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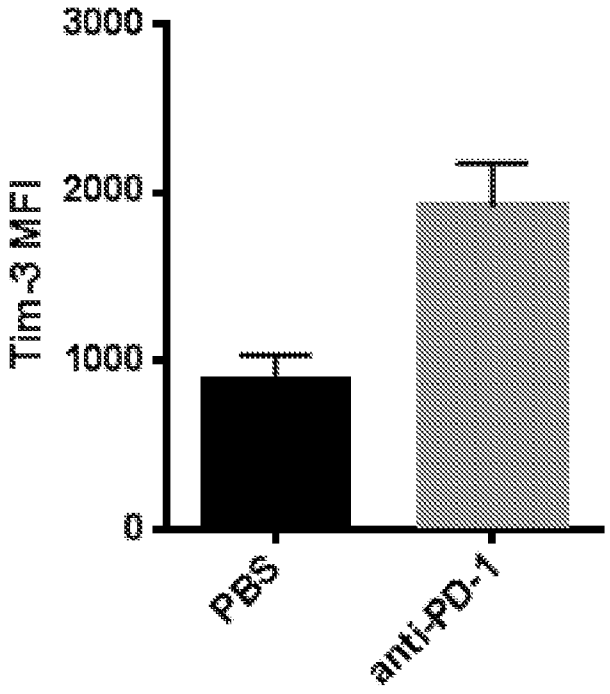
(54) Title: ANTIBODIES SPECIFICALLY BINDING PD-1 AND THEIR USES

(57) Abstract: The present invention relates to antibodies specifically binding PD-1, polynucleotides encoding the antibodies or fragments, and methods of making and using the foregoing.

Figure 1A.



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ANTIBODIES SPECIFICALLY BINDING PD-1 AND THEIR USES

SEQUENCE LISTING

This application contains a Sequence Listing submitted via EFS-Web, the entire content incorporated herein by reference in its entirety. The ASCII text file, created on 28 October 2016, is named JBI5071WOPCT_ST25.txt and is 418 kilobytes in size.

FIELD OF THE INVENTION

The present invention relates antibodies specifically binding PD-1, polynucleotides encoding the antibodies or fragments, and methods of making and using the foregoing.

BACKGROUND OF THE INVENTION

The immune system is tightly controlled by a network of costimulatory and co-inhibitory ligands and receptors. These molecules provide secondary signals for T cell activation and provide a balanced network of positive and negative signals to maximize immune responses against infection and tumors, while limiting immunity to self (Wang *et al.,* (Epub Mar. 7, 2011) *J Exp Med* 208(3):577-92; Lepenies *et al.*, (2008) *Endocr Metab Immune Disord Drug Targets* 8:279-288).

Immune checkpoint therapy, targeting co-inhibitory pathways in T cells to promote antitumor immune responses, has led to advances in clinical care of cancer patients.

PD-1 is a negative immune checkpoint molecule that suppresses $CD4^+$ and $CD8^+$ T cell functions in the tumor microenvironment (TME). PD-1 engagement with its ligands (PD-L1 and PD-L2) drives T cell anergy and exhaustion in tumors by inhibiting multiple pathways downstream of the T cell receptor signaling, resulting in decreased T cell survival, growth and proliferation, compromised effector function, and altered metabolism. Preclinical studies have demonstrated that the PD-1 pathway blockade can reverse T cell exhaustion and stimulate anti-tumor immunity.

The PD-1 pathway hence contributes to downregulation of T cell functions in the (TME) and evasion of tumors via immune destruction. In the TME, exhausted T cells, in addition to expressing high levels of PD-1, express other inhibitory receptors including CTLA-4, TIM-3, LAG-3, CD244, TIGIT and CD160 (see e.g., Pauken & Wherry; 2015, *Trends in Immunology* 36(4): 265–276).

TIM-3 is a transmembrane receptor that is expressed on Th1 (T helper 1) $CD4^+$ cells and cytotoxic $CD8^+$ T cells that secrete IFN-γ. TIM-3 is generally not expressed on naïve T cells but rather upregulated on activated, effector T cells. TIM-3 has a role in regulating immunity and tolerance *in vivo* (see Hastings et al., (2009) *Eur J Immunol* 39(9):2492-501).

PD-1 antibodies have been described for example in: U.S. Patent Nos. 5,897,862 and 7,488,802, and in Int. Patent Publ. Nos. WO2004/004771, WO2004/056875, WO2006/121168, WO2008/156712, WO2010/029435, WO2010/036959, WO2011/110604, WO2012/145493, WO2014/194302, WO2014/206107, WO2015/036394, WO2015/035606, WO2015/085847, WO2015/112900 and WO2015/112805.

TIM-3 antibodies have been described for example in: Monney *et al., Nature* (2002) 415(6871):536-41, and in Int. Patent Publ. Nos. WO2011/155607, WO2013/006490 and WO2015/117002.

Combinations with TIM-3 antibody and a PD-L1 antibody have been evaluated in for example in Int. Patent Publ. No. WO2011/159877.

While anti-PD-1/PD-L1 antibodies are demonstrating encouraging clinical responses in patients with multiple solid tumors, the response rates are still fairly low, about 15% - 20% in pretreated patients (Swaika *et al., (*2015) *Mol Immunol*. doi: 10.1016/j.molimm.2015.02.009).

Therefore, there is a need for new therapeutics that inhibit the immunosuppressive activity of checkpoint inhibitors such as PD-1 and TIM-3, to be used for cancer immunotherapy and treatment of other conditions that would benefit from enhancement of an immune response, such as chronic infections.

BRIEF SUMMARY OF THE INVENTION

The invention provides an isolated antagonistic antibody specifically binding PD-1, comprising a heavy chain complementarity determining region 1 (HCDR1), a HCDR2 and a HCDR3 of SEQ ID NOs: 82, 83 and 84, respectively, or SEQ ID NOs: 82, 83 and 85, respectively.

The invention also provides an isolated antagonistic antibody specifically binding PD-1, comprising a heavy chain complementarity determining region 1 (HCDR1), a HCDR2 and a HCDR3 of SEQ ID NOs: 82, 83 and 84, respectively, and a light chain complementarity determining region 1 (LCDR1), a LCDR2 and a LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.

The invention also provides an isolated antagonistic antibody specifically binding PD-1, comprising a heavy chain complementarity determining region 1 (HCDR1), a HCDR2 and a HCDR3 of SEQ ID NOs: 82, 83 and 85, respectively, and a light chain complementarity determining region 1 (LCDR1), a LCDR2 and a LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.

The invention also provides an isolated antagonistic antibody specifically binding PD-1, comprising certain HCDR1, HCDR2, HCDR3, LCDR1, LCDR2 and LCDR3 amino acid sequences as described herein.

The invention also provides an isolated antagonistic antibody specifically binding PD-1, comprising certain VH and VL amino acid sequences as described herein.

The invention also provides an immunoconjugate comprising the antibody or antigen-binding portion thereof of the invention linked to a therapeutic agent or to an imaging agent.

The invention also provides a pharmaceutical composition comprising the antibody of the invention and a pharmaceutically accepted carrier.

The invention also provides a polynucleotide encoding the antibody VH, the antibody VL or the antibody VH and the antibody VL of the invention.

The invention also provides a vector comprising the polynucleotide encoding the antibody VH, the antibody VL or the antibody VH and the VL of the invention.

The invention also provides a host cell comprising the vector of the invention.

The invention also provides a method of producing the antibody of the invention, comprising culturing the host cell of the invention in conditions that the antibody is expressed, and recovering the antibody produced by the host cell.

The invention also provides a method of treating a cancer in a subject, comprising administering a therapeutically effective amount of the isolated antibody of the invention to the subject in need thereof for a time sufficient to treat the cancer.

The invention also provides a method of enhancing an immune response in a subject, comprising administering a therapeutically effective amount of the isolated antibody of the invention to the subject in need thereof for a time sufficient to enhance the immune response.

The invention also provides an anti-idiotypic antibody binding to the antibody of the invention.

The invention also provides a kit comprising the antibody of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1A shows that TIM-3 surface expression is elevated in tumors after treatment with anti-PD-1 antibodies. Balb/c mice with established CT26 colon carcinoma tumors were treated biweekly with anti-PD-1 antibody or vehicle. Tumors were harvested at day 22 and TIM-3 expression was evaluated on tumor-infiltrating T cells using flow cytometry. MFI: mean fluorescent intensity. PBS: control

Figure 1B shows that TIM-3 surface expression is elevated on tumor infiltrated lymphocytes (TIL) after treatment with anti-PD-1 antibodies. Balb/c mice with established MC38 colon carcinoma tumors were treated biweekly with anti-PD-1 antibody or vehicle. Geometric mean fluorescent intensity (gMFI) of TIM-3 expression on total CD8 TIL population is shown in vehicle treated (PBS) or anti-PD-1 antibody treated (PD-1) animals. p=0.003 vehicle vs anti-PD-1 antibody treated groups.

Figure 1C shows the relative frequency of TIM-3$^+$ CD8 cells of total CD8$^+$ TILs in MC38 tumors harvested from mice treated with vehicle (PBS) or anti-PD-1 antibody (PD-1). p=0.045 vehicle vs anti-PD-1 antibody treated groups.

Figure 2A shows that CD137 surface expression (gMFI) is elevated on TILs in MC38 colon carcinoma tumors in animals treated with anti-PD-1 antibodies (PD-1 group) when compared to vehicle treated (PBS) group. p=0.005 vehicle vs anti-PD-1 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 2B shows that the relative frequency of CD137$^+$ CD8 cells of total CD8+ TILs in is elevated in MC38 colon carcinoma tumors in animals treated with anti-PD-1 antibodies (PD-1 group) when compared to vehicle treated (PBS) group. p=0.0475 vehicle vs anti-PD-1 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 3A shows that OX40 surface expression (gMFI) is elevated on TILs in MC38 colon carcinoma tumors in animals treated with anti-PD-1 antibodies (PD-1 group) when compared to vehicle treated (PBS) group. p=0.0013 vehicle vs anti-PD-1 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 3B shows that the relative frequency of OX40$^+$ CD8 cells of total CD8$^+$ TILs in is elevated in MC38 colon carcinoma tumors in animals treated with anti-PD-1 antibodies (PD-1 group) when compared to vehicle treated (PBS) group. p=0.03 vehicle vs anti-PD-1 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 4A shows that GITR surface expression (gMFI) is elevated on TILs in MC38 colon carcinoma tumors in animals treated with anti-PD-1 antibodies (PD-1 group) when compared to vehicle treated (PBS) group. p=0.0004 vehicle vs anti-PD-1 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 4B shows that the relative frequency of $GITR^+$ CD8 cells of total $CD8^+$ TILs in is elevated in MC38 colon carcinoma tumors in animals treated with anti-PD-1 antibodies (PD-1 group) when compared to vehicle treated (PBS) group. p=0.0015 vehicle vs anti-PD-1 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 5 shows that treatment with anti-TIM-3 antibodies after anti-PD-1 antibody treatment further induces antigen-specific immune response. The antibodies were tested in the CMV assay using PBMCs from CMV positive donors, in which antigen-specific immune responses were induced with pp65 peptide pools. The cells were treated for 5 days with anti-PD-1 antibody PD1B244, re-stimulated, and treated for 24 hours with anti-TIM-3 antibody TM3B105. Immune response was determined by measuring increases in IFN-γ secretion. IgG2s Iso: IgG2sigma isotype control. CMV: sample treated with cytomegalovirus p65 peptides in the absence of antibodies.

Figure 6 shows the HCDR1 sequences of select anti-PD-1 antibodies and the HCDR1 genus sequence.

Figure 7 shows the HCDR2 sequences of select anti-PD-1 antibodies and the HCDR2 genus sequence.

Figure 8 shows the HCDR3 sequences of select anti-PD-1 antibodies and the first HCDR3 genus sequence.

Figure 9 shows the HCDR3 sequences of select anti-PD-1 antibodies and the second HCDR3 genus sequence.

Figure 10 shows the LCDR1 sequences of select anti-PD-1 antibodies and the LCDR1 genus sequence.

Figure 11 shows the LCDR2 sequences of select anti-PD-1 antibodies and the LCDR2 genus sequence.

Figure 12 shows the LCDR3 sequences of select anti-PD-1 antibodies and the LCDR3 genus sequence.

Figure 13 shows the HCDR1 sequences of select anti-TIM-3 antibodies and the HCDR1 genus sequence. The genus sequence was determined by generating molecular models for all Fv (VH/VL pairs) in MOE (CCG, Montreal) using a default protocol for antibody modeling.

For CDRs that have different lengths, these structural models were aligned based upon the structurally conserved regions and the structurally equivalent CDRs positions were identified.

Figure 14 shows the HCDR2 sequences of select anti-TIM-3 antibodies and the HCDR2 genus sequence. The HCDR2 genus sequence was generated as described for Figure 10.

Figure 15 shows the HCDR3 sequences of select anti-TIM-3 antibodies and the first HCDR3 genus sequence. The HCDR3 genus sequence was generated as described for Figure 10.

Figure 16 shows the LCDR1 sequences of select anti-TIM-3 antibodies and the LCDR1 genus sequence. The LCDR1 genus sequence was generated as described for Figure 10.

Figure 17 shows the LCDR2 sequences of select anti-TIM-3 antibodies and the LCDR2 genus sequence. The LCDR2 genus sequence was generated as described for Figure 10.

Figure 18 shows the LCDR3 sequences of select anti-TIM-3 antibodies and the LCDR3 genus sequence. The LCDR3 genus sequence was generated as described for Figure 10.

Figure 19A shows that TIGIT surface expression (gMFI) is elevated on TILs in MC38 colon carcinoma tumors in animals treated with anti-TIM-3 antibodies (TIM-3 group) when compared to vehicle treated (PBS) group. $p=0.0181$ vehicle vs anti-TIM-3 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 19B shows that the relative frequency of TIGIT+ CD8 cells of total CD8+ TILs in is elevated in MC38 colon carcinoma tumors in animals treated with anti-TIM-3 antibodies (TIM-3 group) when compared to vehicle treated (PBS) group. $p=0.0475$ vehicle vs anti-TIM-3 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 20A shows that TIGIT surface expression (gMFI) is elevated on TILs in CT26 colon carcinoma tumors in animals treated with anti-TIM-3 antibodies (TIM-3 group) when compared to vehicle treated (PBS) group. $p<0.001$ vehicle vs anti-TIM-3 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 20B shows that the relative frequency of TIGIT+ CD8 cells of total CD8+ TILs in is elevated in CT26 colon carcinoma tumors in animals treated with anti-TIM-3 antibodies (TIM-3 group) when compared to vehicle treated (PBS) group. $p=0.0105$ vehicle vs anti-TIM-3 antibody treated groups. Each point represents one mouse. Data are representative of at least 2 independent experiments.

Figure 21 shows upregulation of TIM-3 expression on peripheral T cells in melanoma patients PBMCs from treatment naïve melanoma patients stimulated with melanoma antigen peptide pools (NY-ESO, gp100, MART-1) in the presence or absence of anti-PD-1

or anti-TIM-3 function blocking antibodies. Expression of TIM-3 was determined by flow cytometry on restimulated cells on day 6.

Figure 22A shows that TM3B403 treatment increases frequency of activated NK cells in IL-2 stimulated human PBMCs. IgG2s: Isotype control. NK cell activation was assessed as percentage (%) of CD69 expressing cells in the stimulated PBMCs.

Figure 22B shows that TM3B403 treatment increases frequency of activated NK cells in IL-2 stimulated human PBMCs. IgG2s: Isotype control. NK cell activation was assessed as percentage (%) of CD25 expressing cells in the stimulated PBMCs.

DETAILED DESCRIPTION OF THE INVENTION

All publications, including but not limited to patents and patent applications, cited in this specification are herein incorporated by reference as though fully set forth.

It is to be understood that the terminology used herein is for the purpose of describing particular embodiments only and is not intended to be limiting. Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which the invention pertains.

Although any methods and materials similar or equivalent to those described herein may be used in the practice for testing of the present invention, exemplary materials and methods are described herein. In describing and claiming the present invention, the following terminology will be used.

As used in this specification and the appended claims, the singular forms "a," "an," and "the" include plural referents unless the content clearly dictates otherwise. Thus, for example, reference to "a cell" includes a combination of two or more cells, and the like.

"Specific binding" or "specifically binds" or "binds" refers to an antibody binding to an antigen or an epitope within the antigen with greater affinity than for other antigens. Typically, the antibody binds to the antigen or the epitope within the antigen with an equilibrium dissociation constant (K_D) of about $1x10^{-8}$ M or less, for example about $1x10^{-9}$ M or less, about $1x10^{-10}$ M or less, about $1x10^{-11}$ M or less, or about $1x10^{-12}$ M or less, typically with the K_D that is at least one hundred fold less than its K_D for binding to a non-specific antigen (e.g., BSA, casein). The dissociation constant may be measured using standard procedures. Antibodies that specifically bind to the antigen or the epitope within the antigen may, however, have cross-reactivity to other related antigens, for example to the same antigen from other species (homologs), such as human or monkey, for example *Macaca fascicularis* (cynomolgus, cyno), *Pan troglodytes* (chimpanzee, chimp) or

Callithrix jacchus (common marmoset, marmoset). While a monospecific antibody specifically binds one antigen or one epitope, a bispecific antibody specifically binds two distinct antigens or two distinct epitopes.

"Antibodies" is meant in a broad sense and includes immunoglobulin molecules including monoclonal antibodies including murine, human, humanized and chimeric monoclonal antibodies, antigen-binding fragments, bispecific or multispecific antibodies, dimeric, tetrameric or multimeric antibodies, single chain antibodies, domain antibodies and any other modified configuration of the immunoglobulin molecule that comprises an antigen binding site of the required specificity. "Full length antibodies" are comprised of two heavy (H) chains and two light (L) chains inter-connected by disulfide bonds as well as multimers thereof (for example IgM). Each heavy chain is comprised of a heavy chain variable region (VH) and a heavy chain constant region (comprised of domains CH1, hinge CH2 and CH3). Each light chain is comprised of a light chain variable region (VL) and a light chain constant region (CL). The VH and the VL regions may be further subdivided into regions of hypervariability, termed complementarity determining regions (CDR), interspersed with framework regions (FR). Each VH and VL is composed of three CDRs and four FR segments, arranged from amino-terminus to carboxy-terminus in the following order: FR1, CDR1, FR2, CDR2, FR3, CDR3, and FR4.

"Complementarity determining regions (CDR)" are "antigen binding sites" in an antibody. CDRs may be defined using various terms: (i) Complementarity Determining Regions (CDRs), three in the VH (HCDR1, HCDR2, HCDR3) and three in the VL (LCDR1, LCDR2, LCDR3) are based on sequence variability (Wu and Kabat, (1970) *J Exp Med* 132:211-50; Kabat *et al.,* Sequences of Proteins of Immunological Interest, 5th Ed. Public Health Service, National Institutes of Health, Bethesda, Md., 1991). (ii) "Hypervariable regions", "HVR", or "HV", three in the VH (H1, H2, H3) and three in the VL (L1, L2, L3) refer to the regions of an antibody variable domains which are hypervariable in structure as defined by Chothia and Lesk (Chothia and Lesk, (1987) *Mol Biol* 196:901-17). The International ImMunoGeneTics (IMGT) database (http://www_imgt_org) provides a standardized numbering and definition of antigen-binding sites. The correspondence between CDRs, HVs and IMGT delineations is described in Lefranc *et al.,* (2003) *Dev Comparat Immunol* 27:55-77. The term "CDR", "HCDR1", "HCDR2", "HCDR3", "LCDR1", "LCDR2" and "LCDR3" as used herein includes CDRs defined by any of the methods described *supra,* Kabat, Chothia or IMGT, unless otherwise explicitly stated in the specification.

Immunoglobulins may be assigned to five major classes, IgA, IgD, IgE, IgG and IgM, depending on the heavy chain constant domain amino acid sequence. IgA and IgG are further sub-classified as the isotypes IgA1, IgA2, IgG1, IgG2, IgG3 and IgG4. Antibody light chains of any vertebrate species may assigned to one of two clearly distinct types, namely kappa (κ) and lambda (λ), based on the amino acid sequences of their constant domains.

"Antibody fragments" or "antigen-binding portion" refers to a portion of an immunoglobulin molecule that retains the antigen binding properties of the parental full length antibody. Exemplary antigen-binding portions are heavy chain complementarity determining regions (HCDR) 1, 2 and 3, light chain complementarity determining regions (LCDR) 1, 2 and 3, a heavy chain variable region (VH), a light chain variable region (VL), Fab, F(ab')2, Fd and Fv fragments as well as domain antibodies (dAb) consisting of either one VH or VL domain. VH and VL domains may be linked together via a synthetic linker to form various types of single chain antibody designs where the VH/VL domains may pair intramolecularly, or intermolecularly in those cases when the VH and VL domains are expressed by separate single chain antibody constructs, to form a monovalent antigen binding site, such as single chain Fv (scFv) or diabody; described for example in Int. Patent Publ. Nos. WO1998/44001, WO1988/01649, WO1994/13804 and WO1992/01047.

"Monoclonal antibody" refers to an antibody population with single amino acid composition in each heavy and each light chain, except for possible well known alterations such as removal of C-terminal lysine from the antibody heavy chain. Monoclonal antibodies typically bind one antigenic epitope, except that multispecific monoclonal antibodies bind two or more distinct antigens or epitopes. Bispecific monoclonal antibodies bind two distinct antigenic epitopes. Monoclonal antibodies may have heterogeneous glycosylation within the antibody population. Monoclonal antibodies may be monospecific or multispecific, or monovalent, bivalent or multivalent. A multispecific antibody, such as a bispecific antibody or a trispecific antibody is included in the term monoclonal antibody.

"Isolated antibody" refers to an antibody or antibody fragment that is substantially free of other antibodies having different antigenic specificities (e.g., an isolated antibody specifically binding PD-1 is substantially free of antibodies that specifically bind antigens other than PD-1). An isolated antibody specifically binding TIM-3 is substantially free of antibodies that specifically bind antigens other than TIM-3. In case of bispecific PD-1/TIM-3 antibodies, the bispecific antibody specifically binds both PD-1 and TIM-3, and is substantially free of antibodies that specifically bind antigens other that PD-1 and TIM-

3. "Isolated antibody" encompasses antibodies that are isolated to a higher purity, such as antibodies that are 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% pure.

"Humanized antibodies" refers to antibodies in which at least one CDR is derived from non-human species and the variable region frameworks are derived from human immunoglobulin sequences. Humanized antibodies may include intentionally introduced mutations in the framework regions so that the framework may not be an exact copy of expressed human immunoglobulin or germline gene sequences.

"Human antibody" refers to an antibody having heavy and light chain variable regions in which both the framework and all 6 CDRs are derived from sequences of human origin. If the antibody contains a constant region or a portion of the constant region, the constant region also is derived from sequences of human origin.

Human antibody comprises heavy or light chain variable regions that are "derived from" sequences of human origin if the variable regions of the antibody are obtained from a system that uses human germline immunoglobulin or rearranged immunoglobulin genes. Such exemplary systems are human immunoglobulin gene libraries displayed on phage, and transgenic non-human animals such as mice or rats carrying human immunoglobulin loci as described herein. "Human antibody" may contain amino acid differences when compared to the human germline immunoglobulin or rearranged immunoglobulin genes due to for example naturally occurring somatic mutations or intentional introduction of substitutions into the framework or antigen binding site, or both. Typically, "human antibody" is at least about 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% identical in amino acid sequence to an amino acid sequence encoded by human germline immunoglobulin or rearranged immunoglobulin genes. In some cases, "human antibody" may contain consensus framework sequences derived from human framework sequence analyses, for example as described in Knappik *et al.*, (2000) *J Mol Biol* 296:57-86, or synthetic HCDR3 incorporated into human immunoglobulin gene libraries displayed on phage, for example as described in Shi *et al.,* (2010) *J Mol Biol* 397:385-96, and in Int. Patent Publ. No. WO2009/085462.

Human antibodies derived from human immunoglobulin sequences may be generated using systems such as phage display incorporating synthetic CDRs and/or synthetic frameworks, or may be subjected to *in vitro* mutagenesis to improve antibody properties, resulting in antibodies that are not expressed by the human antibody germline repertoire *in vivo.*

"Recombinant" refers to antibodies and other proteins that are prepared, expressed, created or isolated by recombinant means.

"Epitope" refers to a portion of an antigen to which an antibody specifically binds. Epitopes typically consist of chemically active (such as polar, non-polar or hydrophobic) surface groupings of moieties such as amino acids or polysaccharide side chains and may have specific three-dimensional structural characteristics, as well as specific charge characteristics. An epitope may be composed of contiguous and/or discontiguous amino acids that form a conformational spatial unit. For a discontiguous epitope, amino acids from differing portions of the linear sequence of the antigen come in close proximity in 3-dimensional space through the folding of the protein molecule. Antibody "epitope" depends on the methodology used to identify the epitope.

"Multispecific" refers to an antibody that specifically binds at least two distinct antigens or two distinct epitopes within the antigens, for example three, four or five distinct antigens or epitopes.

"Bispecific" refers to an antibody that specifically binds two distinct antigens or two distinct epitopes within the same antigen. The bispecific antibody may have cross-reactivity to other related antigens, for example to the same antigen from other species (homologs), such as human or monkey, for example *Macaca fascicularis* (cynomolgus, cyno), *Pan troglodytes* (chimpanzee, chimp) or *Callithrix jacchus* (common marmoset, marmoset), or may bind an epitope that is shared between two or more distinct antigens.

"Variant" refers to a polypeptide or a polynucleotide that differs from a reference polypeptide or a reference polynucleotide by one or more modifications for example, substitutions, insertions or deletions.

"Vector" refers to a polynucleotide capable of being duplicated within a biological system or that can be moved between such systems. Vector polynucleotides typically contain elements, such as origins of replication, polyadenylation signal or selection markers, that function to facilitate the duplication or maintenance of these polynucleotides in a biological system. Examples of such biological systems may include a cell, virus, animal, plant, and reconstituted biological systems utilizing biological components capable of duplicating a vector. The polynucleotide comprising a vector may be DNA or RNA molecules or a hybrid of these.

"Expression vector" refers to a vector that can be utilized in a biological system or in a reconstituted biological system to direct the translation of a polypeptide encoded by a polynucleotide sequence present in the expression vector.

"Polynucleotide" refers to a synthetic molecule comprising a chain of nucleotides covalently linked by a sugar-phosphate backbone or other equivalent covalent chemistry. cDNA is a typical example of a polynucleotide.

"Polypeptide" or "protein" refers to a molecule that comprises at least two amino acid residues linked by a peptide bond to form a polypeptide. Small polypeptides of less than 50 amino acids may be referred to as "peptides".

PD-1 refers to human programmed cell death protein 1, PD-1. PD-1 is also known as CD279 or PDCD1. The amino acid sequence of the mature human PD-1 (without signal sequence) is shown in **SEQ ID NO: 1**. The extracellular domain spans residues 1-150, the transmembrane domain spans residues 151-171 and the cytoplasmic domain spans residues 172-268 of SEQ ID NO: 1. Throughout the specification, " the extracellular domain of human PD-1 "huPD1-ECD" refers to protein having amino acid sequence of residues 1-149 of SEQ ID NO: 1, and shown in **SEQ ID NO:2**. "PD-1" in the specification refers to human mature PD-1, unless explicitly stated to the contrary.

TIM-3 refers to human hepatitis A virus cellular receptor 2, also called HAVCR2. The amino acid sequence of the mature human TIM-3 (without signal sequence) is shown **in SEQ ID NO: 138.** The extracellular domain spans residues 1-181, the transmembrane domain spans residues 182-202 and the cytoplasmic domain spans residues 203-280 of SEQ ID NO: 138. Throughout the specification, "the extracellular domain of human TIM-3 "huTIM-3-ECD" refers to protein having amino acid sequence of residues 1-179 of SEQ ID NO: 138, and shown in **SEQ ID NO: 89.** TIM-3 in the specification refers to human mature TIM-3, unless explicitly stated to the contrary.

"In combination with" means that two or more therapeutics are administered to a subject together in a mixture, concurrently as single agents or sequentially as single agents in any order.

"Overexpress", "overexpressed" and "overexpressing" is used interchangeably and refers to a sample such as a cancer cell, malignant cell or cancer tissue that has measurably higher levels of PD-1, TIM-3, PD-L1, PD-L2 or TIM-3 ligand when compared to a reference sample. The overexpression may be caused by gene amplification or by increased transcription or translation. Expression and overexpression of protein in the sample may be measured using well know assays using for example ELISA, immunofluorescence, flow cytometry or radioimmunoassay on live or lysed cells. Expression and overexpression of a polynucleotide in the sample may be measured for example using fluorescent *in situ* hybridization, Southern blotting, or PCR techniques. A protein or a polynucleotide is overexpressed when the level of the protein or the

polynucleotide in the sample at least 1.5-fold higher or statistically significant when compared to the reference sample. Selection of the reference sample is known.

"Sample" refers to a collection of similar fluids, cells, or tissues isolated from a subject, as well as fluids, cells, or tissues present within a subject. Exemplary samples are biological fluids such as blood, serum and serosal fluids, plasma, lymph, urine, saliva, cystic fluid, tear drops, feces, sputum, mucosal secretions of the secretory tissues and organs, vaginal secretions, ascites fluids such as those associated with non-solid tumors, fluids of the pleural, pericardial, peritoneal, abdominal and other body cavities, fluids collected by bronchial lavage, liquid solutions contacted with a subject or biological source, for example, cell and organ culture medium including cell or organ conditioned medium, lavage fluids and the like, tissue biopsies, fine needle aspirations or surgically resected tumor tissue.

A "cancer cell" or a "tumor cell" refers to a cancerous, pre-cancerous or transformed cell, either *in vivo*, *ex vivo*, or in tissue culture, that has spontaneous or induced phenotypic changes. These changes do not necessarily involve the uptake of new genetic material. Although transformation may arise from infection with a transforming virus and incorporation of new genomic nucleic acid, uptake of exogenous nucleic acid or it can also arise spontaneously or following exposure to a carcinogen, thereby mutating an endogenous gene. Transformation/cancer is exemplified by morphological changes, immortalization of cells, aberrant growth control, foci formation, proliferation, malignancy, modulation of tumor specific marker levels, invasiveness, tumor growth in suitable animal hosts such as nude mice, and the like, *in vitro*, *in vivo*, and *ex vivo* (Freshney, Culture of Animal Cells: A Manual of Basic Technique (3rd ed. 1994)).

"About" means within an acceptable error range for the particular value as determined by one of ordinary skill in the art, which will depend in part on how the value is measured or determined, i.e., the limitations of the measurement system. Unless explicitly stated otherwise within the Examples or elsewhere in the Specification in the context of a particular assay, result or embodiment, "about" means within one standard deviation per the practice in the art, or a range of up to 5%, whichever is larger.

"Bispecific PD-1/TIM-3 antibody", "PD-1/TIM-3 antibody", "bispecific anti-PD-1/TIM-3 antibody" or "anti-PD-1/TIM-3 antibody" refers to a molecule comprising at least one binding domain specifically binding PD-1 and at least one binding domain specifically binding TIM-3. The domains specifically binding PD-1 and TIM-3 are typically VH/VL pairs. The bispecific anti-PD-1/TIM-3 antibody may be monovalent in terms of its binding to either PD-1 or TIM-3.

"Valent" refers to the presence of a specified number of binding sites specific for an antigen in a molecule. As such, the terms "monovalent", "bivalent", "tetravalent", and "hexavalent" refer to the presence of one, two, four and six binding sites, respectively, specific for an antigen in a molecule.

"An antigen specific $CD4^+$ or $CD8^+$ T cell" refers to a $CD4^+$ or $CD8^+$ T cell activated by a specific antigen, or immunostimulatory epitope thereof.

"CD137" (also called tumor necrosis factor receptor superfamily member 9, TNFRSF9, 4-1BBL) refers to a human CD137 molecule having the amino acid sequence shown in SEQ ID NO: **281.**

SEQ ID NO: 281
MGNSCYNIVATLLLVLNFERTRSLQDPCSNCPAGTFCDNNRNQICSPCPPNSFSSA
GGQRTCDICRQCKGVFRTRKECSSTSNAECDCTPGFHCLGAGCSMCEQDCKQGQ
ELTKKGCKDCCFGTFNDQKRGICRPWTNCSLDGKSVLVNGTKERDVVCGPSPAD
LSPGASSVTPPAPAREPGHSPQIISFFLALTSTALLFLLFFLTLRFSVVKRGRKKLLYI
FKQPFMRPVQTTQEEDGCSCRFPEEEEGGCEL

"TIGIT" (also called T-cell immunoreceptor with Ig and ITIM domains) refers to human TIGIT molecule having the amino acid sequence shown in SEQ ID NO: **301.**

SEQ ID NO: 301
MMTGTIETTGNISAEKGGSIILQCHLSSTTAQVTQVNWEQQDQLLAICNADLGWHI
SPSFKDRVAPGPGLGLTLQSLTVNDTGEYFCIYHTYPDGTYTGRIFLEVLESSVAEH
GARFQIPLLGAMAATLVVICTAVIVVVALTRKKKALRIHSVEGDLRRKSAGQEEW
SPSAPSPPGSCVQAEAAPAGLCGEQRGEDCAELHDYFNVLSYRSLGNCSFFTETG

"Agonist" refers to a molecule that, when bound to a cellular protein, induces at least one reaction or activity that is induced by a natural ligand of the protein. The molecule is an agonist when the at least one reaction or activity is induced by at least about 30%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or 100% greater than the at least one reaction or activity induced in the absence of the agonist (*e.g.*, negative control), or when the induction is statistically significant when compared to the induction in the absence of the agonist. Agonist may be an antibody, a soluble ligand, or a small molecule. An exemplary agonist is an agonistic antibody that specifically binds a T cell activating molecule.

"Antagonist" refers to a molecule that, when bound to a cellular protein, suppresses at least one reaction or activity that is induced by a natural ligand of the protein. A molecule is an antagonist when the at least one reaction or activity is suppressed by at least about 30%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, or 100% more than the at least one reaction or activity suppressed in the absence of the antagonist (*e.g.,* negative control), or when the suppression is statistically significant when compared to the suppression in the absence of the antagonist. Antagonist may be an antibody, a soluble ligand, a small molecule, a DNA or RNA such as siRNA. Exemplary antagonists are an antagonistic antibody specifically binding PD-1, an antagonistic antibody specifically binding TIM-3, an antagonistic bispecific PD-1/TIM-3 antibody or an antagonistic antibody specifically binding a T cell inhibitory molecule. A typical reaction or activity that is induced by PD-1 binding to its receptor PD-L1 or PD-L2 may be reduced antigen-specific $CD4^+$ or $CD8^+$ cell proliferation or reduced interferon-γ (IFN-γ) production by T cells, resulting in suppression of immune responses against for example tumor. A typical reaction or activity that is induced by TIM-3 binding to its receptor, such as galectin-9, may be reduced antigen specific $CD4^+$ or $CD8^+$ cell proliferation, reduced IFN-γ production by T cells, or reduced CD137 surface expression on $CD4^+$ or $CD8^+$ cells, resulting in suppression of immune responses against for example tumor. Similarly, a typical reaction or activity that is induced by a T cell inhibitory molecule is immunosuppression. Hence, an antagonistic PD-1 antibody specifically binding PD-1, an antagonistic antibody specifically binding TIM-3, an antagonistic bispecific PD-1/TIM-3 antibody, or an antagonistic antibody specifically binding a T cell inhibitory molecule induces immune responses by inhibiting the inhibitory pathways.

"Subject" includes any human or nonhuman animal. "Nonhuman animal" includes all vertebrates, *e.g.,* mammals and non-mammals, such as nonhuman primates, sheep, dogs, cats, horses, cows chickens, amphibians, reptiles, etc. Except when noted, the terms "patient" or "subject" are used interchangeably.

The numbering of amino acid residues in the antibody constant region throughout the specification is according to the EU index as described in Kabat *et al.,* Sequences of Proteins of Immunological Interest, 5th Ed. Public Health Service, National Institutes of Health, Bethesda, MD. (1991), unless otherwise explicitly stated.

Conventional one and three-letter amino acid codes are used herein as shown in **Table 1.**

Table 1.

Amino acid	Three-letter code	One-letter code
Alanine	Ala	A
Arginine	Arg	R
Asparagine	Asn	N
Aspartate	Asp	D
Cysteine	Cys	C
Glutamate	Gln	E
Glutamine	Glu	Q
Glycine	Gly	G
Histidine	His	H
Isoleucine	Ile	I
Lysine	Lys	K
Methionine	Met	M
Phenylalanine	Phe	F
Proline	Pro	P
Serine	Ser	S
Threonine	Thr	T
Tryptophan	Trp	W
Tyrosine	Tyr	Y
Valine	Val	V

Compositions of matter

The present invention provides antagonistic antibodies specifically binding PD-1, antagonistic antibodies specifically binding TIM-3, and antagonistic bispecific PD-1/TIM-3 antibodies. The present invention provides polypeptides and polynucleotides encoding the antibodies of the invention or complementary nucleic acids thereof, vectors, host cells, and methods of making and using them.

Antagonistic antibodies specifically binding PD-1

PD-1, upon ligand engagement, suppresses T cell functions through multiple mechanisms (Pauken & Wherry (2015) *Trends in Immunology* 36(4): 265–276). PD-1 engagement directly inhibits T cell receptor (TCR) signaling through co-localization with the TCR and subsequent induction of dephosphorylation of TCR proximal signaling molecules, inhibition of Ras/MEK/ERK pathway leading to inhibition of the cell cycle progression and T cell proliferation, inhibition of cell growth and survival and reprogramming of T cell metabolism through suppression of PI3K/AKT pathway, leading to the upregulation of the BATF transcription factor, and modulation of development, maintenance and function of regulatory T cells. PD-1 has also been proposed to increase T cell motility and to limit duration of interaction between T cells and target cells, thereby reducing the extent of T cell activation (Honda *et al.,* (2014) *Immunity* 40(2):235-47).

Tumors have co-opted the PD-1 pathway to downregulate T cell function in the tumor microenvironment (TME) and to evade immune destruction. In the TME, under conditions of persistent antigen and inflammation, T cells become exhausted, or dysfunctional, and progressively lose their effector function and proliferative capacity. Exhausted T cells express high levels of PD-1, often together with other inhibitory receptors such as TIM-3 or LAG-3 (Pauken & Wherry (2015) *Trends in Immunology* 36(4): 265–276). One of the PD-1 ligands, PD-L1, is also upregulated in various tumors. PD-L1 expression occurs on the cancer cells themselves and/or infiltrating immune cells, including tumor associated macrophages, dendritic cells, fibroblasts and activated T cells (Chen *et al.,* 2012 *Clin Cancer Res* 18(24):6580-7). In this setting, PD-1 engagement is hypothesized to limit anti-tumor T cell responses and lead to immune evasion. Recent studies have shown that a higher frequency and level of PD-1 expression occurs on tumor infiltrating lymphocytes (TILs) in multiple solid tumors. Importantly, these PD-1$^+$ TILs are functionally impaired, as evidenced by lower proliferation and effector functions (Pauken & Wherry; 2015, *Trends in Immunology* 36(4): 265–276) These data support the hypothesis that PD-1 mediates immune suppression in the TME.

T cell exhaustion in tumors is reversible, at least partially, by PD-1 pathway blockade. Anti-PD-1/PD-L1 antibodies have been shown to enhance T cell function and lead to improved anti-tumor immunity in a number of preclinical tumor models. PD-1/PD-L1 antibodies have also shown encouraging clinical responses in multiple solid tumors, with 20-40% overall response rate (ORR) in melanoma, 10-24% in non-small cell lung cancer (NSCLC), 12-31% in renal cell carcinoma (RCC), 24-52% in bladder cancer, and 20% in head and neck cancer (Swaika *et al.,* (2015) *Mol Immunol* 67(2 Pt A):4-17).

The invention provides an isolated antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof comprising a heavy chain complementarity determining region 1 (HCDR1), a HCDR2 and a HCDR3 of SEQ ID NOs: 82, 83 and 84, respectively, or SEQ ID NOs: 82, 83 and 85, respectively.

The invention also provides an isolated antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof comprising a light chain complementarity determining region 1 (LCDR1), a LCDR2 and a LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.

The invention also provides an isolated antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof comprising the HCDR1, the HCDR2 and the HCDR3 of SEQ ID NOs: 82, 83 and 84, respectively, and the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.

The invention also provides an isolated antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof comprising the HCDR1, the HCDR2 and the HCDR3 of SEQ ID NOs: 82, 83 and 85, respectively, and the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.

SEQ ID NOs: 82, 83, 84, 85, 86, 87 and 88 represent the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 genus sequences of affinity-matured variants of antagonistic antibodies specifically binding PD-1 having similar HCDR1, HCDR2, LCDR1, LCDR2 and LCDR3 sequences, and two similar HCDR3 groups of sequences. Antibodies within the genus bind PD-1 with the K_D of less than about $1x10^{-7}$ M, such as less than about $1x10^{-8}$ M, for example less than about $1x10^{-9}$ M, or for example less than about $1x10^{-10}$ M. Exemplary such antibodies are antibodies having the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 amino acid sequences of antibodies PD1B114, PD1B149, PD1B160, PD1B162, PD1B164, PD1B11, PD1B183, PD1B184, PD1B185, PD1B187, PD1B71, PD1B177, PD1B70, PD1B175, PD1B194, PD1B195, PD1B196, PD1B197, PD1B198, PD1B199, PD1B200, PD1B201 and PD1B244 as described herein.

SEQ ID NO: 82

$X_1YX_2IX_3$,

wherein

X_1 is S or D;

X_2 is V or A; and

X_3 is H or S.

SEQ ID NO: 83
GIIPIX_4X_5TANYAQKFQG,
wherein
X_4 is Y or F; and
X_5 is G or D.

SEQ ID NO: 84
PGLAAAYDTGX_6LDY,
wherein
X_6 is N or S.

SEQ ID NO: 85
G$X_7X_8X_9X_{10}$TGX_{11}LDY,
wherein
X_7 is T or Y;
X_8 is L or V;
X_9 is D or R;
X_{10} is R or A; and
X_{11} is H or M.

SEQ ID NO: 86
RASQSV$X_{12}X_{13}$YLA,
wherein
X_{12} is S, R or D; and
X_{13} is S or N.

SEQ ID NO: 87
DASX_{14}RAT,
wherein
X_{14} is N, D, Y, S or T.

SEQ ID NO: 88
QQR$X_{15}X_{16}$WPLT,
wherein

X_{15} is S, N, G, E, D, W or A; and
X_{16} is N, Y, E or A.

In some embodiments, the isolated antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof has one, two, three, four or five of the following properties:

a) enhances an activation of antigen specific $CD4^+$ or $CD8^+$ T cells in a dose dependent manner, wherein the activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1;
b) binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;
c) binds human PD-1 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;
d) binds cynomolgus PD-1 with the K_D of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C, or
e) binds cynomolgus PD-1 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

Exemplary such antibodies are PD-1 antibodies PD1B114, PD1B149, PD1B160, PD1B162, PD1B164, PD1B11, PD1B183, PD1B184, PD1B185, PD1B187, PD1B71, PD1B177, PD1B70, PD1B175, PD1B194, PD1B195, PD1B196, PD1B197, PD1B198, PD1B199, PD1B200, PD1B201 and PD1B244 as described herein.

In some embodiments, the isolated antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof enhances an activation of antigen specific $CD4^+$ or $CD8^+$ T cells in a dose dependent manner, wherein the activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1, and binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the isolated antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof enhances an activation of antigen specific $CD4^+$ or $CD8^+$ T cells in dose dependent manner, wherein the activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1, and binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 10 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the isolated antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof enhances an activation of antigen specific $CD4^+$ or $CD8^+$ T cells in dose dependent manner, wherein the activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1, and binds cynomolgus PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25ºC.

In some embodiments, the isolated antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof enhances an activation of antigen specific $CD4^+$ or $CD8^+$ T cells in dose dependent manner, wherein the activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1, and binds cynomolgus PD-1 with an equilibrium dissociation constant (K_D) of less than about 10 nM, wherein the K_D is measured using ProteOn XPR36 system at +25ºC.

Activation of antigen specific $CD4^+$ or $CD8^+$ T cells may be assessed by measuring increased T cell proliferation in a Mixed Lymphocyte Reaction (MLR) assay, increased interferon-γ (IFN-γ) secretion in the MLR assay, increased TNF-α secretion in the MLR assay, increased IFN-γ secretion in a cytomegalovirus antigen assay (CMV assay) or increased TNF-α secretion in the CMV assay using known protocols and those described in Example 1. Antibodies of the invention enhance the activation of antigen specific $CD4^+$ or $CD8^+$ T when the measured T cell functionality is increased by the antibodies of the invention in a dose-dependent manner.

The affinity of an antibody to human or cynomolgus PD-1 may be determined experimentally using any suitable method. Such methods may utilize ProteOn XPR36, Biacore 3000 or KinExA instrumentation, ELISA or competitive binding assays known to those skilled in the art. The measured affinity of a particular antibody/ PD-1 interaction may vary if measured under different conditions (e.g., osmolarity, pH). Thus, measurements of affinity and other binding parameters (e.g., K_D, K_{on}, K_{off}) are typically made with standardized conditions and a standardized buffer, such as the buffer described herein. Skilled in the art will appreciate that the internal error for affinity measurements for example using Biacore 3000 or ProteOn (measured as standard deviation, SD) may typically be within 5-33% for measurements within the typical limits of detection. Therefore the term "about" in the context of K_D reflects the typical standard deviation in the assay. For example, the typical SD for a K_D of $1x10^{-9}$ M is up to $\pm 0.33x10^{-9}$ M.

In some embodiments, the antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof comprises the HCDR1, the HCDR2 and the HCDR3

contained within a heavy chain variable region (VH) of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47 or 48, wherein the HCDR1, the HCDR2 and the HCDR3 are defined by Chothia, Kabat, or IMGT.

In some embodiments, the antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof of the invention comprises the LCDR1, the LCDR2 and the LCDR3 contained within a light chain variable region (VL) of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61 or 62, wherein the LCDR1, the LCDR2 and the LCDR are defined by Chothia, Kabat, or IMGT.

In some embodiments, the antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof of the invention comprises

the HCDR1 of SEQ ID NOs: 10, 11 or 12;

the HCDR2 of SEQ ID NOs: 13, 14 or 15; and

the HCDR3 of SEQ ID NOs: 16, 17, 18 or 19.

In some embodiments, the antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof of the invention comprises

the LCDR1 of SEQ ID NOs: 20, 21, 22, 23, 24 or 25;

the LCDR2 of SEQ ID NOs: 26, 27, 28, 29 or 30; and

the LCDR3 of SEQ ID NOs: 31, 32, 33, 34, 35, 36, 37, 38, 39, or 40.

In some embodiments, the antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof of the invention comprises

the HCDR1 of SEQ ID NOs: 10, 11 or 12;

the HCDR2 of SEQ ID NOs: 13, 14 or 15;

the HCDR3 of SEQ ID NOs: 16, 17, 18 or 19;

the LCDR1 of SEQ ID NOs: 20, 21, 22, 23, 24 or 25;

the LCDR2 of SEQ ID NOs: 26, 27, 28, 29 or 30; and

the LCDR3 of SEQ ID NOs: 31, 32, 33, 34, 35, 36, 37, 38, 39 or 40.

In some embodiments, the antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof of the invention comprises the HCDR1, the HCDR2 and the HCDR3 of

SEQ ID NOs: 10, 13 and 16, respectively;

SEQ ID NOs: 10, 14 and 16, respectively;

SEQ ID NOs: 10, 13 and 17, respectively;

SEQ ID NOs: 10, 13 and 18, respectively;

SEQ ID NOs: 11, 15 and 18, respectively;

SEQ ID NOs: 10, 13 and 19, respectively;

SEQ ID NOs: 10, 14 and 17, respectively; or

SEQ ID NOs: 12, 13 and 19, respectively.

In some embodiments, the antagonistic antibody specifically binding PD-1 or the antigen-binding portion thereof of the invention comprises the LCDR1, the LCDR2 and the LCDR3 of

SEQ ID NOs: 20, 26 and 31, respectively;

SEQ ID NOs: 21, 26 and 32, respectively;

SEQ ID NOs: 22, 27 and 33, respectively;

SEQ ID NOs: 22, 26 and 34, respectively;

SEQ ID NOs: 23, 28 and 35, respectively;

SEQ ID NOs: 20, 26 and 36, respectively;

SEQ ID NOs: 21, 27 and 37, respectively;

SEQ ID NOs: 23, 26 and 32, respectively;

SEQ ID NOs: 22, 26 and 32, respectively;

SEQ ID NOs: 24, 26 and 38, respectively;

SEQ ID NOs: 20, 29 and 39, respectively;

SEQ ID NOs: 20, 30 and 32, respectively;

SEQ ID NOs: 25, 26 and 40, respectively; or

SEQ ID NOs: 24, 26 and 32, respectively.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, optionally less than about 10 nM, for example less than about 1 nM such as less than about 500 pM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antibody or the antigen-binding portion thereof binds cynomolgous PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, optionally less than about 10 nM, for example less than about 1 nM such as less than about 500 pM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

In some embodiments, the VH and the VL are encoded by polynucleotide sequences of SEQ ID NOs: 196 and 197, respectively.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4/κ isotype, optionally comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 and is an IgG4 isotype, optionally comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 and is an IgG4/κ isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises a heavy chain (HC) of SEQ ID NO: 72 and a light chain (LC) of SEQ ID NO: 73.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG2/κ isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 and is an IgG2/κ isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 and is an IgG2/κ isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The antibody is suitable for use in therapy, for example in treating a melanoma.

The antibody is suitable for use in therapy, for example in treating a lung cancer.

The antibody is suitable for use in therapy, for example in treating non-small cell lung cancer (NSCLC).

The antibody is suitable for use in therapy, for example in treating a squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a non-squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a lung adenocarcinoma.

The antibody is suitable for use in therapy, for example in treating a renal cell carcinoma (RCC).

The antibody is suitable for use in therapy, for example in treating a mesothelioma.

The antibody is suitable for use in therapy, for example in treating a nasopharyngeal carcinoma (NPC).

The antibody is suitable for use in therapy, for example in treating a colorectal cancer.

The antibody is suitable for use in therapy, for example in treating a prostate cancer.

The antibody is suitable for use in therapy, for example in treating a castration-resistant prostate cancer.

The antibody is suitable for use in therapy, for example in treating a stomach cancer.

The antibody is suitable for use in therapy, for example in treating an ovarian cancer.

The antibody is suitable for use in therapy, for example in treating a gastric cancer.

The antibody is suitable for use in therapy, for example in treating a liver cancer.

The antibody is suitable for use in therapy, for example in treating a pancreatic cancer.

The antibody is suitable for use in therapy, for example in treating a thyroid cancer.

The antibody is suitable for use in therapy, for example in treating a squamous cell carcinoma of the head and neck.

The antibody is suitable for use in therapy, for example in treating a carcinomas of the esophagus or gastrointestinal tract.

The antibody is suitable for use in therapy, for example in treating a breast cancer.

The antibody is suitable for use in therapy, for example in treating a fallopian tube cancer.

The antibody is suitable for use in therapy, for example in treating a brain cancer.

The antibody is suitable for use in therapy, for example in treating an urethral cancer.

The antibody is suitable for use in therapy, for example in treating an endometriosis.

The antibody is suitable for use in therapy, for example in treating a cervical cancer.

The antibody is suitable for use in therapy, for example in treating a metastatic lesion of the cancer.

The antibody is suitable for use in therapy, for example in treating a hematological malignancy.

The antibody is suitable for use in therapy, for example in treating a non-Hodgkin's lymphoma.

The antibody is suitable for use in therapy, for example in treating a chronic lymphocytic leukemia.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody specifically binding TIM-3.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with a FGFR inhibitor.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with a vaccine.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding GITR (SEQ ID NO: 271).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding CD137 (SEQ ID NO: 281).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding OX-40 (SEQ ID NO: 279).

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 66, 67, 68, 69, 70 and 71, respectively.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

In some embodiments, the VH and the VL are encoded by polynucleotide sequences of SEQ ID NOs: 198 and 199, respectively.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 63 and the VL of SEQ ID NO: 65.

In some embodiments, the antibody or the antigen-binding portion thereof binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, optionally less than about 10 nM, for example less than about 1 nM such as less than about 100 pM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4/κ isotype, optionally comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 and is an IgG4 isotype, optionally comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 and is an IgG4κ isotype, comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 74 and the LC of SEQ ID NO: 75.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG2/κ isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitution when compared to the wild type IgG2.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 and is an IgG2/κ isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitution when compared to the wild type IgG2.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 and is an IgG2/κ isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitution when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The antibody is suitable for use in therapy, for example in treating a melanoma.

The antibody is suitable for use in therapy, for example in treating a lung cancer.

The antibody is suitable for use in therapy, for example in treating non-small cell lung cancer (NSCLC).

The antibody is suitable for use in therapy, for example in treating a squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a non-squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a lung adenocarcinoma.

The antibody is suitable for use in therapy, for example in treating a renal cell carcinoma (RCC).

The antibody is suitable for use in therapy, for example in treating a mesothelioma.

The antibody is suitable for use in therapy, for example in treating a nasopharyngeal carcinoma (NPC).

The antibody is suitable for use in therapy, for example in treating a colorectal cancer.

The antibody is suitable for use in therapy, for example in treating a prostate cancer.

The antibody is suitable for use in therapy, for example in treating a castration-resistant prostate cancer.

The antibody is suitable for use in therapy, for example in treating a stomach cancer.

The antibody is suitable for use in therapy, for example in treating an ovarian cancer.

The antibody is suitable for use in therapy, for example in treating a gastric cancer.

The antibody is suitable for use in therapy, for example in treating a liver cancer.

The antibody is suitable for use in therapy, for example in treating a pancreatic cancer.

The antibody is suitable for use in therapy, for example in treating a thyroid cancer.

The antibody is suitable for use in therapy, for example in treating a squamous cell carcinoma of the head and neck.

The antibody is suitable for use in therapy, for example in treating a carcinomas of the esophagus or gastrointestinal tract.

The antibody is suitable for use in therapy, for example in treating a breast cancer.

The antibody is suitable for use in therapy, for example in treating a fallopian tube cancer.

The antibody is suitable for use in therapy, for example in treating a brain cancer.

The antibody is suitable for use in therapy, for example in treating an urethral cancer.

The antibody is suitable for use in therapy, for example in treating an endometriosis.

The antibody is suitable for use in therapy, for example in treating a cervical cancer.

The antibody is suitable for use in therapy, for example in treating a metastatic lesion of the cancer.

The antibody is suitable for use in therapy, for example in treating a hematological malignancy.

The antibody is suitable for use in therapy, for example in treating a non-Hodgkin's lymphoma.

The antibody is suitable for use in therapy, for example in treating a chronic lymphocytic leukemia.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody specifically binding TIM-3.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with a FGFR inhibitor.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with a vaccine.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding GITR (SEQ ID NO: 271).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding CD137 (SEQ ID NO: 281).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding OX-40 (SEQ ID NO: 279).

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-biding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the

LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 12, 13, 19, 24, 26 and 38, respectively.

In some embodiments, the antibody or the antigen-biding portion thereof comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-biding portion thereof comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 11, 15, 18, 20, 30 and 32, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60.

In some embodiments, the VH and the VL are encoded by polynucleotide sequences of SEQ ID NOs: 202 and 203, respectively.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 76 and the LC of SEQ ID NO: 77.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-biding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 16, 20, 26 and 31, respectively.

In some embodiments, the antibody or the antigen-biding portion thereof comprises the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 212 and the LC of SEQ ID NO: 213.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-biding portion thereof, comprisingthe HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 16, 21, 26 and 32, respectively.

In some embodiments, the antibody or the antigen-biding portion thereof comprises the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 50.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 214 and the LC of SEQ ID NO: 215.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-biding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 16, 22, 27 and 33, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 51.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 216 and the LC of SEQ ID NO: 217.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-biding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 16, 22, 26 and 34, respectively.

In some embodiments, the antibody or the antigen-biding portion thereof comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 52.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 218 and the LC of SEQ ID NO: 219.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 16, 23, 28 and 35, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 53.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 220 and the LC of SEQ ID NO: 221.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 20, 26 and 31, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 49.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 20, 26 and 36, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 54.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 222 and the LC of SEQ ID NO: 223.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 21, 26 and 32, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 50.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 224 and the LC of SEQ ID NO: 225.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 21, 27 and 37, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 55.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 226 and the LC of SEQ ID NO: 227.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 23, 26 and 32, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 56.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 22, 26 and 32, respectively.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 57.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 228 and the LC of SEQ ID NO: 229.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 18, 20, 26 and 31, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 44 and the VL of SEQ ID NO: 49.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 11, 15, 18, 20, 26 and 31, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 49.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 19, 20, 26 and 31, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 46 and the VL of SEQ ID NO: 49.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 12, 13, 19, 20, 26 and 31, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 49.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 28 and 35, respectively.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 53.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 22, 26 and 34, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 52.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 12, 13, 19, 20, 29 and 39, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 59.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 11, 15, 18, 25, 26 and 40, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 61.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 11, 15, 18, 24, 26 and 32, respectively.

In some embodiments, the antibody of the antigen-binding portion thereof comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 62.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype.

In some embodiments, the antibody is an IgG4 isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47, 48, 63 or 64.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62 or 65.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH of SEQ ID NOs: 41, 42, 43, 44, 45,

46, 47, 48, 63 or 64 and the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62 or 65.

The VH, the VL, the HCDR and the LCDR sequences of exemplary antagonistic antibodies specifically binding PD-1 of the invention are shown in **Table 2.**

Although the embodiments illustrated in the Examples comprise pairs of variable regions, one from a heavy chain and one from a light chain, a skilled artisan will recognize that alternative embodiments may comprise single heavy or light chain variable regions. The single variable region may be used to screen for variable domains capable of forming a two-domain specific antigen-binding fragment capable of, for example, binding to human PD-1. The screening may be accomplished by phage display screening methods using for example hierarchical dual combinatorial approach disclosed in Int. Patent Publ. No. WO1992/01047. In this approach, an individual colony containing either a VH or a VL chain clone is used to infect a complete library of clones encoding the other chain (VL or VH), and the resulting two-chain specific antigen-binding domain is selected in accordance with phage display techniques using known methods and those described herein. Therefore, the individual VH and VL polypeptide chains are useful in identifying additional antibodies specifically binding to human PD-1 using the methods disclosed in Int. Patent Publ. No. WO1992/01047.

In some embodiments, the antagonistic antibody specifically binding PD-1 is a multispecific antibody.

In some embodiments, the antagonistic antibody specifically binding PD-1 is a bispecific antibody.

In some embodiments, antagonistic bispecific antibody specifically binding PD-1 binds PD-L1 (SEQ ID NO: 5), PD-L2 (SEQ ID NO: 8), LAG-3 (SEQ ID NO: 293), TIM-3 (SEQ ID NO: 138), CEACAM-1 (SEQ ID NO: 296), CEACAM-5 (SEQ ID NO: 307), OX-40 (SEQ ID NO: 279), GITR (SEQ ID NO: 271), CD27 (SEQ ID NO: 280), VISTA (SEQ ID NO: 286), CD137 (SEQ ID NO: 281), TIGIT (SEQ ID NO: 301) or CTLA-4 (SEQ ID NO: 292). Bispecific and multispecific antibodies may be generated using methods described herein.

Table 2.

Antibody	SEQ ID NO:							
	HCDR1	HCDR2	HCDR3	LCDR1	LCDR2	LCDR3	VH	VL
PD1B114	10	13	16	20	26	31	41	49
PD1B149	10	13	16	21	26	32	41	50
PD1B160	10	14	16	22	27	33	42	51
PD1B162	10	14	16	22	26	34	42	52
PD1B164	10	14	16	23	28	35	42	53
PD1B11	10	13	17	20	26	31	43	49
PD1B183	10	13	17	20	26	36	43	54
PD1B184	10	13	17	21	26	32	43	50
PD1B185	10	13	17	21	27	37	43	55
PD1B187	10	13	17	23	26	32	43	56
PD1B192	10	13	17	22	26	32	43	57
PD1B71	10	13	18	20	26	31	44	49
PD1B177	11	15	18	20	26	31	45	49
PD1B70	10	13	19	20	26	31	46	49
PD1B175	12	13	19	20	26	31	47	49
PD1B194	10	14	17	23	28	35	48	53
PD1B195	10	14	17	22	26	34	48	52
PD1B196	10	14	17	23	26	32	48	56
PD1B197	12	13	19	24	26	38	47	58
PD1B198	12	13	19	20	29	39	47	59
PD1B199	11	15	18	20	30	32	45	60
PD1B200	11	15	18	25	26	40	45	61
PD1B201	11	15	18	24	26	32	45	62
PD1B131	66	67	68	69	70	71	63	65
PD1B132	66	67	68	69	70	71	64	65

Homologous antibodies

Variants of the antagonistic antibodies specifically binding PD-1 or the antigen-binding portion thereof of the invention comprising the VH, the VL or the VH and the VL amino acid sequences shown in **Table 2, Table 21 and Table 22** are within the scope of

the invention. For example, variants may comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions in the VH and/or the VL as long as the homologous antibodies retain or have improved functional properties when compared to the parental antibodies. In some embodiments, the sequence identity may be about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% to a VH or the VL amino acid sequence of the invention. Optionally, any variation of the variant compared to the parental antibody is not within the CDRs of the variant.

The invention also provides an antagonistic antibody specifically binding PD-1or an antigen-binding portion thereof, comprising the VH of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47, 48, 63 or 64, the VH optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62 or 65, the VL optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising

the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49;

the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 50;

the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 51;

the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 52;

the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 53;

the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 49;

the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 54;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 50;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 55;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 56;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 57;
the VH of SEQ ID NO: 44 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 46 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 53;
the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 52;
the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58;
the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 59;
the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60;
the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 61;
the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 62; or
the VH of SEQ ID NO: 63 and the VL of SEQ ID NO: 65, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH having the amino acid sequence at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47, 48, 64 or 65. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VL having the amino acid sequence at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62 or 65. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH having the amino acid sequence at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47, 48, 63 or 64 and the VL having the amino acid sequence at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to

the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62 or 65. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH and the VL of SEQ ID NOs:

41 and 49, respectively;

41 and 50, respectively;

42 and 51, respectively;

42 and 52, respectively;

42 and 53, respectively;

43 and 49, respectively;

43 and 54, respectively;

43 and 50, respectively;

43 and 55, respectively;

43 and 56, respectively;

43 and 57, respectively;

44 and 49, respectively;

45 and 49, respectively;

46 and 49, respectively;

47 and 49, respectively;

48 and 53, respectively;

48 and 52, respectively;

47 and 58, respectively;

47 and 59, respectively;

45 and 60, respectively;

45 and 61, respectively;

45 and 62, respectively; or

63 and 65, respectively. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The homologous antagonistic antibodies specifically binding PD-1 or the antigen-binding portions thereof of the invention have one, two, three, four or five of the following properties:

a) enhance an activation of antigen specific $CD4^+$ or $CD8^+$ T cells in a dose dependent manner, wherein the activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1;

b) bind human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;

c) bind human PD-1 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;

d) bind cynomolgus PD-1 of SEQ ID NO: 3 with the K_D of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C, or

e) bind cynomolgus PD-1 of SEQ ID NO: 3 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antibody enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells in a dose dependent manner, wherein activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1, and binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antibody enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells in dose dependent manner, wherein activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1, and binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 10 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

The percent identity between the two sequences is a function of the number of identical positions shared by the sequences (i.e., % identity = number of identical positions/total number of positions ×100), taking into account the number of gaps, and the length of each gap, which need to be introduced for optimal alignment of the two sequences.

The percent identity between two amino acid sequences may be determined using the algorithm of E. Meyers and W. Miller (*Comput Appl Biosci* 4:11-17 (1988)) which has been incorporated into the ALIGN program (version 2.0), using a PAM120 weight residue table, a gap length penalty of 12 and a gap penalty of 4. In addition, the percent identity between two amino acid sequences may be determined using the Needleman and Wunsch (*J Mol Biol* 48:444-453 (1970)) algorithm which has been incorporated into the GAP program in the GCG software package (available at http://_www_gcg_com), using either a Blossum 62 matrix or a PAM250 matrix, and a gap weight of 16, 14, 12, 10, 8, 6, or 4 and a length weight of 1, 2, 3, 4, 5, or 6.

Antibodies with conservative modifications

The invention also provides antagonistic antibodies specifically binding PD-1 or antigen-binding portions thereof comprising the VH comprising the HCDR1, the HCDR2 and the HCDR3 sequences and the VL comprising the LCDR1, the LCDR2 and the LCDR3 sequences, wherein one or more of the CDR sequences comprise specified amino acid sequences based on the antibodies described herein (e.g., antibodies shown in **Table 2, Table 21 and Table 22,** or conservative modifications thereof, and wherein the antibodies retain the desired functional properties of the parental antagonistic antibodies specifically binding PD-1 of the invention.

The antibodies with conservative modifications have one, two, three, four or five of the following properties:

a) enhance an activation of antigen specific $CD4^+$ or $CD8^+$ T cells in dose dependent manner, wherein the activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1;

b) bind human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;

c) bind human PD-1 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;

d) bind cynomolgus PD-1of SEQ ID NO: 3 with the K_D of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C, or

e) bind cynomolgus PD-1of SEQ ID NO: 3 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antibody enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells in dose dependent manner, wherein activation is measured using a

cytomegalovirus antigen recall assay (CMV assay) as described in Example 1, and binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25ºC.

In some embodiments, the antibody enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells in dose dependent manner, wherein activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1, and binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 10 nM, wherein the K_D is measured using ProteOn XPR36 system at +25ºC.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 66, 67, 68, 69, 70 and 71, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of

SEQ ID NOs: 10, 13, 16, 20, 26 and 31, respectively;
SEQ ID NOs: 10, 13, 16, 21, 26 and 32, respectively;
SEQ ID NOs: 10, 14, 16, 22, 27 and 33, respectively;
SEQ ID NOs: 10, 14, 16, 22, 26 and 34, respectively;
SEQ ID NOs: 10, 14, 16, 23, 28 and 35, respectively;
SEQ ID NOs: 10, 13, 17, 20, 26 and 31, respectively;
SEQ ID NOs: 10, 13, 17, 20, 26 and 36, respectively;
SEQ ID NOs: 10, 13, 17, 21, 26 and 32, respectively;
SEQ ID NOs: 10, 13, 17, 21, 27 and 37, respectively;
SEQ ID NOs: 10, 13, 17, 23, 26 and 32, respectively;
SEQ ID NOs: 10, 13, 17, 22, 26 and 32, respectively;
SEQ ID NOs: 10, 13, 18, 20, 26 and 31, respectively;
SEQ ID NOs: 11, 15, 18, 20, 26 and 31, respectively;
SEQ ID NOs: 10, 13, 19, 20, 26 and 31, respectively;
SEQ ID NOs: 12, 13, 19, 20, 26 and 31, respectively;
SEQ ID NOs: 10, 14, 17, 23, 28 and 35, respectively;

SEQ ID NOs: 10, 14, 17, 22, 26 and 34, respectively;

SEQ ID NOs: 12, 13, 19, 24, 26 and 38, respectively;

SEQ ID NOs: 12, 13, 19, 20, 29 and 39, respectively;

SEQ ID NOs: 11, 15, 18, 20, 30 and 32, respectively;

SEQ ID NOs: 11, 15, 18, 25, 26 and 40, respectively;

SEQ ID NOs: 11, 15, 18, 24, 26 and 32, respectively, and conservative modifications thereof.

"Conservative modification" refers to amino acid modifications that do not significantly affect or alter the binding characteristics of the antibody containing the amino acid sequences. Conservative modifications include amino acid substitutions, additions and deletions. Conservative substitutions are those in which the amino acid is replaced with an amino acid residue having a similar side chain. The families of amino acid residues having similar side chains are well defined and include amino acids with acidic side chains (for example, aspartic acid, glutamic acid), basic side chains (for example, lysine, arginine, histidine), nonpolar side chains (for example, alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine), uncharged polar side chains (for example, glycine, asparagine, glutamine, cysteine, serine, threonine, tyrosine, tryptophan), aromatic side chains (for example, phenylalanine, tryptophan, histidine, tyrosine), aliphatic side chains (for example, glycine, alanine, valine, leucine, isoleucine, serine, threonine), amide (for example, asparagine, glutamine), beta-branched side chains (for example, threonine, valine, isoleucine) and sulfur-containing side chains (cysteine, methionine). Furthermore, any native residue in the polypeptide may also be substituted with alanine, as has been previously described for alanine scanning mutagenesis (MacLennan *et al.*, Acta Physiol. Scand. Suppl. 643:55-67, 1998; Sasaki *et al.*, Adv. Biophys. 35:1-24, 1998). Amino acid substitutions to the antibodies of the invention may be made by well-known methods for example by PCR mutagenesis (US Pat. No. 4,683,195). Alternatively, libraries of variants may be generated using known methods, for example using random (NNK) or non-random codons, for example DVK codons, which encode 11 amino acids (Ala, Cys, Asp, Glu, Gly, Lys, Asn, Arg, Ser, Tyr, Trp). The resulting antibody variants may be tested for their characteristics using assays described herein.

Antagonistic antibodies specifically binding TIM-3

T-cell immunoglobulin domain and mucin domain 3 (TIM-3, also known as Hepatitis A virus cellular receptor 2 (HAVCR2)) is a co-inhibitory immune checkpoint receptor that has been proposed to negatively regulate both adaptive and innate immune

responses. TIM-3 is expressed on specific subsets of $CD4^+$ and $CD8^+$ T cells and functions to limit the duration and magnitude of T cell responses.

Multiple lines of evidence support the inhibitory role of TIM-3 in regulating T cell responses. Tim-3-deficient mice exhibit defects in the induction of both antigen-specific and transplantation tolerance, consistent with TIM-3 inhibiting effector T cells during normal immune responses (Sabatos *et al.,* (2003) *Nat Immunol* 4(11):1102-1110, Sanchez-Fueyo *et al.,* (2003) *Nat Immunol* 4(11):1093-1101). Anti-TIM-3 antibodies exacerbate experimental autoimmune encephalomyelitis (EAE) in animal models (Monney *et al.,* (2002) *Nature* 415(6871):536-541). TIM-3 has been shown to be a critical driver of the dysfunctional or exhausted T cell state that occurs in chronic infection and cancer (Sakuishi, K. and A. C. Anderson (2014). Tim-3 Regulation of Cancer Immunity. Tumor-Induced Immune Suppression. D. I. Gabrilovich and A. A. Hurwitz, Springer New York: 239-261).

Blockade of TIM-3 has been shown to restore activity in effector cells, such as cytokine secretion and proliferation. In virally exhausted cell populations, e.g., cells infected with HCV, TIM-3-expressing cells ($TIM\text{-}3^+$ cells) express less TNF-α and IFN-γ cytokines than TIM-3 negative cells in both effector cell populations, $CD4^+$ and $CD8^+$ T cells (Golden-Mason *et al.,* (2009) *J Virol* 83:9122). Blockade of TIM-3 restored proliferation in $CD8^+$ T cells from an HIV patient, or in cells that recapitulated viral exhaustion (Jones *et al.,* (2008) *J Exp Med* 205:2763), or proliferation and IFN-γ and/or TNF-α secretion in NY-ESO-1 specific T cells from PBMCs from metastatic patients (Fourcade *et al., (*2010) *J Exp Med* 207:2175). $TIM\text{-}3^+$ T cells have been found to be concentrated in tumors, and contribute to the immunosuppressive tumor environment (Sakuishi *et al.,* (2013) *Oncoimmunology,* 2:e23849).

Blockade of TIM-3 (partially alone and additively or synergistically in combination with PD-1 pathway blockade) has shown anti-tumor efficacy in several preclinical cancer models, including CT26 colon carcinoma (Sakuishi *et al.,* (2010) *J Exp Med* 207(10):2187-94), WT3 sarcoma and TRAMP-C1 prostate carcinoma (Ngiow *et al.,* (2011) *Cancer Res* 71(10):3540-3551).

The mechanisms through which TIM-3 inhibits T cell responses are not fully understood. The cytoplasmic tail of TIM-3 contains multiple tyrosine residues (Ferris *et al.,* (2014) *J Immunol* 193(4): 1525-1530) but lacks inhibitory signaling motifs such as ITIMs or ITSMs that are found in the PD-1 intracellular tail. The Src family tyrosine kinases Fyn and Lck have been shown to bind to TIM-3, although the exact consequences of these interactions remain to be confirmed *in vivo*. Two opposing models have been

proposed for how TIM-3 regulates T cell signaling. On one hand, TIM-3 has been postulated to negatively regulate TCR signaling by recruiting a phosphatase to the immunological synapse, and de-phosphorylating Lck (Clayton, *et al.,* (2014) *J Immunol* 192(2):782-791). In contrast, TIM-3 has also been proposed to enhance TCR signaling and paradoxically drive T cells towards a more exhausted state, through increased activation of NFAT activity and NFκB signaling.

In addition to expression on effector T cells, TIM-3 is also expressed on regulatory T cells (T-regs) and has been shown to mark a suppressive T-reg subset in tumors. Analyses using both primary human cells and mouse preclinical models have shown that TIM-3$^+$ T-regs are more effective at inhibiting T helper1 (Th1) and T helper 17 (Th17) T cell responses than TIM-3$^-$ T-regs (Gautron *et al.,* (2014) *Eur J Immunol* 44(9): 2703-2711; Sakuishi *et al.,* (2013) *Oncoimmunology,* 2:e23849). Since TIM-3 is expressed on highly suppressive Tregs, it can directly inhibit CD4$^+$ and CD8$^+$ T cell responses. In addition, TIM-3$^+$ Tregs express high levels of IL-10, which has been proposed to drive exhaustion of effector T cells in the TME as an additional indirect mechanism of suppressing anti-tumor immune responses (Sakuishi *et al.,* (2013) *Oncoimmunology,* 2:e23849).

TIM-3 is expressed on several innate immune cell types, including monocytes/macrophages, dendritic cells, and NK cells. Existing data are consistent with a suppressive role for TIM-3 in these different cell types.

TIM-3 is constitutively expressed by circulating CD14$^+$ monocytes in healthy donors, and its expression on peripheral monocytes is significantly increased in patients with chronic inflammation and cancer (Rong *et al.,* (2014) *Tissue Antigens* 83(2):76-81). TIM-3 levels are also upregulated on macrophages that infiltrate hepatocellular carcinoma (HCC) tumors, compared to macrophages from adjacent tissues, and is proposed to play a role in driving the polarization of macrophages to an M2 tumor-promoting phenotype.

Recently, TIM-3 was reported to be expressed on dendritic cells that infiltrate mouse tumors. In this setting, interaction of TIM-3 with HMBG1 was proposed to suppress innate immunity by interfering with the recognition of and response to immunostimulatory nucleic acid (Chiba *et al.,* (2012) *Immunol* 13(9): 832-842). TIM-3 is also constitutively expressed on NK cells in peripheral blood. A recent study showed that NK cells from advanced melanoma patients express high levels of TIM-3 on peripheral NK cells. Importantly, TIM-3$^+$ NK cells were functionally exhausted and anti-TIM-3 blockade was able to reverse the exhaustion and enhance NK cell functionality (da Silva *et al.,* (2014) *Cancer Immunol Res* 2(5): 410-422).

TIM-3 binds ligands galectin-9 (Gal-9), phosphatidylserine (PtdSer), HMGB1 and CEACAM-1. S-type lectin galectin-9 can inhibit TIM-3-associated Th1 effector function and induce apoptosis on TIM-3-expressing T cells in murine models. PtdSer usually resides on the intracellular side of the plasma membrane, but is flipped to the extracellular side during apoptosis. PtdSer binds a preserved cleft in all three human TIM family members (TIM-1, 3, 4). Inhibition of PtdSer binding to TIM-3 may activate T-cell response. Galectin-9 is secreted by tumor cells and can contribute to evasion from anti-tumor immunity. DNA alarmin HMGB1, for which TIM-3 may act as a "sink," can prevent the HMGB1/RAGE interactions that stimulate innate immunity. CEACAM-1 can interact with TIM-3 both in cis as a heterodimer on T cells and in trans as a ligand. Interaction between CEACAM-1 and TIM-3 may help mediate block immune response signaling. Co-blockade of TIM-3 and CEACAM-1 in CT26 colon carcinoma showed similar efficacy to that seen for co-blockade of PD-L1 and TIM-3.

Thus, blockade of TIM-3 using the antibodies of the invention described herein that inhibit TIM-3 function may improve the immune response against infection and anti-tumor immunity.

The invention also provides an isolated antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, wherein the antibody inhibits binding of TIM-3 to galectin-9.

Inhibition of binding of TIM-3 to galectin-9 by the antibodies of the invention may be assessed using competition ELISA. In an exemplary assay, 1 μg/ml recombinant human Fc-TIM-3 is bound on wells of microtiter plates, the wells are washed and blocked, and 10 μg/ml of the test antibody is added. Without washing, 7.5 μg/ml galectin-9 is added into the wells and incubated for 30 min, after which 0.5 μg/ml anti-galectin-9-biotin antibody is added and incubated for 30 min. The plates are washed and 0.5 μg/mL neutravidin-HRP conjugate polyclonal antibody is added and incubated for 30 minutes. The plates are washed and POD Chemiluminescence substrate added immediately prior to reading the luminescence signal. Antibodies of the invention inhibit binding of TIM-3 to galectin-9 when the binding of galectin-9 is reduced by at least about 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95% or 100% using an assay described herein and in Example 1. Exemplary antibodies that inhibit TIM-3 binding to galectin-9 are antibodies TM3B103, TM3B105, TM3B107, TM3B108, TM3B109, TM3B113, TM3B189, TM3B190 and TM3B196.

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells.

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof enhances an activation of antigen specific $CD4^+$ or $CD8^+$ T cells, wherein the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant enhancement of CD137 surface expression on antigen specific $CD4^+$ or $CD8^+$ T cells according to methods described in Example 14.

Use of CD137 as a marker of antigen specific $CD8^+$ and $CD4^+$ T cells that expand in response to CMV antigen stimulation allowed the detection of the functional effects of the antagonistic TIM-3 antibodies of the invention.

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof binds TIM-3 within TIM-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261).

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof binds TIM-3 within TIM-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261) and residues 50-56 (DERDVNY) (SEQ ID NO: 262).

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof binds TIM-3 within TIM-3 residues 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263).

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof binds TIM-3 within TIM-3 residues 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263) and residues 50-56 (DERDVNY) SEQ ID NO: 262.

"Within" means that 80% or more of the epitope residues the antibody binds to reside within the recited amino acid stretches, and that up to 20% of the epitope residues the antibody binds to reside outside of the recited amino acid stretches.

The Tim-3 epitope the antibody binds to may be resolved for example using hydrogen/deuterium exchange (H/D exchange) or by analyzing a crystal structure of the antibody in complex with TIM-3. The epitope residues are those which are protected by the antibody by at least 5% difference in deuteration levels through H/D exchange or those surface exposed amino acid residues determined to bind the antibody in a crystal structure of a complex of the antibody and TIM-3. In the crystal structure of a complex of the

antibody and TIM-3, the epitope residues are those TIM-3 residues that reside within 4 Å distance or less from any of the antibody CDR residues.

In an H/D exchange assay, TIM-3 protein is incubated in the presence or absence of the antibody in deuterated water for predetermined times resulting in deuterium incorporation at exchangeable hydrogen atoms which are unprotected by the antibody, followed by protease digestion of the protein and analyses of the peptide fragments using LC-MS. In an exemplary assay, 5 µL of the test antibody (10 µg) or 5 µL of the complex of TIM-3 and the test antibody (10 and 7.35 µg, respectively) is incubated with 120 µL deuterium oxide labeling buffer (50mM phosphate, 100mM sodium chloride at pH 7.4) for 0 sec, 60 sec, 300 sec, 1800 sec, 7200 sec, and 14400 sec. Deuterium exchange is quenched by adding 63 µL of 5 M guanidine hydrochloride and final pH is 2.5. The quenched sample is subjected to on-column pepsin/protease type XIII digestion and LC-MS analysis. For pepsin/protease type XIII digestion, 5 µg of the samples in 125 µL control buffer (50mM phosphate, 100mM sodium chloride at pH 7.4) are denatured by adding 63 µL of 5 M guanidine hydrochloride (final pH is 2.5) and incubating the mixture for 3 min. Then, the mixture is subjected to on-column pepsin/protease type XIII digestion and the resultant peptides analyzed using an UPLC-MS system comprised of a Waters Acquity UPLC coupled to a Q Exactive™ Hybrid Quadrupole-Orbitrap Mass Spectrometer (Thermo). Raw MS data is processed using HDX WorkBench, software for the analysis of H/D exchange MS data. The deuterium levels are calculated using the average mass difference between the deuteriated peptide and its native form (t_0). Peptide identification is done through searching MS/MS data against the TIM-3 sequence with Mascot. The mass tolerance for the precursor and product ions is 20 ppm and 0.05 Da, respectively.

For X-ray crystallography, TIM-3 and the test antibody are expressed and purified using standard protocols. The TIM-3/test antibody complex is incubated overnight at 4°C, concentrated, and separated from the uncomplexed species using size-exclusion chromatography. The complex is crystallized by the vapor-diffusion method from various known test solutions for example solutions containing PEG3350, ammonium citrate and 2-(N-Morpholino)ethanesulfonic acid (MES).

Antibodies binding within Tim-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261), 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263) and/or 50-56 (DERDVNY) (SEQ ID NO: 262) may be generated by isolating antibodies binding TIM-3 using phage display libraries, selecting those antibodies that compete with the reference antibody TM3B105 (VH of SEQ ID NO: 146 and VL of SEQ ID NO: 156) or TM3B291

(VH of SEQ ID NO: 172 and VL of SEQ ID NO: 173) for binding to TIM-3 by 100%, and confirming the epitope of the generated antibodies by solving the crystal structure of the antibody/TIM-3 complex. Alternatively, mice, rats or rabbits may be immunized using peptides encompassing residues 32-47, 90-102 and/or 50-56 of TIM-3 and the generated antibodies may be evaluated for their binding within the recited region.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof comprising a heavy chain complementarity determining region 1 (HCDR1), a HCDR2 and a HCDR3 of SEQ ID NOs: 164, 165 and 166, respectively.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof comprising a light chain complementarity determining region 1 (LCDR1), LCDR2 and LCDR3 of SEQ ID NOs: 167, 168 and 169 respectively.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof comprising the HCDR1, the HCDR2 and the HCDR3 of SEQ ID NOs: 164, 165 and 166, respectively, and the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 167, 168 and 169 respectively.

SEQ ID NOs: 164, 165, 166, 167, 168 and 169 represent the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 genus sequences of TIM-3 antagonists derived from phage display libraries. The genus sequences were generated based on structural models that resulted in the sequence alignments given in **Figure 13, Figure 14, Figure 15, Figure 16, Figure 17** and **Figure 18** and summarized herein.

SEQ ID NO: 164

$X_{17}YX_{18}MX_{19}$,

wherein

X_{17} is N, S, G or D;

X_{18} is W or A; and

X_{19} is S or H.

SEQ ID NO: 165

$X_{20}IX_{21}X_{22}SGGSX_{23}YYADSVKG$,

wherein

X_{20} is A or V;

X_{21} is S or K;

X_{22} is G or Y; and

X_{23} is T or K.

SEQ ID NO: 166

$X_{24}X_{25}X_{26}X_{27}X_{28}X_{29}X_{30}X_{31}DY$,

wherein

X_{24} is D, S, N, G or E;

X_{25} is H, P, E, T or L;

X_{26} is W, E, N or deleted;

X_{27} is D, P or deleted;

X_{28} is P, Y, D or deleted;

X_{29} is N, A, D, G or deleted;

X_{30} is F, P, R, W or V; and

X_{31} is L or F.

SEQ ID NO: 167

$X_{32}X_{33}SQSVX_{34}X_{35}X_{36}X_{37}X_{38}X_{39}X_{40}X_{41}X_{42}LA$,

wherein

X_{32} is R or K;

X_{33} is A or S;

X_{34} is S, N or L;

X_{35} is S, A, N or deleted;

X_{36} is S or deleted;

X_{37} is S or deleted;

X_{38} is N or deleted;

X_{39} is N or deleted;

X_{40} is K or deleted;

X_{41} is S, D or N; and

X_{42} is Y or T.

SEQ ID NO: 168

$X_{43}ASX_{44}RX_{45}X_{46}$,

wherein

X_{43} is G, D, W or T;

X_{44} is S, N or T;

X_{45} is A or E; and

X_{46} is T or S.

SEQ ID NO: 169

$QQX_{47}X_{48}X_{49}X_{50}PX_{51}T$ (SEQ ID NO: 169),

wherein

X_{47} is Y, G or S;

X_{48} is G or Y;

X_{49} is S, H or T;

X_{50} is S, A or T; and

X_{51} is L, I or W.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof comprising the HCDR1, the HCDR2 and the HCDR3 contained within a heavy chain variable region (VH) of SEQ ID NOs: 145, 146, 147, 148 or 149, wherein the HCDR1, the HCDR2 and the HCDR3 are defined by Chothia, Kabat, or IMGT.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof comprising the LCDR1, the LCDR2 and the LCDR3 contained within a light chain variable region (VL) of SEQ ID NOs: 155, 156, 157 or 158, wherein the LCDR1, the LCDR2 and the LCDR3 are defined by Chothia, Kabat, or IMGT.

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof comprises

the HCDR1 of SEQ ID NOs: 90, 91, 92 or 93;

the HCDR2 of SEQ ID NOs: 99, 100 or 101; and

the HCDR3 of SEQ ID NOs: 107, 108, 109, 110 or 111.

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof of the invention comprises

the LCDR1 of SEQ ID NOs: 117, 118, 119 or 120;

the LCDR2 of SEQ ID NOs: 126, 127, 128 or 129; and

the LCDR3 of SEQ ID NOs: 135, 136, 137 or 139.

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof comprises

the HCDR1 of SEQ ID NOs: 90, 91, 92 or 93;

the HCDR2 of SEQ ID NOs: 99, 100 or 101;

the HCDR3 of SEQ ID NOs: 107, 108, 109, 110 or 111;

the LCDR1 of SEQ ID NOs: 117, 118, 119 or 120;

the LCDR2 of SEQ ID NOs: 126, 127, 128 or 129; or

the LCDR3 of SEQ ID NOs: 135, 136, 137 or 139.

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof comprises the HCDR1, the HCDR2 and the HCDR3 of

SEQ ID NOs: 90, 99 and 107, respectively;

SEQ ID NOs: 91, 99 and 108, respectively;

SEQ ID NOs: 91, 99 and 109, respectively;

SEQ ID NOs: 92, 100 and 110, respectively; or

SEQ ID NOs: 93, 101 and 111, respectively;

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof comprises the LCDR1, the LCDR2 and the LCDR3 of

SEQ ID NOs: 117, 126 and 135, respectively;

SEQ ID NOs: 118, 127 and 136, respectively;

SEQ ID NOs: 119, 128 and 137, respectively; or

SEQ ID NOs: 120, 129 and 139, respectively.

In some embodiments, the antagonistic antibody specifically binding TIM-3 or the antigen-binding portion thereof comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of

SEQ ID NOs: 90, 99, 107, 117, 126 and 135, respectively;

SEQ ID NOs: 91, 99, 108, 118, 127 and 136, respectively;

SEQ ID NOs: 91, 99, 109, 119, 128 and 137, respectively;

SEQ ID NOs: 92, 100, 110, 117, 126 and 135, respectively; or

SEQ ID NOs: 93, 101, 111, 120, 129 and 139, respectively.

The invention also provides an isolated antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2 and, the HCDR3 of SEQ ID NOs: 164, 165 and 108, respectively, and the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 118, 168 and 169 respectively.

The invent ion also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 91, 99, 108, 118, 127 and 136, respectively.

In some embodiments, the antibody specifically binding TIM-3 or the antigen-binding portion thereof binds TIM-3 within TIM-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261).

In some embodiments, the antibody specifically binding TIM-3 or the antigen-binding portion thereof binds TIM-3 within TIM-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261) and residues 50-56 (DERDVNY) SEQ ID NO: 262.

In some embodiments, the antibody specifically binding TIM-3 or the antigen-binding portion thereof inhibits binding of TIM-3 to galectin-9.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV3-23 (SEQ ID NO: 174) and a light chain framework derived from IGKV3-11 (SEQ ID NO: 171).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

In some embodiments, the VH and the VL are encoded by polynucleotide sequences of SEQ ID NOs: 204 and 205, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells, wherein activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant enhancement of CD137 surface expression on antigen specific $CD4^+$ or $CD8^+$ T cells according to methods described in Example 14.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4/κ isotype, optionally comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156 and is an IgG4 isotype, optionally comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156 and is an IgG4κ isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG2/κ isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156 and is an IgG2/κ isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156 and is an IgG2/κ isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 78 and the LC of SEQ ID NO: 79.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 240 and the LC of SEQ ID NO: 79.

SEQ ID NO: 78

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMSWVRQAPGKGLEWVSAISGSG
GSTYYADSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCAKSPYAPLDYWGQ
GTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTS
GVHTFPAVLQSSGLYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKTVERKCC
VECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQFNWYVD
GVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKT
ISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNY
KTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG
K

SEQ ID NO: 79

EIVLTQSPATLSLSPGERATLSCRASQSVNDYLAWYQQKPGQAPRLLIYDA
SNRATGIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQGGHAPITFGQGTKVEIKR
TVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESV
TEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 240

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMSWVRQAPGKGLEWVSAISGSG
GSTYYADSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCAKSPYAPLDYWGQ
GTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTS
GVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDKTVERKCC
VECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQFNWYVD
GVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKT
ISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNY
KTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG
K

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The antibody is suitable for use in therapy, for example in treating a melanoma.

The antibody is suitable for use in therapy, for example in treating a lung cancer.

The antibody is suitable for use in therapy, for example in treating non-small cell lung cancer (NSCLC).

The antibody is suitable for use in therapy, for example in treating a squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a non-squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a lung adenocarcinoma.

The antibody is suitable for use in therapy, for example in treating a renal cell carcinoma (RCC).

The antibody is suitable for use in therapy, for example in treating a mesothelioma.

The antibody is suitable for use in therapy, for example in treating a nasopharyngeal carcinoma (NPC).

The antibody is suitable for use in therapy, for example in treating a colorectal cancer.

The antibody is suitable for use in therapy, for example in treating a prostate cancer.

The antibody is suitable for use in therapy, for example in treating a castration-resistant prostate cancer.

The antibody is suitable for use in therapy, for example in treating a stomach cancer.

The antibody is suitable for use in therapy, for example in treating an ovarian cancer.

The antibody is suitable for use in therapy, for example in treating a gastric cancer.

The antibody is suitable for use in therapy, for example in treating a liver cancer.

The antibody is suitable for use in therapy, for example in treating a pancreatic cancer.

The antibody is suitable for use in therapy, for example in treating a thyroid cancer.

The antibody is suitable for use in therapy, for example in treating a squamous cell carcinoma of the head and neck.

The antibody is suitable for use in therapy, for example in treating a carcinomas of the esophagus or gastrointestinal tract.

The antibody is suitable for use in therapy, for example in treating a breast cancer.

The antibody is suitable for use in therapy, for example in treating a fallopian tube cancer.

The antibody is suitable for use in therapy, for example in treating a brain cancer.

The antibody is suitable for use in therapy, for example in treating an urethral cancer.

The antibody is suitable for use in therapy, for example in treating an endometriosis.

The antibody is suitable for use in therapy, for example in treating a cervical cancer.

The antibody is suitable for use in therapy, for example in treating a metastatic lesion of the cancer.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody that specifically binds PD-1.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with the antagonistic antibody that specifically binds PD-1 comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with the antagonistic antibody that specifically binds PD-1 comprising the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with the antagonistic antibody that specifically binds PD-1 comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with the antagonistic antibody that specifically binds PD-1 comprising the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an antagonistic antibody specifically binding TIGIT (SEQ ID NO: 301).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with a FGFR inhibitor.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with a vaccine.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding GITR (SEQ ID NO: 271).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding CD137 (SEQ ID NO: 281).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding OX-40 (SEQ ID NO: 279).

The antibody is suitable for use in therapy in a subject who is being treated or who has been treated with an antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab)).

The antibody is suitable for use in therapy in a subject who is being treated or who has been treated with an antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The antibody is suitable for use in therapy in a subject who is refractory to treatment with the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab)).

The antibody is suitable for use in therapy in a subject who is refractory to treatment with the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The antibody is suitable for use in therapy in a subject who has a relapsed tumor after treatment with the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab).

The antibody is suitable for use in therapy in a subject who has a relapsed tumor after treatment with the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 97, 105, 115, 124, 133 and 143, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV5-51 (SEQ ID NO: 179) and a light chain framework derived from IGKV1-39 (SEQ ID NO: 182).

In some embodiments, the antibody comprises the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

In some embodiments, the VH and the VL are encoded by polynucleotide sequences of SEQ ID NOs: 206 and 207, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells, wherein the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant enhancement of CD137 surface expression on antigen specific $CD4^+$ or $CD8^+$ T cells according to methods described in Example 14.

In some embodiments, the antibody specifically binding TIM-3 or the antigen-binding portion thereof binds TIM-3 within TIM-3 residues 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263).

In some embodiments, the antibody specifically binding TIM-3 or the antigen-binding portion thereof binds TIM-3 within TIM-3 residues 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263) and residues 50-56 (DERDVNY) SEQ ID NO: 262.

In some embodiments, the antibody specifically binding TIM-3 or the antigen-binding portion thereof inhibits binding of TIM-3 to galectin-9.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4/κ isotype, optionally comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173 and is an IgG4 isotype, optionally comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173 and is an IgG4κ isotype comprising the S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG2/κ isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173 and is an IgG2/κ isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody comprises the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173 and is an IgG2/κ isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody comprises the HC of SEQ ID NO: 80 and the LC of SEQ ID NO: 81.

SEQ ID NO: 80

EVQLVQSGAEVKKPGESLKISCKGSGYSFTSYWMQWVRQMPGKGLEWMGAIYP
GDGDIRYTQNFKGQVTISADKSISTAYLQWSSLKASDTAMYYCARWEKSTTVVQ
RNYFDYWGQGTTVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTV

SWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKV
DKTVERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDP
EVQFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVS
NKGLPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVE
WESNGQPENNYKTTPPMLDSDGSFFLYSRLTVDKSRWQQGNVFSCSVMHEALHN
HYTQKSLSLSPGK

SEQ IN NO: 81

DIQMTQSPSSLSASVGDRVTITCKASENVGTFVSWYQQKPGKAPKLLIYGASNRY
TGVPSRFSGSGSGTDFTLTISSLQPEDFATYYCGQSYSYPTFGQGTKLEIKRTVAAP
SVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDS
KDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

In some embodiments, the antibody is a bispecific antibody, such as a bispecific PD-1/TIM-3 antibody.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The antibody is suitable for use in therapy, for example in treating a melanoma.

The antibody is suitable for use in therapy, for example in treating a lung cancer.

The antibody is suitable for use in therapy, for example in treating non-small cell lung cancer (NSCLC).

The antibody is suitable for use in therapy, for example in treating a squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a non-squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a lung adenocarcinoma.

The antibody is suitable for use in therapy, for example in treating a renal cell carcinoma (RCC).

The antibody is suitable for use in therapy, for example in treating a mesothelioma.

The antibody is suitable for use in therapy, for example in treating a nasopharyngeal carcinoma (NPC).

The antibody is suitable for use in therapy, for example in treating a colorectal cancer.

The antibody is suitable for use in therapy, for example in treating a prostate cancer.

The antibody is suitable for use in therapy, for example in treating a castration-resistant prostate cancer.

The antibody is suitable for use in therapy, for example in treating a stomach cancer.

The antibody is suitable for use in therapy, for example in treating an ovarian cancer.

The antibody is suitable for use in therapy, for example in treating a gastric cancer.

The antibody is suitable for use in therapy, for example in treating a liver cancer.

The antibody is suitable for use in therapy, for example in treating a pancreatic cancer.

The antibody is suitable for use in therapy, for example in treating a thyroid cancer.

The antibody is suitable for use in therapy, for example in treating a squamous cell carcinoma of the head and neck.

The antibody is suitable for use in therapy, for example in treating a carcinomas of the esophagus or gastrointestinal tract.

The antibody is suitable for use in therapy, for example in treating a breast cancer.

The antibody is suitable for use in therapy, for example in treating a fallopian tube cancer.

The antibody is suitable for use in therapy, for example in treating a brain cancer.

The antibody is suitable for use in therapy, for example in treating an urethral cancer.

The antibody is suitable for use in therapy, for example in treating an endometriosis.

The antibody is suitable for use in therapy, for example in treating a cervical cancer.

The antibody is suitable for use in therapy, for example in treating a metastatic lesion of the cancer.

The antibody is suitable for use in therapy, for example in treating a hematological malignancy.

The antibody is suitable for use in therapy, for example in treating an acute lymphoblastic leukemia (ALL).

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with an antagonistic antibody that specifically binds PD-1.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with the antagonistic antibody that specifically binds PD-1 comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with the antagonistic antibody that specifically binds PD-1 comprising the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with the antagonistic antibody that specifically binds PD-1 comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60.

The antibody is suitable for use in therapy, for example in treating a cancer, in combination with the antagonistic antibody that specifically binds PD-1 comprising the VH of SEQ ID NO: 65 and the VL of SEQ ID NO: 65.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an antagonistic antibody specifically binding TIGIT (SEQ ID NO: 301).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with a FGFR inhibitor.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with a vaccine.

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding GITR (SEQ ID NO: 271).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding CD137 (SEQ ID NO: 281).

The antibody is suitable for use in therapy, for example in treating cancer, such as a solid tumor, in combination with an agonistic antibody specifically binding OX-40 (SEQ ID NO: 279).

The antibody is suitable for use in therapy in a subject who is being treated or who has been treated with an antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab)).

The antibody is suitable for use in therapy in a subject who is being treated or who has been treated with an antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The antibody is suitable for use in therapy in a subject who is refractory to treatment with the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab)).

The antibody is suitable for use in therapy in a subject who is refractory to treatment with the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The antibody is suitable for use in therapy in a subject who has a relapsed tumor after treatment with the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab).

The antibody is suitable for use in therapy in a subject who has a relapsed tumor after treatment with the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 90, 99, 107, 117, 126 and 135, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV3-23 (SEQ ID NO: 174) and a light chain framework derived from IGKV3-20 (SEQ ID NO: 180).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155.

In some embodiments, the VH and the VL are encoded by polynucleotide sequences of SEQ ID NOs: 208 and 209, respectively.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 91, 99, 109, 119, 128 and 137,

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV3-23 (SEQ ID NO: 174) and a light chain framework derived from IGKV4-1 (SEQ ID NO: 181).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 148 and the VL of SEQ ID NO: 157.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 92, 100, 110, 117, 126 and 135, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV3-23 (SEQ ID NO: 174) and a light chain framework derived from IGKV3-20 (SEQ ID NO: 180).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 147 and the VL of SEQ ID NO: 155.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 93, 101, 111, 120, 129 and 139, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV3-23 (SEQ ID NO: 174) and a light chain framework derived from IGKV3-20 (SEQ ID NO: 180).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 149 and the VL of SEQ ID NO: 158.

In some embodiments, the VH and the VL are encoded by polynucleotide sequences of SEQ ID NOs: 201 and 211, respectively.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 94, 102, 112, 121, 130 and 140, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV1-02 (SEQ ID NO: 175) and a light chain framework derived from IGKV4-1 (SEQ ID NO: 181).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 150 and the VL of SEQ ID NO: 159.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 95, 103, 113, 122, 131 and 141, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV4-30-4 (SEQ ID NO: 176) and a light chain framework derived from IGKV1-39 (SEQ ID NO: 182).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 151 and the VL of SEQ ID NO: 160.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 96, 104, 114, 123, 132 and 142, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV1-03 (SEQ ID NO: 177) and a light chain framework derived from IGKV1-33 (SEQ ID NO: 183).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 152 and the VL of SEQ ID NO: 161.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 97, 105, 115, 124, 133 and 143, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV1-03 (SEQ ID NO: 177) and a light chain framework derived from IGKV1-39 (SEQ ID NO: 182).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 98, 106, 116, 125, 134 and 144, respectively.

In some embodiments, the antibody or the antigen-binding portion thereof comprises a heavy chain framework derived from IGHV2-26 (SEQ ID NO: 178) and a light chain framework derived from IGKV4-1 (SEQ ID NO: 181).

In some embodiments, the antibody or the antigen-binding portion thereof comprises the VH of SEQ ID NO: 154 and the VL of SEQ ID NO: 163.

In some embodiments, the antibody or the antigen-binding portion thereof enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells, wherein activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant

enhancement of CD137 surface expression on antigen specific $CD4^+$ or $CD8^+$ T cells according to methods described in Example 14.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

The VH, the VL, the HCDR and the LCDR sequences of exemplary antagonistic antibodies specifically binding TIM-3 of the invention are shown in **Table 3.**

Although the embodiments illustrated in the Examples comprise pairs of variable regions, one from a heavy chain and one from a light chain, a skilled artisan will recognize that alternative embodiments may comprise single heavy or light chain variable regions. The single variable region may be used to screen for variable domains capable of forming a two-domain specific antigen-binding fragment capable of, for example, binding to human TIM-3. The screening may be accomplished by phage display screening methods similarly as described herein.

In some embodiments, the antagonistic antibody specifically binding TIM-3 is a multispecific antibody.

In some embodiments, the antagonistic antibody specifically binding TIM-3 is a bispecific antibody.

In some embodiments, the bispecific or the multispecific antibody binds PD-1 (SEQ ID NO: 1), PD-L1 (SEQ ID NO: 5), PD-L2 (SEQ ID NO: 8), LAG-3 (SEQ ID NO: 293), CEACAM-1 (SEQ ID NO: 296), CEACAM-5 (SEQ ID NO: 307), NKG2D (SEQ ID NO: 282), or TIGITI (SEQ ID NO: 301). Bispecific and multispecific antibodies may be generated using methods described herein.

Table 3.

mAb name	SEQ ID NO:							
	HCDR1	HCDR2	HCDR3	LCDR1	LCDR2	LCDR3	VH	VL
TM3B103	90	99	107	117	126	135	145	155
TM3B105	91	99	108	118	127	136	146	156

TM3B109	91	99	109	119	128	137	148	157
TM3B108	92	100	110	117	126	135	147	155
TM3B113	93	101	111	120	129	139	149	158
TM3B189	94	102	112	121	130	140	150	159
TM3B190	95	103	113	122	131	141	151	160
TM3B193	96	104	114	123	132	142	152	161
TM3B195	97	105	115	124	133	143	153	162
TM3B196	98	106	116	125	134	144	154	163
TM3B291	97	105	115	124	133	143	172	173

Homologous antibodies

Variants of the antagonistic antibodies specifically binding TIM-3 of the invention comprising VH or VL amino acid sequences shown in **Table 3, Table 36** and **Table 37** are within the scope of the invention. For example, variants may comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions in the VH and/or the VL as long as the homologous antibodies retain or have improved functional properties when compared to the parental antibodies. In some embodiments, the sequence identity may be about 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% to a VH or the VL amino acid sequence of the invention.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 148 and the VL of SEQ ID NO: 157, wherein the VH, the VL or both the VH and the VL optionally

comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 147 and the VL of SEQ ID NO: 155, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 149 and the VL of SEQ ID NO: 158, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 150 and the VL of SEQ ID NO: 159, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 151 and the VL of SEQ ID NO: 160, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 152 and the VL of SEQ ID NO: 161, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 153 and the VL

of SEQ ID NO: 162, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof,, comprising the VH of SEQ ID NO: 154 and the VL of SEQ ID NO: 163, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions. Optionally, any substitutions are not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH having the amino acid sequence at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NOs: 145, 146, 147, 148, 149, 150, 151, 152, 153, 154 or 172. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VL having the amino acid sequence at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VL of SEQ IS NOs: 155, 156, 157, 158, 159, 160, 161, 162, 163 or 173. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH having the amino acid sequence at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NOs: 145, 146, 147, 148, 149, 150, 151, 152, 153, 154 or 172 and the VL having the amino acid sequence at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VL of SEQ ID NOs: 155, 156, 157, 158, 159, 160, 161, 162, 163 or 173. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 148 and the VL of SEQ ID NO: 157. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 147 and the VL of SEQ ID NO: 155. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 149 and the VL of SEQ ID NO: 158. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 150 and the VL of SEQ ID NO: 159. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 151 and the VL of SEQ ID NO: 160. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 152 and the VL of SEQ ID NO: 161. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 154 and the VL of SEQ ID NO: 163. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL having the amino acid sequences at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173. Optionally, any variation from the sequences of the SEQ ID NOs is not within the CDRs.

The homologous antibodies of the invention described herein have substantially similar functionality when compared to the parental TIM-3 antibodies.

Antagonistic antibodies specifically binding TIM-3 of the invention with conservative modifications

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH comprising the HCDR1, the HCDR2 and the HCDR3 sequences and the VL comprising the LCDR1, the LCDR2 and the LCDR3 sequences, wherein one or more of the CDR sequences comprise specified amino acid sequences based on the antibodies described herein (e.g., antibodies shown in **Table 3, Table 36** or **Table 37** or conservative modifications thereof, and wherein the antibodies retain the desired functional properties of the parental antagonistic antibodies specifically binding TIM-3 of the invention.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3,

the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 90, 99, 107, 117, 126 and 135, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 91, 99, 108, 118, 127 and 136, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 91, 99, 109, 119, 128 and 137, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 92, 100, 110, 117, 126 and 135, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 93, 101, 111, 120, 129 and 139, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 94, 102, 112, 121, 130 and 140, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 95, 103, 113, 122, 131 and 141, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 96, 104, 114, 123, 132 and 142, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 97, 105, 115, 124, 133 and 143, respectively, and conservative modifications thereof.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 98, 106, 116, 125, 134 and 144, respectively, and conservative modifications thereof.

"Conservative modification" refers to modifications as described herein.

Antagonistic antibodies specifically binding TIM-3 of the invention with specific framework sequences

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH and the VL derived from particular human germline immunoglobulin sequences.

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV3-23 (SEQ ID NO: 174), IGHV1-02 (SEQ ID NO: 175), IGHV4-30-4 (SEQ ID NO: 176), IGHV1-03 (SEQ ID NO: 177), IGHV2-26 (SEQ ID NO: 178) or IGHV5-51 (SEQ ID NO: 179).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VL framework derived from IGKV3-20 (A27) (SEQ ID NO: 180), IGKV3-11 (L6) (SEQ ID NO: 171), IGKV4-1 (B3) (SEQ ID NO: 181), IGKV1-39) (O12) (SEQ ID NO: 182) or IGKV1-33 (O18) (SEQ ID NO: 183).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV3-23 (SEQ ID NO: 174) and the VL framework derived from IGKV3-20 (SEQ ID NO: 180).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV3-23 (SEQ ID NO: 174) and the VL framework derived from IGKV3-11 (SEQ ID NO: 171).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV3-23 (SEQ ID NO: 174) and the VL framework derived from IGKV4-1 (SEQ ID NO: 181).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from

IGHV1-02 (SEQ ID NO: 175) and the VL framework derived from IGKV4-1 (SEQ ID NO: 181).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV4-30-4 (SEQ ID NO: 176) and the VL framework derived from IGKV1-39 (SEQ ID NO: 182).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV1-03 (SEQ ID NO: 177) and the VL framework derived from IGKV1-33 (SEQ ID NO: 183).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV1-03 (SEQ ID NO: 177) and the VL framework derived from IGKV1-39 (SEQ ID NO: 182).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV2-26 (SEQ ID NO: 178) and the VL framework derived from IGKV4-1 (SEQ ID NO: 181).

The invention also provides an antagonistic antibody specifically binding TIM-3 or an antigen-binding portion thereof, comprising the VH framework derived from IGHV5-51 (SEQ ID NO: 179) and the VL framework derived from IGKV1-39 (SEQ ID NO: 182).

The antibodies of the invention comprising heavy or light chain variable regions "derived from" a particular framework or germline sequence refer to antibodies obtained from a system that uses human germline immunoglobulin genes, such as from transgenic mice or from phage display libraries as discussed herein. An antibody that is "derived from" a particular framework or germline sequence may contain amino acid differences as compared to the sequence it was derived from, due to, for example, naturally-occurring somatic mutations or intentional substitutions.

Exemplary antagonistic antibodies specifically binding TIM-3 having certain VH and VL framework sequences are shown in **Table 38.**

Bispecific anti-PD-1/TIM-3 antibodies

The invention also provides antagonistic bispecific PD-1/TIM-3 antibodies.

The invention also provides an isolated antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention enhances activation of antigen-specific $CD4^+$ or $CD8^+$ T cells.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention enhances activation of antigen-specific $CD4^+$ or $CD8^+$ T cells, wherein enhanced activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant increase of CD137 surface expression on antigen-specific $CD4^+$ or $CD8^+$ T cells.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention inhibits TIM-3 binding to galectin-9.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention

binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;

binds human PD-1 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;

binds cynomolgus PD-1 with the K_D of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C; or

binds cynomolgus PD-1 with the K_D of less than about 1 nM;

wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention enhances an activation of antigen-specific $CD4^+$ or $CD8^+$ T cells, wherein the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant increase of CD137 surface expression on antigen-specific $CD4^+$ or $CD8^+$ T cells and binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention enhances the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells, wherein the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant increase of CD137 surface expression on antigen-specific $CD4^+$ or $CD8^+$ T cells, and binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention enhances the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells, wherein the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant increase of CD137 surface expression on antigen-specific $CD4^+$ or $CD8^+$ T cells and binds cynomolgus PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention enhances the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells, wherein the activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant increase of CD137 surface expression on antigen-specific $CD4^+$ or $CD8^+$ T cells, and binds cynomolgus PD-1 with an equilibrium dissociation constant (K_D) of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

The antagonistic bispecific PD-1/TIM-3 antibodies of the invention described herein may be evaluated for their ability to enhance antigen specific $CD4^+$ or $CD8^+$ T cell activation, to inhibit TIM-3 binding to galectin-9, and binding kinetics to human or cynomolgus PD-1 or TIM-3 may be assessed using methods described herein.

For example, CD137 may be used as a marker for activation of antigen specific $CD4^+$ or $CD8^+$ T cells. CD137 surface expression may be measured on T cells cultured in the presence or in the absence of a test antibody, such as the bispecific PD-1/TIM-3 antibody, using anti-CD137 antibody and a secondary antibody conjugated for example to a fluorescent dye. The statistically significant difference in the obtained signal on T cells cultured in the presence or in the absence of the test antibody is evaluated.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention binds TIM-3 within TIM-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261).

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention binds TIM-3 within TIM-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261) and residues 50-56 (DERDVNY) SEQ ID NO: 262.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention binds TIM-3 within TIM-3 residues 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263).

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody of the invention binds TIM-3 within TIM-3 residues 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263) and residues 50-56 (DERDVNY) SEQ ID NO: 262.

In some embodiments, the first domain comprises a heavy chain complementarity determining region (HCDR) 1 a HCDR2 and a HCDR3 of SEQ ID NOs: 82, 83 and 84, respectively.

In some embodiments, the first domain comprises the HCDR1, the HCDR2 and the HCDR3 of SEQ ID NOs: 82, 83 and 85, respectively.

In some embodiments, the first domain comprises a light chain complementarity determining regions (LCDR) 1, a LCDR2 and a LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.

In some embodiments, the first domain comprises the HCDR1, the HCDR2 and the HCDR3 of SEQ ID NOs: 82, 83 and 84, respectively, and the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.

In some embodiments, the first domain comprises the HCDR1, the HCDR2 and the HCDR3 of SEQ ID NOs: 82, 83 and 85, respectively, and the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.

In some embodiments, the second domain comprises the HCDR1, the HCDR2 and the HCDR3 amino acid sequences of SEQ ID NOs: 164, 165 and 166, respectively.

In some embodiments, the second domain comprises the LCDR1, the LCDR2 and the LCDR3 amino acid sequences of SEQ ID NOs: 167, 168 and 169, respectively.

In some embodiments, the second domain comprises the HCDR1, the HCDR2 and the HCDR3 amino acid sequences of SEQ ID NOs: 164, 165 and 166, respectively, and the LCDR1, the LCDR2 and the LCDR3 amino acid sequences of SEQ ID NOs: 167, 168 and 169 respectively.

In some embodiments, the first domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of

SEQ ID NOs: 10, 13, 16, 20, 26 and 31, respectively;

SEQ ID NOs: 10, 13, 16, 21, 26 and 32, respectively;

SEQ ID NOs: 10, 14, 16, 22, 27 and 33, respectively;

SEQ ID NOs: 10, 14, 16, 22, 26 and 34, respectively;

SEQ ID NOs: 10, 14, 16, 23, 28 and 35, respectively;

SEQ ID NOs: 10, 13, 17, 20, 26 and 31, respectively;

SEQ ID NOs: 10, 13, 17, 20, 26 and 36, respectively;

SEQ ID NOs: 10, 13, 17, 21, 26 and 32, respectively;

SEQ ID NOs: 10, 13, 17, 21, 27 and 37, respectively;

SEQ ID NOs: 10, 13, 17, 23, 26 and 32, respectively;

SEQ ID NOs: 10, 13, 17, 22, 26 and 32, respectively;

SEQ ID NOs: 10, 13, 18, 20, 26 and 31, respectively;

SEQ ID NOs: 11, 15, 18, 20, 26 and 31, respectively;

SEQ ID NOs: 10, 13, 19, 20, 26 and 31, respectively;

SEQ ID NOs: 12, 13, 19, 20, 26 and 31, respectively;

SEQ ID NOs: 10, 14, 17, 23, 28 and 35, respectively;

SEQ ID NOs: 10, 14, 17, 22, 26 and 34, respectively;

SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively;

SEQ ID NOs: 12, 13, 19, 24, 26 and 38, respectively;

SEQ ID NOs: 12, 13, 19, 20, 29 and 39, respectively;

SEQ ID NOs: 11, 15, 18, 20, 30 and 32, respectively;

SEQ ID NOs: 11, 15, 18, 25, 26 and 40, respectively;

SEQ ID NOs: 11, 15, 18, 24, 26 and 32, respectively; or

SEQ ID NOs: 66, 67, 68, 69, 70 and 71, respectively.

In some embodiments, the second domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of

SEQ ID NOs: 90, 99, 107, 117, 126 and 135, respectively;

SEQ ID NOs: 91, 99, 108, 118, 127 and 136, respectively;

SEQ ID NOs: 91, 99, 109, 119, 128 and 137, respectively;

SEQ ID NOs: 92, 100, 110, 117, 126 and 135, respectively;

SEQ ID NOs: 93, 101, 111, 120, 129 and 139, respectively;

SEQ ID NOs: 94, 102, 112, 121, 130 and 140, respectively;

SEQ ID NOs: 95, 103, 113, 122, 131 and 141, respectively;

SEQ ID NOs: 96, 104, 114, 123, 132 and 142, respectively;

SEQ ID NOs: 97, 105, 115, 124, 133 and 143, respectively; or

SEQ ID NOs: 98, 106, 116, 125, 134 and 144, respectively.

In some embodiments, the first domain comprises the VH of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47, 48, 63 or 64, the VH optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen conservative amino acid substitutions. Optionally, any substitutions are not within the CDRs.

In some embodiments, the first domain comprises the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62 or 65, the VL optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen conservative amino acid substitutions. Optionally, any substitutions are not within the CDRs.

In some embodiments, the first domain comprises the VH of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47, 48, 63 or 64 and the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62 and 65, the VH, the VL, or the VH and the VL optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen conservative amino acid substitutions. Optionally, any substitutions are not within the CDRs.

In some embodiments, the second domain comprises the VH of SEQ ID NOs: 145, 146, 147, 148, 149, 150, 151, 152, 153, 154 or 172, the VH optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen conservative amino acid substitutions. Optionally, any substitutions are not within the CDRs.

In some embodiments, the second domain comprises the VL of SEQ IS NOs: 155, 156, 157, 158, 159, 160, 161, 162, 163 or 173, the VL optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen conservative amino acid substitutions. Optionally, any substitutions are not within the CDRs.

In some embodiments, the second domain comprises the VH of SEQ ID NOs: 145, 146, 147, 148, 149, 150, 151, 152, 153, 154 or 172 and the VL of SEQ ID NOs: 155, 156, 157, 158, 159, 160, 161, 162, 163 or 173, the VH and the VL optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen conservative amino acid substitutions. Optionally, any substitutions are not within the CDRs.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 50.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 51.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 52.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 53.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 49.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 54.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 50.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 55.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 56.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 57.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 44 and the VL of SEQ ID NO: 49.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 49.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 46 and the VL of SEQ ID NO: 49.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 49.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 53.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 52.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 59.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 61.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 62.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 63 and the VL of SEQ ID NO: 65.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 148 and the VL of SEQ ID NO: 157.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 147 and the VL of SEQ ID NO: 155.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 149 and the VL of SEQ ID NO: 158.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 150 and the VL of SEQ ID NO: 159.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 151 and the VL of SEQ ID NO: 160.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 152 and the VL of SEQ ID NO: 161.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 154 and the VL of SEQ ID NO: 163.

In some embodiments, the second domain comprises the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

The invention also provides an isolated antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3, wherein the first domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively, and the second domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 91, 99, 108, 118, 127 and 136, respectively.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody binds TIM-3 within TIM-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261).

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody binds TIM-3 within TIM-3 residues 32-47 (WGKGACPVFECGNVVL) (SEQ ID NO: 261) and residues 50-56 (DERDVNY) SEQ ID NO: 262.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody inhibits TIM-3 binding to galectin-9.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 and the second domain comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising a F405L and/or a K409R substitution.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4 isotype comprising a F405L and a K409R substitution.

In some embodiments, the antibody is an IgG4 isotype comprising a heavy chain substitution S228P when compared to the wild type IgG4.

In some embodiments, the isolated antagonistic bispecific PD-1/TIM-3 antibody comprises a first heavy chain (HC1) a first light chain (LC1), a second heavy chain (HC2) and a second light chain (LC2) of SEQ ID NOs: 241, 188, 245 or 194, respectively.

In some embodiments, the isolated antagonistic bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 186, 188, 191 or 194, respectively.

In some embodiments, the isolated antagonistic bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 186, 188, 248 or 194, respectively.

In some embodiments, the isolated antagonistic bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 243, 188, 246 or 194, respectively.

The antibody is suitable for use in therapy, for example in treating a cancer.

The antibody is suitable for use in therapy, for example in treating a solid tumor.

The antibody is suitable for use in therapy, for example in treating a melanoma.

The antibody is suitable for use in therapy, for example in treating a lung cancer.

The antibody is suitable for use in therapy, for example in treating a non-small cell lung cancer (NSCLC)

The antibody is suitable for use in therapy, for example in treating a squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a non-squamous NSCLC.

The antibody is suitable for use in therapy, for example in treating a lung adenocarcinoma.

The antibody is suitable for use in therapy, for example in treating a renal cell carcinoma (RCC).

The antibody is suitable for use in therapy, for example in treating a mesothelioma.

The antibody is suitable for use in therapy, for example in treating a nasopharyngeal carcinoma (NPC).

The antibody is suitable for use in therapy, for example in treating a colorectal cancer.

The antibody is suitable for use in therapy, for example in treating a prostate cancer.

The antibody is suitable for use in therapy, for example in treating a castration-resistant prostate cancer.

The antibody is suitable for use in therapy, for example in treating a stomach cancer.

The antibody is suitable for use in therapy, for example in treating an ovarian cancer.

The antibody is suitable for use in therapy, for example in treating a gastric cancer.

The antibody is suitable for use in therapy, for example in treating a liver cancer.

The antibody is suitable for use in therapy, for example in treating pancreatic cancer.

The antibody is suitable for use in therapy, for example in treating a thyroid cancer.

The antibody is suitable for use in therapy, for example in treating a squamous cell carcinoma of the head and neck.

The antibody is suitable for use in therapy, for example in treating a carcinomas of the esophagus or gastrointestinal tract.

The antibody is suitable for use in therapy, for example in treating a breast cancer.

The antibody is suitable for use in therapy, for example in treating a fallopian tube cancer.

The antibody is suitable for use in therapy, for example in treating a brain cancer.

The antibody is suitable for use in therapy, for example in treating an urethral cancer.

The antibody is suitable for use in therapy, for example in treating an endometriosis.

The antibody is suitable for use in therapy, for example in treating a cervical cancer.

The antibody is suitable for use in therapy, for example in treating a metastatic lesion of the cancer.

The antibody is suitable for use in therapy in a subject who is being treated or who has been treated with anti-PD-1 antibody comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab)).

The antibody is suitable for use in therapy in a subject who is being treated or who has been treated with anti-PD-1 antibody comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The antibody is suitable for use in therapy in a subject who is refractory to treatment with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab)).

The antibody is suitable for use in therapy in a subject who is refractory to treatment with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The antibody is suitable for use in therapy in a subject who has a relapsed tumor after treatment with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231. (e.g. KEYTRUDA® (pembrolizumab).

The antibody is suitable for use in therapy in a subject who has a relapsed tumor after treatment with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233. (e.g. OPDIVO® (nivolumab)).

The invention also provides an isolated antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3, wherein the first domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 66, 67, 68, 69, 70 and 71, respectively, and the second domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs 97, 105, 115, 124, 133 and 143, respectively.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody binds TIM-3 within TIM-3 residues 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263).

In some embodiments, the bispecific PD-1/TIM-3 antibody binds TIM-3 within TIM-3 residues 90-102 (RIQIPGIMNDEKF) (SEQ ID NO: 263) and residues 50-56 (DERDVNY) SEQ ID NO: 262.

In some embodiments, the bispecific PD-1/TIM-3 antibody inhibits binding of TIM-3 to galectin-9.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 and the second domain comprises the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising a F405L and/or a K409R substitution.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4 isotype comprising a F405L and a K409R substitution.

In some embodiments, the antibody is an IgG4 isotype comprising a heavy chain substitution S228P when compared to the wild type IgG4.

In some embodiments, the isolated bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 187, 189, 190 and 193, respectively.

The invention also provides an isolated antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain

specifically binding TIM-3, wherein the first domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 66, 67, 68, 69, 70 and 71, respectively, and the second domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 91, 99, 108, 118, 127 and 136, respectively.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 and the second domain comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising a F405L and/or a K409R substitution.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4 isotype comprising a F405L and a K409R substitution.

In some embodiments, the antibody is an IgG4 isotype comprising a heavy chain substitution S228P when compared to the wild type IgG4.

In some embodiments, the isolated antagonistic bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 187, 189, 191 and 194, respectively.

In some embodiments, the isolated bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 242, 189, 246 and 194, respectively.

The invention also provides an isolated antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3, wherein the first domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively, and the second domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 97, 105, 115, 124, 133 and 143, respectively.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 and the second domain comprises the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising a F405L and/or a K409R substitution.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4 isotype comprising a F405L and a K409R substitution.

In some embodiments, the antibody is an IgG4 isotype comprising a heavy chain substitution S228P when compared to the wild type IgG4.

In some embodiments, the isolated antagonistic bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 186, 188, 192 and 195, respectively.

In some embodiments, the isolated antagonistic bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 241, 188, 244 and 195, respectively.

In some embodiments, the isolated antagonistic bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 243, 188, 247 and 195, respectively.

In some embodiments, the antibody enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells, wherein activation of antigen-specific $CD4^+$ or $CD8^+$ T cells is assessed by measuring a statistically significant enhancement of CD137 surface expression on antigen specific $CD4^+$ or $CD8^+$ T cells according to methods described in Example 14.

The invention also provides an isolated antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3, wherein the first domain comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively, and the second domain comprises the HCDR1, the HCDR2, the

HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 97, 105, 115, 124, 133 and 143, respectively.

In some embodiments, the first domain comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 and the second domain comprises the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 156.

In some embodiments, the antibody is an IgG1 isotype.

In some embodiments, the antibody is an IgG2 isotype.

In some embodiments, the antibody is an IgG2 isotype comprising a F405L and/or a K409R substitution.

In some embodiments, the antibody is an IgG2 isotype, optionally comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2.

In some embodiments, the antibody is an IgG3 isotype.

In some embodiments, the antibody is an IgG4 isotype, optionally comprising a S228P substitution when compared to the wild type IgG4.

In some embodiments, the antibody is an IgG4 isotype comprising a F405L and a K409R substitution.

In some embodiments, the antibody is an IgG4 isotype comprising a heavy chain substitution S228P when compared to the wild type IgG4.

In some embodiments, the isolated bispecific PD-1/TIM-3 antibody comprises the HC1, the LC1, the HC2 and the LC2 of SEQ ID NOs: 186, 188, 190 and 193, respectively.

Exemplary antagonistic bispecific PD-1/TIM-3 antibodies of the invention having certain VH, VL, HCDR and LCDR sequences as shown in **Table 4 and Table 5.**

Table 4.

mAb	PD-1 binding arm SEQ ID NOs:							
	VH	VL	HCDRs			LCDRs		
			1	2	3	1	2	3
PTBB14	48	56	10	14	17	23	26	32
PTBB15	48	56	10	14	17	23	26	32
PTBB16	64	65	66	67	68	69	70	71
PTBB17	64	65	66	67	68	69	70	71

PTBB24	48	56	10	14	17	23	26	32
PTBB30	48	56	10	14	17	23	26	32
PTBB27	48	56	10	14	17	23	26	32
PTBB28	48	56	10	14	17	23	26	32
PTBB18	64	65	66	67	68	69	70	71
PTBB20	48	56	10	14	17	23	26	32
PTBB21	48	56	10	14	17	23	26	32

Table 5.

mAb	TIM-3 binding arm SEQ ID NOs:							
	VH	VL	HCDRs			LCDR2		
			1	2	3	1	2	3
PTBB14	153	162	97	105	115	124	133	143
PTBB15	146	156	91	99	108	118	127	136
PTBB16	153	162	97	105	115	124	133	143
PTBB17	146	156	91	99	108	118	127	136
PTBB24	172	173	97	105	115	124	133	143
PTBB30	146	156	91	99	108	118	127	136
PTBB27	172	173	97	105	115	124	133	143
PTBB28	146	156	91	99	108	118	127	136
PTBB18	146	156	91	99	108	118	127	136
PTBB20	146	156	91	99	108	118	127	136
PTBB21	172	173	97	105	115	124	133	143

Engineered and modified antibodies

The antibodies of the invention may further be engineered to generate modified antibodies with similar or altered properties when compared to the parental antibodies.

The VH, the VL, the VH and the VL, the constant regions, VH framework, VL framework, or any or all of the six CDRs may be engineered in the antibodies of the invention.

"The antibodies of the invention" as used herein refers to the antagonistic antibodies specifically binding PD-1, the antagonistic antibodies specifically binding TIM-3, and the antagonistic bispecific PD-1/TIM-3 antibodies comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3 (e.g. bispecific PD-1/TIM-3 antibodies) as described herein.

The antibodies of the invention may be engineered by CDR grafting. One or more CDR sequences of the antibodies of the invention described herein may be grafted to a different framework sequence. CDR grafting may be done using known methods and methods described herein.

In some embodiments, the antagonistic antibodies specifically binding PD-1 or the bispecific PD-1/TIM-3 antibodies of the invention comprise the VH that comprises the HDCR1 of SEQ ID NOs: 10, 11 or 12, the HCDR2 of SEQ ID NOs: 13, 14 or 15, the HCDR3 of SEQ ID NOs: 16, 17, 18 or 19, and the VL that comprises the LCDR1 of SEQ ID NOs: 20, 21, 22, 23, 24 or 25, the LCDR2 of SEQ ID NOs: 26, 27, 28, 29 or 30, and/or the LCDR3 of SEQ ID NOs: 31, 32, 33, 34, 35, 36, 37, 38, 39 or 40, wherein the VH framework is derived from the VH framework other than VH1-69 (SEQ ID NO: 170) and the VL framework is derived from the VL framework other than IGKV3-11 (SEQ ID NO: 171).

In some embodiments, the antagonistic antibodies specifically binding TIM-3 or the bispecific PD-1/TIM-3 antibodies of the invention comprise the HDCR1 of SEQ ID NOs: 90, 91, 92, 93, 94, 95, 96, 97 or 98, the HCDR2 of SEQ ID NOs: 99, 100, 101, 102, 10, 104, 105 or 106, the HCDR3 of SEQ ID NOs: 107, 108, 109, 110, 111, 112, 113, 114, 115 or 116, and the VL that comprises the LCDR1 of SEQ ID NOs: 117, 118, 119, 120, 121, 122, 123, 124 or 125, the LCDR2 of SEQ ID NOs: 126, 127, 128, 129, 130, 131, 132, 133 or 134, and/or the LCDR3 of SEQ ID NOs: 135, 136, 137, 139, 140, 141, 142, 143 or 144, wherein the VH framework is derived from the human VH germline gene sequences other than those of IGHV3-23 (SEQ ID NO: 174), IGHV1-02 (SEQ ID NO: 175), IGHV4-30-4 (SEQ ID NO: 176), IGHV1-03 (SEQ ID NO: 177), IGHV2-26 (SEQ ID NO: 178) or IGHV5-51 (SEQ ID NO: 179), and the VL framework is derived from the human VL germline gene sequences other than those of IGKV3-20 (A27) (SEQ ID NO: 180), IGKV3-11 (L6) (SEQ ID NO: 171), IGKV4-1 (B3) (SEQ ID NO: 181), IGKV1-39 (O12) (SEQ ID NO: 182) or IGKV1-33 (O18) (SEQ ID NO: 183).

The framework sequences to be used may be obtained from public DNA databases or published references that include germline antibody gene sequences. For example, germline DNA and the encoded protein sequences of human heavy and light chain variable region genes may be found at IMGT®, the international ImMunoGeneTics information system® (http://_www-imgt_org). Framework sequences that may be used to replace the existing framework sequences in the antibodies of the invention may be those that show the highest percent identity to the parental frameworks over the entire length of the VH or the VL, or over the length of the FR1, FR2, FR3 and FR4. In addition, suitable frameworks may further be selected based on the VH and the VL CDR1 and CDR2 lengths or identical LCDR1, LCDR2, LCDR3, HCDR1 and HCDR2 canonical structure. Suitable frameworks may be selected using known methods, such as human framework adaptation described in U.S. Patent No. 8,748,356 or superhumanization described in U.S. Patent No. 7,709, 226.

The framework sequences of the parental and engineered antibodies may further be modified, for example by backmutations to restore and/or improve binding of the generated antibody to the antigen as described for example in U.S. Patent No. 6,180,370. The framework sequences of the parental or engineered antibodies may further be modified by mutating one or more residues within the framework region, or within one or more CDR regions, to remove T-cell epitopes to thereby reduce the potential immunogenicity of the antibody. This approach is also referred to as "deimmunization" and described in further detail in U.S. Patent Publ. No. US20070014796.

The CDR residues of the antibodies of the invention may be mutated to improve affinity of the antibodies to PD-1, TIM-3, or PD-1 and TIM-3.

The CDR residues of the antibodies of the invention may be mutated for example to minimize risk of post-translational modifications. Amino acid residues of putative motifs for deamination (NS), acid-catalyzed hydrolysis (DP), isomerization (DS), or oxidation (W) may be substituted with any of the naturally occurring amino acids to mutagenize the motifs, and the resulting antibodies may be tested for their functionality and stability using methods described herein.

Fc substitutions may be made to the antibodies of the invention to modulate antibody effector functions and pharmacokinetic properties. In traditional immune function, the interaction of antibody-antigen complexes with cells of the immune system results in a wide array of responses, ranging from effector functions such as antibody-dependent cytotoxicity, mast cell degranulation, and phagocytosis to immunomodulatory signals such as regulating lymphocyte proliferation and antibody secretion. All of these

interactions are initiated through the binding of the Fc domain of antibodies or immune complexes to specialized cell surface receptors on hematopoietic cells. The diversity of cellular responses triggered by antibodies and immune complexes results from the structural heterogeneity of the three Fc receptors: FcγRI (CD64), FcγRII (CD32), and FcγRIII (CD16). FcγRI (CD64), FcγRIIA (CD32A) and FcγRIII (CD16) are "activating Fcγ receptors" (i e, immune system enhancing); FcγRIIB (CD32B) is an inhibiting Fcγ receptor" (i.e., immune system dampening). Binding to the FcRn receptor modulates antibody half-life.

In some embodiments, the antagonistic antibodies of the invention comprise at least one substitution in an Fc region

In some embodiments, the antagonistic antibodies of the invention comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or

Fc positions that may be substituted to modulate antibody half-life are those described for example in Dall'Acqua *et al.*, (2006) *J Biol Chem* 281:23514–240, Zalevsky *et al.,* (2010) *Nat Biotechnol* 28:157-159, Hinton *et al.,* (2004) *J Biol Chem* 279(8):6213-6216, Hinton *et al.,* (2006) *J Immunol* 176:346-356, Shields *et al.*(2001) *J Biol Chem* 276:6591-6607, Petkova *et al.,* (2006). *Int Immunol* 18:1759-1769, Datta-Mannan *et al.,* (2007) *Drug Metab Dispos*, 35:86-94, 2007, Vaccaro *et al.,* (2005) *Nat Biotechnol* 23:1283-1288, Yeung *et al.,* (2010) *Cancer Res,* 70:3269-3277 and Kim *et al.,* (1999) *Eur J Immunol* 29: 2819, and include positions 250, 252, 253, 254, 256, 257, 307, 376, 380, 428, 434 and 435. Exemplary substitutions that may be made singularly or in combination are substitutions T250Q, M252Y, I253A, S254T, T256E, P257I, T307A, D376V, E380A, M428L, H433K, N434S, N434A, N434H, N434F, H435A and H435R. Exemplary singular or combination substitutions that may be made to increase the half-life of the antibody are substitutions M428L/N434S, M252Y/S254T/T256E, T250Q/M428L, N434A and T307A/E380A/N434A. Exemplary singular or combination substitutions that may be made to reduce the half-life of the antibody are substitutions H435A, P257I/N434H, D376V/N434H, M252Y/S254T/T256E/H433K/N434F, T308P/N434A and H435R.

In some embodiments, the antibodies of the invention comprise at least one substitution in the antibody Fc at amino acid position 250, 252, 253, 254, 256, 257, 307, 376, 380, 428, 434 or 435.

In some embodiments, the antibodies of the invention comprise at least one substitution in the antibody Fc selected from the group consisting of T250Q, M252Y, I253A, S254T, T256E, P257I, T307A, D376V, E380A, M428L, H433K, N434S, N434A, N434H, N434F, H435A and H435R.

In some embodiments, the antibodies of the invention comprise at least one substitution in the antibody Fc selected from the group consisting of M428L/N434S, M252Y/S254T/T256E, T250Q/M428L, N434A, T307A/E380A/N434A, H435A, P257I/N434H, D376V/N434H, M252Y/S254T/T256E/H433K/N434F, T308P/N434A and H435R.

In some embodiments, the antibodies of the invention comprise at least one substitution in the antibody Fc that reduces binding of the antibody to an activating Fcγ receptor (FcγR) and/or reduces Fc effector functions such as C1q binding, complement dependent cytotoxicity (CDC), antibody-dependent cell-mediated cytotoxicity (ADCC) or phagocytosis (ADCP).

Fc positions that may be substituted to reduce binding of the antibody to the activating FcγR and subsequently to reduce effector function are those described for example in Shields *et al.,* (2001) *J Biol Chem* 276:6591-6604, Intl. Patent Publ. No. WO2011/066501, U.S. Patent Nos. 6,737,056 and 5,624,821, Xu *et al.,* (2000) *Cell Immunol,* 200:16-26, Alegre *et al.,* (1994) *Transplantation* 57:1537-1543, Bolt *et al.,* (1993) *Eur J Immunol* 23:403-411, Cole *et al.,* (1999) *Transplantation,* 68:563-571, Rother *et al.,* (2007) *Nat Biotechnol* 25:1256-1264, Ghevaert *et al.,* (2008) *J Clin Invest* 118:2929-2938, An *et al.,* (2009) mAbs, 1:572-579) and include positions 214, 233, 234, 235, 236, 237, 238, 265, 267, 268, 270, 295, 297, 309, 327, 328, 329, 330, 331 and 365. Exemplary substitutions that may be made singularly or in combination are substitutions K214T, E233P, L234V, L234A, deletion of G236, V234A, F234A, L235A, G237A, P238A, P238S, D265A, S267E, H268A, H268Q, Q268A, N297A, A327Q, P329A, D270A, Q295A, V309L, A327S, L328F, A330S and P331S in IgG1, IgG2, IgG3 or IgG4. Exemplary combination substitutions that result in antibodies with reduced ADCC are substitutions L234A/L235A on IgG1, V234A,/G237A/ P238S/H268A/V309L/A330S/P331S on IgG2, F234A/L235A on IgG4, S228P/F234A/ L235A on IgG4, N297A on all Ig isotypes, V234A/G237A on IgG2, K214T/E233P/ L234V/L235A/G236-deleted/A327G/P331A/D365E/L358M on IgG1, H268Q/V309L/ A330S/P331S on IgG2, S267E/L328F on IgG1, L234F/L235E/D265A on IgG1, L234A/L235A/G237A/P238S/H268A/A330S/P331S on IgG1, S228P/F234A/L235A/G237A/P238S on IgG4, and S228P/F234A/L235A/G236-deleted/G237A/P238S on IgG4. Hybrid IgG2/4 Fc domains may also be used, such as Fc with residues 117-260 from IgG2 and residues 261-447 from IgG4.

Well-known S228P substitution may be made in IgG4 antibodies to enhance IgG4 stability.

In some embodiments, the antibodies of the invention comprise a substitution in at least one residue position 214, 233, 234, 235, 236, 237, 238, 265, 267, 268, 270, 295, 297, 309, 327, 328, 329, 330, 331 or 365, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise at least one substitution selected from the group consisting of K214T, E233P, L234V, L234A, deletion of G236, V234A, F234A, L235A, G237A, P238A, P238S, D265A, S267E, H268A, H268Q, Q268A, N297A, A327Q, P329A, D270A, Q295A, V309L, A327S, L328F, A330S and P331S, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a substitution in at least one residue position 228, 234, 235, 237, 238, 268, 330 or 331, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a S228P substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a V234A substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a F234A substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a G237A substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a P238S substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a H268A substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a Q268A substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise an A330S substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise a P331S substitution, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise L234A, L235A, G237A, P238S, H268A, A330S and P331S substitutions, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise F234A, L235A, G237A, P238S and Q268A substitutions, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise L234A, L235A or L234A and L235A substitutions, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise F234A, L235A or F234A and L235A substitutions, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise S228P, F234A and L235A substitutions, wherein residue numbering is according to the EU Index.

In some embodiments, the antibodies of the invention comprise at least one substitution in an antibody Fc that enhances binding of the antibody to an Fcγ receptor (FcγR) and/or enhances Fc effector functions such as C1q binding, complement dependent cytotoxicity (CDC), antibody-dependent cell-mediated cytotoxicity (ADCC) or phagocytosis (ADCP).

In addition to their immunomodulatory activity, the PD-1 or the TIM-3 antibodies of the invention may kill tumor cells expressing PD-1 and/or TIM-3 directly via antibody-mediated effector functions, for example by ADCC, ADCP or CDC.

Fc positions that may be substituted to increase binding of the antibody to the activating Fcγ and/or enhance antibody effector functions are those described for example in U.S. Patent No. 6,737,056, U.S. Patent Publ. No. 2015/0259434, Shields *et al.,* (2001) *J Biol Chem* 276:6591-6604, Lazar *et al.,* (2006) *Proc Natal Acad Sci*, 103:4005-4010, Stavenhagen et al., (2007) *Cancer Res* 67:8882-8890, Richards *et al.,* (2008) *Mol Cancer Ther* 7:2517-2527, Diebolder *et al., Science;* published online March 13, 2014; doi:10.1126/science.1248943, and include positions 236, 239, 243, 256, 290, 292, 298, 300, 305, 312, 326, 330, 332, 333, 334, 345, 360, 339, 378, 396 or 430 (residue numbering according to the EU index). Exemplary substitutions that may be made singularly or in combination are G236A, S239D, F243L, T256A, K290A, R292P, S298A, Y300L, V305L, K326A, A330K, I332E, E333A, K334A, A339T and P396L. Exemplary combination substitutions that result in antibodies with increased ADCC or ADCP are substitutions S239D/I332E, S298A/E333A/K334A, F243L/R292P/Y300L,

F243L/R292P/Y300L/P396L, F243L/R292P/Y300L/V305I/P396L and G236A/S239D/I332E on IgG1.

Fc positions that may be substituted to enhance CDC of the antibody are those described for example in Int. Patent Appl. WO2014/108198, Idusogie *et al.,* (2001) *J Immunol* 166:2571-2575 and Moore *et al.,* (2010) Mabs, 2:181-189, and include positions 267, 268, 324, 326, 333, 345 and 430. Exemplary substitutions that may be made singularly or in combination are substitutions S267E, H268F, S324T, K326A, K326W, E333A, E345K, E345Q, E345R, E345Y, E430S, E430F and E430T. Exemplary combination substitutions that result in antibodies with increased CDC are substitutions K326A/E333A, K326W/E333A, H268F/S324T, S267E/H268F, S267E/S324T and S267E/H268F/S324T on IgG1.

"Antibody-dependent cellular cytotoxicity", "antibody-dependent cell-mediated cytotoxicity" or "ADCC" is a mechanism for inducing cell death that depends upon the interaction of antibody-coated target cells with effector cells possessing lytic activity, such as natural killer cells, monocytes, macrophages and neutrophils via Fc gamma receptors (FcγR) expressed on effector cells. For example, NK cells express FcγRIIIa, whereas monocytes express FcγRI, FcγRII and FcγRIIIa. Death of the antibody-coated target cell, such as PD-1 or TIM-3 expressing cells, occurs as a result of effector cell activity through the secretion of membrane pore-forming proteins and proteases. To assess ADCC activity of the antibody of the invention described herein, the antibody may be added to TIM-3 or PD-1 expressing cells in combination with immune effector cells, which may be activated by the antigen antibody complexes resulting in cytolysis of the target cell. Cytolysis may be detected by the release of label (e.g. radioactive substrates, fluorescent dyes or natural intracellular proteins) from the lysed cells. Exemplary effector cells for such assays include peripheral blood mononuclear cells (PBMC) and NK cells. Exemplary target cells include cells expressing TIM-3 or PD-1 either endogenously or recombinantly. In an exemplary assay, target cells are used with a ratio of 1 target cell to 50 effector cells. Target cells are pre-labeled with BATDA (PerkinElmer) for 20 minutes at 37°C, washed twice and resuspended in DMEM, 10% heat-inactivated FBS, 2mM L-glutamine (all from Invitrogen). Target ($1x10^4$ cells) and effector cells ($0.5x10^6$ cells) are combined and 100 µl of cells are added to the wells of 96-well U-bottom plates. An additional 100 µl is added with or without the test antibodies. The plates are centrifuged at 200g for 3 minutes, incubated at 37°C for 2 hours, and then centrifuged again at 200g for 3 minutes. A total of 20 µl of supernatant is removed per well and cell lysis is measured by the

addition of 200 µl of the DELPHIA Europium-based reagent (PerkinElmer). Data is normalized to maximal cytotoxicity with 0.67% Triton X-100 (Sigma Aldrich) and minimal control determined by spontaneous release of BATDA from target cells in the absence of any antibody. The antibody of the invention may induce ADCC by about 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95% or 100% .

"Antibody-dependent cellular phagocytosis" ("ADCP") refers to a mechanism of elimination of antibody-coated target cells by internalization by phagocytic cells, such as macrophages or dendritic cells. ADCP may be evaluated by using monocyte-derived macrophages as effector cells and Daudi cells (ATCC® CCL-213™) or B cell leukemia or lymphoma or tumor cells expressing TIM-3 or PD-1 as target cells engineered to express GFP or other labeled molecule. Effector:target cell ratio may be for example 4:1. Effector cells may be incubated with target cells for 4 hours with or without the antibody of the invention. After incubation, cells may be detached using accutase. Macrophages may be identified with anti-CD11b and anti-CD14 antibodies coupled to a fluorescent label, and percent phagocytosis may be determined based on % GFP fluorescence in the $CD11^{+}CD14^{+}$ macrophages using standard methods. The antibody of the invention may induce ADCP by about 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95% or 100% .

"Complement-dependent cytotoxicity", or "CDC", refers to a mechanism for inducing cell death in which the Fc effector domain of a target-bound antibody binds and activates complement component C1q which in turn activates the complement cascade leading to target cell death. Activation of complement may also result in deposition of complement components on the target cell surface that facilitate ADCC by binding complement receptors (e.g., CR3) on leukocytes. CDC of TIM-3 or PD-1 expressing cells may be measured for example by plating Daudi cells at 1×10^{5} cells/well (50 µl/well) in RPMI-B (RPMI supplemented with 1% BSA), adding 50 µl of test antibodies to the wells at final concentration between 0-100 µg/ml, incubating the reaction for 15 min at room temperature, adding 11 µl of pooled human serum to the wells, and incubation the reaction for 45 min at 37° C. Percentage (%) lysed cells may be detected as % propidium iodide stained cells in FACS assay using standard methods. Antibodies of the invention may induce CDC by about 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95% or 100% .

The ability of antibodies of the invention described herein to induce ADCC may be enhanced by engineering their oligosaccharide component. Human IgG1 or IgG3 are

N-glycosylated at Asn297 with the majority of the glycans in the well-known biantennary G0, G0F, G1, G1F, G2 or G2F forms. Antibodies produced by non-engineered CHO cells typically have a glycan fucose content of about at least 85%. The removal of the core fucose from the biantennary complex-type oligosaccharides attached to the Fc regions enhances the ADCC of antibodies via improved FcγRIIIa binding without altering antigen binding or CDC activity. Such mAbs may be achieved using different methods reported to lead to the successful expression of relatively high defucosylated antibodies bearing the biantennary complex-type of Fc oligosaccharides such as control of culture osmolality (Konno *et al.,* (2012) *Cytotechnology* 64:249-65), application of a variant CHO line Lec13 as the host cell line (Shields *et al.,* (2002) *J Biol Chem* 277:26733-26740), application of a variant CHO line EB66 as the host cell line (Olivier *et al., MAbs* ;2(4), 2010; Epub ahead of print; PMID:20562582), application of a rat hybridoma cell line YB2/0 as the host cell line (Shinkawa *et al.,* (2003) *J Biol Chem* 278:3466-3473), introduction of small interfering RNA specifically against the α 1,6-fucosyltrasferase (*FUT8)* gene (Mori *et al.,* (2004) *Biotechnol Bioeng* 88:901-908), or coexpression of β-1,4-*N*-acetylglucosaminyltransferase III and Golgi α-mannosidase II or a potent alpha-mannosidase I inhibitor, kifunensine (Ferrara *et al.,* (2006) *J Biol Chem* 281:5032-5036, Ferrara *et al.,* (2006) *Biotechnol Bioeng* 93:851-861; Xhou *et al.,* (2008) *Biotechnol Bioeng* 99:652-65).

In some embodiments, the antibodies of the invention comprise at least one substitution in the antibody Fc that enhances effector function of the antibody.

In some embodiments, the antibodies of the invention comprise at least one substitution in the antibody Fc at amino acid position 236, 239, 243, 256, 267, 268, 290, 292, 298, 300, 305, 312, 324, 326, 330, 332, 333, 334, 345, 360, 339, 378, 396 or 430.

In some embodiments, the antibodies of the invention comprise at least one substitution in the antibody Fc selected from the group consisting of G236A, S239D, F243L, T256A, K290A, R292P, S298A, Y300L, V305L, K326A, A330K, I332E, E333A, K334A, A339T, P396L, S267E, H268F, S324T, K326A, K326W, E333A, E345K, E345Q, E345R, E345Y, E430S, E430F and E430T.

In some embodiments, the antibodies of the invention comprise at least one substitution in the antibody Fc selected from the group consisting of S239D/I332E, S298A/E333A/K334A, F243L/R292P/Y300L, F243L/R292P/Y300L/P396L, F243L/R292P/Y300L/V305I/P396L, G236A/S239D/I332E, K326A/E333A, K326W/E333A, H268F/S324T, S267E/H268F, S267E/S324T and S267E/H268F/S324T.

In some embodiments, the antibodies of the invention have a biantennary glycan structure with fucose content of about between 0% to about 15%, for example 15%, 14%, 13%, 12%, 11% 10%, 9%, 8%, 7%, 6%, 5%, 4%, 3%, 2%, 1% or 0%.

In some embodiments, the antibodies of the invention have a biantennary glycan structure with fucose content of about 50%, 40%, 45%, 40%, 35%, 30%, 25%, 20%, 15%, 14%, 13%, 12%, 11% 10%, 9%, 8%, 7%, 6%, 5%, 4%, 3%, 2%, 1% or 0%.

Substitutions in the Fc and reduced fucose content may enhance the ADCC activity of the antagonistic antibodies specifically binding TIM-3 or PD-1 of the invention. TIM-3 or PD-1 antibodies with enhanced ADCC, ADCP and/or CDC activity may be useful in the treatment of patients with TIM-3 and/or PD-1 expressing tumors, including heme malignancies.

"Fucose content" means the amount of the fucose monosaccharide within the sugar chain at Asn297. The relative amount of fucose is the percentage of fucose-containing structures related to all glycostructures. These may be characterized and quantified by multiple methods, for example: 1) using MALDI-TOF of N-glycosidase F treated sample (e.g. complex, hybrid and oligo- and high-mannose structures) as described in Intl. Patent Publ. No. WO2008/077546; 2) by enzymatic release of the Asn297 glycans with subsequent derivatization and detection/ quantitation by HPLC (UPLC) with fluorescence detection and/or HPLC-MS (UPLC-MS); 3) intact protein analysis of the native or reduced mAb, with or without treatment of the Asn297 glycans with Endo S or other enzyme that cleaves between the first and the second GlcNAc monosaccharides, leaving the fucose attached to the first GlcNAc; 4) digestion of the mAb to constituent peptides by enzymatic digestion (e.g., trypsin or endopeptidase Lys-C), and subsequent separation, detection and quantitation by HPLC-MS (UPLC-MS) or 5) separation of the mAb oligosaccharides from the mAb protein by specific enzymatic deglycosylation with PNGase F at Asn 297. The oligosaccharides released may be labeled with a fluorophore, separated and identified by various complementary techniques which allow fine characterization of the glycan structures by matrix-assisted laser desorption ionization (MALDI) mass spectrometry by comparison of the experimental masses with the theoretical masses, determination of the degree of sialylation by ion exchange HPLC (GlycoSep C), separation and quantification of the oligosaccharide forms according to hydrophilicity criteria by normal-phase HPLC (GlycoSep N), and separation and quantification of the oligosaccharides by high performance capillary electrophoresis-laser induced fluorescence (HPCE-LIF).

"Low fucose" or "low fucose content" refers to antibodies with fucose content of about 0% - 15%.

"Normal fucose" or 'normal fucose content" refers to antibodies with fucose content of about over 50%, typically about over 60%, 70%, 80% or over 85%.

The antibodies of the invention may be post-translationally modified by processes such as glycosylation, isomerization, deglycosylation or non-naturally occurring covalent modification such as the addition of polyethylene glycol moieties (pegylation) and lipidation. Such modifications may occur *in vivo* or *in vitro*. For example, the antibodies of the invention described herein may be conjugated to polyethylene glycol (PEGylated) to improve their pharmacokinetic profiles. Conjugation may be carried out by techniques known to those skilled in the art. Conjugation of therapeutic antibodies with PEG has been shown to enhance pharmacodynamics while not interfering with function (Knigh *et al.,* (2004) *Platelets* 15:409-18; Leong *et al.,* (2001) *Cytokine* 16:106-19; Yang *et al.,* (2003) *Protein Eng* 16:761-70).

Antibodies of the invention may be modified to improve stability, selectivity, cross-reactivity, affinity, immunogenicity or other desirable biological or biophysical property are within the scope of the invention. Stability of an antibody is influenced by a number of factors, including (1) core packing of individual domains that affects their intrinsic stability, (2) protein/protein interface interactions that have impact upon the HC and LC pairing, (3) burial of polar and charged residues, (4) H-bonding network for polar and charged residues; and (5) surface charge and polar residue distribution among other intra- and inter-molecular forces (Worn *et al.,* (2001) *J Mol Biol* 305:989-1010). Potential structure destabilizing residues may be identified based upon the crystal structure of the antibody or by molecular modeling in certain cases, and the effect of the residues on antibody stability may be tested by generating and evaluating variants harboring mutations in the identified residues. One of the ways to increase antibody stability is to raise the thermal transition midpoint (T_m) as measured by differential scanning calorimetry (DSC). In general, the protein T_m is correlated with its stability and inversely correlated with its susceptibility to unfolding and denaturation in solution and the degradation processes that depend on the tendency of the protein to unfold (Remmele *et al.,* (2000) *Biopharm* 13:36-46). A number of studies have found correlation between the ranking of the physical stability of formulations measured as thermal stability by DSC and physical stability measured by other methods (Gupta *et al.,* (2003) *AAPS PharmSci* 5E8; Zhang *et al.,* (2004) *J Pharm Sci* 93:3076-89; Maa *et al.,* (1996) *Int J Pharm* 140:155-68; Bedu-Addo *et al.*, (2004) *Pharm Res* 21:1353-61; Remmele *et al.*, (1997) Pharm Res 15:200-8).

Formulation studies suggest that a Fab T_m has implication for long-term physical stability of a corresponding mAb.

C-terminal lysine (CTL) may be removed from injected antibodies by endogenous circulating carboxypeptidases in the blood stream (Cai *et al.*, (2011) *Biotechnol Bioeng* 108:404-412). During manufacturing, CTL removal may be controlled to less than the maximum level by control of concentration of extracellular Zn^{2+}, EDTA or EDTA – Fe^{3+} as described in U.S. Patent Publ. No. US20140273092. CTL content in antibodies can be measured using known methods.

In some embodiments, the antibodies of the invention have a C-terminal lysine content of about 10% to about 90%, about 20% to about 80%, about 40% to about 70%, about 55% to about 70%, or about 60%.

In some embodiments, the antibodies of the invention have a C-terminal lysine content of about 0%, 5%, 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90% or 100%.

Methods of generating homologous antibodies, antibodies with conservative modifications, and engineered and modified antibodies

The antibodies of the invention that have altered amino acid sequences when compared to the parental antibodies may be generated using standard cloning and expression technologies. For example, site-directed mutagenesis or PCR-mediated mutagenesis may be performed to introduce the mutation(s) and the effect on antibody binding or other property of interest, may be evaluated using well known methods and the methods described herein in the Examples.

Antibody allotypes

The antibody of the invention may be an IgG1, IgG2, IgG3 or IgG4 isotype.

In some embodiments, the antibody of the invention is an IgG1 isotype.

In some embodiments, the antibody of the invention is an IgG2 isotype.

In some embodiments, the antibody of the invention is an IgG3 isotype.

In some embodiments, the antibody of the invention is an IgG4 isotype.

Immunogenicity of therapeutic antibodies is associated with increased risk of infusion reactions and decreased duration of therapeutic response (Baert *et al.*, (2003) *N Engl J Med* 348:602-08). The extent to which therapeutic antibodies induce an immune response in the host may be determined in part by the allotype of the antibody (Stickler *et al.*, (2011) *Genes and Immunity* 12:213-21). Antibody allotype is related to amino acid sequence variations at specific locations in the constant region sequences of the antibody.

Table 6 shows select IgG1, IgG2 and IgG4 allotypes.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention are of G2m(n), G2m(n-), G2m(n)/(n-), nG4m(a), G1m(17) or G1m(17,1) allotype.

In some embodiments, the antagonistic antibodies specifically binding TIM-3 of the invention are of G2m(n), G2m(n-), G2m(n)/(n-), nG4m(a), G1m(17) or G1m(17,1) allotype.

In some embodiments, the bispecific PD-1/TIM-3 antibodies of the invention are of G2m(n), G2m(n-), G2m(n)/(n-), nG4m(a), G1m(17) or G1m(17,1) allotype.

Table 6.

Allotype	Amino acid residue at position of diversity (residue numbering: EU Index)							
	IgG2		IgG4		IgG1			
	189	282	309	422	214	356	358	431
G2m(n)	T	M						
G2m(n-)	P	V						
G2m(n)/(n-)	T	V						
nG4m(a)			L	R				
G1m(17)					K	E	M	A
G1m(17,1)					K	D	L	A

Anti-idiotypic antibodies

The present invention provides an anti-idiotypic antibody binding to the antibody of the invention.

The invention also provides an anti-idiotypic antibody specifically binding to the anti-PD-1 antibody of the invention.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 50.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 51.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 52.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 53.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 49.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 54.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 50.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 55.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 56.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 57.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 44 and the VL of SEQ ID NO: 49.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 49.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 46 and the VL of SEQ ID NO: 49.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 49.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 53.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 52.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 59.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 61.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 62.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 63 and the VL of SEQ ID NO: 65.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

The invention also provides an anti-idiotypic antibody specifically binding the antagonistic antibody specifically binding TIM-3 of the invention.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 148 and the VL of SEQ ID NO: 157.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 147 and the VL of SEQ ID NO: 155.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 149 and the VL of SEQ ID NO: 158.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 150 and the VL of SEQ ID NO: 159.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 151 and the VL of SEQ ID NO: 160.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 152 and the VL of SEQ ID NO: 161.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 154 and the VL of SEQ ID NO: 163.

In some embodiments, the kit comprises the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

In some embodiments, the kit comprises the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

The invention also provides an anti-idiotypic antibody specifically binding the antibody comprising the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

In some embodiments, the anti-idiotypic antibody is used for detecting the level of the therapeutic antibodies (e.g. anti-PD-1, anti-TIM-3 or the bispecific PD-1/TIM-3 antibodies of the invention described herein) in a sample.

An anti-idiotypic (Id) antibody is an antibody which recognizes the antigenic determinants (*e.g.* the paratope or CDRs) of the antibody. The Id antibody may be antigen-blocking or non-blocking. The antigen-blocking Id may be used to detect the free antibody in a sample (e.g. anti-PD-1, anti-TIM-3 or the bispecific PD-1/TIM-3 antibody of the invention described herein). The non-blocking Id may be used to detect the total antibody (free, partially bond to antigen, or fully bound to antigen) in a sample. An Id antibody may be prepared by immunizing an animal with the antibody to which an anti-Id is being prepared.

An anti-Id antibody may also be used as an immunogen to induce an immune response in yet another animal, producing a so-called anti-anti-Id antibody. An anti-anti-Id may be epitopically identical to the original mAb, which induced the anti-Id. Thus, by using antibodies to the idiotypic determinants of a mAb, it is possible to identify other clones expressing antibodies of identical specificity. Anti-Id antibodies may be varied (thereby producing anti-Id antibody variants) and/or derivatized by any suitable technique, such as those described elsewhere herein with respect to the antibodies specifically binding PD-1 or TIM-3, or the bispecific PD-1/TIM-3 antibodies.

Immunoconjugates

An "immunoconjugate" refers to the antibody of the invention conjugated to one or more heterologous molecule(s).

In some embodiments, the antibody of the invention is conjugated to one or more cytotoxic agents or an imaging agent.

Exemplary cytotoxic agents include chemotherapeutic agents or drugs, growth inhibitory agents, toxins (e.g., protein toxins, enzymatically active toxins of bacterial, fungal, plant, or animal origin, or fragments thereof), and radionuclides.

The cytotoxic agent may be one or more drugs, such as to a mayatansinoid (see, e.g., U.S. Patent No. 5,208,020, 5,416,06), an auristatin such as monomethylauristatin drug moieties DE and DF (MMAE and MMAF) (see, e.g., U.S. Patent Nos. 5,635,483 and

5,780,588, and 7,498,298), a dolastatin, a calicheamicin or derivative thereof (see, e.g., U.S. Patent Nos. 5,712,374, 5,714,586, 5,739, 116, 5,767,285, 5,770,701, 5,770,710, 5,773,001, and 5,877,296; Hinman *et al.,* (1993) *Cancer Res* 53:3336-3342; and Lode *et al.,*(1998) *Cancer Res* 58:2925-2928); an anthracycline such as daunomycin or doxorubicin (see, e.g., Kratz *et al.,* (2006) *Current Med. Chem* 13:477-523; Jeffrey *et al.,* (2006) Bioorganic & *Med Chem Letters* 16:358-362; Torgov *et al.,* (2005) *Bioconj Chem* 16:717-721; Nagy *et* al., (2000) *Proc Natl Acad Sci* USA 97:829-834; Dubowchik et al, Bioorg. & Med. Chem. Letters 12: 1529-1532 (2002); King *et al.,* (2002) *J Med Chem* 45:4336-4343; and U.S. Patent No. 6,630,579), methotrexate, vindesine, a taxane such as docetaxel, paclitaxel, larotaxel, tesetaxel, and ortataxel.

The cytotoxic agent may also be an enzymatically active toxin or fragment thereof, such as diphtheria A chain, nonbinding active fragments of diphtheria toxin, exotoxin A chain (from Pseudomonas aeruginosa), ricin A chain, modeccin A chain, alpha-sarcin, Aleurites fordii proteins, dianthins, Phytolacca americana proteins (PAPI, PAPII, and PAP-S), momordica charantia inhibitor, curcin, crotin, sapaonaria officinalis inhibitor, gelonin, mitogellin, restrictocin, phenomycin, enomycin, and the tricothecenes.

The cytotoxic agent or an imaging agent may also be a radionuclide. Exemplary radionuclides include Ac-225, At-211, 1-131, I-125, Y-90, Re-186, Re-188, Sm-153, Bi-212, P-32, Pb-212 and radioactive isotopes of Lu. When the radioconjugate is used for detection, it may comprise a radioactive atom for scintigraphic studies, for example Tc-99m or I-123, or a spin label for nuclear magnetic resonance (NMR) imaging (also known as magnetic resonance imaging, mri), such as I-123, I-131, In-111, F-19, C-13, N-15 or O-17.

Conjugates of the antibodies of the invention and the heterologous molecule may be made using a variety of bifunctional protein coupling agents such as N-succinimidyl-3-(2-pyridyldithio) propionate (SPDP), succinimidyl-4-(N-maleimidomethyl) cyclohexane-l-carboxylate (SMCC), iminothiolane (IT), bifunctional derivatives of imidoesters (such as dimethyl adipimidate HQ), active esters (such as disuccinimidyl suberate), aldehydes (such as glutaraldehyde), bis-azido compounds (such as bis (p-azidobenzoyl) hexanediamine), bis-diazonium derivatives (such as bis-(p-diazoniumbenzoyl)-ethylenediamine), diisocyanates (such as toluene 2,6-diisocyanate), and bis-active fluorine compounds (such as l,5-difluoro-2,4-dinitrobenzene). For example, a ricin immunotoxin may be prepared as described in Vitetta *et al.,* (1987) *Science* 238: 1098. Carbon- 14-labeled l-isothiocyanatobenzyl-3-methyldiethylene triaminepentaacetic acid (MX- DTPA) is an exemplary chelating agent for conjugation of radionucleotide to the antibody. See,

e.g., W094/11026. The linker may be a "cleavable linker" facilitating release of a cytotoxic drug in the cell. For example, an acid-labile linker, peptidase-sensitive linker, photolabile linker, dimethyl linker or disulfide-containing linker (Chari *et al.,* (1992) *Cancer Res* 52: 127-131; U.S. Patent No. 5,208,020) may be used.

Conjugates of the antibodies of the invention and the heterologous molecule may be prepared with cross-linker reagents such as BMPS, EMCS, GMBS, HBVS, LC-SMCC, MBS, MPBH, SBAP, SIA, SIAB, SMCC, SMPB, SMPH, sulfo-EMCS, sulfo-GMBS, sulfo-KMUS, sulfo-MBS, sulfo-SIAB, sulfo-SMCC, and sulfo- SMPB, and SVSB (succinimidyl-(4-vinylsulfone)benzoate) which are commercially available (e.g., from Pierce Biotechnology, Inc., Rockford, IL., U.S.A).

The invention also provides an immunoconjugate comprising the antagonistic antibody specifically binding PD-1 of the invention linked to a therapeutic agent or an imaging agent.

The invention also provides an immunoconjugate comprising the antagonistic antibody specifically binding TIM-3 of the invention linked to a therapeutic agent or an imaging agent.

The invention also provides an immunoconjugate comprising the bispecific PD-1/TIM-3 antibody of the invention linked to a therapeutic agent or an imaging agent.

Generation of monospecific antibodies of the invention

In some embodiments, the antibodies of the invention are human.

In some embodiments, the antibodies of the invention are humanized.

Monospecific antibodies of the invention described herein (e.g. antibodies specifically binding PD-1 or TIM-3) may be generated using various technologies. For example, the hybridoma method of Kohler and Milstein, *Nature* 256:495, 1975 may be used to generate monoclonal antibodies. In the hybridoma method, a mouse or other host animal, such as a hamster, rat or monkey, is immunized with human or cyno PD-1 or TIM-3 or fragments of PD-1 or TIM-3, such as the extracellular domain of PD-1 or TIM-3, followed by fusion of spleen cells from immunized animals with myeloma cells using standard methods to form hybridoma cells (Goding, Monoclonal Antibodies: Principles and Practice, pp.59-103 (Academic Press, 1986)). Colonies arising from single immortalized hybridoma cells are screened for production of antibodies with desired properties, such as specificity of binding, cross-reactivity or lack thereof, and affinity for the antigen.

Various host animals may be used to produce the antibodies of the invention. For example, Balb/c mice may be used to generate mouse anti-human PD-1 or TIM-3 antibodies. The antibodies made in Balb/c mice and other non-human animals may be humanized using various technologies to generate more human-like sequences.

Exemplary humanization techniques including selection of human acceptor frameworks are known and include CDR grafting (U.S. Patent No. 5,225,539), SDR grafting (U.S. Patent No. 6,818,749), Resurfacing (Padlan, (1991) *Mol Immunol* 28:489-499), Specificity Determining Residues Resurfacing (U.S. Patent Publ. No. 2010/0261620), human framework adaptation (U.S. Patent No. 8,748,356) or superhumanization (U.S. Patent No. 7,709, 226). In these methods, CDRs of parental antibodies are transferred onto human frameworks that may be selected based on their overall homology to the parental frameworks, based on similarity in CDR length, or canonical structure identity, or a combination thereof.

Humanized antibodies may be further optimized to improve their selectivity or affinity to a desired antigen by incorporating altered framework support residues to preserve binding affinity (backmutations) by techniques such as those described in Int. Patent Publ. Nos. WO1090/007861 and WO1992/22653, or by introducing variation at any of the CDRs for example to improve affinity of the antibody.

Transgenic animals, such as mice or rats carrying human immunoglobulin (Ig) loci in their genome may be used to generate human antibodies against a target protein, and are described in for example U.S. Patent No. 6,150,584, Int. Patent Publ. No. WO99/45962, Int. Patent Publ. Nos. WO2002/066630, WO2002/43478, WO2002/043478 and WO1990/04036, Lonberg *et al* (1994) *Nature* 368:856-9; Green *et al* (1994) *Nature Genet.* 7:13-21; Green & Jakobovits (1998) *Exp. Med.* 188:483-95; Lonberg and Huszar (1995) *Int Rev Immunol* 13:65-93; Bruggemann *et al.,* (1991) *Eur J Immunol* 21:1323- 1326; Fishwild *et al.,* (1996) *Nat Biotechnol* 14:845-851; Mendez *et al.,* (1997) *Nat Genet* 15:146-156; Green (1999) *J Immunol Methods* 231:11-23; Yang *et al.,* (1999) *Cancer Res* 59:1236-1243; Brüggemann and Taussig (1997) *Curr Opin Biotechnol* 8:455-458. The endogenous immunoglobulin loci in such animal may be disrupted or deleted, and at least one complete or partial human immunoglobulin locus may be inserted into the genome of the antimal using homologous or non-homologous recombination, using transchromosomes, or using minigenes. Companies such as Regeneron (http://_www_regeneron_com), Harbour Antibodies (http://_www_harbourantibodies_com), Open Monoclonal Technology, Inc. (OMT) (http://_www_omtinc_net), KyMab (http://_www_kymab_com), Trianni

(http://_www.trianni_com) and Ablexis (http://_www_ablexis_com) may be engaged to provide human antibodies directed against a selected antigen using technologies as described above.

Human antibodies may be selected from a phage display library, where the phage is engineered to express human immunoglobulins or portions thereof such as Fabs, single chain antibodies (scFv), or unpaired or paired antibody variable regions (Knappik *et al.*, (2000) *J Mol Biol* 296:57-86; Krebs *et al.*, (2001) *J Immunol Meth* 254:67-84; Vaughan *et al.*, (1996) *Nature Biotechnology* 14:309-314; Sheets *et al.*, (1998) *PITAS (USA)* 95:6157-6162; Hoogenboom and Winter (1991) *J Mol Biol* 227:381; Marks *et al.*, (1991) *J Mol Biol* 222:581). The antibodies of the invention may be isolated for example from phage display library expressing antibody heavy and light chain variable regions as fusion proteins with bacteriophage pIX coat protein as described in Shi *et al.*, (2010) *J Mol Biol* 397:385-96, and Int. Patent Publ. No. WO09/085462). The libraries may be screened for phage binding to human and/or cyno PD-1 or TIM-3 and the obtained positive clones may be further characterized, the Fabs isolated from the clone lysates, and expressed as full length IgGs. Such phage display methods for isolating human antibodies are described in for example: U.S. Patent Nos. 5,223,409, 5,403,484, 5,571,698, 5,427,908, 5, 580,717, 5,969,108, 6,172,197, 5,885,793; 6,521,404; 6,544,731; 6,555,313; 6,582,915 and 6,593,081.

Preparation of immunogenic antigens and monoclonal antibody production may be performed using any suitable technique, such as recombinant protein production. The immunogenic antigens may be administered to an animal in the form of purified protein, or protein mixtures including whole cells or cell or tissue extracts, or the antigen may be formed *de novo* in the animal's body from nucleic acids encoding said antigen or a portion thereof.

Generation of bispecific PD-1/TIM-3 antibodies of the invention

The bispecific PD-1/TIM-3 antibodies of the invention (*e.g.* the bispecific antibodies comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3) may be generated by combining PD-1 binding VH/VL domains with TIM-3 binding VH/VL domains isolated and characterized herein. Alternatively, the bispecific PD-1/TIM-3 antibodies may be engineered using VH/VL domains from publicly available monospecific anti-PD-1 and anti-TIM-3 antibodies, and/or by mix-matching the PD-1 or TIM-3 binding VH/VL domains identified herein with publicly available PD-1 or TIM-3 binding VH/VL domains.

Exemplary anti-PD-1 antibodies that may be used to engineer bispecific PD-1/TIM-3 molecules are for example those described in U.S. Patent Nos. 5,897,862 and 7,488,802, and in Int. Patent Publ. Nos. WO2004/004771, WO2004/056875, WO2006/121168, WO2008/156712, WO2010/029435, WO2010/036959, WO2011/110604, WO2012/145493, WO2014/194302, WO2014/206107, WO2015/036394, WO2015/035606, WO2015/085847, WO2015/112900 and WO2015/112805. For example, the VH/VL domains of KEYTRUDA® (pembrolizumab) and OPDIVO® (nivolumab) may be used. These PD-1 VH/VL domains may be incorporated into bispecific antibodies comprising TIM-3 binding VH/VL domains described herein and in **Table 3**. For example, the VH/VL domains of the TIM-3 antibodies TM3B103, TM3B105, TM3B107, TM3B108, TM3B109, TM3B113, TM3B189, TM3B190 and TM3B196 described herein may be used to generate bispecific PD-1/TIM-3 antibodies.

Similarly, exemplary anti-TIM-3 antibodies that may be used to engineer bispecific PD-1/TIM-3 molecules are for example those described in Int. Patent Publ. Nos. WO2011/155607, WO2013/006490, and WO2015/117002. These TIM-3 VH/VL domains may be incorporated into bispecific antibodies comprising PD-1 binding VH/VL domains described herein and in **Table 2**. For example, the VH/VL domains of the PD-1 antibodies PD1B114, PD1B149, PD1B160, PD1B162, PD1B164, PD1B11, PD1B183, PD1B184, PD1B185, PD1B187, PD1B192, PD1B71, PD1B177, PD1B70, PD1B175, PD1B194, PD1B195, PD1B196, PD1B197, PD1B198, PD1B199, PD1B200, PD1B201, PD1B131 and PD1B132 described herein may be used to generate bispecific PD-1/TIM-3 antibodies.

The generated bispecific PD-1/TIM-3 antibodies may be tested for their binding to PD-1 and TIM-3, and for their desired functional characteristics, such as enhancement of activation of antigen specific $CD4^+$ and $CD4^+$ T cells using methods described herein.

Bispecific antibodies of the invention comprise antibodies having a full length antibody structure.

Full length bispecific antibodies may be generated for example using Fab arm exchange (e.g., half molecule exchange, exchanging on heavy chain – light chain pair) between two monospecific bivalent antibodies by introducing mutations at the heavy chain CH3 interface in each half-molecule to favor heterodimer formation of two antibody half-molecules having distinct specificity either *in vitro* in cell-free environment or using co-expression. The Fab arm exchange reaction is the result of a disulfide-bond isomerization reaction and dissociation-association of CH3 domains. The heavy chain disulfide bonds in

the hinge regions of the parental monospecific antibodies are reduced. The resulting free cysteines of one of the parental monospecific antibodies form an inter heavy-chain disulfide bond with cysteine residues of a second parental monospecific antibody molecule and simultaneously CH3 domains of the parental antibodies release and reform by dissociation-association. The CH3 domains of the Fab arms may be engineered to favor heterodimerization over homodimerization. The resulting product is a bispecific antibody having two Fab arms r half molecules which each bind a distinct epitope. Mutations F405L in one heavy chain and K409R in the other heavy chain may be used in case of IgG1 antibodies. For IgG2 antibodies, a wild-type IgG2 and a IgG2 antibody with F405L and R409K substitutions may be used. To generate bispecific antibodies, first monospecific bivalent antibody and the second monospecific bivalent antibody are engineered to have a F405L or a K409R mutation in the Fc region, the antibodies are incubated together under reducing conditions sufficient to allow the cysteines in the hinge region to undergo disulfide bond isomerization; thereby generating the bispecific antibody by Fab arm exchange. The incubation conditions may optimally be restored to non-reducing. Exemplary reducing agents that may be used are 2- mercaptoethylamine (2-MEA), dithiothreitol (DTT), dithioerythritol (DTE), glutathione, tris(2 carboxyethyl)phosphine (TCEP), L-cysteine and beta- mercaptoethanol. For example, incubation for at least 90 min at a temperature of at least 20°C in the presence of at least 25 mM 2-MEA or in the presence of at least 0.5 mM dithiothreitol at a pH of from 5-8, for example at pH of 7.0 or at pH of 7.4 may be used.

Bispecific antibodies may also be generated using designs such as the Knob-in-Hole (Genentech), CrossMAbs (Roche) and the electrostatically-matched (Chugai, Amgen, NovoNordisk, Oncomed), the LUZ-Y (Genentech), the Strand Exchange Engineered Domain body (SEEDbody)(EMD Serono), and the Biclonic (Merus).

The "knob-in-hole" strategy (see, e.g., Intl. Publ. No. WO 2006/028936) may be used to generate full length bispecific antibodies of the invention. Briefly, selected amino acids forming the interface of the CH3 domains in human IgG can be mutated at positions affecting CH3 domain interactions to promote heterodimer formation. An amino acid with a small side chain (hole) is introduced into a heavy chain of an antibody specifically binding a first antigen and an amino acid with a large side chain (knob) is introduced into a heavy chain of an antibody specifically binding a second antigen. After co-expression of the two antibodies, a heterodimer is formed as a result of the preferential interaction of the heavy chain with a "hole" with the heavy chain with a "knob". Exemplary CH3 substitution pairs forming a knob and a hole are (expressed as modified position in the first

CH3 domain of the first heavy chain/ modified position in the second CH3 domain of the second heavy chain): T366Y/F405A, T366W/F405W, F405W/Y407A, T394W/Y407T, T394S/Y407A, T366W/T394S, F405W/T394S and T366W/T366S_L368A_Y407V.

The CrossMAb technology may be used to generate full length bispecific antibodies of the invention. CrossMAbs, in addition to utilizing the "knob-in-hole" strategy to promoter Fab arm exchange, have in one of the half arms the CH1 and the CL domains exchanged to ensure correct light chain pairing of the resulting bispecific antibody (see e.g. U.S. Patent No. 8,242,247).

Other cross-over strategies may be used to generate full length bispecific antibodies of the invention by exchanging variable or constant, or both domains between the heavy chain and the light chain or within the heavy chain in the bispecific antibodies, either in one or both arms. These exchanges include for example VH-CH1 with VL-CL, VH with VL, CH3 with CL and CH3 with CH1 as described in Int. Patent Publ. Nos. WO2009/080254, WO2009/080251, WO2009/018386 and WO2009/080252.

Other strategies such as promoting heavy chain heterodimerization using electrostatic interactions by substituting positively charged residues at one CH3 surface and negatively charged residues at a second CH3 surface may be used, as described in US Patent Publ. No. US2010/0015133; US Patent Publ. No. US2009/0182127; US Patent Publ. No. US2010/028637 or US Patent Publ. No. US2011/0123532. In other strategies, heterodimerization may be promoted by following substitutions (expressed as modified positions in the first CH3 domain of the first heavy chain/ modified position in the second CH3 domain of the second heavy chain): L351Y_F405A_Y407V/T394W, T366I_K392M_T394W/F405A_Y407V, T366L_K392M_T394W/F405A_Y407V, L351Y_Y407A/T366A_K409F, L351Y_Y407A/T366V_K409F, Y407A/T366A_K409F, or T350V_L351Y_F405A_Y407V/T350V_T366L_K392L_T394W as described in U.S. Patent Publ. No. US2012/0149876 or U.S. Patent Publ. No. US2013/0195849.

LUZ-Y technology may be utilized to generate bispecific antibodies of the invention. In this technology, a leucine zipper is added into the C terminus of the CH3 domains to drive the heterodimer assembly from parental mAbs that is removed post-purification as described in Wranik *et al.,* (2012) *J Biol Chem* 287(52): 42221-9.

SEEDbody technology may be utilized to generate bispecific antibodies of the invention. SEEDbodies have, in their constant domains, select IgG residues substituted with IgA residues to promote heterodimerization as described in U.S. Patent No. US20070287170.

Mutations are typically made at the DNA level to a molecule such as the constant domain of the antibody using standard methods.

The antibodies of the invention may be engineered into various well known antibody formats.

In some embodiments, the bispecific antibodies include recombinant IgG-like dual targeting molecules, wherein the two sides of the molecule each contain the Fab fragment or part of the Fab fragment of at least two different antibodies; IgG fusion molecules, wherein full length IgG antibodies are fused to an extra Fab fragment or parts of Fab fragment; Fc fusion molecules, wherein single chain Fv molecules or stabilized diabodies are fused to heavy-chain constant-domains, Fc-regions or parts thereof; Fab fusion molecules, wherein different Fab-fragments are fused together; ScFv- and diabody-based and heavy chain antibodies (e.g., domain antibodies, nanobodies) wherein different single chain Fv molecules or different diabodies or different heavy-chain antibodies (e.g. domain antibodies, nanobodies) are fused to each other or to another protein or carrier molecule.

Polynucleotides, vectors and host cells

The invention also provides an antagonistic antibody that specifically binds PD-1, TIM-3 or PD-1 and TIM-3 having certain VH and VL sequences, wherein the antibody VH is encoded by a first polynucleotide and the antibody VL is encoded by a second polynucleotide. The polynucleotide may be a complementary deoxynucleic acid (cDNA), and may be codon optimized for expression in suitable host. Codon optimization is a well-known technology.

The invention also provides an isolated polynucleotide encoding the VH of the antibody of the invention, the VL of the antibody of the invention, the heavy chain of the antibody of the invention or the light chain of the antibody of the invention.

The invention also provides an isolated polynucleotide encoding the VH, the VL, or the VH and the VL of the antagonistic antibody specifically binding PD-1 of the invention.

The invention also provides an isolated polynucleotide encoding the VH of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47, 48, 63 or 64.

The invention also provides an isolated polynucleotide encoding the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62 or 65.

The invention also provides an isolated polynucleotide comprising the polynucleotide sequence of SEQ ID NOs: 196, 197, 198, 199, 200, 201, 202 or 203.

The invention also provides an isolated polynucleotide encoding the VH, the VL, or the VH and the VL of the antagonistic antibody specifically binding TIM-3 of the invention.

The invention also provides an isolated polynucleotide encoding the VH of SEQ ID NOs: 145, 146, 147, 148, 149, 150, 151, 152, 153, 154 or 172.

The invention also provides an isolated polynucleotide encoding the VL of SEQ ID NOs: 155, 156, 157, 158, 159, 160, 161, 162, 163 or 173.

The invention also provides an isolated polynucleotide comprising the polynucleotide sequence of SEQ ID NOs: 204, 205, 206, 207, 208, 209, 210 or 211.

The invention also provides an isolated polynucleotide encoding the HC1, the LC1, the HC2 or the LC2 of the antagonistic bispecific PD-1/TIM-3 antibody of the invention.

The invention also provides an isolated polynucleotide encoding the HC1 of SEQ ID NOs: 186, 187, 241, 242 or 243.

The invention also provides an isolated polynucleotide encoding the LC1 of SEQ ID NOs: 188 or 189.

The invention also provides an isolated polynucleotide encoding the HC2 of SEQ ID NOs: 190, 191, 192, 244, 245, 246, 247 or 248.

The invention also provides an isolated polynucleotide encoding the LC2 of SEQ ID NOs: 193, 194 or 195.

The invention also provides an isolated polynucleotide comprising the polynucleotide sequence of SEQ ID NOs: 253, 254, 255, 256, 257, 258, 259 and 260.

SEQ ID NO: 196 (PD1H170)

CAGGTGCAGCTGGTGCAGAGCGGCGCGGAAGTGAAAAAACCGGGCAGCAGCG
TGAAAGTGAGCTGCAAAGCGAGCGGCGGCACCTTTAGCAGCTATGCGATTAG
CTGGGTGCGCCAGGCGCCGGGCCAGGGCCTGGAATGGATGGGCGGCATTATT
CCGATTTTTGACACCGCGAACTATGCGCAGAAATTTCAGGGCCGCGTGACCAT
TACCGCGGATGAAAGCACCAGCACCGCGTATATGGAACTGAGCAGCCTGCGC
AGCGAAGATACCGCGGTGTATTATTGCGCGCGCCCTGGTCTCGCTGCGGCTTA
TGATACTGGTTCCTTGGACTATTGGGGCCAGGGCACCCTGGTGACCGTGAGCA
GC

SEQ ID NO: 197 (PD1L148)

GAAATTGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGAGCCCGGGCGAAC
GCGCGACCCTGAGCTGCCGCGCGAGCCAGAGCGTTCGCTCCTACCTGGCGTGG
TATCAGCAGAAACCGGGCCAGGCGCCGCGCCTGCTGATCTACGACGCGAGCA
ATCGTGCGACCGGCATTCCGGCGCGCTTTAGCGGCTCCGGTAGCGGCACCGAT
TTTACCCTGACCATTAGCAGCCTGGAACCGGAAGATTTTGCGGTGTATTATTGC
CAGCAACGTAATTATTGGCCGCTGACCTTTGGCCAGGGCACCAAAGTGGAAAT
TAAA

SEQ ID NO: 198 (PD1H129)
GAAGTGCAGCTGGTGGAATCTGGCGGCGGACTGGTGCAGCCTGGCGGATCTCT
GAGACTGAGCTGTGCCGCCAGCGGCTTCGCCTTCAGCAGATACGACATGAGCT
GGGTGCGCCAGGCCCCTGGCAAAGGACTGGAAAGCGTGGCCTACATCTCTGG
CGGAGGCGCCAACACCTACTACCTGGACAACGTGAAGGGCCGGTTCACCATC
AGCCGGGACAACGCCAAGAACAGCCTGTACCTGCAGATGAACTCCCTGCGGG
CCGAGGACACCGCCGTGTACTATTGCGCCTCCCCCTACCTGAGCTACTTCGAC
GTGTGGGGCCAGGGCACACTCGTGACCGTGTCATCT

SEQ ID NO: 199 (PD1L62)
GAGATCGTGATGACCCAGAGCCCTGCCACCCTGTCCGTGTCTCCAGGCGAAAG
AGCCACCCTGAGCTGCAGAGCCAGCCAGAGCCTGAGCGACTACCTGCACTGGT
ATCAGCAGAAGCCCGGCCAGGCCCCCAGACTGCTGATCAAGTCTGCCAGCCA
GTCCATCAGCGGCATCCCCGCCAGATTTTCTGGCAGCGGCTCCGGCACCGAGT
TCACCCTGACAATCAGCAGCCTGCAGAGCGAGGACTTCGCCGTGTACTACTGC
CAGAACGGCCACAGCTTCCCTTACACCTTCGGCCAGGGCACCAAGCTGGAAAT
CAAG

SEQ ID NO: 200 (PD1H163)
CAGGTGCAGCTGGTGCAGAGCGGCGCGGAAGTGAAAAAACCGGGCAGCAGCG
TGAAAGTGAGCTGCAAAGCGAGCGGCGGCACCTTCAAGTCCTATGTGATTCAT
TGGGTGCGCCAGGCGCCGGGCCAGGGCCTGGAATGGATGGGCGGTATTATCC
CAATTTTTGGCACCGCCAATTATGCGCAGAAATTTCAGGGCCGCGTGACCATT
ACCGCTGATGAAAGCACCAGCACCGCGTATATGGAACTGAGCAGCCTGCGCA
GCGAAGATACCGCGGTGTATTATTGCGCGCGCGGTTATGTGCGGGCTACGGGC
ATGTTGGACTATTGGGGCCAGGGCACCCTGGTGACCGTGAGCAGC

SEQ ID NO: 201 (PD1L185)
GAAATTGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGAGCCCGGGCGAAC
GCGCGACCCTGAGCTGCCGCGCGAGCCAGAGCGTTAGCAATTATCTGGCGTGG
TATCAGCAGAAACCGGGCCAGGCGCCGCGCCTGCTGATCTACGACGCCAGCA
ATCGCGCGACCGGCATTCCGGCGCGCTTTAGCGGCTCCGGTAGCGGCACCGAT
TTTACCCTGACCATTAGCAGCCTGGAACCGGAAGATTTTGCGGTGTATTATTGC
CAGCAACGTGCATATTGGCCGCTGACCTTTGGCCAGGGCACCAAAGTGGAAAT
TAAA

SEQ ID NO: 202 (PD1H164)
CAGGTGCAGCTGGTGCAGAGCGGCGCGGAAGTGAAAAAACCGGGCAGCAGCG
TGAAAGTGAGCTGCAAAGCGAGCGGCGGCACCTTCAGCGATTATGTGATTTCC
TGGGTGCGCCAGGCGCCGGGCCAGGGCCTGGAATGGATGGGCGGTATTATCC
CGATTTACGGGACCGCTAACTATGCGCAGAAATTTCAGGGCCGCGTGACCATT
ACCGCTGATGAAAGCACCAGCACCGCGTATATGGAACTGAGCAGCCTGCGCA
GCGAAGATACCGCGGTGTATTATTGCGCGCGCGGTACCCTCGACCGGACCGGG
CATTTGGACTATTGGGGCCAGGGCACCCTGGTGACCGTGAGCAGC

SEQ ID NO: 203 (PD1L86)
GAAATTGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGAGCCCGGGCGAAC
GCGCGACCCTGAGCTGCCGCGCGAGCCAGAGCGTCTCCTCCTACCTTGCGTGG
TATCAGCAGAAACCGGGCCAGGCGCCGCGCCTGCTGATCCACGACGCCTCTAC
GCGTGCGACCGGCATTCCGGCGCGCTTTAGCGGCTCCGGTAGCGGCACCGATT
TTACCCTGACCATTAGCAGCCTGGAACCGGAAGATTTTGCGGTGTATTATTGC
CAGCAACGTAATTATTGGCCGCTCACCTTTGGCCAGGGCACCAAAGTGGAAAT
TAAA

SEQ ID NO: 204 (TM3H24)
GAAGTGCAGCTGCTGGAAAGCGGCGGCGGCCTGGTGCAGCCGGGCGGCAGCC
TGCGCCTGAGCTGCGCGGCAAGCGGCTTTACCTTTAGCAGCTATGCGATGAGC
TGGGTGCGCCAGGCGCCGGGCAAAGGCCTGGAATGGGTGAGCGCGATTAGCG
GCAGCGGCGGCAGCACCTATTATGCGGATAGCGTGAAAGGCCGCTTTACCATT
AGCCGCGATAACAGCAAAAACACCCTGTATCTGCAGATGAACAGCCTGCGCG
CGGAAGATACCGCGGTGTATTATTGCGCGAAATCCCCGTACGCGCCCTTGGAC
TATTGGGGCCAGGGCACCCTGGTGACCGTGAGCAGC

SEQ ID NO: 205 (TM3L33)
GAAATTGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGAGCCCGGGCGAAC
GCGCGACCCTTAGCTGCCGTGCAAGTCAGAGTGTGAACGACTACCTGGCGTGG
TATCAGCAGAAACCGGGCCAGGCGCCGCGCCTGCTGATTTATGATGCGAGCAA
CCGCGCGACCGGCATTCCGGCGCGCTTTAGCGGCAGCGGCAGCGGCACCGATT
TTACCCTGACCATTAGCAGCCTGGAACCGGAAGATTTTGCGGTGTATTATTGC
CAGCAGGGTGGTCACGCGCCGATCACCTTTGGCCAGGGCACCAAAGTGGAAA
TTAAA

SEQ ID NO: 206 (TM3H162)
GAAGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAGCCTGGCGAGAGCC
TGAAGATCAGCTGCAAGGGCAGCGGCTACAGCTTCACCAGCTACTGGATGCA
GTGGGTGCGCCAGATGCCTGGCAAGGGCCTGGAATGGATGGGCGCCATCTATC
CCGGCGACGGCGACATCAGATACACCCAGAACTTCAAGGGCCAAGTGACCAT
CAGCGCCGACAAGAGCATCAGCACCGCCTACCTGCAGTGGTCCAGCCTGAAG
GCCAGCGACACCGCCATGTACTACTGTGCCAGATGGGAGAAGTCCACCACCGT
GGTGCAGCGGAACTACTTCGACTACTGGGGCCAGGGCACCACAGTGACCGTGT
CTAGT

SEQ ID NO: 207 (TM3L85)
GACATCCAGATGACCCAGAGCCCCAGCAGCCTGTCTGCCAGCGTGGGCGACA
GAGTGACCATCACATGCAAGGCCAGCGAGAACGTGGGCACCTTCGTGTCCTGG
TATCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGGCGCCAGCA
ACAGATACACCGGCGTGCCCAGCAGATTCAGCGGCTCTGGCAGCGGCACCGA
CTTCACCCTGACCATCTCTAGCCTGCAGCCCGAGGACTTCGCCACCTACTACTG
CGGCCAGAGCTACAGCTACCCCACCTTTGGCCAGGGCACCAAGCTGGAAATCA
AG

SEQ ID NO: 208 (TM3H21)
GAAGTGCAGCTGCTGGAAAGCGGCGGCGGCCTGGTGCAGCCGGGCGGCAGCC
TGCGCCTGAGCTGCGCGGCGAGCGGCTTTACCTTTAGCAACTATTGGATGAGC
TGGGTGCGCCAGGCGCCGGGCAAAGGCCTGGAATGGGTGAGCGCGATTAGCG
GCAGCGGCGGCAGCACCTATTATGCGGATAGCGTGAAAGGCCGCTTTACCATT
AGCCGCGATAACAGCAAAAACACCCTGTATCTGCAGATGAACAGCCTGCGCG

CGGAAGATACCGCGGTGTATTATTGCGCGAAAGATCATTGGGATCCCAATTTT
TTGGACTATTGGGGCCAGGGCACCCTGGTGACCGTGAGCAGC

SEQ ID NO: 209 (PH9L1)
GAAATTGTGCTGACCCAGAGCCCGGGCACCCTGAGCCTGAGCCCGGGCGAAC
GCGCGACCCTGAGCTGCCGCGCGAGCCAGAGCGTGAGCAGCAGCTATCTGGC
GTGGTATCAGCAGAAACCGGGCCAGGCGCCGCGCCTGCTGATTTATGGCGCGA
GCAGCCGCGCGACCGGCATTCCGGATCGCTTTAGCGGCAGCGGCAGCGGCAC
CGATTTTACCCTGACCATTAGCCGCCTGGAACCGGAAGATTTTGCGGTGTATT
ATTGCCAGCAGTATGGCAGCAGCCCGCTGACCTTTGGCCAGGGCACCAAAGTG
GAAATTAAA

SEQ ID NO: 210 (TM3H65)
GAAGTGCAGCTGCTGGAAAGCGGCGGCGGCCTGGTGCAGCCGGGCGGCAGCC
TGCGCCTGAGCTGCGCGGCGAGCGGCTTTACCTTTAGCGACTATTGGATGAGC
TGGGTGCGCCAGGCGCCGGGCAAAGGCCTGGAATGGGTGAGCGTGATCAAGT
ATAGCGGTGGCTCCAAATATTATGCGGATAGCGTGAAAGGCCGCTTTACCATT
AGCCGCGATAACAGCAAAAACACCCTGTATCTGCAGATGAACAGCCTGCGCG
CGGAAGATACCGCGGTGTATTATTGCGCGAAAGAGCTGGAGGGGGTGTTCGA
CTATTGGGGCCAGGGCACCCTGGTGACCGTGAGCAGC

SEQ ID NO: 211 (TM3L12)
GAAATTGTGCTGACCCAGAGCCCGGGCACCCTGAGCCTGAGCCCGGGCGAAC
GCGCGACCCTGAGCTGCCGCGCGAGCCAGAGCGTTAGCAATAGCACTCTGGC
GTGGTATCAGCAGAAACCGGGCCAGGCGCCGCGCCTGCTGATTTATACTGCGA
GCAGCCGCGCGACCGGCATTCCGGATCGCTTTAGCGGCAGCGGCAGCGGCAC
CGATTTTACCCTGACCATTAGCCGCCTGGAACCGGAAGATTTTGCGGTGTATT
ATTGCCAGCAGTCTTACACATCTCCGTGGACTTTTGGCCAGGGCACCAAAGTG
GAAATTAAA

The polynucleotide sequences encoding the VH or the VL or an antigen-binding fragment thereof of the antibodies of the invention, or the heavy chain and the light chain of the antibodies of the invention may be operably linked to one or more regulatory elements, such as a promoter or enhancer, that allow expression of the nucleotide sequence in the intended host cell. The polynucleotide may be a cDNA.

The invention also provides a vector comprising the polynucleotide of the invention. Such vectors may be plasmid vectors, viral vectors, vectors for baculovirus expression, transposon based vectors or any other vector suitable for introduction of the synthetic polynucleotide of the invention into a given organism or genetic background by any means. For example, polynucleotides encoding light and/or heavy chain variable regions of the antibodies of the invention, optionally linked to constant regions, are inserted into expression vectors. The light and/or heavy chains may be cloned in the same or different expression vectors. The DNA segments encoding immunoglobulin chains may be operably linked to control sequences in the expression vector(s) that ensure the expression of immunoglobulin polypeptides. Such control sequences include signal sequences, promoters (e.g. naturally associated or heterologous promoters), enhancer elements, and transcription termination sequences, and are chosen to be compatible with the host cell chosen to express the antibody. Once the vector has been incorporated into the appropriate host, the host is maintained under conditions suitable for high level expression of the proteins encoded by the incorporated polynucleotides.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 196 and 197.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 198 and 199.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 200 and 201.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 202 and 203.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 204 and 205.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 206 and 207.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 208 and 209.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 210 and 211.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 253 and 254.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 255 and 256.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 257 and 258.

In some embodiments, the vector comprises the polynucleotide of SEQ ID NO: 259 and 260.

Suitable expression vectors are typically replicable in the host organisms either as episomes or as an integral part of the host chromosomal DNA. Commonly, expression vectors contain selection markers such as ampicillin-resistance, hygromycin-resistance, tetracycline resistance, kanamycin resistance or neomycin resistance to permit detection of those cells transformed with the desired DNA sequences.

Suitable promoter and enhancer elements are known in the art. For expression in a eukaryotic cell, exemplary promoters include light and/or heavy chain immunoglobulin gene promoter and enhancer elements; cytomegalovirus immediate early promoter; herpes simplex virus thymidine kinase promoter; early and late SV40 promoters; promoter present in long terminal repeats from a retrovirus; mouse metallothionein-I promoter; and various known tissue specific promoters. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art.

Exemplary vectors that may be used are Bacterial: pBs, phagescript, PsiX174, pBluescript SK, pBs KS, pNH8a, pNH16a, pNH18a, pNH46a (Stratagene, La Jolla, Calif., USA); pTrc99A, pKK223-3, pKK233-3, pDR540, and pRIT5 (Pharmacia, Uppsala, Sweden). Eukaryotic: pWLneo, pSV2cat, pOG44, PXR1, pSG (Stratagene) pSVK3, pBPV, pMSG and pSVL (Pharmacia), pEE6.4 (Lonza) and pEE12.4 (Lonza).

The invention also provides a host cell comprising one or more vectors of the invention. "Host cell" refers to a cell into which a vector has been introduced. It is understood that the term host cell is intended to refer not only to the particular subject cell but to the progeny of such a cell, and also to a stable cell line generated from the particular subject cell. Because certain modifications may occur in succeeding generations due to either mutation or environmental influences, such progeny may not be identical to the parent cell, but are still included within the scope of the term "host cell" as used herein. Such host cells may be eukaryotic cells, prokaryotic cells, plant cells or archeal cells. *Escherichia coli*, bacilli, such as *Bacillus subtilis*, and other enterobacteriaceae, such as Salmonella, Serratia, and various Pseudomonas species are examples of prokaryotic host cells. Other microbes, such as yeast, are also useful for expression. Saccharomyces (for example, S. cerevisiae) and Pichia are examples of suitable yeast host cells. Exemplary eukaryotic cells may be of mammalian, insect, avian or other animal origins. Mammalian eukaryotic cells include immortalized cell lines such as hybridomas or myeloma cell lines

such as SP2/0 (American Type Culture Collection (ATCC), Manassas, VA, CRL-1581), NS0 (European Collection of Cell Cultures (ECACC), Salisbury, Wiltshire, UK, ECACC No. 85110503), FO (ATCC CRL-1646) and Ag653 (ATCC CRL-1580) murine cell lines. An exemplary human myeloma cell line is U266 (ATTC CRL-TIB-196). Other useful cell lines include those derived from Chinese Hamster Ovary (CHO) cells such as CHOK1SV (Lonza Biologics, Walkersville, MD), Potelligent® CHOK2SV (Lonza), CHO-K1 (ATCC CRL-61) or DG44.

The invention also provides a method of producing an antibody of the invention comprising culturing the host cell of the invention in conditions that the antibody is expressed, and recovering the antibody produced by the host cell. Methods of making antibodies and purifying them are well known in the art. Once synthesized (either chemically or recombinantly), the whole antibodies, their dimers, individual light and/or heavy chains, or other antibody fragments such as VH and/ or VL, may be purified according to standard procedures, including ammonium sulfate precipitation, affinity columns, column chromatography, high performance liquid chromatography (HPLC) purification, gel electrophoresis, and the like (see generally Scopes, Protein Purification (Springer- Verlag, N.Y., (1982)). A subject antibody may be substantially pure, for example, at least about 80% to 85% pure, at least about 85% to 90% pure, at least about 90% to 95% pure, or at least about 98% to 99%, or more, pure, for example, free from contaminants such as cell debris, macromolecules, etc. other than the subject antibody.

The polynucleotide sequences of the invention may be incorporated into vectors using standard molecular biology methods. Host cell transformation, culture, antibody expression and purification are done using well known methods. Another embodiment of the invention is a method of producing the antagonistic antibody specifically binding PD-1 of the invention, comprising:

incorporating the first polynucleotide encoding the VH of the antibody and the second polynucleotide encoding the VL of the antibody into an expression vector;

transforming a host cell with the expression vector;

culturing the host cell in culture medium under conditions wherein the VL and the VH are expressed and form the antibody; and

recovering the antibody from the host cell or culture medium.

Another embodiment of the invention described herein is a method of producing the antagonistic antibody specifically binding TIM-3 of the invention, comprising:

incorporating the first polynucleotide encoding the VH of the antibody and the second polynucleotide encoding the VL of the antibody into an expression vector;

transforming a host cell with the expression vector;

culturing the host cell in culture medium under conditions wherein the VL and the VH are expressed and form the antibody; and

recovering the antibody from the host cell or culture medium.

The polynucleotides encoding certain VH or VL sequences of the invention described herein, and in some embodiments of each and every one of the numbered embodiments listed below, may be incorporated into vectors using standard molecular biology methods. Host cell transformation, culture, antibody expression and purification are done using well known methods.

Pharmaceutical compositions/Administration

The invention provides pharmaceutical compositions comprising the antibodies of the invention and a pharmaceutically acceptable carrier. For therapeutic use, the antibodies of the invention may be prepared as pharmaceutical compositions containing an effective amount of the antibody as an active ingredient in a pharmaceutically acceptable carrier. "Carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the antibody of the invention is administered. Such vehicles may be liquids, such as water and oils, including those of petroleum, animal, vegetable or synthetic origin, such as peanut oil, soybean oil, mineral oil, sesame oil and the like. For example, 0.4% saline and 0.3% glycine may be used. These solutions are sterile and generally free of particulate matter. They may be sterilized by conventional, well-known sterilization techniques (*e.g.*, filtration). The compositions may contain pharmaceutically acceptable auxiliary substances as required to approximate physiological conditions such as pH adjusting and buffering agents, stabilizing, thickening, lubricating and coloring agents, etc. The concentration of the antibodies of the invention in such pharmaceutical formulation may vary, from less than about 0.5%, usually to at least about 1% to as much as 15 or 20% by weight and may be selected primarily based on required dose, fluid volumes, viscosities, etc., according to the particular mode of administration selected. Suitable vehicles and formulations, inclusive of other human proteins, e.g., human serum albumin, are described, for example, in e.g. Remington: The Science and Practice of Pharmacy, 21st Edition, Troy, D.B. ed., Lipincott Williams and Wilkins, Philadelphia, PA 2006, Part 5, Pharmaceutical Manufacturing pp 691-1092, See especially pp. 958-989.

The mode of administration for therapeutic use of the antibodies of the invention may be any suitable route that delivers the antibody to the host, such as parenteral administration, e.g., intradermal, intramuscular, intraperitoneal, intravenous or

subcutaneous, pulmonary, transmucosal (oral, intranasal, intravaginal, rectal), using a formulation in a tablet, capsule, solution, powder, gel, particle; and contained in a syringe, an implanted device, osmotic pump, cartridge, micropump; or other means appreciated by the skilled artisan, as well known in the art. Site specific administration may be achieved by for example intratumoral, intrarticular, intrabronchial, intraabdominal, intracapsular, intracartilaginous, intracavitary, intracelial, intracerebellar, intracerebroventricular, intracolic, intracervical, intragastric, intrahepatic, intracardial, intraosteal, intrapelvic, intrapericardiac, intraperitoneal, intrapleural, intraprostatic, intrapulmonary, intrarectal, intrarenal, intraretinal, intraspinal, intrasynovial, intrathoracic, intrauterine, intravascular, intravesical, intralesional, vaginal, rectal, buccal, sublingual, intranasal, or transdermal delivery.

The antibodies of the invention may be administered to a subject by any suitable route, for example parentally by intravenous *(i.v.)* infusion or bolus injection, intramuscularly or subcutaneously or intraperitoneally. *i.v.* infusion may be given over for example 15, 30, 60, 90, 120, 180, or 240 minutes, or from 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11 or 12 hours.

The dose given to a subject is sufficient to alleviate or at least partially arrest the disease being treated ("therapeutically effective amount") and may be sometimes 0.005 mg to about 100 mg/kg, e.g. about 0.05 mg to about 30 mg/kg or about 5 mg to about 25 mg/kg, or about 4 mg/kg, about 8 mg/kg, about 16 mg/kg or about 24 mg/kg , or for example about 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10 mg/kg, but may even higher, for example about 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 30, 40, 50, 60, 70, 80, 90 or 100 mg/kg.

A fixed unit dose may also be given, for example, 50, 100, 200, 500 or 1000 mg, or the dose may be based on the patient's surface area, e.g., 500, 400, 300, 250, 200, or 100 mg/m^2. Usually between 1 and 8 doses, (e.g., 1, 2, 3, 4, 5, 6, 7 or 8) may be administered to treat the patient, but 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20 or more doses may be given.

The administration of the antibodies of the invention may be repeated after one day, two days, three days, four days, five days, six days, one week, two weeks, three weeks, one month, five weeks, six weeks, seven weeks, two months, three months, four months, five months, six months or longer. Repeated courses of treatment are also possible, as is chronic administration. The repeated administration may be at the same dose or at a different dose. For example, the antibodies of the invention may be administered at 8 mg/kg or at 16 mg/kg at weekly interval for 8 weeks, followed by administration at 8 mg/kg or at 16 mg/kg every two weeks for an additional 16 weeks,

followed by administration at 8 mg/kg or at 16 mg/kg every four weeks by intravenous infusion.

For example, the antibodies of the invention may be provided as a daily dosage in an amount of about 0.1-100 mg/kg, such as 0.5, 0.9, 1.0, 1.1, 1.5, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 40, 45, 50, 60, 70, 80, 90 or 100 mg/kg, per day, on at least one of day 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, or 40, or alternatively, at least one of week 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19 or 20 after initiation of treatment, or any combination thereof, using single or divided doses of every 24, 12, 8, 6, 4, or 2 hours, or any combination thereof.

The antibodies of the invention, may also be administered prophylactically in order to reduce the risk of developing cancer, delay the onset of the occurrence of an event in cancer progression, and/or reduce the risk of recurrence when a cancer is in remission.

The antibodies of the invention may be lyophilized for storage and reconstituted in a suitable carrier prior to use. This technique has been shown to be effective with conventional protein preparations and well known lyophilization and reconstitution techniques can be employed.

Methods and Uses

The antibodies of the invention have *in vitro* and *in vivo* diagnostic, as well as therapeutic and prophylactic utilities. For example, the antibodies of the invention may be administered to cells in culture, *in vitro* or *ex vivo*, or to a subject to treat, prevent, and/or diagnose a variety of disorders, such as cancers and infectious disorders.

The invention provides a method of modifying an immune response in a subject comprising administering to the subject the antibody of the invention for a time sufficient to modify the immune response.

In some embodiments, the immune response is enhanced, stimulated or up-regulated.

In some embodiments described herein, the subject is a human patient.

In some embodiments described herein, the subject is a human patient in need of enhancement of the immune response.

In some embodiments, the subject is immunocompromised.

In some embodiments, the subject is at risk of being immunocompromised. Immunocompromised subject may be undergoing, or has undergone a chemotherapeutic or radiation therapy.

In some embodiment, the subject is or is at risk of being immunocompromised as a result of an infection.

The antibodies of the invention are suitable for treating a subject having a disorder that may be treated by augmenting T-cell mediated immune responses.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention described herein is PD1B114, PD1B149, PD1B160, PD1B162, PD1B164, PD1B11, PD1B183, PD1B184, PD1B185, PD1B187, PD1B71, PD1B177, PD1B70, PD1B175, PD1B194, PD1B195, PD1B196, PD1B197, PD1B198, PD1B199, PD1B200, PD1B201, PD1B243, PD1B244, PD1B131 or PD1B132. The VH and the VL amino acid sequences of these antibodies are shown in **Table 2.**

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention described herein is TM3B103, TM3B105, TM3B109, TM3B108, TM3B113, TM3B189, TM3B190, TM3B193, TM3B195, TM3B196 or TM3B291. The VH and the VL amino acid sequences of these antibodies are shown in **Table 3.**

In some embodiments, the bispecific PD-1/TIM-3 antibody used in the methods of the invention is PTBB14, PTBB15, PTBB16, PTBB17, PTBB24, PTBB30, PTBB27, PTBB28, PTBB18, PTBB20 or PTBB21. The HC1, the LC1, the HC2 and the LC2 amino acid sequences of these antibodies are shown in **Table 41** and **Table 42.**

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 50.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 51.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 52.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 53.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 49.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 54.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 50.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 55.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 56.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 57.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 44 and the VL of SEQ ID NO: 49.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 49.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 46 and the VL of SEQ ID NO: 49.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 49.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 53.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 52.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 59.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 61.

In some embodiments, the antagonistic antibody specifically binding PD-1 used in the methods of the invention comprises the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 62.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 148 and the VL of SEQ ID NO: 157.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 147 and the VL of SEQ ID NO: 155.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 149 and the VL of SEQ ID NO: 158.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 150 and the VL of SEQ ID NO: 159.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 151 and the VL of SEQ ID NO: 160.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 152 and the VL of SEQ ID NO: 161.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 154 and the VL of SEQ ID NO: 163.

In some embodiments, the antagonistic antibody specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 in the first domain, and the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162 in the second domain.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 in the first domain, and the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156 in the second domain.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 in the first domain, and the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162 in the second domain.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3 used in the methods of the invention, comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 in the first domain, and the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156 in the second domain.

In some embodiments, the antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3 used in the methods of the invention comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 in the first domain, and the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173 in the second domain.

Cancer

Blockade of PD-1 may enhance an immune response to cancerous cells in a subject. The ligand for PD-1, PD-L1, is abundantly expressed in a variety of human cancers (Dong *et al.,* (2002) *Nat Med* 8:787-9). The interaction between PD-1 and PD-L1 can result in a decrease in tumor infiltrating lymphocytes, a decrease in T-cell receptor mediated proliferation, and/or immune evasion by the cancerous cells (Dong *et al.,* (2003) *J Mol Med* 81:281-7; Blank *et al.,* (2005) *Cancer Immunol Immunother* 54:307-314; Konishi *et al.,* (2004) *Clin Cancer Res* 10:5094-100). Immune suppression may be reversed by inhibiting the local interaction of PD-1 to PD-L1; the effect is additive when the interaction of PD-1 to the second PD-1 ligand, PD-L2, is blocked as well (Iwai *et al.,* (2002) *PorcNatl Acad Sci* 99:12293-7; Brown *et al.,* (2003) *J Immunol* 170:1257-66). Thus, inhibition of PD-1 may result in augmenting an immune response.

TIM-3 is a coinhibitory protein expressed on activated T helper 1 (Thl) $CD4^+$ and cytotoxic $CD8^+$ T cells that secrete IFN-γ. TIM-3 is co-expressed on PD-1+ exhausted T cells as shown in preclinical models of cancer and viral exhaustion. Co-blockade of these pathways may restore effector T cell function (e.g., IFN-γ secretion, proliferation) in several models as well as human PBMCs derived from metastatic melanoma patients and patients with HIV or HCV. TIM-3 is also enriched on Foxp3+ regulatory T cells and Tregs co-expressing TIM-3, LAG3 and CTLA4 have been shown to be highly efficient

suppressors of effector T cells (Teff) (Galuton *et al.,* (2014) *Eur J Immunol* 44(9):2703-11). TIM-3 expression has been correlated with poorer prognosis in NSCLC (Zhuang *et al.,* (2012) Am J Clin Pathol 137(6):978-85). Lymphocytes from tumor tissues of ovarian, colorectal, cervical and hepatocellular carcinoma patients exhibit higher proportion of TIM-3$^+$ CD4 T cells, which cells have impaired capacity to produce ILF-γ (Yan *et al.,* (2013) *PLoS One* 8(3):e58006).

The invention also provides a method of inhibiting growth of tumor cells in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody specifically binding PD-1 of the invention for a time sufficient to inhibit growth of tumor cells...

The invention also provides a method of inhibiting growth of tumor cells in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody specifically binding TIM-3 of the invention for a time sufficient to inhibit growth of tumor cells.

The invention also provides a method of inhibiting growth of tumor cells in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic bispecific PD-1/TIM-3 antibody of the invention for a time sufficient to inhibit growth of tumor cells.

The invention also provides a method of treating a cancer by administering to the subject in need thereof a therapeutically effective amount of the antagonistic antibody specifically binding PD-1 of the invention for a time sufficient to treat the cancer.

The invention also provides a method of treating a cancer by administering to the subject in need thereof a therapeutically effective amount of the antagonistic antibody specifically binding TIM-3 of the invention for a time sufficient to treat the cancer.

The invention also provides a method of treating a cancer by administering to the subject in need thereof a therapeutically effective amount of the bispecific PD-1/TIM-3 antibody of the invention for a time sufficient to treat the cancer.

Exemplary antibodies that may be used are antagonistic antibodies specifically binding PD-1, antagonistic antibodies specifically binding TIM-3, and antagonistic bispecific PD-1/TIM-3 antibodies PD1B114, PD1B149, PD1B160, PD1B162, PD1B164, PD1B11, PD1B183, PD1B184, PD1B185, PD1B187, PD1B71, PD1B177, PD1B70, PD1B175, PD1B194, PD1B195, PD1B196, PD1B197, PD1B198, PD1B199, PD1B200, PD1B201, TM3B103, TM3B105, TM3B109, TM3B108, TM3B113, TM3B189, TM3B190, TM3B193, TM3B195, TM3B196, TM3B291, PTBB14, PTBB15, PTBB16,

PTBB17, PTBB24, PTBB30, PTBB27, PTBB28, PTBB18, PTBB20 and PTBB21 having the VH and the VL amino acid sequence and characteristics as described herein.

Cancer may be a hyperproliferative condition or disorder, a solid tumor, a hematological malignancy, a soft tissue tumor, or a metastatic lesion.

"Cancer" is meant to include all types of cancerous growths or oncogenic processes, metastatic tissues or malignantly transformed cells, tissues, or organs, irrespective of histopathology type or stage of invasiveness. Examples of cancers include solid tumors, hematological malignancies, soft tissue tumors, and metastatic lesions. Exemplary solid tumors include malignancies, e.g., sarcomas, and carcinomas (including adenocarcinomas and squamous cell carcinomas) of the various organ systems, such as those affecting liver, lung, breast, lymphoid, gastrointestinal (e.g., colon), genitourinary tract (e.g., renal, urothelial cells), prostate and pharynx. Adenocarcinomas include malignancies such as most colon cancers, a rectal cancer, a renal-cell carcinoma, a liver cancer, a non-small cell carcinoma of the lung, a cancer of the small intestine and a cancer of the esophagus. Squamous cell carcinomas include malignancies, e.g., in the lung, esophagus, skin, head and neck region, oral cavity, anus, and cervix.

In some embodiments, the cancer is a melanoma.

Metastatic lesions of the aforementioned cancers may also be treated or prevented using the methods and antibodies of the invention described herein.

Exemplary cancers whose growth may be inhibited or reduced using the antibodies of the invention include cancers that may be responsive to immunotherapy. Exemplary such cancers include a melanoma, a renal cancer, a prostate cancer, a breast cancer, a colon cancer, a gastrointestinal cancer, a stomach cancer, an esophageal cancer, a lung cancer, a metastatic malignant melanoma, a clear cell carcinoma, a hormone refractory prostate adenocarcinoma, a non-small cell lung cancer or cancer of the head and neck. Refractory or recurrent malignancies may be treated using the antibodies of the invention described herein.

Exemplary other cancers that may be treated with the antibodies of the invention ae an anal cancer, a basal cell carcinoma, a biliary tract cancer, a bladder cancer, a bone cancer, brain and CNS cancers, a carcinoma of the fallopian tubes, carcinoma of the vagina, a carcinoma of the vulva, a cutaneous or intraocular malignant melanoma, a astro-esophageal cancer, a testicular cancer, an ovarian cancer, a pancreatic cancer, a rectal cancer, an uterine cancer, a primary CNS lymphoma; a neoplasm of the central nervous system (CNS), a cervical cancer, a choriocarcinoma, a rectum cancer, a connective tissue cancer, a cancer of the digestive system, an endometrial cancer, an eye cancer; an intra-

epithelial neoplasm, a kidney cancer, a larynx cancer, a liver cancer; a small cell lung cancer, a neuroblastoma, an oral cavity cancer (e.g., lip, tongue, mouth, and pharynx), a nasopharyngeal cancer, a retinoblastoma, a rhabdomyosarcoma, a cancer of the respiratory system, a sarcoma, a thyroid cancer, a cancer of the urinary system, a hepatocarcinoma, a cancer of the anal region, a carcinoma of the fallopian tubes, a carcinoma of the vagina, a carcinoma of the vulva, a cancer of the small intestine, a cancer of the endocrine system, a cancer of the parathyroid gland, a cancer of the adrenal gland, a sarcoma of soft tissue, a cancer of the urethra, a cancer of the penis, solid tumors of childhood, a tumor angiogenesis, a spinal axis tumor, a brain stem glioma, a pituitary adenoma, Kaposi's sarcoma, Merkel cell cancer, an epidermoid cancer, a squamous cell cancer, an environmentally induced cancers including those induced by asbestos, as well as other carcinomas and sarcomas, and combinations of said cancers.

Exemplary hematological malignancies that may be treated with the antibodies of the invention include leukemias, lymphomas and myeloma, such as a precursor B-cell lymphoblastic leukemia/lymphoma and a B-cell non-Hodgkin's lymphoma, an acute promyelocytic leukemia, an acute lymphoblastic leukemia (ALL), a B-cell chronic lymphocytic leukemia(CLL)/small lymphocytic lymphoma (SLL), a B-cell acute lymphocytic leukemia, a B-cell prolymphocytic leukemia, a lymphoplasmacytic lymphoma, a mantle cell lymphoma (MCL), a follicular lymphoma (FL), including low-grade, intermediate- grade and high-grade FL, a cutaneous follicle center lymphoma, a marginal zone B-cell lymphoma (MALT type, nodal and splenic type), a hairy cell leukemia, a diffuse large B-cell lymphoma (DLBCL), Burkitt's lymphoma (BL), a plasmacytoma, a multiple myeloma (MM), a plasma cell leukemia, a post-transplant lymphoproliferative disorder, Waldenstrom's macroglobulinemia, plasma cell disorders, an anaplastic large-cell lymphoma (ALCL), a T-cell acute lymphocytic leukemia, a primary systemic amyloidosis (e.g. light chain amyloidosis), a pro-lymphocytic/myelocytic leukemia, an acute myeloid leukemia (AML), a chronic myeloid leukemia (CML), a large granular lymphocytic (LGL) leukemia, a NK-cell leukemia and Hodgkin’s lymphoma.

“Plasma cell disorder” refers to disorders characterized by clonal plasma cells, and includes a multiple myeloma, a light chain amyloidosis and Waldenstrom’s macroglobulinemia. Light chain amyloidosis and Waldenstrom’s macroglobulinemia can arise independently from multiple myeloma. They may also present simultaneously with multiple myeloma, and develop either before or after the development of multiple myeloma.

Exemplary B-cell non-Hodgkin's lymphomas are a lymphomatoid granulomatosis, a primary effusion lymphoma, an intravascular large B-cell lymphoma, a mediastinal large B-cell lymphoma, heavy chain diseases (including γ, μ, and a disease), lymphomas induced by therapy with immunosuppressive agents, such as cyclosporine-induced lymphoma, and methotrexate-induced lymphoma.

Patients having cancer including metastatic cancer that express PD-L1 may be treated with the antibodies of the invention. The cancer may be a melanoma, a renal cell carcinoma, a squamous non-small cell lung cancer (NSCLC), a non-squamous NSCLC, a colorectal cancer, a castration-resistant prostate cancer, an ovarian cancer, a gastric cancer, an adenocarcinoma (ACA), a squamous cell carcinoma (SCC), a hepatocellular carcinoma (HCC), a pancreatic carcinoma, a squamous cell carcinoma of the head and neck, carcinomas of the esophagus, gastrointestinal tract and breast.

Patients having cancer that expresses TIM-3 may be treated with the antibodies of the invention. TIM-3-expressing cancers include a cervical cancer, a lung cancer, a NSCLC, an acute myeloid leukemia (AML), a diffuse large B cell lymphoma (DLBCL), a melanoma, a renal cancer, a renal cell carcinoma (RCC), a kidney clear cell carcinoma, a kidney papillary cell carcinoma, a metastatic renal cell carcinoma, a squamous cell carcinoma, an esophageal squamous cell carcinoma, a nasopharyngeal carcinoma, a colorectal cancer, a breast cancer (e.g., a breast cancer that does not express one, two or all of estrogen receptor, progesterone receptor, or Her2/neu, e.g., a triple negative breast cancer), a mesothelioma, a hepatocellular carcinoma, and an ovarian cancer. The TIM-3-expressing cancer may be a metastatic cancer.

In some embodiments, the subject has a solid tumor.

In some embodiments, the subject has a hematological malignancy.

In some embodiments, the solid tumor is a melanoma.

In some embodiments, the solid tumor is a lung cancer.

In some embodiments, the solid tumor is a non-small cell lung cancer (NSCLC).

In some embodiments, the solid tumor is a squamous non-small cell lung cancer (NSCLC).

In some embodiments, the solid tumor is a non-squamous NSCLC.

In some embodiments, the solid tumor is a lung adenocarcinoma.

In some embodiments, the solid tumor is a renal cell carcinoma (RCC).

In some embodiments, the solid tumor is a mesothelioma.

In some embodiments, the solid tumor is a nasopharyngeal carcinoma (NPC).

In some embodiments, the solid tumor is a colorectal cancer.

In some embodiments, the solid tumor is a prostate cancer.

In some embodiments, the solid tumor is castration-resistant prostate cancer.

In some embodiments, the solid tumor is a stomach cancer.

In some embodiments, the solid tumor is an ovarian cancer.

In some embodiments, the solid tumor is a gastric cancer.

In some embodiments, the solid tumor is a liver cancer.

In some embodiments, the solid tumor is pancreatic cancer.

In some embodiments, the solid tumor is a thyroid cancer.

In some embodiments, the solid tumor is a squamous cell carcinoma of the head and neck.

In some embodiments, the solid tumor is a carcinomas of the esophagus or gastrointestinal tract.

In some embodiments, the solid tumor is a breast cancer.

In some embodiments, the solid tumor is a fallopian tube cancer.

In some embodiments, the solid tumor is a brain cancer.

In some embodiments, the solid tumor is an urethral cancer.

In some embodiments, the solid tumor is a genitourinary cancer.

In some embodiments, the solid tumor is an endometriosis.

In some embodiments, the solid tumor is a cervical cancer.

In some embodiments, the solid tumor is a metastatic lesion of the cancer.

In some embodiments, the hematological malignancy is a lymphoma, a myeloma or a leukemia.

In some embodiments, the hematological malignancy is a B cell lymphoma.

In some embodiments, the hematological malignancy is Burkitt's lymphoma.

In some embodiments, the hematological malignancy is Hodgkin's lymphoma.

In some embodiments, the hematological malignancy is a non-Hodgkin's lymphoma.

In some embodiments, the hematological malignancy is a myelodysplastic syndrome.

In some embodiments, the hematological malignancy is an acute myeloid leukemia (AML).

In some embodiments, the hematological malignancy is a chronic myeloid leukemia (CML).

In some embodiments, the hematological malignancy is a chronic myelomoncytic leukemia (CMML).

In some embodiments, the hematological malignancy is a multiple myeloma (MM).

In some embodiments, the hematological malignancy is a plasmacytoma.

In some embodiments, the subject has a tumor that expresses PD-L1.

In some embodiments, the subject has tumor-infiltrating T lymphocytes (TILs) in the tumor tissue.

In some embodiments, the subject has PD-1^{+}TIM-3^{+} TILs in the tumor tissue.

In some embodiments, the subject has increased number of PD-1^{+}TIM-3^{+} tumor-infiltrating T lymphocytes (TILs) in the tumor tissue.

"Increased number" refers to statistically significant increase in a subject when compared to a control. "Increased number" for example refers to statistically significant increase in the number of TILs in a subject (e.g. patient) pre- and post-treatment with a PD-1 antibody or other therapeutic.

In some embodiments, the subject has increased expression or activity of interferon-gamma (IFN-γ).

In some embodiments, the subject has been treated with an anti-PD-1 antibody.

In some embodiments, the subject is refractory to treatment with the anti-PD-1 antibody.

In some embodiments, the subject has a relapsed tumor after treatment with the anti-PD-1 antibody.

In some embodiments, the subject has been treated with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231 (e.g. KEYTRUDA® (pembrolizumab)).

In some embodiments, the subject has been treated with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233 (e.g. OPDIVO® (nivolumab)).

In some embodiments, the subject is refractory to treatment with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231 (e.g. KEYTRUDA® (pembrolizumab)).

In some embodiments, the subject is refractory to treatment with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233 (e.g. OPDIVO® (nivolumab)).

In some embodiments, the subject has a relapsed tumor after treatment with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231 (e.g. KEYTRUDA® (pembrolizumab).

In some embodiments, the subject has a relapsed tumor after treatment with the anti-PD-1 antibody comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233 (e.g. OPDIVO® (nivolumab)).

SEQ ID NO: 230
QVQLVQSGVEVKKPGASVKVSCKASGYTFTNYYMYWVRQAPGQGLEWMGG
INPSNGGTNFNEKFKNRVTLTTDSSTTTAYMELKSLQFDDTAVYYCARRDYRFDM
GFDYWGQGTTVTVSS

SEQ ID NO: 231
EIVLTQSPATLSLSPGERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLAS
YLESGVPARFSGSGSGTDFTLTISSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK

SEQ ID NO: 232
QVQLVESGGGVVQPGRSLRLDCKASGITFSNSGMHWVRQAPGKGLEWVAVIWY
DGSKRYYADSVKGRFTISRDNSKNTLFLQMNSLRAEDTAVYYCATNDDYWGQG
TLVTVSS

SEQ ID NO: 233
EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQSSNWPRTFGQGTKVEIK

In some embodiments, the subject has been treated or is being treated with a PD-L1 antibody.

In some embodiments, the subject is refractory to treatment with the PD-L1 antibody.

In some embodiments, the subject has a relapsed tumor after treatment with the PD-L1 antibody.

In some embodiments, the subject is refractory or relapsed after treatment with the PD-L1 antibody durvalumab (MEDI-4736). Durvalumab comprises the VH of SEQ ID NO: 234 and the VL of SEQ ID NO: 235.

SEQ ID NO: 234

EVQLVESGGG LVQPGGSLRLSCAASGFTFSRYWMSWVRQAPGKGLEWVAN
IKQDGSEKYYVDSVKGRFTISRDNAKNSLYLQMNSLRAEDTAVYYCAREG
GWFGELAFDYWGQGTLVTVSS

SEQ ID NO: 235
EIVLTQSPGTLSLSPGERATLSCRASQRVSSSYLAWYQQK PGQAPRLLIY
DASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSLPWTFG
QGTKVEIK

In some embodiments, the subject is refractory or relapsed after treatment with the PD-L1 antibody atezolizumab.
Atezolizumab comprises the VH of SEQ ID NO: 236 and the VL of SEQ ID NO: 237.

SEQ ID NO: 236
EVQLVESGGGLVQPGGSLRLSCAASGFTFSDSWIHWVRQAPGKGLEWVAW
ISPYGGSTYYADSVKGRFTISADTSKNTAYLQMNSLRAEDTAVYYCARRH
WPGGFDYWGQGTLVTVSS

SEQ ID NO: 237
DIQMTQSPSSLSASVGDRVTITCRASQDVSTAVAWYQQKPGKAPKLLIYS
ASFLYSGVPSRFSGSGSGTDFTLTISSLQPEDFATYYCQQYLYHPATFGQ
GTKVEIK

In some embodiments, the subject is refractory or relapsed after treatment with the PD-L1 antibody avelumab.
Avelumab comprises the VH of SEQ ID NO: 238 and the VL of SEQ ID NO: 239.

SEQ ID NO: 238
EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYIMMWVRQAPGKGLEWVSS
IYPSGGITFYADTVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCARIK
LGTVTTVDYWGQGTLVTVSS

SEQ ID NO: 239
QSALTQPASVSGSPGQSITISCTGTSSDVGGYNYVSWYQQHPGKAPKLMI
YDVSNRPSGVSNRFSGSKSGNTASLTISGLQAEDEADYYCSSYTSSSTRV

FGTGTKVTVL

In some embodiments, the subject is refractory or relapsed after treatment with the PD-L1 antibody MDX-1105.

In some embodiments, the subject has been treated or is being treated with a PD-L2 antibody.

In some embodiments described herein, the subject is refractory to treatment with a PD-L2 antibody.

In some embodiments, the subject has a relapsed tumor after treatment with a PD-L2 antibody.

Various qualitative and/or quantitative methods may be used to determine relapse or refractory nature of the disease. Symptoms that may be associated with relapse or resistance are, for example, a decline or plateau of the well-being of the patient or re-establishment or worsening of various symptoms associated with solid tumors, and/or the spread of cancerous cells in the body from one location to other organs, tissues or cells.

TIM-3 expression was found herein to be elevated in $CD8^+$ T cells isolated from tumors after anti-PD-1 antibody treatment. Therefore, therapeutic administration of antagonistic antibodies specifically binding TIM-3 or antagonistic bispecific PD-1/TIM-3 antibodies described herein to a subject who has already received or is receiving anti-PD-1 antibody therapy, is refractory to the anti-PD-1 antibody treatment or has relapsed after or during the anti-PD-1 antibody treatment may improve the clinical outcome of the patients.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody specifically binding TIM-3 of the invention, wherein the subject is being treated or has been treated with an anti-PD-1 antibody.

In some embodiments, the antagonistic antibody specifically binding TIM-3 comprises the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156, wherein the subject is being treat or has been treated with the anti-PD-1 antibody KEYTRUDA® (pembrolizumab) comprising the VH of SEQ ID NO: 230 and the VL of SEQ ID NO: 231.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody

specifically binding TIM-3 comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156, wherein the subject is being treat or has been treated with the anti-PD-1 antibody OPDIVO® (nivolumab) comprising the VH of SEQ ID NO: 232 and the VL of SEQ ID NO: 233.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody specifically binding TIM-3 of the invention, wherein the subject is being treated or has been treated with an anti-PD-L1 antibody.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody specifically binding TIM-3 of the invention, wherein the subject is being treated or has been treated with an anti-PD-L2 antibody.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic bispecific PD-1/TIM-3 antibody the invention, wherein the subject is being treated or has been treated with an anti-PD-1 antibody.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic bispecific PD-1/TIM-3 antibody the invention, wherein the subject is being treated or has been treated with an anti-PD-L1 antibody.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic bispecific PD-1/TIM-3 antibody the invention, wherein the subject is being treated or has been treated with an anti-PD-L2 antibody.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 for a time sufficient to treat the cancer.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject a therapeutically effective amount of the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65 for a time sufficient to treat the cancer.

Any of the PD-1, TIM-3 or bispecific PD-1/TIM-3 antibodies of the invention described herein may be used in the methods of the invention.

"Treat" or "treatment" refers to therapeutic treatment wherein the object is to slow down (lessen) an undesired physiological change or disease, such as the development or spread of tumor or tumor cells, or to provide a beneficial or desired clinical outcome during treatment. Beneficial or desired clinical outcomes include alleviation of symptoms, diminishment of extent of disease, stabilized (i.e., not worsening) state of disease, delay or slowing of disease progression, lack of metastasis, amelioration or palliation of the disease state, and remission (whether partial or total), whether detectable or undetectable. "Treatment" may also mean prolonging survival as compared to expected survival if a subject was not receiving treatment. Those in need of treatment include those subjects already with the undesired physiological change or diseases well as those subjects prone to have the physiological change or disease.

A "therapeutically effective amount" refers to an amount effective, at dosages and for periods of time necessary, to achieve a desired therapeutic result. A therapeutically effective amount of the antibody of the invention may vary according to factors such as the disease state, age, sex, and weight of the individual, and the ability of the antibody of the invention to elicit a desired response in the individual. Exemplary indicators of an effective therapeutic or combination of therapeutics include, for example, improved well-being of the patient, reduction in a tumor burden, arrested or slowed growth of a tumor, and/or absence of metastasis of cancer cells to other locations in the body.

Combination therapies for cancer treatment

The antibodies of the invention may be administered in combination with a second therapeutic agent.

The antibodies of the invention may be administered in combination with one, two, three, four, five or six additional therapeutic agents.

Any of the antagonistic antibodies specifically binding PD-1, antagonistic antibodies specifically binding TIM-3 or antagonistic bispecific PD-1/TIM-3 antibodies of the invention may be used in combination with a second therapeutic agent.

Any of the antagonistic antibodies specifically binding PD-1, antagonistic antibodies specifically binding TIM-3 or antagonistic bispecific PD-1/TIM-3 antibodies of the invention may be used in combination with one, two, three, four, five or six additional therapeutic agents.

"In combination with" refers to administering of the antibodies of the invention and at least one second therapeutic agent concurrently as single agents or sequentially as

single agents in any order. In general, each agent will be administered at a dose and/or on a time schedule determined for that agent.

In some embodiments, the second therapeutic agent modulates activity of a molecule involved in the cancer-immunity cycle, e.g. a molecule involved in stimulatory or inhibitory pathways functioning in release of cancer cell antigens, cancer antigen presentation, T cell priming and activation, trafficking of T cells to tumors, infiltration of T cells into tumors, recognition of cancer cells by T cells, and killing of cancer cells. The cancer-immunity cycle is described in Chen and Mellman (2013) *Immunity* 39:1-10. In some embodiments, the second therapeutic agend modulates activity of a molecule involved in regulation of activity of T regulatory cells (Treg), co-stimulatory or co-inhibitory ligands expressed on tumors, activating or inhibitory receptors on natural killer (NK) cells, or immunosuppressive factors in the tumor microenvironment. Combination cancer immunotherapies are described in Manoney *et al.,* (2015) *Nature Reviews* 14:561-584.

The second therapeutic agent typically enhances the activity of stimulatory molecules and suppresses the activity of inhibitory molecules, as is well known. Thus, "modulate" refers to the enhancement of immune response by the second therapeutic agent, wheatear the agent itself is agonist or antagonist of a specific molecule.

In some embodiments, the antibodies of the invention are administered in combination with an inhibitor of a T cell inhibitory molecule.

In some embodiments, the antibodies of the invention are administered in combination with an inhibitor of a T cell inhibitory molecule PD-1, PD-L1, PD-L2, VISTA, BTNL2, B7-H3, B7-H4, HVEM, HHLA2, CTLA-4, LAG-3, TIM-3, BTLA, CD160, CEACAM-1, LAIR1, TGFβ, IL-10, Siglec family protein, KIR, CD96, TIGIT, NKG2A, CD112, CD47, SIRPA or CD244.

In some embodiments, KIR is KIR2DL1, KIR2DL2 or KIR2DL3.

Inhibition of inhibitory molecules may be performed by inhibition at the DNA, RNA or protein level. In some embodiments, an inhibitory nucleic acid (e.g., a dsRNA, siRNA or shRNA) is used to inhibit expression of the inhibitory molecule.

In some embodiments, the inhibitor of the inhibitory molecule is a soluble ligand of the inhibitory molecule.

In some embodiments, the inhibitor of the inhibitory molecule is an antagonistic antibody specifically binding the inhibitory molecule.

In some embodiments, the inhibitor of the inhibitory molecule is CTLA-4-Fc or TIM-3-Fc fusion protein.

In some embodiments, the inhibitor of the inhibitory molecule is an antibody or an antibody fragment that binds PD-1, PD-L1, PD-L2, VISTA, BTNL2, B7-H3, B7-H4, HVEM, HHLA2, CTLA-4, LAG-3, TIM-3, BTLA, CD160, CEACAM-1, LAIR1, TGFβ, IL-10, Siglec family protein, KIR, CD96, TIGIT, NKG2A, CD112, CD47, SIRPA or CD244.

Exemplary anti-PD-1 antibodies that may be used in the methods of the invention are those described herein and in U.S. Patent Nos. 5,897,862 and 7,488,802, and in Int. Patent Publ. Nos. WO2004/004771, WO2004/056875, WO2006/121168, WO2008/156712, WO2010/029435, WO2010/036959, WO2011/110604, WO2012/145493, WO2014/194302, WO2014/206107, WO2015/036394, WO2015/035606, WO2015/085847, WO2015/112900 and WO2015/112805. Exemplary anti-PD1 antibodies include KEYTRUDA® (pembrolizumab) and OPDIVO® (nivolumab).

In some embodiments, the antibodies of the invention are administered in combination with a soluble PD-1 ligand.

In some embodiments, the soluble PD-1 ligand is soluble PD-L1 or soluble PD-L2 fused to an Fc.

In some embodiments, the soluble PD-1 ligand is AMP-224.

In some embodiments, the antibodies of the invention are administered in combination with an anti-PD-L1 antibody, or antigen-binding fragments thereof.

Exemplary PD-L1 antibodies that may be used in the methods of the invention are antibodies MDPL3280A (Genentech/Roche) and other human monoclonal antibodies disclosed in U.S. Patent No. 7,943,743 and U.S Patent Publ. No. 20120039906. Other anti-PD-L1 binding agents include YW243.55.S70 (heavy and light chain variable regions are shown in SEQ ID NOs 20 and 21 in WO2010/077634) and MDX-1105 (also referred to as BMS-936559, and, e.g., anti-PD-L1 binding agents disclosed in WO2007/005874). The VH and the VL sequences of anti-PD-L1 antibodies durvalumab, atezolimumab and avelumab that may be used are disclosed herein.

Exemplary PD-L2 antibodies that may be used in the methods of the invention are those described in U.S. Patent Nos. 8,080,636, 8,188,238, U.S. Patent Publ. No. 20110271358 and Int. Patent Publ. No. WO2012145493.

Exemplary B7-H4 antibodies that may be used in the methods of the invention are those described in U.S. Patent Nos. 7,888,477, 8,609,816, 7,931,896, European Patent No. 1817055, U.S. Patent Publ. No. US20140037551and US2014029486, and Int. Patent Publ. Nos. WO2014/100483 and WO2014/159835.

Exemplary anti-CTLA-4 antibodies that may be used in the methods of the invention are ipilimumab (MDX-010, CAS No. 477202-00-9) and tremelimumab (IgG2 monoclonal antibody available from Pfizer, formerly known as ticilimumab, CP-675,206).

Exemplary anti-LAG-3 antibodies that may be used in the methods of the invention are those described for example in Int. Patent Publ. Nos. WO2008/132601 and WO2010/019570.

Exemplary anti-CEACAM-1 antibodies that may be used in the methods of the invention are those described in U.S. Patent No. 8,598,322 and in U.S. Patent Publ. Nos. US2004/0047858, US20140271618 and US20120100158. Without wishing to be bound by any particular theory, CEACAM-1 has been described as a ligand and partner of TIM-3 (see e.g., Int. Patent Publ. No. WO2014/022332). Synergistic *in vivo* effect of the combination of anti-TIM-3 and anti-CEACAM-1 antibodies have been detected in xenograft cancer models (see e.g., Int. Patent Publ. No. WO2014/022332). Tumors may use CEACAM-1 to inhibit the immune system. Therefore, anti- CEACAM-1 antibodies may be used in combination with the antibodies of the invention described herein.

Exemplary anti-LAIR1 antibodies that may be used in the methods of the invention are those described in U.S. Patent No. 6,479,638 and Int. Patent Publ. No. WO2010/078580.

Exemplary anti-CD96 antibodies that may be used in the methods of the invention are those described in Int. Patent Publ. No. WO2015/024060.

Exemplary anti-TIM-3 antibodies that may be used in the methods of the invention are those described herein and in Int. Patent Publ. Nos. WO2011/155607, WO2013/006490 and WO2015/117002.

Exemplary anti-TIGIT antibodies that may be used in the methods of the invention are those described in U.S. Patent Publ. Nos. US20140056890 and US20150216970. An exemplary anti-TIGIT antibody is RG-6058 (MTIG-7192A).

TIGIT expression was found herein to be elevated in $CD8^+$ T cells isolated from tumors after anti-TIM-3 antibody treatment in animal models of cancer. Therefore, therapeutic administration of antagonistic antibodies specifically binding TIGIT to a subject who has already received or is receiving anti-TIM-3 antibody therapy, is refractory to the anti-TIM-3 antibody treatment or has relapsed after or during the anti-TIM-3 antibody treatment may improve the clinical outcome of the patients.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject in need thereof a therapeutically effective amount of an

antagonistic antibody that specifically binds TIM-3 and an antagonistic antibody that specifically binds TIGIT for a time sufficient to treat the cancer.

In some embodiments, the antagonistic antibody that specifically binds TIGIT is administered after administration of the antagonistic antibody specifically binding TIM-3.

In some embodiments, the antagonistic antibody that specifically binds TIGIT and the antagonistic antibody specifically binding TIM-3 are administered concurrently as single agents or sequentially as single agents in any order.

Exemplary anti-BTLA antibodies that may be used in the methods of the invention are those described in U.S. Patent Nos. 8,546,541, 7,479,544, 8,188,232, 8,247,537, 8,563,694 and in Int. Patent Publ. No. WO2014184360.

Exemplary anti-HVEM antibodies that may be used in the methods of the invention are those described in U.S. Patent Publ. No. US20110280866.

Exemplary CD47 antibodies that may be used in the methods of the invention are those described in U.S. Patent No. 8,101,719.

Exemplary CD244 antibodies that may be used in the methods of the invention include those described in U.S. Patent No. 5,688,690.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-TIM-3 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-PD-L1 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-PD-L2 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-VISTA antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the

antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-BTNL2 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-B7-H3 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-B7-H4 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-HVEM antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-HLA2 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CTLA-4 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-LAG-3 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-TIM-3 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-BTLA antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD160 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CEACAM-1 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-LAIR1 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-TGFβ antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-IL-10 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-TIGIT antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-KIR antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-NKG2A antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the

antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD112 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD47 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-SIRPA antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD244antibody or antigen-binding fragment thereof.

The immune inhibitory molecules may regulate or synergistically regulate T-cell functions to promote tumoral immune escape. Therefore, combination therapies with two or more inhibitors of the inhibitory molecules may provide an improved therapy to a patient when compared to monotherapy alone.

In some embodiments, the antibodies of the invention are administered in combination with an activator of an activating molecule.

In some embodiments, the antibodies of the invention are administered in combination with an activator of an activating molecule CD86, CD80, CD28, ICOS, ICOS ligand, TMIGD2, CD40, GITR ligand, 4-1BB ligand, OX40 ligand, CD70, CD40L, TNFRSF25, LIGHT, GITR, OX-40, CD27, CD137, NKG2D, CD48, CD226 or MICA.

Activation of activating molecules may be performed using for example soluble ligands or ligand derivatives of the activating molecules, peptides or agonistic antibodies.

In some embodiments, the activator of the activating molecule is a soluble ligand of the T cell activating molecule.

In some embodiments, the activator of the activating molecule is an agonistic antibody specifically binding the activating molecule.

Exemplary anti-CD40 antibodies that may be used in the methods of the invention include CP-870,893 and humanized S2C6 described in U.S. Patent No. 7,288,251 (antibody 21.4.1) and U.S. Patent No. 8,303,955, respectively, and anti-CD40 antibodies described in Int. Patent Publ. Nos. WO2001/056603, WO2001/083755, WO2013/034904 and WO2014/070934.

Exemplary GITR agonists include, e.g., GITR fusion proteins and anti-GITR antibodies (e.g., bivalent anti-GITR antibodies), such as, a GITR fusion protein described in U.S. Patent No. 6,111,090, European Patent No. 090505B1, U.S. Patent No. 8,586,023, Int. Patent. Publ. Nos. WO2010/003118 and WO2011/090754, or an anti-GITR antibody described in U.S. Patent Nos. 7,025,962, 7,812,135, 8,388,967, 8,591,886 and 7,618,632, European Patent Nos.1947183 and 1866339, or Int. Patent Publ. Nos. WO2011/028683, WO2013/039954, WO2005/007190, WO2007/133822, WO2005/055808, WO1999/40196, WO2001/03720, WO1999/20758, WO2006/083289, WO2005/115451 and WO2011/051726.

GITR expression was found herein to be elevated in $CD8^+$ T cells isolated from tumors after anti-PD-1 antibody treatment in animal models of cancer. The restoration of GITR expression on TILs by anti-PD-1 treatment supports that combination therapy with anti-GITR and anti-PD-l antibodies may improve the clinical outcome of the patients.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject in need thereof a therapeutically effective amount of an antagonistic antibody that specifically binds PD-1 and an agonistic antibody that specifically binds GITR for a time sufficient to treat the cancer.

In some embodiments, the agonistic antibody that specifically binds GITR is administered after administration of the antagonistic antibody specifically binding PD-1.

In some embodiments, the agonistic antibody that specifically binds GITR and the antagonistic antibody specifically binding PD-1 are administered concurrently as single agents or sequentially as single agents in any order.

Exemplary OX40 antibodies that may be used in the methods of the invention include those described in U.S. Patent Nos. 8,133,983, 7,960,515, U.S. Patent Publ. No. 20130280275 and Int. Patent Publ. Nos. WO2013028231 and WO2014148895.

An exemplary OX40 antibody that may be used in the methods of the invention is an antibody comprising the VH of SEQ ID NO: 309 and the VL of SEQ ID NO: 310.

Another exemplary OX40 antibody that may be used in the methods of the invention is an antibody comprising the VH of SEQ ID NO: 311 and the VL of SEQ ID NO: 312.

OX40 expression was found herein to be elevated in $CD8^+$ T cells isolated from tumors after anti-PD-1 antibody treatment in animal models of cancer. The restoration of OX40 expression on TILs by anti-PD-1 treatment supports that combination therapy with anti-OX40 and anti-PD-l antibodies may improve the clinical outcome of the patients.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject in need thereof a therapeutically effective amount of an antagonistic antibody that specifically binds PD-1 and an agonistic antibody that specifically binds OX40 for a time sufficient to treat the cancer.

In some embodiments, the agonistic antibody that specifically binds OX40 is administered after administration of the antagonistic antibody specifically binding PD-1.

In some embodiments, the agonistic antibody that specifically binds OX40 and the antagonistic antibody specifically binding PD-1 are administered concurrently as single agents or sequentially as single agents in any order.

Exemplary CD70 antibodies that may be used in the methods of the invention include those described in U.S. Patent Publ. No. US20130336976.

Exemplary TNFRSF25 antibodies that may be used in the methods of the invention include those described in U.S. Patent No. 7,708,996.

Exemplary CD27 antibodies that may be used in the methods of the invention include those described in U.S. Patent Publ. No. US20130336976.

Exemplary CD137 antibodies that may be used in the methods of the invention include those described in U.S. Patent Nos. 6,974,863, 6,303,121, 7,138,500, 7,288,638, 8,716,452, 8,821,867 and in U.S. Patent Publ. No. US20130149301.

CD137 expression was found herein to be elevated in $CD8^+$ T cells isolated from tumors after anti-PD-1 antibody treatment in animal models of cancer. The restoration of CD137 expression on TILs by anti-PD-1 treatment supports that combination therapy with anti-CD137 and anti-PD-l antibodies may improve the clinical outcome of the patients.

The invention also provides a method of treating a cancer in a subject, comprising administering to the subject in need thereof a therapeutically effective amount of an antagonistic antibody that specifically binds PD-1 and an agonistic antibody that specifically binds CD137 for a time sufficient to treat the cancer.

In some embodiments, the agonistic antibody that specifically binds CD137 is administered after administration of the antagonistic antibody specifically binding PD-1.

In some embodiments, the agonistic antibody that specifically binds CD137 and the antagonistic antibody specifically binding PD-1 are administered concurrently as single agents or sequentially as single agents in any order.

Exemplary NKG2D antibodies that may be used in the methods of the invention include those described in U.S. Patent Publ. No. US20110150870.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the

antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD86 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD80 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD28 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-ICOS antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-ICOS ligand antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-TMIGD2 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD40 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-GITR ligand antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-4-1BB ligand antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-OX40 ligand antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD70 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD40L antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-TNFRSF25 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-LIGHT antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-GITR antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-OX40 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD27 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the

antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD137 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-NKG2D antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD48 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-CD226 antibody or antigen-binding fragment thereof.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with an anti-MICA antibody or antigen-binding fragment thereof.

The combination of antibodies recited herein can be administered separately, e.g., as separate antibodies, or linked, e.g., as a bispecific or trispecific antibody molecule.

The efficacy of the combinations described herein may be tested in animal models known in the art.

Antibodies of the invention described herein may be administered in combination with a vaccine.

Exemplary vaccines are immunogenic agents, such as cancerous cells, purified tumor antigens (including recombinant proteins, antigen epitopes, peptides and carbohydrate molecules), tumor antigens delivered to a patient via gene therapy, cells, and cells transfected with genes encoding immune stimulating cytokines. Exemplary vaccines that may be used include peptides of melanoma antigens, such as peptides of gp100, MAGE antigens, Trp-2, MART1 and/or tyrosinase, or tumor cells transfected to express the cytokine GM-CSF, DNA-based vaccines, RNA-based vaccines, and viral transduction-based vaccines, peptides or prostate antigens or peptides of lung cancer antigens. The cancer vaccine may be prophylactic or therapeutic.

Many experimental strategies for vaccination against tumors have been devised (see Rosenberg, S., 2000, Development of Cancer Vaccines, ASCO Educational Book

Spring: 60-62; Logothetis, C., 2000, ASCO Educational Book Spring: 300-302; Khayat, D. 2000, ASCO Educational Book Spring: 414-428; Foon, K. 2000, ASCO Educational Book Spring: 730-738; see also Restifo, N. and Sznol, M., Cancer Vaccines, Ch. 61, pp. 3023-3043 in DeVita, V. et al. (eds.), 1997, Cancer: Principles and Practice of Oncology. Fifth Edition). In one of these strategies, a vaccine is prepared using autologous or allogeneic tumor cells. These cellular vaccines have been shown to be most effective when the tumor cells are transduced to express GM-CSF. GM-CSF has been shown to be a potent activator of antigen presentation for tumor vaccination (Dranoff *et al.,* (1993) *Proc Natl Acad Sci* U.S.A. 90: 3539-43).

The antibodies of the invention described herein may be administered in combination with one or a collection of recombinant proteins and/or peptides expressed in or on a tumor in order to generate an immune response to these proteins. These proteins are normally viewed by the immune system as self-antigens and are therefore tolerant to them. The tumor antigen may also include the protein telomerase, which is required for the synthesis of telomeres of chromosomes and which is expressed in more than 85% of human cancers and in only a limited number of somatic tissues (Kim *et al.,* (1994) *Science* 266: 2011-2013). Tumor antigens may also be "neo-antigens" expressed in or on cancer cells as a result of somatic mutations that alter protein sequence or create fusion proteins between two unrelated sequences (e.g., bcr-abl in the Philadelphia chromosome), or idiotype from B cell tumors. The tumor antigens may be antigen epitopes of prostate specific antigen (PSA), mesothelin, prostate-specific membrane antigen (PSMA), synovial sarcoma X2 (SSX2), NKX3.1, prostatic acidic phosphatase (PAP), or epidermal growth factor receptors, or peptides specific for variants of EGFR such as the well-known EGFRvIII overexpressed on tumor cells.

Other tumor vaccines may include the proteins from viruses implicated in human cancers such a Human Papilloma Viruses (HPV), Hepatitis Viruses (HBV and HCV) and Kaposi's Herpes Sarcoma Virus (KHSV), and Epstein-Barr virus (EBV). Another form of tumor specific antigens which may be used in combination with the antibodies of the invention described herein is purified heat shock proteins (HSP) isolated from the tumor tissue itself. HSP contain fragments of proteins from the tumor cells and are highly efficient at delivery to antigen presenting cells for eliciting tumor immunity (Suot and Srivastava (1995) *Science* 269:1585-1588; Tamura *et al.,* (1997) *Science* 278:117-120).

Dendritic cells (DC) are potent antigen presenting cells that may be used to prime antigen-specific responses. DC's may be produced *ex vivo* and loaded with various protein and peptide antigens as well as tumor cell extracts (Nestle *et al.,* (1998) *Nature Medicine*

4: 328-332). DCs may also be transduced by genetic means to express these tumor antigens. DCs have also been fused directly to tumor cells for the purposes of immunization (Kugler *et al.,* (2000) *Nature Medicine* 6:332-336). As a method of vaccination, DC immunization may be effectively combined with the antibodies of the invention described herein to activate more potent anti-tumor responses.

In some embodiments, the vaccine is a polypeptide or a fragment thereof, or a DNA or a RNA encoding the polypeptide or fragment thereof expressed on tumor cells.

In some embodiments, the polypeptide or fragment thereof expressed on tumor cells is PSMA.

In some embodiments, the polypeptide or fragment thereof expressed on tumor cells is mesothelin.

In some embodiments, the polypeptide or fragment thereof expressed on tumor cells is EGFR or EGFR variant such as EGFRvIII.

In some embodiments, the polypeptide or fragment thereof expressed on tumor cells is PAP.

In some embodiments, the polypeptide or fragment thereof expressed on tumor cells is synovial sarcoma X2 (SSX2).

In some embodiments, the polypeptide or fragment thereof expressed on tumor cells is NKX3.1.

In some embodiments, the tumor cells are melanoma, lung cancer, squamous non-small cell lung cancer (NSCLC), non-squamous NSCLC, colorectal cancer, prostate cancer, castration-resistant prostate cancer, ovarian cancer, gastric cancer, liver cancer, pancreatic cancer, thyroid cancer, squamous cell carcinoma of the head and neck, carcinomas of the esophagus or gastrointestinal tract or breast cancer cells.

In some embodiments, the antibodies of the invention are administered in combination with a renal carcinoma (RCC) vaccine.

In some embodiments, the antibodies of the invention are administered in combination with a lung cancer vaccine.

In some embodiments, the antibodies of the invention are administered in combination with a prostate cancer vaccine.

In some embodiments, the antibodies of the invention are administered in combination with a lung cancer vaccine.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in

combination with a tumor vaccine comprising a peptide fragment of EGFR or EGFRvIII, or a vector encoding the peptide fragment of EGFR or EGFRvIII.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with a tumor vaccine comprising a peptide fragment of mesothelin, or a vector encoding the peptide fragment of mesothelin.

In some embodiments, the antagonistic antibodies specifically binding PD-1 of the invention, the antagonistic antibodies specifically binding TIM-3 or the invention, or the antagonistic bispecific PD-1/TIM-3 antibodies of the invention are administered in combination with a tumor vaccine comprising a peptide fragment of prostate specific antigen, or a vector encoding the peptide fragment of prostate specific antigen.

Suitable vectors that may be used in the methods of the invention are well known and include lentiviral vectors, adenoviral vectors, minimal nucleic acid vector (MNAV), vaccinia virus, flow pox virus, Alpha virus-derived VRP, Saccharomyces cerevisiae, MVA, Listeria moonocytogenes, pVAX-based plasmid, see e.g. Pol *et al.,* (2014) *Oncoimmunology* 1(3):e28185.

The antibodies of the invention may be administered in combination with a standard of care cancer treatment.

The antibodies of the invention described herein may be administered in combination with a standard of care cancer chemotherapeutic regimes. In these instances, it may be possible to reduce the dose of chemotherapeutic reagent administered (Mokyr *et al.,* (1998) *Cancer Research* 58: 5301-5304).

In some embodiments, the antibodies of the invention may be administered in combination with one or more of other antibody molecules, chemotherapy, other anti-cancer therapy (e.g., targeted anti-cancer therapies, or oncolytic drugs), cytotoxic agents, cytokines, surgical and/or radiation procedures.

Exemplary cytotoxic agents that may be administered in combination with the antibodies of the invention include antimicrotubule agents, topoisomerase inhibitors, anti-metabolites, mitotic inhibitors, alkylating agents, anthracyclines, vinca alkaloids, intercalating agents, agents capable of interfering with a signal transduction pathway, agents that promote apoptosis, proteosome inhibitors, and radiation (e.g., local or whole body irradiation).

Standard of care therapeutics include anastrozole (Arimidex®), bicalutamide (Casodex®), bleomycin sulfate (Blenoxane®), busulfan (Myleran®), busulfan injection

(Busulfex®), capecitabine (Xeloda®), N4-pentoxycarbonyl-5-deoxy-5-fluorocytidine, carboplatin (Paraplatin®), carmustine (BiCNU®), chlorambucil (Leukeran®), cisplatin (Platinol®), cladribine (Leustatin®), cyclophosphamide (Cytoxan® or Neosar®), cytarabine, cytosine arabinoside (Cytosar-U®), cytarabine liposome injection (DepoCyt®), dacarbazine (DTIC-Dome®), dactinomycin (Actinomycin D, Cosmegan), daunorubicin hydrochloride (Cerubidine®), daunorubicin citrate liposome injection (DaunoXome®), dexamethasone, docetaxel (Taxotere®), doxorubicin hydrochloride (Adriamycin®, Rubex®), etoposide (Vepesid®), fludarabine phosphate (Fludara®), 5-fluorouracil (Adrucil®, Efudex®), flutamide (Eulexin®), tezacitibine, Gemcitabine (difluorodeoxycitidine), hydroxyurea (Hydrea®), Idarubicin (Idamycin®), ifosfamide (IFEX®), irinotecan (Camptosar®), L-asparaginase (ELSPAR®), leucovorin calcium, melphalan (Alkeran®), 6-mercaptopurine (Purinethol®), methotrexate (Folex®), mitoxantrone (Novantrone®), paclitaxel (Taxol®), phoenix (Yttrium90/MX-DTPA), pentostatin, polifeprosan 20 with carmustine implant (Gliadel®), tamoxifen citrate (Nolvadex®), teniposide (Vumon®), 6-thioguanine, thiotepa, tirapazamine (Tirazone®), topotecan hydrochloride for injection (Hycamptin®), vinblastine (Velban®), vincristine (Oncovin®), vinorelbine (Navelbine®), Ibrutinib, idelalisib, and brentuximab vedotin.

Exemplary alkylating agents include, nitrogen mustards, ethylenimine derivatives, alkyl sulfonates, nitrosoureas and triazenes: uracil mustard (Aminouracil Mustard®, Chlorethaminacil®, Demethyldopan®, Desmethyldopan®, Haemanthamine®, Nordopan®, Uracil Nitrogen Mustard®, Uracillost®, Uracilmostaza®, Uramustin®, Uramustine®), chlormethine (Mustargen®), cyclophosphamide (Cytoxan®, Neosar®, Clafen®, Endoxan®, Procytox®, Revimmune™), ifosfamide (Mitoxana®), melphalan (Alkeran®), chlorambucil (Leukeran®), pipobroman (Amedel®, Vercyte®), triethylenemelamine (Hemel®, Hexalen®, Hexastat®), triethylenethiophosphoramine, temozolomide (Temodar®), thiotepa (Thioplex®), busulfan (Busilvex®, Myleran®), carmustine (BiCNU®), lomustine (CeeNU®) and streptozocin (Zanosar®). Additional exemplary alkylating agents include, oxaliplatin (Eloxatin®), temozolomide (Temodar® and Temodal®), dactinomycin (also known as actinomycin-D, Cosmegen®), altretamine (also known as hexamethylmelamine (HMM), Hexalen®), bendamustine (Treanda®), carboplatin (Paraplatin®), lomustine (also known as CCNU, CeeNU®), cisplatin (also known as CDDP, Platinol® and Platinol®-AQ), chlorambucil (Leukeran®), prednumustine, procarbazine (Matulane®),and thiotepa (also known as thiophosphoamide, TESPA and TSPA, Thioplex®).

Exemplary anthracyclines include, e.g., doxorubicin (Adriamycin® and Rubex®); bleomycin (Lenoxane®), daunorubicin (dauorubicin hydrochloride, daunomycin, and rubidomycin hydrochloride, Cerubidine®), daunorubicin liposomal (daunorubicin citrate liposome, DaunoXome®), mitoxantrone (DHAD, Novantrone®), epirubicin (Ellence™), idarubicin (Idamycin®, Idamycin PFS®), mitomycin C (Mutamycin®), geldanamycin, herbimycin, ravidomycin, and desacetylravidomycin.

Exemplary vinca alkaloids that may be used in combination with the antibodies of the invention include vinorelbine tartrate (Navelbine®), vincristine (Oncovin®), and vindesine (Eldisine®), vinblastine (also known as vinblastine sulfate, vincaleukoblastine and VLB, Alkaban-AQ® and Velban®) and vinorelbine (Navelbine®).

Exemplary proteosome inhibitors that may be used in combination with the antibodies of the invention are bortezomib (Velcade®); carfilzomib (Kyprolis®), ixazomib (Ninlaro®), marizomib (NPI-0052) and delanzomib (CEP-18770).

In some embodiments, the antibodies of the invention are administered in combination with a tyrosine kinase inhibitor (e.g., a receptor tyrosine kinase (RTK) inhibitor). Exemplary tyrosine kinase inhibitor include an epidermal growth factor (EGF) pathway inhibitor (e.g., an epidermal growth factor receptor (EGFR) inhibitor), a vascular endothelial growth factor (VEGF) pathway inhibitor (e.g., a vascular endothelial growth factor receptor (VEGFR) inhibitor (e.g., a VEGFR-1 inhibitor, a VEGFR-2 inhibitor, a VEGFR-3 inhibitor), a platelet derived growth factor (PDGF) pathway inhibitor (e.g., a platelet derived growth factor receptor (PDGFR) inhibitor (e.g., a PDGFR-β inhibitor), a RAF-1 inhibitor, a KIT inhibitor and a RET inhibitor. In some embodiments, the second therapeutic is axitinib (AG013736), bosutinib (SKI-606), cediranib (RECENTIN™, AZD2171), dasatinib (SPRYCEL®, BMS-354825), erlotinib (TARCEVA®), gefitinib (IRESSA®), imatinib (Gleevec®, CGP57148B, STI-571), lapatinib (TYKERB®, TYVERB®), lestaurtinib (CEP-701), neratinib (HKI-272), nilotinib (TASIGNA®), semaxanib (semaxinib, SU5416), sunitinib (SUTENT®, SU11248), toceranib (PALLADIA®), vandetanib (ZACTIMA®, ZD6474), vatalanib (PTK787, PTK/ZK), trastuzumab (HERCEPTIN®), bevacizumab (AVASTIN®), rituximab (RITUXAN®), cetuximab (ERBITUX®), panitumumab (VECTIBIX®), ranibizumab (Lucentis®), nilotinib (TASIGNA®), sorafenib (NEXAVAR®), alemtuzumab (CAMPATH®), gemtuzumab ozogamicin (MYLOTARG®), ENMD-2076, PCI-32765, AC220, dovitinib lactate (TKI258, CHIR-258), BIBW 2992 (TOVOK™), SGX523, PF-04217903, PF-02341066, PF-299804, BMS-777607, ABT-869, MP470, BIBF 1120 (VARGATEF®), AP24534, JNJ-26483327, MGCD265, DCC-2036, BMS-690154, CEP-11981, tivozanib

(AV-951), OSI-930, MM-121, XL-184, XL-647, XL228, AEE788, AG-490, AST-6, BMS-599626, CUDC-101, PD153035, pelitinib (EKB-569), vandetanib (zactima), WZ3146, WZ4002, WZ8040, ABT-869 (linifanib), AEE788, AP24534 (ponatinib), AV-951 (tivozanib), axitinib, BAY 73-4506 (regorafenib), brivanib alaninate (BMS-582664), brivanib (BMS-540215), cediranib (AZD2171), CHIR-258 (dovitinib), CP 673451, CYC116, E7080, Ki8751, masitinib (AB1010), MGCD-265, motesanib diphosphate (AMG-706), MP-470, OSI-930, pazopanib hydrochloride, PD173074, Sorafenib Tosylate (Bay 43-9006), SU 5402, TSU-68 (SU6668), vatalanib, XL880 (GSK1363089, EXEL-2880). Selected tyrosine kinase inhibitors are chosen from sunitinib, erlotinib, gefitinib, or sorafenib. In some embodiments, the EGFR inhibitor is a bispecific EGFRc-Met antibody (EM-1 mAb) comprising the heavy and the light chains of SEQ ID NOs: 249, 250, 251 and 252 (US2014/0141000).

In some embodiments, the antibodies of the invention are administered in combination with Vascular Endothelial Growth Factor (VEGF) receptor inhibitors, including bevacizumab (Avastin®), axitinib (Inlyta®), brivanib alaninate (BMS-582664, (S)—((R)-1-(4-(4-Fluoro-2-methyl-1H-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy)propan-2-yl)2-aminopropanoate), sorafenib (Nexavar®); Pazopanib (Votrient®), sunitinib malate (Sutent®), cediranib (AZD2171, CAS 288383-20-1), vargatef (BIBF1120, CAS 928326-83-4), foretinib (GSK1363089), telatinib (BAY57-9352, CAS 332012-40-5), apatinib (YN968D1, CAS 811803-05-1), imatinib (Gleevec®), ponatinib (AP24534, CAS 943319-70-8), tivozanib (AV951, CAS 475108-18-0), regorafenib (BAY73-4506, CAS 755037-03-7), vatalanib dihydrochloride (PTK787, CAS 212141-51-0), brivanib (BMS-540215, CAS 649735-46-6), vandetanib (Caprelsa® or AZD6474), motesanib diphosphate (AMG706, CAS 857876-30-3, N-(2,3-dihydro-3,3-dimethyl-1H-indol-6-yl)-2-[(4-pyridinylmethyl)amino]-3-pyridinecarboxamide, described in PCT Publication No. WO 02/066470), dovitinib dilactic acid (TKI258, CAS 852433-84-2), linfanib (ABT869, CAS 796967-16-3); Cabozantinib (XL184, CAS 849217-68-1), lestaurtinib (CAS 111358-88-4); N-[5-[[[5-(1,1-Dimethylethyl)-2-oxazolyl]methyl]thio]-2-thiazolyl]-4-piperidinecarboxamide (BMS38703, CAS 345627-80-7); (3R,4R)-4-Amino-1-((4-((3-methoxyphenyl)amino)pyrrolo[2,1-f][1,2,4]triazin-5-yl)methyl)piperidin-3-ol (BMS690514); N-(3,4-Dichloro-2-fluorophenyl)-6-methoxy-7-[[(3aα,5β,6aα)-octahydro-2-methylcyclopenta[c]pyrrol-5-yl]methoxy]-4-quinazolinamine (XL647, CAS 781613-23-8); 4-Methyl-3-[[1-methyl-6-(3-pyridinyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl]amino]-N-

[3-(trifluoromethyl)phenyl]-benzamide (BHG712, CAS 940310-85-0); and aflibercept (Eylea®).

In some embodiments, the antibodies of the invention are administered in combination with a PI3K inhibitor. In one embodiment, the PI3K inhibitor is an inhibitor of delta and gamma isoforms of PI3K. Exemplary PI3K inhibitors that may be used are described in, e.g., WO 2010/036380, WO 2010/006086, WO 09/114870, WO 05/113556, GSK 2126458, GDC-0980, GDC-0941, Sanofi XL147, XL756, XL147, PF-46915032, BKM 120, CAL-101, CAL 263, SF1126, PX-886, and a dual PI3K inhibitor (e.g., Novartis BEZ235).

In some embodiments, the antibodies of the invention are administered in combination with a mTOR inhibitor, e.g., one or more mTOR inhibitors chosen from one or more of rapamycin, temsirolimus (TORISEL®), AZD8055, BEZ235, BGT226, XL765, PF-4691502, GDC0980, SF1126, OSI-027, GSK1059615, KU-0063794, WYE-354, Palomid 529 (P529), PF-04691502, or PKI-587. ridaforolimus (formally known as deferolimus, (1R,2R,4S)-4-[(2R)-2 [(1R,9S,12S,15R,16E,18R,19R,21R,23S,24E,26E,28Z,30S,32S,35R)-1,18-dihydroxy-19,30-dimethoxy-15,17,21,23,29,35-hexamethyl-2,3,10,14,20-pentaoxo-11,36-dioxa-4-azatricyclo[30.3.1.04,9] hexatriaconta-16,24,26,28-tetraen-12-yl]propyl]-2-methoxycyclohexyl dimethylphosphinate, also known as AP23573 and MK8669, and described in PCT Publication No. WO 03/064383); everolimus (Afinitor® or RAD001); rapamycin (AY22989, Sirolimus®); simapimod (CAS 164301-51-3); emsirolimus, (5-{2,4-Bis[(3S)-3-methylmorpholin-4-yl]pyrido[2,3-d]pyrimidin-7-yl}-2-methoxyphenyl)methanol (AZD8055); 2-Amino-8-[trans-4-(2-hydroxyethoxy)cyclohexyl]-6-(6-methoxy-3-pyridinyl)-4-methyl-pyrido[2,3-d]pyrimidin-7(8H)-one (PF04691502, CAS 1013101-36-4); and N2-[1,4-dioxo-4-[[4-(4-oxo-8-phenyl-4H-1-benzopyran-2-yl)morpholinium-4-yl]methoxy]butyl]-L-arginylglycyl-L-α-aspartylL-serine- (SEQ ID NO: 237), inner salt (SF1126, CAS 936487-67-1), and XL765.

In some embodiments, the antibodies of the invention are administered in combination with a BRAF inhibitor, e.g., GSK2118436, RG7204, PLX4032, GDC-0879, PLX4720, and sorafenib tosylate (Bay 43-9006).

In some embodiments, the antibodies of the invention are administered in combination with a MEK inhibitor.

In some embodiments, the antibodies of the invention are administered in combination with a JAK2 inhibitor, e.g., CEP-701, INCB18424, CP-690550 (tasocitinib).

In some embodiments, the antibodies of the invention are administered in combination with paclitaxel or a paclitaxel agent, e.g., TAXOL®, protein-bound paclitaxel (e.g., ABRAXANE®). Exemplary paclitaxel agents include nanoparticle albumin-bound paclitaxel (ABRAXANE, marketed by Abraxis Bioscience), docosahexaenoic acid bound-paclitaxel (DHA-paclitaxel, Taxoprexin, marketed by Protarga), polyglutamate bound-paclitaxel (PG-paclitaxel, paclitaxel poliglumex, CT-2103, XYOTAX, marketed by Cell Therapeutic), the tumor-activated prodrug (TAP), ANG105 (Angiopep-2 bound to three molecules of paclitaxel, marketed by ImmunoGen), paclitaxel-EC-1 (paclitaxel bound to the erbB2-recognizing peptide EC-1; see Li et al., Biopolymers (2007) 87:225-230), and glucose-conjugated paclitaxel (e.g., 2′-paclitaxel methyl 2-glucopyranosyl succinate, see Liu *et al.,* (2007) *Bioorganic & Medicinal Chemistry Letters* 17:617-620).

In some embodiments, the antibodies of the invention are administered in combination with a cellular immunotherapy (e.g., Provenge (e.g., Sipuleucel)), and optionally in combination with cyclophosphamide.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of pancreatic cancer include a chemotherapeutic agent, e.g., paclitaxel or a paclitaxel agent (e.g., a paclitaxel formulation such as TAXOL, an albumin-stabilized nanoparticle paclitaxel formulation (e.g., ABRAXANE) or a liposomal paclitaxel formulation); gemcitabine (e.g., gemcitabine alone or in combination with AXP107-11); other chemotherapeutic agents such as oxaliplatin, 5-fluorouracil, capecitabine, rubitecan, epirubicin hydrochloride, NC-6004, cisplatin, docetaxel (e.g., TAXOTERE), mitomycin C, ifosfamide; interferon; tyrosine kinase inhibitor (e.g., EGFR inhibitor (e.g., erlotinib, panitumumab, cetuximab, nimotuzumab); HER2/neu receptor inhibitor (e.g., trastuzumab); dual kinase inhibitor (e.g., bosutinib, saracatinib, lapatinib, vandetanib); multikinase inhibitor (e.g., sorafenib, sunitinib, XL184, pazopanib); VEGF inhibitor (e.g., bevacizumab, AV-951, brivanib); radioimmunotherapy (e.g., XR303); cancer vaccine (e.g., GVAX, survivin peptide); COX-2 inhibitor (e.g., celecoxib); IGF-1 receptor inhibitor (e.g., AMG 479, MK-0646); mTOR inhibitor (e.g., everolimus, temsirolimus), IL-6 inhibitor (e.g., CNTO 328); cyclin-dependent kinase inhibitor (e.g., P276-00, UCN-01); Altered Energy Metabolism-Directed (AEMD) compound (e.g., CPI-613); HDAC inhibitor (e.g., vorinostat); TRAIL receptor 2 (TR-2) agonist (e.g., conatumumab); MEK inhibitor (e.g., AS703026, selumetinib, GSK1120212); Raf/MEK dual kinase inhibitor (e.g., RO5126766), Notch signaling inhibitor (e.g., MK0752), monoclonal antibody-antibody fusion protein (e.g., L19IL2), curcumin; HSP90 inhibitor (e.g., tanespimycin, STA-9090), rIL-2; denileukin diftitox; topoisomerase 1 inhibitor (e.g.,

irinotecan, PEP02); statin (e.g., simvastatin), Factor VIIa inhibitor (e.g., PCI-27483), AKT inhibitor (e.g., RX-0201), hypoxia-activated prodrug (e.g., TH-302), metformin hydrochloride, gamma-secretase inhibitor (e.g., R04929097), ribonucleotide reductase inhibitor (e.g., 3-AP), immunotoxin (e.g., HuC242-DM4), PARP inhibitor (e.g., KU-0059436, veliparib), CTLA-4 inhbitor (e.g., CP-675,206, ipilimumab), AdV-tk therapy, proteasome inhibitor (e.g., bortezomib (Velcade), NPI-0052), thiazolidinedione (e.g., pioglitazone), NPC-1C; Aurora kinase inhibitor (e.g., R763/AS703569), CTGF inhibitor (e.g., FG-3019), siG12D LODER and radiation therapy (e.g., tomotherapy, stereotactic radiation, proton therapy), surgery, and a combination thereof. In certain embodiments, a combination of paclitaxel or a paclitaxel agent, and gemcitabine can be used with the antibodies of the invention.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of small cell lung cancer (SCLC) include approved drugs for treatment of SCLC such as methotrexate (Folex®, Mexate®), everolimus (Afinitor®), doxorubicin hydrochloride, etoposide phosphate (Etopophos®), topotecan hydrochloride (Hycamtin®), mechlorethamine hydrochloride (Mustargen®), topotecan hydrochloride. Other therapeutic agents that may be used are carboplatin, cisplatin, oxaliplatin, irinotecan, gemcitabine, liposomal SN-38, bendamustine, temozolomide, belotecan, NK012, FR901228, flavopiridol), tyrosine kinase inhibitor (e.g., EGFR inhibitor (e.g., erlotinib, gefitinib, cetuximab, panitumumab), multikinase inhibitor (e.g., sorafenib, sunitinib), VEGF inhibitor (e.g., bevacizumab, vandetanib), cancer vaccine (e.g., GVAX); Bcl-2 inhibitor (e.g., oblimersen sodium, ABT-263), proteasome inhibitor (e.g., bortezomib (Velcade), NPI-0052), paclitaxel or a paclitaxel agent; docetaxel, IGF-1 receptor inhibitor (e.g., AMG 479), HGF/SF inhibitor (e.g., AMG 102, MK-0646), chloroquine, Aurora kinase inhibitor (e.g., MLN8237), radioimmunotherapy (e.g., TF2), HSP90 inhibitor (e.g., tanespimycin, STA-9090), mTOR inhibitor (e.g., everolimus), Ep-CAM/CD3-bispecific antibody (e.g., MT110), CK-2 inhibitor (e.g., CX-4945), HDAC inhibitor (e.g., belinostat), SMO antagonist (e.g., BMS 833923), peptide cancer vaccine, and radiation therapy (e.g., intensity-modulated radiation therapy (IMRT), hypofractionated radiotherapy, hypoxia-guided radiotherapy), surgery, and combinations thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of non-small cell lung cancer include approved drugs for treatment of NSCLC including methotrexate (Folex®, Mexate®), paclitaxel (Abraxane®), afatinib (Gilotrif®), everolimus (Afinitor®), alectinib (Alecensa®), pemetrexed disodium (Alimta®), bevacizumab (Avastin®), carboplatin, ceritinib (Zykadia®), crizotinib

(Xalkori®), ramucirumab (Cyramza®), docetaxel, everolimus (Afinitor®),gefitinib (Iressa®), afatinib dimaleate (Gilotrif®), gemcitabine hydrochloride (Gmezar®), pembrolizumab (Keytruda®), mechlorethamine hydrochloride (Mustargen®), vinorelbine tartrate (Navelbine®), necitumumab (Portrazza®), nivolumab (Opdivo®), osimertinib, paclitaxel (Taxol®), carboplatin, pemetrexed disodium, ramucirumab (Cyramza®), osimertinib (Tagrisso®). Other therapeutic agents that may be used are vinorelbine, cisplatin, docetaxel, pemetrexed disodium, etoposide, gemcitabine, carboplatin, liposomal SN-38, TLK286, temozolomide, topotecan, pemetrexed disodium, azacitidine, irinotecan, tegafur-gimeracil-oteracil potassium, sapacitabine), tyrosine kinase inhibitor (e.g., EGFR inhibitor (e.g., erlotinib, gefitinib, cetuximab, panitumumab, necitumumab, PF-00299804, nimotuzumab, RO5083945), MET inhibitor (e.g., PF-02341066, ARQ 197), PI3K kinase inhibitor (e.g., XL147, GDC-0941), Raf/MEK dual kinase inhibitor (e.g., RO5126766), PI3K/mTOR dual kinase inhibitor (e.g., XL765), SRC inhibitor (e.g., dasatinib), dual inhibitor (e.g., BIBW 2992, GSK1363089, ZD6474, AZD0530, AG-013736, lapatinib, MEHD7945A, linifanib), multikinase inhibitor (e.g., sorafenib, sunitinib, pazopanib, AMG 706, XL184, MGCD265, BMS-690514, R935788), VEGF inhibitor (e.g., endostar, endostatin, bevacizumab, cediranib, BIBF 1120, axitinib, tivozanib, AZD2171), cancer vaccine (e.g., BLP25 liposome vaccine, GVAX, recombinant DNA and adenovirus expressing L523S protein), Bcl-2 inhibitor (e.g., oblimersen sodium), proteasome inhibitor (e.g., bortezomib, carfilzomib, NPI-0052, MLN9708), paclitaxel or a paclitaxel agent, docetaxel, IGF-1 receptor inhibitor (e.g., cixutumumab, MK-0646, OSI 906, CP-751,871, BIIB022), hydroxychloroquine, HSP90 inhibitor (e.g., tanespimycin, STA-9090, AUY922, XL888), mTOR inhibitor (e.g., everolimus, temsirolimus, ridaforolimus), Ep-CAM/CD3-bispecific antibody (e.g., MT110), CK-2 inhibitor (e.g., CX-4945), HDAC inhibitor (e.g., MS 275, LBH589, vorinostat, valproic acid, FR901228), DHFR inhibitor (e.g., pralatrexate), retinoid (e.g., bexarotene, tretinoin), antibody-drug conjugate (e.g., SGN-15), bisphosphonate (e.g., zoledronic acid), cancer vaccine (e.g., belagenpumatucel-L), low molecular weight heparin (LMWH) (e.g., tinzaparin, enoxaparin), GSK1572932A, melatonin, talactoferrin, dimesna, topoisomerase inhibitor (e.g., amrubicin, etoposide, karenitecin), nelfinavir, cilengitide, ErbB3 inhibitor (e.g., MM-121, U3-1287), survivin inhibitor (e.g., YM155, LY2181308), eribulin mesylate, COX-2 inhibitor (e.g., celecoxib), pegfilgrastim, Polo-like kinase 1 inhibitor (e.g., BI 6727), TRAIL receptor 2 (TR-2) agonist (e.g., CS-1008), CNGRC peptide (SEQ ID NO: 225)-TNF alpha conjugate, dichloroacetate (DCA), HGF inhibitor (e.g., SCH 900105), SAR240550, PPAR-gamma agonist (e.g., CS-7017), gamma-secretase inhibitor (e.g., RO4929097), epigenetic therapy

(e.g., 5-azacitidine), nitroglycerin, MEK inhibitor (e.g., AZD6244), cyclin-dependent kinase inhibitor (e.g., UCN-01), cholesterol-Fus1, antitubulin agent (e.g., E7389), farnesyl-OH-transferase inhibitor (e.g., lonafarnib), immunotoxin (e.g., BB-10901, SS1 (dsFv) PE38), fondaparinux, vascular-disrupting agent (e.g., AVE8062), PD-L1 inhibitor (e.g., MDX-1105, MDX-1106), beta-glucan, NGR-hTNF, EMD 521873, MEK inhibitor (e.g., GSK1120212), epothilone analog (e.g., ixabepilone), kinesin-spindle inhibitor (e.g., 4SC-205), telomere targeting agent (e.g., KML-001), P70 pathway inhibitor (e.g., LY2584702), AKT inhibitor (e.g., MK-2206), angiogenesis inhibitor (e.g., lenalidomide), Notch signaling inhibitor (e.g., OMP-21M18), EGFR/c-Met bispecific antibody EM-1 as described in US2014/0141000A1 , radiation therapy, surgery, and combinations thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of ovarian cancer include approved drugs for treatment of ovarian cancer, such as melphalan (Alkeran®), bevacizumab (Avastin®), carboplatin, cyclophosphamide (Clafen®, Cytoxan®), clisplatin, doxorubicin hydrochloride, gemcitabine hydrochloride (Gemzar®), topotecan hydrochloride (Hycamtin®), Olaparib (Lynparza®), carboplatin, cisplatin, paclitaxel (Taxol®), thiotepa and topotecan hydrochloride. Other therapeutic agents that may be used are, ifosfamide, olaparib, oxaliplatin, pemetrexed disodium, SJG-136, etoposide, decitabine; immunotherapy (e.g., APC8024, oregovomab, OPT-821), tyrosine kinase inhibitor (e.g., EGFR inhibitor (e.g., erlotinib), dual inhibitor (e.g., E7080), multikinase inhibitor (e.g., AZD0530, JI-101, sorafenib, sunitinib, pazopanib), VEGF inhibitor (e.g., bevacizumab, BIBF 1120, cediranib, AZD2171), PDGFR inhibitor (e.g., IMC-3G3), paclitaxel, topoisomerase inhibitor (e.g., karenitecin, Irinotecan), HDAC inhibitor (e.g., valproate, vorinostat), folate receptor inhibitor (e.g., farletuzumab), angiopoietin inhibitor (e.g., AMG 386), epothilone analog (e.g., ixabepilone), proteasome inhibitor (e.g., carfilzomib), IGF-1 receptor inhibitor (e.g., OSI 906, AMG 479), PARP inhibitor (e.g., veliparib, AG014699, iniparib, MK-4827), Aurora kinase inhibitor (e.g., MLN8237, ENMD-2076), angiogenesis inhibitor (e.g., lenalidomide), DHFR inhibitor (e.g., pralatrexate), radioimmunotherapeutic agent (e.g., Hu3S193), statin (e.g., lovastatin), topoisomerase 1 inhibitor (e.g., NKTR-102), cancer vaccine (e.g., p53 synthetic long peptides vaccine, autologous OC-DC vaccine), mTOR inhibitor (e.g., temsirolimus, everolimus), BCR/ABL inhibitor (e.g., imatinib), ET-A receptor antagonist (e.g., ZD4054), TRAIL receptor 2 (TR-2) agonist (e.g., CS-1008), HGF/SF inhibitor (e.g., AMG 102), EGEN-001, Polo-like kinase 1 inhibitor (e.g., BI 6727), gamma-secretase inhibitor (e.g., RO4929097), Wee-1 inhibitor (e.g., MK-1775), antitubulin agent (e.g., vinorelbine, E7389), immunotoxin (e.g., denileukin diftitox), SB-

485232, vascular-disrupting agent (e.g., AVE8062), integrin inhibitor (e.g., EMD 525797), kinesin-spindle inhibitor (e.g., 4SC-205), revlimid, HER2 inhibitor (e.g., MGAH22), ErrB3 inhibitor (e.g., MM-121), radiation therapy, and combinations thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of a myeloma include one or more of chemotherapy or other anti-cancer agents (e.g., thalidomide analogs, e.g., lenalidomide), HSCT (Cook, (2008) J *Manag Care Pharm.* 14(7 Suppl):19-25), an anti-TIM-3 antibody (Hallett *et al,.* (2011) *J of American Society for Blood and Marrow Transplantation* 17(8):1133-145), tumor antigen-pulsed dendritic cells, fusions (e.g., electrofusions) of tumor cells and dendritic cells, or vaccination with immunoglobulin idiotype produced by malignant plasma cells (reviewed in Yi (2009) *Cancer J* 15(6):502-10).

Exemplary therapeutics agents that may be used in combination with the antibodies of the invention for treatment of a renal cancer, e.g., a renal cell carcinoma (RCC) or metastatic RCC include drugs approved for treatment of RCC, including everolimus (Afinitor®), aldesleukin, bevacizumab (Avastin®), axitinib (Inlyta®), cabozantinib-S-Malate (Cabometyx®), aldesleukin (Proleukin®), lenvatinib mesylate (Lenvima®), sorafenib tosylate (Nexavar®), nivolumab (Opdivo®), pazopanib hydrochloride, sorafenib tosylate, sunitinib (Sutent®), temsirolimus (Torisel®) and pazopanib hydrochloride (Votrient®). Other therapeutics that may be used are a targeted agent (e.g., a VEGF inhibitor such as a monoclonal antibody to VEGF, e.g., bevacizumab, a VEGF tyrosine kinase inhibitor such as sorafenib, axitinib and pazopanib.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of a chronic myelogenous leukemia (AML) include a chemotherapeutic (e.g., cytarabine, hydroxyurea, clofarabine, melphalan, thiotepa, fludarabine, busulfan, etoposide, cordycepin, pentostatin, capecitabine, azacitidine, cyclophosphamide, cladribine, topotecan), tyrosine kinase inhibitor (e.g., BCR/ABL inhibitor (e.g., imatinib, nilotinib), dual inhibitor (e.g., dasatinib, bosutinib), multikinase inhibitor (e.g., DCC-2036, ponatinib, sorafenib, sunitinib, RGB-286638), interferon alfa, steroids, apoptotic agent (e.g., omacetaxine mepesuccinat), immunotherapy (e.g., allogeneic CD4+ memory Th1-like T cells/microparticle-bound anti-CD3/anti-CD28, autologous cytokine induced killer cells (CIK), AHN-12), CD52 targeting agent (e.g., alemtuzumab), HSP90 inhibitor (e.g., tanespimycin, STA-9090, AUY922, XL888), mTOR inhibitor (e.g., everolimus), SMO antagonist (e.g., BMS 833923), ribonucleotide reductase inhibitor (e.g., 3-AP), JAK-2 inhibitor (e.g., INCB018424), hydroxychloroquine, retinoid (e.g., fenretinide), cyclin-dependent kinase inhibitor (e.g., UCN-01), HDAC inhibitor (e.g.,

belinostat, vorinostat, JNJ-26481585), PARP inhibitor (e.g., veliparib), MDM2 antagonist (e.g., RO5045337), Aurora B kinase inhibitor (e.g., TAK-901), radioimmunotherapy (e.g., actinium-225-labeled anti-CD33 antibody HuM 195), Hedgehog inhibitor (e.g., PF-04449913), STAT3 inhibitor (e.g., OPB-31121), KB004, cancer vaccine (e.g., AG858), bone marrow transplantation, stem cell transplantation, radiation therapy, and combinations thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of a chronic lymphocytic leukemia (CLL) include a chemotherapeutic agent (e.g., fludarabine, cyclophosphamide, doxorubicin, vincristine, chlorambucil, bendamustine, chlorambucil, busulfan, gemcitabine, melphalan, pentostatin, mitoxantrone, 5-azacytidine, pemetrexed disodium), tyrosine kinase inhibitor (e.g., EGFR inhibitor (e.g., erlotinib), BTK inhibitor (e.g., PCI-32765 (ibrutinib), multikinase inhibitor (e.g., MGCD265, RGB-286638), CD-20 targeting agent (e.g., rituximab, ofatumumab, RO5072759, LFB-R603), CD52 targeting agent (e.g., alemtuzumab), prednisolone, darbepoetin alfa, lenalidomide, Bcl-2 inhibitor (e.g., ABT-263), immunotherapy (e.g., allogeneic $CD4^+$ memory Th1-like T cells/microparticle-bound anti-CD3/anti-CD28, autologous cytokine induced killer cells (CIK), HDAC inhibitor (e.g., vorinostat, valproic acid, LBH589, JNJ-26481585, AR-42), XIAP inhibitor (e.g., AEG35156), CD-74 targeting agent (e.g., milatuzumab), mTOR inhibitor (e.g., everolimus), AT-101, immunotoxin (e.g., CAT-8015, anti-Tac(Fv)-PE38 (LMB-2)), CD37 targeting agent (e.g., TRU-016), radioimmunotherapy (e.g., 131-tositumomab), hydroxychloroquine, perifosine, SRC inhibitor (e.g., dasatinib), thalidomide, PI3K delta inhibitor (e.g., CAL-101), retinoid (e.g., fenretinide), MDM2 antagonist (e.g., RO5045337), plerixafor, Aurora kinase inhibitor (e.g., MLN8237, TAK-901), proteasome inhibitor (e.g., bortezomib), CD-19 targeting agent (e.g., MEDI-551, MOR208), MEK inhibitor (e.g., ABT-348), JAK-2 inhibitor (e.g., INCB018424), hypoxia-activated prodrug (e.g., TH-302), paclitaxel or a paclitaxel agent, HSP90 inhibitor, AKT inhibitor (e.g., MK2206), HMG-CoA inhibitor (e.g., simvastatin), GNKG186, radiation therapy, bone marrow transplantation, stem cell transplantation, and a combination thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of an acute lymphocytic leukemia (ALL) include a chemotherapeutic agent (e.g., prednisolone, dexamethasone, vincristine, asparaginase, daunorubicin, cyclophosphamide, cytarabine, etoposide, thioguanine, mercaptopurine, clofarabine, liposomal annamycin, busulfan, etoposide, capecitabine, decitabine, azacitidine, topotecan, temozolomide), tyrosine kinase inhibitor (e.g., BCR/ABL inhibitor

(e.g., imatinib, nilotinib), ON 01910.Na, multikinase inhibitor (e.g., sorafenib), CD-20 targeting agent (e.g., rituximab), CD52 targeting agent (e.g., alemtuzumab), HSP90 inhibitor (e.g., STA-9090), mTOR inhibitor (e.g., everolimus, rapamycin), JAK-2 inhibitor (e.g., INCB018424), HER2/neu receptor inhibitor (e.g., trastuzumab), proteasome inhibitor (e.g., bortezomib), methotrexate, asparaginase, CD-22 targeting agent (e.g., epratuzumab, inotuzumab), immunotherapy (e.g., autologous cytokine induced killer cells (CIK), AHN-12), blinatumomab, cyclin-dependent kinase inhibitor (e.g., UCN-01), CD45 targeting agent (e.g., BC8), MDM2 antagonist (e.g., RO5045337), immunotoxin (e.g., CAT-8015, DT2219ARL), HDAC inhibitor (e.g., JNJ-26481585), JVRS-100, paclitaxel or a paclitaxel agent, STAT3 inhibitor (e.g., OPB-31121), PARP inhibitor (e.g., veliparib), EZN-2285, radiation therapy, steroid, bone marrow transplantation, stem cell transplantation, or a combination thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of an acute myeloid leukemia (AML) include a chemotherapeutic agent (e.g., cytarabine, daunorubicin, idarubicin, clofarabine, decitabine, vosaroxin, azacitidine, clofarabine, ribavirin, CPX-351, treosulfan, elacytarabine, azacitidine), tyrosine kinase inhibitor (e.g., BCR/ABL inhibitor (e.g., imatinib, nilotinib), ON 01910.Na, multikinase inhibitor (e.g., midostaurin, SU 11248, quizartinib, sorafinib), immunotoxin (e.g., gemtuzumab ozogamicin), DT388IL3 fusion protein, HDAC inhibitor (e.g., vorinostat, LBH589), plerixafor, mTOR inhibitor (e.g., everolimus), SRC inhibitor (e.g., dasatinib), HSP90 inhbitor (e.g., STA-9090), retinoid (e.g., bexarotene, Aurora kinase inhibitor (e.g., BI 811283), JAK-2 inhibitor (e.g., INCB018424), Polo-like kinase inhibitor (e.g., BI 6727), cenersen, CD45 targeting agent (e.g., BC8), cyclin-dependent kinase inhibitor (e.g., UCN-01), MDM2 antagonist (e.g., RO5045337), mTOR inhibitor (e.g., everolimus), LY573636-sodium, ZRx-101, MLN4924, lenalidomide, immunotherapy (e.g., AHN-12), histamine dihydrochloride, radiation therapy, bone marrow transplantation, stem cell transplantation, and a combination thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of a multiple myeloma (MM) include a chemotherapeutic agent (e.g., melphalan, amifostine, cyclophosphamide, doxorubicin, clofarabine, bendamustine, fludarabine, adriamycin, SyB L-0501), thalidomide, lenalidomide, dexamethasone, prednisone, pomalidomide, proteasome inhibitor (e.g., bortezomib, carfilzomib, MLN9708), cancer vaccine (e.g., GVAX), CD-40 targeting agent (e.g., SGN-40, CHIR-12.12), perifosine, zoledronic acid, Immunotherapy (e.g., MAGE-A3, NY-ESO-1, HuMax-CD38), HDAC inhibitor (e.g., vorinostat, LBH589, AR-42), aplidin, cycline-

dependent kinase inhibitor (e.g., PD-0332991, dinaciclib), arsenic trioxide, CB3304, HSP90 inhibitor (e.g., KW-2478), tyrosine kinase inhibitor (e.g., EGFR inhibitor (e.g., cetuximab), multikinase inhibitor (e.g., AT9283), VEGF inhibitor (e.g., bevacizumab), plerixafor, MEK inhibitor (e.g., AZD6244), IPH2101, atorvastatin, immunotoxin (e.g., BB-10901), NPI-0052, radioimmunotherapeutic (e.g., yttrium Y 90 ibritumomab tiuxetan), STAT3 inhibitor (e.g., OPB-31121), MLN4924, Aurora kinase inhibitor (e.g., ENMD-2076), IMGN901, ACE-041, CK-2 inhibitor (e.g., CX-4945), an anti-CD38 antibody (e.g. DARZALEX® (daratumumab), radiation therapy, bone marrow transplantation, stem cell transplantation, and a combination thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of a prostate cancer are approved drugs for treatment of the prostate cancer, such as abiraterone acetate (Zytiga®), bicalutamide (Casodex®), cabazitaxel (Jevtana®), conjugated estrogens (Premarin®), stradiol (Estrace®), estradiol valerate (Delestrogen®), estrogens, esterified (Menest®), degarelix (Firmagon®), docetaxel (Taxotere®), enzalutamide (Xtandi®), flutamide, goserelin acetate (Zoladex®), Cabazitaxel (Jevtana®), leuprolide acetate (Lupron®), mitoxantrone hydrochloride, nilutamide (Nilandron®) Sipuleucel-T (Provenge®) and radium 223 dichloride (Xofigo®). Other drugs that may be used include a chemotherapeutic agent (e.g., carboplatin, fludarabine), hormonal therapy (e.g., cyproterone acetate, ketoconazole, aminoglutethimide, abarelix, degarelix, leuprolide, triptorelin, buserelin), tyrosine kinase inhibitor (e.g., dual kinase inhibitor (e.g., lapatanib), multikinase inhibitor (e.g., sorafenib, sunitinib), VEGF inhibitor (e.g., bevacizumab), TAK-700, cancer vaccine (e.g., BPX-101, PEP223), lenalidomide, TOK-001, IGF-1 receptor inhibitor (e.g., cixutumumab), TRC105, Aurora A kinase inhibitor (e.g., MLN8237), proteasome inhibitor (e.g., bortezomib), OGX-011, radioimmunotherapy (e.g., HuJ591-GS), HDAC inhibitor (e.g., valproic acid, SB939, LBH589), hydroxychloroquine, mTOR inhibitor (e.g., everolimus), dovitinib lactate, diindolylmethane, efavirenz, OGX-427, genistein, IMC-3G3, bafetinib, CP-675,206, radiation therapy, surgery, or a combination thereof.

Exemplary therapeutic agents that may be used in combination with the antibodies of the invention for treatment of a head and neck squamous cell carcinoma (HNSCC) include methotrexate (Folex®, Mexate®), bleomycin (Blenoxane®), docetaxel (Taxotere®), erbitux (Cetuximab®), hydroxyurea (Hydrea®) or pembrolizumab (Keytruda®),

In some embodiments, the antibodies of the invention are administered in combination with a TLR agonist.

In some embodiments, the TLR3 agonist is TLR4 agonist.

In some embodiments, the TLR3 agonist is a TLR7/8 agonist.

Exemplary TLR agonists are Pam3Cys, a TLR-1/2 agonist; CFA, a TLR-2 agonist; MALP2, a TLR-2 agonist; Pam2Cys, a TLR-2 agonist; FSL-1, a TLR-2 agonist; Hib-OMPC, a TLR-2 agonist; polyribosinic:polyribocytidic acid (Poly I:C), a TLR-3 agonist; polyadenosine-polyuridylic acid (poly AU), a TLR-3 agonist; Polyinosinic-Polycytidylic acid stabilized with poly-L-lysine and carboxymethylcellulose (Hiltonol®), a TLR-3 agonist; monophosphoryl lipid A (MPL), a TLR-4 agonist; LPS, a TLR-4 agonist; bacterial flagellin, a TLR-5 agonist; sialyl-Tn (STn), a carbohydrate associated with the MUCI mucin on a number of human cancer cells and a TLR-4 agonist; imiquimod, a TLR-7 agonist; resiquimod, a TLR-7/8 agonist; loxoribine, a TLR-7/8 agonist; and unmethylated CpG dinucleotide (CpG-ODN), a TLR-9 agonist.

Exemplary TLR4 agonists are agonistic antibodies specifically binding TLR4.

In some embodiments described herein, the antibodies of the invention are administered in combination with an antibody that bids CSF-1R

Exemplary antibodies that bind CSF-1R are those described in Int. Patent Publ. No. WO2013132044.

In some embodiments described herein, the antibodies of the invention are administered in combination with LXRβ agonist.

In some embodiments described herein, the antibodies of the invention are administered in combination with a DR4 agonist.

In some embodiments described herein, the antibodies of the invention are administered in combination with a DR5 agonist.

Suitable DR4 and DR5 agonists are described for example in Int. Patent Publ. No. WO2014159562.

In some embodiments described herein, the antibodies of the invention are administered in combination with an anti-galectin 1 antibody.

Exemplary anti-galectin 1 antibodies that may be used in combination with the antibodies of the invention are those described in Int. Patent Publ. No. WO2015013389.

In some embodiment described herein, the antibodies of the invention are administered in combination with a BTK inhibitor.

In some embodiments, the BTK inhibitor is IMBRUVICA® (ibrutinib).

In some embodiments described herein, the antibodies of the invention are administered in combination with an anti-HER2 antibody.

In some embodiments described herein, the antibodies of the invention are administered in combination with an anti-CD20 antibody.

In some embodiments, the antibodies of the invention are administered in conjunction with (e.g., before, simultaneously or following) bone marrow transplantation, T cell ablative therapy using chemotherapy agents such as, fludarabine, external-beam radiation therapy (XRT), cyclophosphamide, and/or antibodies such as OKT3 or CAMPATH. In some embodiments, the antibodies of the invention may be administered following B-cell ablative therapy such as agents that react with CD20, e.g., Rituxan. For example, in one embodiment, subjects may undergo standard treatment with high dose chemotherapy followed by peripheral blood stem cell transplantation. In certain embodiments, following the transplant, subjects receive the antibodies of the invention.

In some embodiments described herein, the antibodies of the invention are administered before or following surgery.

In some embodiments described herein, the antibodies of the invention are administered in combination with radiation therapy.

Radiation therapy may be administered using various methods, including external-beam therapy, internal radiation therapy, implant radiation, stereotactic radiosurgery, systemic radiation therapy, radiotherapy and permanent or temporary interstitial brachytherapy. External-beam therapy involves three dimensional, conformal radiation therapy where the field of radiation is designed, local radiation (e.g., radiation directed to a preselected target or organ), or focused radiation. Focused radiation may be selected from stereotactic radiosurgery, fractionated stereotactic radiosurgery or intensity-modulated radiation therapy. Focused radiation may have particle beam (proton), cobalt-60 (photon) linear accelerator (x-ray) as a radiation source (see e.g. WO 2012/177624). "Brachytherapy," refers to radiation therapy delivered by a spatially confined radioactive material inserted into the body at or near a tumor or other proliferative tissue disease site, and includes exposure to radioactive isotopes (e.g., At-211, I-131, I-125, Y-90, Re-186, Re-188, Sm-153, Bi-212, P-32, and radioactive isotopes of Lu). Suitable radiation sources for use as a cell conditioner include both solids and liquids. The radiation source can be a radionuclide, such as I-125, I-131, Yb-169, Ir-192 as a solid source, I-125 as a solid source, or other radionuclides that emit photons, beta particles, gamma radiation, or other therapeutic rays. The radioactive material may also be a fluid made from any solution of radionuclide(s), e.g., a solution of I-125 or I-131, or a radioactive fluid can be produced using a slurry of a suitable fluid containing small particles of solid radionuclides, such as Au-198, Y-90. The radionuclide(s) may be embodied in a gel or radioactive micro spheres.

In some embodiments, the antibodies of the invention are administered in combination with decarbazine for the treatment of melanoma. Without being bound by any particular theory, the combined use of PD-1 and/or TIM-3 blockade and chemotherapy is believed to be facilitated by cell death that is a consequence of the cytotoxic action of most chemotherapeutic compounds, which can result in increased levels of tumor antigen in the antigen presentation pathway. Other combination therapies that may result in synergy with PD-1 and/or TIM-3 blockade through cell death are radiation, surgery, and hormone deprivation. Each of these protocols creates a source of tumor antigen in the host. Angiogenesis inhibitors may also be combined with PD-1 and/or TIM-3 blockade. Inhibition of angiogenesis leads to tumor cell death which may feed tumor antigen into host antigen presentation pathways.

The monospecific PD-1 and/or TIM-3 antibodies of the invention may also be used in combination with bispecific antibodies. Bispecific antibodies may be used to target two separate antigens. For example anti-Fc receptor/anti-tumor antigen (e.g., Her-2/neu) bispecific antibodies have been used to target macrophages to sites of tumor. Bispecific targeting may more effectively activate tumor specific responses. The T cell arm of these responses would be augmented by the use of PD-1 and/or TIM-3 blockade. Alternatively, antigen may be delivered directly to DCs by the use of bispecific antibodies which bind to tumor antigen and a dendritic cell specific cell surface marker.

The antibodies of the invention may be used in unconjugated forms or conjugated to a second agent, e.g., a cytotoxic drug, radioisotope, or a protein, e.g., a protein toxin or a viral protein. The antibody molecules may be used to deliver a variety of therapeutic agents, e.g., a cytotoxic moiety, e.g., a therapeutic drug, a radioisotope, molecules of plant, fungal, or bacterial origin, or biological proteins (e.g., protein toxins) or particles (e.g., a recombinant viral particles, e.g.; via a viral coat protein), or mixtures thereof.

Infectious Diseases

The invention also provides a method of treating a subject that has been exposed to particular toxins or pathogen with the antibodies of the invention for a time sufficient to treat the subject.

The invention also provides a method of treating a subject having an infectious disease, comprising administering a therapeutically efficient amount of the antibody of the invention to the subject in need thereof for a time sufficient to treat the infectious disease.

The invention also provides a method of treating a subject having a viral infection, comprising administering a therapeutically efficient amount of the antibody of the invention to the subject in need thereof for a time sufficient to treat the viral infection.

The invention also provides a method of treating a subject having a bacterial infection, comprising administering a therapeutically efficient amount of the antibody of the invention to the subject in need thereof for a time sufficient to treat the bacterial infection.

The invention also provides a method of treating a subject having a fungal infection, comprising administering a therapeutically efficient amount of the antibody of the invention to the subject in need thereof for a time sufficient to treat the fungal infection.

In the treatment of infection (e.g., acute and/or chronic), administration of the antibodies of the invention may be combined with conventional treatments in addition to or in lieu of stimulating natural host immune defenses to infection. Natural host immune defenses to infection include inflammation, fever, antibody-mediated host defense, T-lymphocyte-mediated host defenses, including lymphokine secretion and cytotoxic T-cells (especially during viral infection), complement mediated lysis and opsonization (facilitated phagocytosis), and phagocytosis. The ability of the antibodies of the invention to reactivate dysfunctional T-cells would be useful to treat chronic infections, in particular those in which cell-mediated immunity is important for complete recovery.

Similar to its application to tumors as discussed above, antibodies of the invention may be used alone, or as an adjuvant, in combination with vaccines, to stimulate the immune response to pathogens, toxins, and self-antigens. Examples of pathogens for which this therapeutic approach may be useful include pathogens for which there is currently no effective vaccine, or pathogens for which conventional vaccines are less than completely effective. These include HIV, Hepatitis (A, B, & C), Influenza, Herpes, Giardia, Malaria, Leishmania, Staphylococcus aureus and Pseudomonas Aeruginosa. PD-1 and/or TIM-3 blockade may be useful against established infections by agents such as HIV that present altered antigens over the course of the infections. These novel epitopes are recognized as foreign at the time of administration of the antibodies of the invention, thus provoking a strong T cell response that is not dampened by negative signals through PD-1 or TIM-3.

Viruses

For infections resulting from viral causes, the antibodies of the invention may be combined with standard therapies for treating viral infections. Such standard therapies vary depending upon type of virus, although in almost all cases, administration of human serum containing antibodies (e.g., IgA, IgG) specific to the virus can be effective.

Exemplary pathogenic viruses causing infections that may be treatable by the antibodies of the invention include HIV, hepatitis (A, B, or C), herpes virus (e.g., VZV, HSV-1, HAV-6, HSV-II, and CMV, Epstein Barr virus), adenovirus, influenza virus, flaviviruses, echovirus, rhinovirus, coxsackie virus, cornovirus, respiratory syncytial virus, mumps virus, rotavirus, measles virus, rubella virus, parvovirus, vaccinia virus, HTLV virus, dengue virus, papillomavirus, molluscum virus, poliovirus, rabies virus, JC virus and arboviral encephalitis virus.

In some embodiments, the virus infection is an influenza virus infection. Influenza infection can result in fever, cough, myalgia, headache and malaise, which often occur in seasonal epidemics. Influenza is also associated with a number of postinfectious disorders, such as encephalitis, myopericarditis, Goodpasture's syndrome, and Reye's syndrome. Influenza infection also suppresses normal pulmonary antibacterial defenses, such that patients recovering from influenza have an increased risk of developing bacterial pneumonia. Influenza viral surface proteins show marked antigenic variation, resulting from mutation and recombination. Thus, cytolytic T lymphocytes are the host's primary vehicle for the elimination of virus after infection. Influenza is classified into three primary types: A, B and C. Influenza A is unique in that it infects both humans and many other animals (e.g., pigs, horses, birds and seals) and is the principal cause of pandemic influenza. A cell can be infected by two different influenza A strains, the segmented RNA genomes of two parental virus types mix during replication to create a hybrid replicant, resulting in new epidemic strains. Influenza B does not replicate in animals and thus has less genetic variation and influenza C has only a single serotype.

Most conventional therapies are palliatives of the symptoms resulting from infection, while the host's immune response actually clears the disease. However, certain strains (e.g., influenza A) can cause more serious illness and death. Influenza A may be treated both clinically and prophylactically by the administration of the cyclic amines inhibitors amantadine and rimantadine, which inhibit viral replication. However, the clinical utility of these drugs is limited due to the relatively high incidence of adverse reactions, their narrow anti-viral spectrum (influenza A only), and the propensity of the virus to become resistant. The administration of serum IgG antibody to the major influenza surface proteins, hemagglutinin and neuraminidase can prevent pulmonary

infection, whereas mucosal IgA is required to prevent infection of the upper respiratory tract and trachea. The most effective current treatment for influenza is vaccination with the administration of virus inactivated with formalin or β-propiolactone.

In some embodiments, the infection is a hepatitis infection, e.g., a Hepatitis B or C infection.

Hepatitis B virus (HB-V) is the most infectious known blood borne pathogen. It is a major cause of acute and chronic hepatitis and hepatic carcinoma, as well as life-long, chronic infection. Following infection, the virus replicates in hepatocytes, which also then shed the surface antigen HBsAg. The detection of excessive levels of HBsAg in serum is used as a standard method for diagnosing a hepatitis B infection. An acute infection may resolve or it can develop into a chronic persistent infection. Current treatments for chronic HBV include α-interferon, which increases the expression of class I human leukocyte antigen (HLA) on the surface of hepatocytes, thereby facilitating their recognition by cytotoxic T lymphocytes. Additionally, the nucleoside analogs ganciclovir, famciclovir and lamivudine have also shown some efficacy in the treatment of HBV infection in clinical trials. Additional treatments for HBV include pegylated α-interferon, adenfovir, entecavir and telbivudine. While passive immunity can be conferred through parental administration of anti-HBsAg serum antibodies, vaccination with inactivated or recombinant HBsAg also confers resistance to infection. The antibodies of the invention may be combined with conventional treatments for hepatitis B infections for therapeutic advantage.

Hepatitis C virus (HC-V) infection may lead to a chronic form of hepatitis, resulting in cirrosis. While symptoms are similar to infections resulting from Hepatitis B, in distinct contrast to HB-V, infected hosts can be asymptomatic for 10-20 years. The antibodies of the invention can be administered as a monotherapy, or combined with the standard of care for hepatitis C infection. For example, the antibodies of the invention can be administered with one or more of Sovaldi (sofosbuvir) Olysio (simeprevir), plus ribavirin or pegylated interferon. Although regimens that include Incivek (telaprevir) or Victrelis (boceprevir) plus ribavirin and pegylated interferon are also approved, they are associated with increased side effects and longer duration of treatment.

Conventional treatment for HC-V infection includes the administration of a combination of α-interferon and ribavirin. A promising potential therapy for HC-V infection is the protease inhibitor telaprevir (VX-960). Additional treatments include bavituximab (an antibody that binds anionic phospholipid phosphatidylserine in a B2-glycoprotein I dependent manner, Peregrine Pharmaceuticals), anti-HPV viral coat protein

E2 antibod(y)(ies) (e.g., ATL 6865-Ab68+Ab65, XTL Pharmaceuticals) and Civacir® (polyclonal anti-HCV human immune globulin). The antibodies of the invention may be combined with one or more of these treatments for hepatitis C infections for therapeutic advantage. Protease, polymerase and NS5A inhibitors which may be used in combination with the antibodies of the invention to specifically treat Hepatitis C infection include those described in US 2013/0045202.

In another embodiment, the infection is a measles virus. After an incubation of 9-11 days, hosts infected with the measles virus develop fever, cough, coryza and conjunctivitis. Within 1-2 days, an erythematous, maculopapular rash develop, which quickly spreads over the entire body. Because infection also suppresses cellular immunity, the host is at greater risk for developing bacterial superinfections, including otitis media, pneumonia and postinfectious encephalomyelitis. Acute infection is associated with significant morbidity and mortality, especially in malnourished adolescents.

Treatment for measles includes the passive administration of pooled human IgG, which can prevent infection in non-immune subjects, even if given up to one week after exposure. However, prior immunization with live, attenuated virus is the most effective treatment and prevents disease in more than 95% of those immunized. As there is one serotype of this virus, a single immunization or infection typically results in protection for life from subsequent infection.

In a small proportion of infected hosts, measles can develop into SSPE, which is a chronic progressive neurologic disorder resulting from a persistent infection of the central nervous system. S SPE is caused by clonal variants of measles virus with defects that interfere with virion assembly and budding. For these patients, reactivation of T-cells with the antibodies of the invention so as to facilitate viral clearance would be desirable.

In another embodiment, the infection is HIV. HIV attacks $CD4^+$ cells, including T-lymphocytes, monocyte-macrophages, follicular dendritic cells and Langerhan's cells, and $CD4^+$ helper/inducer cells are depleted. As a result, the host acquires a severe defect in cell-mediated immunity. Infection with HIV results in AIDS in at least 50% of individuals, and is transmitted via sexual contact, administration of infected blood or blood products, artificial insemination with infected semen, exposure to blood-containing needles or syringes and transmission from an infected mother to infant during childbirth.

A host infected with HIV may be asymptomatic, or may develop an acute illness that resembling mononucleosis—fever, headache, sore throat, malaise and rash. Symptoms can progress to progressive immune dysfunction, including persistent fever, night sweats, weight loss, unexplained diarrhea, eczema, psoriasis, seborrheic dermatitis, herpes zoster,

oral candidiasis and oral hairy leukoplakia. Opportunistic infections by a host of parasites are common in patients whose infections develop into AIDS.

Treatments for HIV include antiviral therapies including nucleoside analogs, zidovudine (AST) either alone or in combination with didanosine or zalcitabine, dideoxyinosine, dideoxycytidine, lamidvudine, stavudine; reverse transcriptive inhibitors such as delavirdine, nevirapine, loviride, and proteinase inhibitors such as saquinavir, ritonavir, indinavir and nelfinavir. Treatments for HIV include EDURANT® (rilpivirine). The antibodies of the invention may be combined with conventional treatments for HIV infections for therapeutic advantage.

In another embodiment, the infection is a Cytomegalovirus (CMV) infection. CMV infection is often associated with persistent, latent and recurrent infection. CMV infects and remains latent in monocytes and granulocyte-monocyte progenitor cells. The clinical symptoms of CMV include mononucleosis-like symptoms (i.e., fever, swollen glands, malaise), and a tendency to develop allergic skin rashes to antibiotics. The virus is spread by direct contact. The virus is shed in the urine, saliva, semen and to a lesser extent in other body fluids. Transmission can also occur from an infected mother to her fetus or newborn and by blood transfusion and organ transplants. CMV infection results in general impairment of cellular immunity, characterized by impaired blastogenic responses to nonspecific mitogens and specific CMV antigens and diminished cytotoxic ability.

Treatments of CMV infection include the anti-virals ganciclovir, foscarnet and cidovir, but these drugs are typically only prescribed in immunocompromised patients. The antibodies of the invention described herein may be combined with conventional treatments for cytomegalovirus infections for therapeutic advantage.

In another embodiment, the infection is Epstein-Barr virus (EBV) infection. EBV can establish persistent and latent infections and primarily attacks B cells. Infection with EBV results in the clinical condition of infectious mononucleosis, which includes fever, sore throat, often with exudate, generalized lymphadenopathy and splenomegaly. Hepatitis is also present, which can develop into jaundice.

While typical treatments for EBV infections are palliative of symptoms, EBV is associated with the development of certain cancers such as Burkitt's lymphoma and nasopharyngeal cancer. Thus, clearance of viral infection before the complications develop would be of great benefit. The antibodies of the invention may be combined with conventional treatments for Epstein-Barr virus infections for therapeutic advantage.

In another embodiment, the infection is Herpes simplex virus (HSV) infection. HSV is transmitted by direct contact with an infected host. A direct infection may be

asymptomatic, but typically result in blisters containing infectious particles. The disease manifests as cycles of active periods of disease, in which lesions appear and disappear as the virus latently infects the nerve ganglion for subsequent outbreaks. Lesions may be on the face, genitals, eyes and/or hands. In some case, an infection can also cause encephalitis.

Treatments for herpes infections are directed primarily to resolving the symptomatic outbreaks, and include systemic antiviral medicines such as: acyclovir (e.g., Zovirax®), valaciclovir, famciclovir, penciclovir, and topical medications such as docosanol (Abreva®), tromantadine and zilactin. The clearance of latent infections of herpes would be of great clinical benefit. The antibodies of the invention may be combined with conventional treatments for herpes virus infections for therapeutic advantage.

In another embodiment, the infection is Human T-lymphotrophic virus (HTLV-1, HTLV-2). HTLV is transmitted via sexual contact, breast feeding or exposure to contaminated blood. The virus activates Th1 cells, resulting in their overproliferation and overproduction of Th1 related cytokines (e.g., IFN-γ and TNF-α). This in turn results in a suppression of Th2 lymphocytes and reduction of Th2 cytokine production (e.g., IL-4, IL-5, IL-10 and IL-13), causing a reduction in the ability of an infected host to mount an adequate immune response to invading organisms requiring a Th2-dependent response for clearance (e.g., parasitic infections, production of mucosal and humoral antibodies).

HTLV infections lead to opportunistic infections resulting in bronchiectasis, dermatitis and superinfections with Staphylococcus spp. and Strongyloides spp. resulting in death from polymicrobial sepsis. HTLV infection can also lead directly to adult T-cell leukemia/lymphoma and progressive demyelinating upper motor neuron disease known as HAM/TSP. The clearance of HTLV latent infections would be of great clinical benefit. The antibodies of the invention may be combined with conventional treatments for HTLV infections for therapeutic advantage.

In another embodiment, the infection is Human papilloma virus (HPV). HPV primarily affects keratinocytes and occurs in two forms: cutaneous and genital. Transmission is believed to occur through direct contact and/or sexual activity. Both cutaneous and genital HPV infection can result in warts and latent infections and sometimes recurring infections, which are controlled by host immunity which controls the symptoms and blocks the appearance of warts, but leaves the host capable of transmitting the infection to others.

Infection with HPV can also lead to certain cancers, such as cervical, anal, vulvar, penile and oropharynial cancer. There are no known cures for HPV infection, but current treatment is topical application of Imiquimod, which stimulates the immune system to attack the affected area. The clearance of HPV latent infections would be of great clinical benefit. The antibodies of the invention may be combined with conventional treatments for HPV infections for therapeutic advantage.

Bacterial Infections

Some examples of pathogenic bacteria causing infections that may be treated with the antibodies of the invention include syphilis, chlamydia, rickettsial bacteria, mycobacteria, staphylococci, streptococci, pneumonococci, meningococci and conococci, klebsiella, proteus, serratia, pseudomonas, legionella, diphtheria, salmonella, bacilli, cholera, tetanus, botulism, anthrax, plague, leptospirosis, and Lymes disease bacteria. The antibodies of the invention can be used in combination with existing treatment modalities for the aforesaid infections. For example, treatments for syphilis include penicillin (e.g., penicillin G.), tetracycline, doxycycline, ceftriaxone and azithromycin.

Lyme disease, caused by Borrelia burgdorferi is transmitted into humans through tick bites. The disease manifests initially as a localized rash, followed by flu-like symptoms including malaise, fever, headache, stiff neck and arthralgias. Later manifestations can include migratory and polyarticular arthritis, neurologic and cardiac involvement with cranial nerve palsies and radiculopathy, myocarditis and arrhythmias. Some cases of Lyme disease become persistent, resulting in irreversible damage analogous to tertiary syphilis . Current therapy for Lyme disease includes primarily the administration of antibiotics. Antibiotic-resistant strains may be treated with hydroxychloroquine or methotrexate. Antibiotic refractory patients with neuropathic pain can be treated with gabapentin. Minocycline may be helpful in late/chronic Lyme disease with neurological or other inflammatory manifestations.

Other forms of borreliois, such as those resulting from B. recurentis, B. hermsii, B. turicatae, B. parikeri, B. hispanica, B. duttonii and B. persica, as well leptospirosis (E.g., L. interrogans), typically resolve spontaneously unless blood titers reach concentrations to cause intrahepatic obstruction.

Fungi and Parasites

Some examples of pathogenic fungi causing infections that may be treated with the antibodies of the invention include Candida (albicans, krusei, glabrata, tropicalis, etc.),

Cryptococcus neoformans, Aspergillus (fumigatus, niger, etc.), Genus Mucorales (mucor, absidia, rhizophus), Sporothrix schenkii, Blastomyces dermatitidis, Paracoccidioides brasiliensis, Coccidioides immitis and Histoplasma capsulatum.

Some examples of pathogenic parasites causing infections treatable with the antibodies of the invention described herein include Entamoeba histolytica, Balantidium coli, Naegleriafowleri, Acanthamoeba sp., Giardia lambia, Cryptosporidium sp., Pneumocystis carinii, Plasmodium vivax, Babesia microti, Trypanosoma brucei, Trypanosoma cruzi, Leishmania donovani, Toxoplasma gondi, and Nippostrongylus brasiliensis.

Diagnostic uses and kits

Kits

The invention also provides a kit comprising the antagonistic antibody specifically binding PD-1 of the invention.

The invention also provides a kit comprising the antagonistic antibody specifically binding TIM-3 of the invention.

The invention also provides a kit comprising the antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second domain specifically binding TIM-3 of the invention.

The kit may be used for therapeutic uses and as diagnostic kits.

The kit may be used to detect the presence of PD-1, TIM-3, or PD-1 and TIM-3 in a biological sample.

In some embodiments, the kit comprises the antibody of the invention described herein and reagents for detecting the antibody. The kit can include one or more other elements including: instructions for use; other reagents, e.g., a label, a therapeutic agent, or an agent useful for chelating, or otherwise coupling, an antibody to a label or therapeutic agent, or a radioprotective composition; devices or other materials for preparing the antibody for administration; pharmaceutically acceptable carriers; and devices or other materials for administration to a subject.

In some embodiments, the kit comprises the antibody of the invention in a container and instructions for use of the kit.

In some embodiments, the antibody in the kit is labeled.

In some embodiments, the kit comprises the antagonistic antibody specifically binding PD-1, comprising

the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49;

the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 50;
the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 51;
the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 52;
the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 53;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 54;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 50;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 55;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 56;
the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 57;
the VH of SEQ ID NO: 44 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 46 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 49;
the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 53;
the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 52;
the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56;
the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58;
the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 59;
the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60;
the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 61;
the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 62;
the VH of SEQ ID NO: 63 and the VL of SEQ ID NO: 65; or
the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

In some embodiments, the kit comprises the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

In some embodiments, the kit comprises the antagonistic antibody specifically binding PD-1 comprising the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

In some embodiments, the kit comprises the antagonistic antibody specifically binding TIM-3, comprising

the VH of SEQ ID NO: 145 and the VL of SEQ ID NO: 155;
the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156;
the VH of SEQ ID NO: 148 and the VL of SEQ ID NO: 157;
the VH of SEQ ID NO: 147 and the VL of SEQ ID NO: 155;
the VH of SEQ ID NO: 149 and the VL of SEQ ID NO: 158;

the VH of SEQ ID NO: 150 and the VL of SEQ ID NO: 159;
the VH of SEQ ID NO: 151 and the VL of SEQ ID NO: 160;
the VH of SEQ ID NO: 152 and the VL of SEQ ID NO: 161;
the VH of SEQ ID NO: 153 and the VL of SEQ ID NO: 162;
the VH of SEQ ID NO: 154 and the VL of SEQ ID NO: 163; or
the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

In some embodiments, the kit comprises the antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

In some embodiments, the kit comprises the antagonistic antibody specifically binding TIM-3 comprising the VH of SEQ ID NO: 172 and the VL of SEQ ID NO: 173.

In some embodiments, the kit comprises the antagonistic bispecific PD-1/TIM-3 antibody comprising the HC1, the LC1, the HC2 and the LC2 of

SEQ ID NOs: 186, 188, 190 and 193, respectively;
SEQ ID NOs: 186, 188, 191 and 194, respectively;
SEQ ID NOs: 187, 189, 190 and 193, respectively;
SEQ ID NOs: 187, 189, 191, 194, respectively;
SEQ ID NOs: 186, 188, 192 and 195, respectively;
SEQ ID NOs: 186, 188, 248 and 194, respectively;
SEQ ID NOs: 241, 188, 244, 195, respectively;
SEQ ID NOs: 241, 188, 245, 194, respectively;
SEQ ID NOs: 242, 189, 246, 194, respectively;
SEQ ID NOs: 243, 188, 246, 194, respectively; or
SEQ ID NOs: 243, 188, 247, 195, respectively.

Methods of detecting PD-1, TIM-3 or PD-1 and TIM-3

The invention also provides a method of detecting PD-1 in a sample, comprising obtaining the sample, contacting the sample with the antagonistic antibody specifically binding PD-1 of the invention, and detecting the antibody bound to PD-1 in the sample.

The invention also provides a method of detecting TIM-3 in a sample, comprising obtaining the sample, contacting the sample with the antagonistic antibody specifically binding TIM-3 of the invention, and detecting the antibody bound to TIM-3 in the sample.

The invention also provides a method of detecting PD-1 and TIM-3 in a sample, comprising obtaining the sample, contacting the sample with the antagonistic bispecific PD-1/TIM-3 antibody comprising a first domain specifically binding PD-1 and a second

domain specifically binding TIM-3 of the invention, and detecting the antibody bound to PD-1 and TIM-3 in the sample.

In some embodiments, the sample may be derived from urine, blood, serum, plasma, saliva, ascites, circulating cells, circulating tumor cells, cells that are not tissue associated (*i.e.*, free cells), tissues (*e.g.*, surgically resected tumor tissue, biopsies, including fine needle aspiration), histological preparations, and the like.

The antibodies of the invention bound to PD-1, TIM-3 or PD-1 and TIM-3 may be detected using known methods. Exemplary methods include direct labeling of the antibodies using fluorescent or chemiluminescent labels, or radiolabels, or attaching to the antibodies of the invention a moiety which is readily detectable, such as biotin, enzymes or epitope tags. Exemplary labels and moieties are ruthenium, ^{111}In-DOTA, ^{111}In-diethylenetriaminepentaacetic acid (DTPA), horseradish peroxidase, alkaline phosphatase and beta-galactosidase, poly-histidine (HIS tag), acridine dyes, cyanine dyes, fluorone dyes, oxazin dyes, phenanthridine dyes, rhodamine dyes and Alexafluor® dyes.

The antibodies of the invention may be used in a variety of assays to detect PD-1, TIM-3 or PD-1 and TIM-3 in the sample. Exemplary assays are western blot analysis, radioimmunoassay, surface plasmon resonance, immunoprecipitation, equilibrium dialysis, immunodiffusion, electrochemiluminescence (ECL) immunoassay, immunohistochemistry, fluorescence-activated cell sorting (FACS) or ELISA assay.

Further embodiments of the invention: antibodies specifically binding PD-1

Set out below are certain further embodiments of the invention according to the disclosures elsewhere herein. Features from embodiments of the invention set out above described as relating to the invention disclosed herein also relate to each and every one of these further numbered embodiments.

1) An isolated antagonistic antibody specifically binding PD-1, comprising heavy chain complementarity determining regions (HCDR) 1 (HCDR1) 2 (HCDR2) and 3 (HCDR3) amino acid sequences of SEQ ID NOs: 82, 83 and 84, respectively, or SEQ ID NOs: 82, 83 and 85, respectively.
2) The antibody according to embodiment 1, comprising light chain complementarity determining regions (LCDR) 1 (LCDR1), 2 (LCDR2) and 3 (LCDR3) amino acid sequences of SEQ ID NOs: 86, 87 and 88, respectively.
3) The antibody according to embodiment 1 or 2, wherein the antibody has one, two, three, four or five of the following properties:

a) enhances activation of antigen specific $CD4^+$ or $CD8^+$ T cells in dose dependent manner, wherein activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1;
b) binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;
c) binds human PD-1 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;
d) binds cynomolgus PD-1 with the K_D of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C, or
e) binds cynomolgus PD-1 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.

4) The antibody according to any one of embodiments 1-3, comprising the HCDR1, the HCDR2 and the HCDR3 contained within a heavy chain variable region (VH) having the amino acid sequence of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47 or 48, wherein the HCDRs are defined by Chothia, Kabat, or IMGT.
5) The antibody according to any one of embodiments 1-4, comprising the LCDR1, the LCDR2 and the LCDR3 contained within a light chain variable region (VL) having the amino acid sequence of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61 or 62, wherein the LCDRs are defined by Chothia, Kabat, or IMGT.
6) The antibody according to any one of embodiments 1-5, comprising
a) the HCDR1 of SEQ ID NOs: 10, 11 or 12;
b) the HCDR2 of SEQ ID NOs: 13, 14 or 15; and
c) the HCDR3 of SEQ ID NOs: 16, 17, 18 or 19.
7) The antibody according to any one of embodiments 1-6, comprising
a) the LCDR1 of SEQ ID NOs: 20, 21, 22, 23, 24 or 25;
b) the LCDR2 of SEQ ID NOs: 26, 27, 28, 29 or 30; and
c) the LCDR3 of SEQ ID NOs: 31, 32, 33, 34, 35, 36, 37, 38, 39, or 40.
8) The antibody according to any one of embodiments 1-7, wherein the antibody comprises a heavy chain framework derived from VH1-69 (SEQ ID NO: 170) and a light chain framework derived from IGKV3-11 (SEQ ID NO: 171).
9) The antibody according to any one of embodiments 1-8, comprising the HCDR1, the HCDR2 and the HCDR3 of
a) SEQ ID NOs: 10, 13 and 16, respectively;
b) SEQ ID NOs: 10, 14 and 16, respectively;

c) SEQ ID NOs: 10, 13 and 17, respectively;
d) SEQ ID NOs: 10, 13 and 18, respectively;
e) SEQ ID NOs: 11, 15 and 18, respectively;
f) SEQ ID NOs: 10, 13 and 19, respectively;
g) SEQ ID NOs: 12, 13 and 19, respectively;
h) SEQ ID NOs: 10, 13 and 16, respectively; or
i) SEQ ID NOs: 10, 14 and 17, respectively.

10) The antibody according to any one of embodiments 1-9, comprising the LCDR1, the LCDR2 and the LCDR3 of
a) SEQ ID NOs: 20, 26 and 31, respectively;
b) SEQ ID NOs: 21, 26 and 32, respectively;
c) SEQ ID NOs: 22, 27 and 33, respectively;
d) SEQ ID NOs: 22, 26 and 34, respectively;
e) SEQ ID NOs: 23, 28 and 35, respectively;
f) SEQ ID NOs: 20, 26 and 36, respectively;
g) SEQ ID NOs: 21, 27 and 37, respectively;
h) SEQ ID NOs: 23, 26 and 32, respectively;
i) SEQ ID NOs: 22, 26 and 32, respectively;
j) SEQ ID NOs: 24, 26 and 38, respectively;
k) SEQ ID NOs: 20, 29 and 39, respectively;
l) SEQ ID NOs: 20, 30 and 32, respectively;
m) SEQ ID NOs: 25, 26 and 40, respectively;
n) SEQ ID NOs: 24, 26 and 32, respectively; or
o) SEQ ID NOs: 69, 70 and 71, respectively.

11) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 16, 20, 26 and 31, respectively.

12) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 16, 21, 26 and 32, respectively.

13) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 16, 22, 27 and 33, respectively.

14) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 16, 22, 26 and 34, respectively.

15) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 16, 23, 28 and 35, respectively.

16) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 20, 26 and 31, respectively.

17) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 20, 26 and 36, respectively.

18) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 21, 26 and 32, respectively.

19) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 21, 27 and 37, respectively.

20) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 23, 26 and 32, respectively.

21) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 17, 22, 26 and 32, respectively.

22) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 18, 20, 26 and 31, respectively.

23) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 11, 15, 18, 20, 26 and 31, respectively.

24) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 13, 19, 20, 26 and 31, respectively.

25) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 12, 13, 19, 20, 26 and 31, respectively.

26) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 28 and 35, respectively.

27) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 22, 26 and 34, respectively.

28) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively.

29) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 12, 13, 19, 24, 26 and 38, respectively.

30) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 12, 13, 19, 20, 29 and 39, respectively.

31) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 11, 15, 18, 20, 30 and 32, respectively.

32) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 11, 15, 18, 25, 26 and 40, respectively.

33) The antibody according to any one of embodiments 1-10, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 11, 15, 18, 24, 26 and 32, respectively.

34) The antibody according to any one of embodiments 1-33, comprising a heavy chain variable region (VH) of SEQ ID NOs: 41, 42, 43, 44, 45, 46, 47 or 48, the VH optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen conservative amino acid substitutions.

35) The antibody according to any one of embodiments 1-34, comprising a light chain variable region (VL) of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61 or 62 the VL optionally having one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen conservative amino acid substitutions.

36) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

37) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 50, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

38) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 51, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

39) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 52, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

40) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 53, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

41) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 49, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

42) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 54, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

43) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 50, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

44) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 55, wherein the VH, the VL or both the VH

and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

45) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 56, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

46) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 57, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

47) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 44 and the VL of SEQ ID NO: 49, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

48) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 49, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

49) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 46 and the VL of SEQ ID NO: 49, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

50) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 49, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

51) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 53, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

52) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 52, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

53) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

54) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 58, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

55) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 47 and the VL of SEQ ID NO: 59, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

56) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 60, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

57) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 61, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

58) The antibody according to any one of embodiments 1-35, comprising the VH of SEQ ID NO: 45 and the VL of SEQ ID NO: 62, wherein the VH, the VL or both the VH and the VL optionally comprise one, two, three, four, five, six, seven, eight, nine, ten, eleven, twelve, thirteen, fourteen or fifteen amino acid substitutions.

59) An isolated antagonistic antibody that specifically binds PD-1, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 amino acid sequences of SEQ ID NOs: 66, 67, 68, 69, 70 and 71, respectively.

60) The antibody according to embodiment 59, wherein the antibody comprises the VH of SEQ ID NO: 63 and the VL of SEQ ID NO: 65 or the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

61) The antibody according to any one of embodiments 1-60, wherein the antibody is human or humanized.

62) The antibody according to any one of embodiments 1-61, wherein the antibody is of IgG1, IgG2, IgG3 or IgG4 isotype.

63) The antibody according to any one of embodiments 1-62, comprising one, two, three, four, five, six, seven, eight, nine or ten substitutions in the antibody Fc.

64) The antibody according to any one of embodiments 1-63, wherein the one, two, three, four, five, six, seven, eight, nine or ten substitutions result in an increase in a half-life of the antibody.

65) The antibody according to any one of embodiments 1-64, comprising M252Y, S254T and T256E substitutions, residue numbering according to the EU Index.

66) The antibody according to any one of embodiments 1-65, wherein the one, two, three, four, five, six, seven, eight, nine or ten substitutions result in a reduced binding of the antibody to an activating Fcγ receptor (FcγR).

67) The antibody according to any one of embodiments 1-66, wherein the activating FcγR is FcγRI, FcγRIIa, FcγRIIIa, or FcγRIIIb.

68) The antibody according to any one of embodiments 1-67, comprising

a) L234A, L235A, G237A, P238S, H268A, A330S and P331S substitutions;

b) V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions;

c) F234A, L235A, G237A, P238S and Q268A substitutions;

d) L234A, L235A or L234A and L235A substitutions;

e) F234A, L235A or F234A and L235A substitutions; or

f) V234A substitution, wherein residue numbering is according to the EU Index.

69) The antibody according to any one of embodiments 1-68, comprising S228P substitutions, wherein residue numbering is according to the EU Index.

70) The antibody according to any one of embodiments 1-69, wherein the antibody is bispecific.

71) The antibody according to embodiment 70, wherein the antibody specifically binds PD-L1, PD-L2, LAG-3, TIM-3, CEACAM-1, CEACAM-5, OX-40, GITR, CD27, VISTA or CTLA-4.

72) An immunoconjugate comprising the antibody or antigen-binding portion thereof according to any one of embodiments 1-71 linked to a therapeutic agent or an imaging agent.

73) A pharmaceutical composition comprising the antibody according to any one of embodiments 1-71 or the immunoconjugate according to embodiment 72 and a pharmaceutically accepted carrier.

74) A polynucleotide encoding the antibody VH according to embodiment 34, the antibody VL according embodiment 35, or the antibody VH and the antibody VL according to embodiment 34 or 35.

75) A polynucleotide encoding the antibody VH, the antibody VL, or the antibody VH and the antibody VL according to embodiment 60.

76) A vector comprising the polynucleotide according to embodiment 74 or 75.

77) A host cell comprising the vector according to embodiment 76.

78) A method of producing the antibody according to any one of embodiments 1-71, comprising culturing the host cell according to embodiment 77 in conditions that the antibody is expressed, and recovering the antibody produced by the host cell.

79) The antibody according to any one of embodiments 1-71, the immunoconjugate according to embodiment 72, or the pharmaceutical composition according to embodiment 73 for use in the treatment of cancer.

80) The antibody, the immunoconjugate or the pharmaceutical composition for use according to embodiment 79, wherein the cancer is a solid tumor or a hematological malignancy.

81) The antibody, the immunoconjugate or the pharmaceutical composition for use according to embodiment 80, wherein the solid tumor is a melanoma, a lung cancer, a squamous non-small cell lung cancer (NSCLC), a non-squamous NSCLC, a colorectal cancer, a prostate cancer, a castration-resistant prostate cancer, a stomach cancer, an ovarian cancer, a gastric cancer, a liver cancer, a pancreatic cancer, a thyroid cancer, a squamous cell carcinoma of the head and neck, a carcinoma of the esophagus or gastrointestinal tract, a breast cancer, a fallopian tube cancer, a brain cancer, an urethral cancer, a genitourinary cancer, an endometriosis, a cervical cancer or a metastatic lesion of the cancer.

82) The antibody, the immunoconjugate or the pharmaceutical composition for use according to embodiment 80, where the hematological malignancy is a lymphoma, a myeloma or a leukemia.

83) The antibody according to any one of embodiments 1-71, the immunoconjugate according to embodiment 72, or the pharmaceutical composition according to embodiment 73 for use to enhance an immune response in a subject.

84) The antibody, the immunoconjugate or the pharmaceutical composition for use according to embodiment 83, wherein the subject has cancer or viral infection.

85) The antibody according to any one of embodiments 1-71, the immunoconjugate according to embodiment 72, or the pharmaceutical composition according to

embodiment 73 for use according to any one of embodiments 70-84 in combination with a second therapeutic agent.

86) The antibody according to any one of embodiments 1-71 for use according to embodiment 85, wherein the second therapeutic agent is a standard of care drug for treatment of the solid tumor or the hematological malignancy.

87) The antibody according to any one of embodiments 1-71 for use according to embodiment 85 or 86, wherein the second therapeutic agent is an agonist of a T cell activating molecule.

88) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85-87, wherein the T cell activating molecule is CD86, CD80, CD28, ICOS, ICOS ligand, TMIGD2, CD40, TL1A, GITR ligand, 4-1BB ligand, OX40 ligand, CD70, CD40L, TNFRSF25, LIGHT, GITR, OX-40, CD27, CD137, NKG2D, CD48, CD226 or MICA.

89) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85-88, wherein the agonist is an antibody that specifically binds the T cell activating molecule.

90) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85 or 86, wherein the second therapeutic agent is an inhibitor of a T cell inhibitory molecule.

91) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85, 86 or 90, wherein the T cell inhibitory molecule is PD-1, PD-L1, PD-L2, VISTA, BTNL2, B7-H3, B7-H4, HVEM, HHLA2, CTLA-4, LAG-3, TIM-3, BTLA, CD160, CEACAM-1, LAIR1, TGFβ, IL-10, Siglec family, KIR, CD96, TIGIT, NKG2A, CD112, CD47, SIRPA or CD244.

92) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85, 86, 90 or 91, wherein the inhibitor or the T cell inhibitory molecule is an antibody that specifically binds the T cell inhibitory molecule.

93) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85, 86 or 90-92, wherein the antibody specifically binds TIM-3 and blocks TIM-3 binding to galectin-9.

94) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85, 86 or 90-93, wherein the antibody that specifically binds the T cell inhibitory molecule comprises the VH and the VL of

a) SEQ ID NOs: 145 and 155, respectively;

b) SEQ ID NOs: 146 and 156, respectively;

c) SEQ ID NOs: 148 and 157, respectively;
d) SEQ ID NOs: 147 and 155, respectively;
e) SEQ ID NOs: 149 and 158, respectively;
f) SEQ ID NOs: 150 and 159, respectively;
g) SEQ ID NOs:151 and 160, respectively;
h) SEQ ID NOs: 152 and 161, respectively;
i) SEQ ID NOs: 153 and 162, respectively; or
j) SEQ ID NOs: 154 and 163, respectively.

95) The antibody according to any one of embodiments 1-71 for use according to embodiment 85, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

96) The antibody according to any one of embodiments 1-71 for use according to embodiment 85, wherein the second therapeutic agent is a vaccine.

97) The antibody according to any one of embodiments 1-71 for use according to embodiment 85 or 96, wherein the vaccine is a polypeptide or a fragment thereof, or a DNA or a RNA encoding the polypeptide or fragment thereof expressed on tumor cells.

98) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85, 96 or 97, wherein the tumor cells are melanoma, lung cancer, squamous non-small cell lung cancer (NSCLC), non-squamous NSCLC, colorectal cancer, prostate cancer, castration-resistant prostate cancer, ovarian cancer, gastric cancer, liver cancer, pancreatic cancer, thyroid cancer, squamous cell carcinoma of the head and neck, carcinomas of the esophagus or gastrointestinal tract or breast cancer cells.

99) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85 or 96-98, wherein the polypeptide is PSMA, mesothelin, EGFR or EGFRvIII.

100) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 85-99, wherein the second therapeutic agent is administered simultaneously, sequentially or separately.

101) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 79-100, wherein the subject is treated or is being treated with radiation therapy.

102) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 79-101, wherein the patient has or will undergo surgery.

103) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 79-102, wherein the isolated antibody comprises the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49.

104) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 79-102, wherein the isolated antibody comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively.

105) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 79-102, wherein the isolated antibody comprises the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 66, 67, 68, 69, 70 and 71, respectively.

106) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 79-102, wherein the isolated antibody comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

107) The antibody according to any one of embodiments 1-71 for use according to any one of embodiments 79-102, wherein the isolated antibody comprises the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65.

108) A polynucleotide encoding the VH comprising the HCDR1, the HCDR2 and the HCDR3 of SEQ ID NOs: 10, 14 and 17, respectively.

109) A polynucleotide encoding the VL comprising the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs 23, 26 and 32, respectively.

110) A polynucleotide encoding the VH comprising the HCDR1, the HCDR2 and the HCDR3 of SEQ ID NOs: 10, 14 and 17, respectively, and the VL comprising the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs 23, 26 and 32, respectively.

111) A polynucleotide encoding the VH comprising the amino acid sequence of SEQ ID NO: 48.

112) A polynucleotide encoding the VL comprising the amino acid sequence of SEQ ID NO: 56.

113) A polynucleotide encoding the VH comprising the amino acid sequence of SEQ ID NO: 48 and the VL comprising the amino acid sequence of SEQ ID NO: 56.

114) A vector comprising the polynucleotide vector according to embodiment 108.

115) A vector comprising the polynucleotide vector according to embodiment 109.

116) A vector comprising the polynucleotide vector according to embodiment 110.

117) A vector comprising the polynucleotide vector according to embodiment 111.

118) A vector comprising the polynucleotide vector according to embodiment 112.

119) A vector comprising the polynucleotide vector according to embodiment 113.

120) A host cell comprising the vector according vector according to embodiment 114.

121) A host cell comprising the vector according vector according to embodiment 115.

122) A host cell comprising the vector according vector according to embodiment 116.

123) A host cell comprising the vector according vector according to embodiment 117.

124) A host cell comprising the vector according vector according to embodiment 118.

125) A host cell comprising the vector according vector according to embodiment 119.

126) A method of producing the antibody comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56, comprising culturing the host cell according to embodiment 125 in conditions that the antibody is expressed, and recovering the antibody produced by the host cell.

127) An immunoconjugate comprising the antibody or antigen-binding portion of the antibody comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56, linked to a therapeutic agent or an imaging agent.

128) A pharmaceutical composition comprising the antibody comprising the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56 and a pharmaceutically accepted carrier.

129) An isolated antagonistic antibody specifically binding PD-1, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively, for use in the treatment of cancer.

130) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 wherein the antibody comprises the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

131) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the cancer is a solid tumor.

132) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a melanoma.

133) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a lung cancer.

134) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a squamous non-small cell lung cancer (NSCLC).

135) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a non-squamous NSCLC.

136) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a colorectal cancer.

137) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a prostate cancer.

138) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a castration-resistant prostate cancer.

139) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a stomach cancer.

140) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is an ovarian cancer.

141) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a gastric cancer.

142) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a liver cancer.

143) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a pancreatic cancer.

144) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a thyroid cancer.

145) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a squamous cell carcinoma of the head and neck.

146) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a carcinoma of the esophagus or gastrointestinal tract.

147) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a breast cancer.

148) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a fallopian tube cancer.

149) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a brain cancer.

150) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is an urethral cancer.

151) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130,, wherein the solid tumor is a genitourinary cancer.

152) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is an endometriosis.

153) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a cervical cancer.

154) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 129 or 130, wherein the solid tumor is a metastatic lesion of the cancer.

155) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 131 in combination with a second therapeutic agent.

156) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 132 in combination with a second therapeutic agent.

157) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 133 in combination with a second therapeutic agent.

158) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 134 in combination with a second therapeutic agent.

159) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 135 in combination with a second therapeutic agent.

160) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 135 in combination with a second therapeutic agent.

161) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 137 in combination with a second therapeutic agent.

162) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 138 in combination with a second therapeutic agent.

163) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 139 in combination with a second therapeutic agent.

164) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 140 in combination with a second therapeutic agent.

165) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 141 in combination with a second therapeutic agent.

166) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 142 in combination with a second therapeutic agent.

167) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 143 in combination with a second therapeutic agent.

168) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 144 in combination with a second therapeutic agent.

169) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 145 in combination with a second therapeutic agent.

170) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 146 in combination with a second therapeutic agent.

171) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 147 in combination with a second therapeutic agent.

172) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 148 in combination with a second therapeutic agent.

173) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 149 in combination with a second therapeutic agent.

174) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 150 in combination with a second therapeutic agent.

175) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 151 in combination with a second therapeutic agent.

176) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 152 in combination with a second therapeutic agent.

177) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 153 in combination with a second therapeutic agent.

178) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 154 in combination with a second therapeutic agent.

179) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 155, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

180) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 156, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

181) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 157, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

182) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 158, wherein the second therapeutic agent is an antagonistic antibody

specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

183) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 159, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

184) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 160, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

185) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 161, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

186) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 162, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

187) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 163, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

188) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 164, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

189) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 165, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

190) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 166, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

191) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 167, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

192) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 168, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

193) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 169, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

194) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 170, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

195) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 171, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

196) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 172, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

197) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 173, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

198) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 174, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

199) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 175, wherein the second therapeutic agent is an antagonistic antibody

specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

200) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 176, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

201) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 177, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

202) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 178, wherein the second therapeutic agent is an antagonistic antibody specifically binding TIM-3, comprising the VH of SEQ ID NO: 146 and the VL of SEQ ID NO: 156.

203) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 155, wherein the second therapeutic agent is a vaccine.

204) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 156, wherein the second therapeutic agent is a vaccine.

205) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 157, wherein the second therapeutic agent is a vaccine.

206) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 158, wherein the second therapeutic agent is a vaccine.

207) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 159, wherein the second therapeutic agent is a vaccine.

208) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 160, wherein the second therapeutic agent is a vaccine.

209) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 161, wherein the second therapeutic agent is a vaccine.

210) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 162, wherein the second therapeutic agent is a vaccine.

211) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 163, wherein the second therapeutic agent is a vaccine.

212) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 164, wherein the second therapeutic agent is a vaccine.

213) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 165, wherein the second therapeutic agent is a vaccine.

214) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 166, wherein the second therapeutic agent is a vaccine.

215) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 167, wherein the second therapeutic agent is a vaccine.

216) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 168, wherein the second therapeutic agent is a vaccine.

217) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 169, wherein the second therapeutic agent is a vaccine.

218) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 170, wherein the second therapeutic agent is a vaccine.

219) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 171, wherein the second therapeutic agent is a vaccine.

220) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 172, wherein the second therapeutic agent is a vaccine.

221) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 173, wherein the second therapeutic agent is a vaccine.

222) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 174, wherein the second therapeutic agent is a vaccine.

223) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 175, wherein the second therapeutic agent is a vaccine.

224) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 176, wherein the second therapeutic agent is a vaccine.

225) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 177, wherein the second therapeutic agent is a vaccine.

226) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 178, wherein the second therapeutic agent is a vaccine.

227) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 155, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

228) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 156, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

229) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 157, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

230) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 158, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

231) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 159, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

232) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 160, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

233) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 161, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

234) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 162, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

235) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 163, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

236) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 164, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

237) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 165, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

238) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 166, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

239) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 167, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

240) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 168, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

241) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 169, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

242) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 170, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

243) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 171, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

244) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 172, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

245) The isolated antagonistic antibody specifically binding PD-1 for use according to embodiment 173, wherein the second therapeutic agent is a fibroblast growth factor receptor (FGFR) inhibitor.

The present invention will now be described with reference to the following specific, non-limiting examples.

Example 1. General methods

Purified human mixed lymphocyte reaction (MLR)

A purified human mixed lymphocyte reaction (MLR assay) was used to measure changes in cytokine production induced by addition of test antibodies to co-cultures of $CD4^+$ T cells and dendritic cells.

Peripheral blood mononuclear cells (PBMCs) were isolated from a leukopak (Biological Specialty Corporation) using a Ficoll gradient. $CD4^+$ T cells were then freshly isolated by negative selection from PBMCs using the Miltenyi AutoMACS and $CD4^+$ T cell isolation beads per manufacturer's instructions or were commercially purchased as frozen $CD4^+$ T cells (Hemacare Corporation). One dendritic cell donor (Hemacare Corporation) was used. Post-isolation or thaw, $CD4^+$ T cells and dendritic cells were washed and resuspended in assay media (RMPI1640 media supplemented with 10 % fetal

bovine serum, 1 % penicillin/streptomycin, 1X non-essential amino acids, and 1X sodium pyruvate-Invitrogen). The purified human $CD4^+$ T cells were diluted to $1x10^6$ cells/mL and seeded at 100,000 cells/100 µL/well. Dendritic cells were diluted to 0.1 $x10^6$ cells/mL and seeded at 5,000 cells/50 µL/well in U-bottom plates. Test antibodies or control antibodies were prepared at a 4X concentration in assay media yielding 1X when 50 µL of antibody was added to 150 µL of cells.

10-point serial dilutions of test or control antibodies were added to the wells at a final concentration of: 30, 10, 3.33, 1.11, 0.37, 0.12, 0.04, 0.01, 0.0046 and 00015 nM. $CD4^+$ T cells plus dendritic cells and dendritic cells alone were included as controls to measure basal cytokine secretion. Cells were maintained at 37 °C, 5 %CO_2 for 5 days. On Day 5, 100 µL of tissue culture supernatant was removed from culture plates and transferred to V-bottom plates. Supernatant was frozen at least overnight at -80 °C. Cumulative cytokine production was measured in tissue culture supernatant using Meso Scale Discovery (MSD) Th1/Th2 human cytokine 10-plex plates following manufacturer's protocol. Briefly, MSD plates were blocked with 1% blocker B overnight at 4 °C. The following day, blocker was removed and plates were washed using the Biotek 406 plate washer. An 8-point standard curve were prepared and added in duplicate to the plates. Thawed tissue culture supernatant was added at 25 µL/well, plates were sealed and shaken vigorously for 1.5 hours. Without removing standards or supernatant, 25 µL of detection antibody was added to each well. Plates were sealed, and shaken vigorously for 1.5 hours. Plates were washed, read buffer was added and plates were read using Meso Scale Discovery's plate reader.

Cytokine concentrations were calculated by MSD software. The concentration of cytokine in unknown samples is calculated by comparing the unknown's output signal to the output signal and known cytokine concentrations in the standard curve. Calculated concentrations were uploaded in Spotfire TIBCO software for visualization. After a visual inspection of the data, MAD-median outlier procedure with a threshold of 3.5 was used to identify and exclude outliers on log-transformed data. Robust analysis of the half-maximal effective concentration (Robust EC50) was carried out on each cytokine for each antibody.

CMV assay

A cytomegalovirus antigen recall assay (CMV assay) was used to measure changes in cytokine production induced by addition of test antibodies to cultures of peripheral blood mononuclear cells (PBMCs) with CMV whole antigen (for PD-1

antibodies) or with a pool of 138 15-mer peptides that overlap through the 65 kd phosphoprotein (pp65) (for TIM-3 mAbs and PD1/TIM-3 bispecific mAbs).

Post-thaw, PBMCs (Astarte Biologics and Hemcare Corporation) were washed and resuspended in assay media (RMPI1640 media supplemented with 10 % fetal bovine serum, 1 % penicillin/streptomycin, 1X non-essential amino acids, and 1X sodium pyruvate-Invitrogen). The PBMCs were diluted to$1.5x10^6$ cells/mL and seeded at 150,000 cells/100 µL/well. CMV antigen (Astarte Biologics) was prepared at a 4X concentration of 0.4 µg/mL in assay media yielding 0.1 µg/mL when 50 µL of antigen was added to 100 µL of cells and 50 µL of antibody. Antibodies were prepared at a 4X concentration in assay media yielding 1X when 50 µL of antibody was added to cells and peptide.

Serial dilutions of test antibodies were added to the wells at a final concentration between 150 – 0.001 nM. Cells plus CMV antigen or pp65 pool, cells alone, and isotype control prepared at a final concentration of 50 or 30 nM were included as controls to measure basal cytokine secretion. Cells were maintained at 37 °C, 5 %CO_2 for 6 days. For MSD analysis, on Day 6, 100 µL of tissue culture supernatant was removed from culture plates and transferred to V-bottom plates. Supernatant was frozen at least overnight at -80°C. Cumulative cytokine production was measured in tissue culture supernatant using Meso Scale Discovery (MSD) Th1/Th2 human cytokine 10-plex plates following manufacturer's protocol. Briefly, MSD plates were blocked with 1% blocker B overnight at 4 °C. The following day, blocker was removed and plates were washed using the Biotek 406 plate washer. An 8-point standard curve was prepared and added in duplicate to the plates. Thawed tissue culture supernatant was added at 25 µL/well, plates were sealed and shaken vigorously for 1.5 hours. Without removing standards or supernatant, 25 µL of detection antibody was added to each well. Plates were sealed, and shaken vigorously for 1.5 hours. Plates were washed, read buffer was added and plates were read using Meso Scale Discovery's plate reader.

Cytokine concentrations were calculated by MSD software. The concentration of cytokine in unknown samples is calculated by comparing the unknown's output signal to the output signal and known cytokine concentrations in the standard curve. Calculated concentrations were uploaded in Spotfire TIBCO software for visualization. After a visual inspection of the data, MAD-median outlier procedure with a threshold of 3.5 was used to identify and exclude outliers on log-transformed data. Robust analysis of the half-maximal effective concentration (Robust EC50) was carried out on each cytokine for each antibody.

For TIM-3 antibodies and PD1/TIM-3 bispecific antibodies, at day 6, after supernatant was collected for MSD analysis, cells were washed once with PBS and

subsequently stained for Live/Dead discrimination and the following cell surface markers: CD3, CD4, CD8, CD137, PD-1 and TIM-3. Flow cytometry was performed on a LSR Fortessa (BD). Data was analyzed using the Flow Jo software. CD137+ cells were identified based on Fluorescence Minus One (FMO) method on viable CMV-treated CD8+ and CD4+ cells.

For the sequential treatment experiments, CMV recall assays were carried out as above with pp65 peptide pool stimulation for six days. On day six, supernatant was removed and cells were restimulated with pp65 pool in the presence of anti-TIM-3 antibodies. Twenty-four hours later, supernatant was removed and IFN-γ levels were measured by MSD, as described above.

PD-1 Ligand inhibition assay

The ligand inhibition assay design was MSD (Mescoscale Discovery) based. A MSD plate was directly coated with ligand (cynoPDL1-ECD, huPDL1-ECD or huPDL2-ECD) and incubated overnight at 4°C. The following day, the coating solution was removed and the plate was blocked. A fixed concentration of biotinylated PD-1 (huPD1-ECD) was pre incubated with antibodies or with an isotype control antibody as a negative control. Depending on the panel of antibodies to be tested, the antibodies were tested as titrations or at a fixed concentration. The MSD plate was washed and the biotinylated PD-1/ antibody mixture was added to the ligand coated MSD plate. The plate was washed and biotinylated PD-1 bound to ligand was detected by ruthenylated streptavidin. Inhibition of PD-1 binding by an antibody resulted in decreased signal in the MSD assay. Maximal biotinylated PD-1 binding in the absence of inhibitor was determined and sometimes used to normalize the data to a percentage of maximal biotinylated PD-1 signal. The mAbs that were positive for inhibition of ligand binding at one concentration were also tested in dose responses for inhibition of various PD-1 ligands.

Jurkat cell binding

Jurkat cells were stimulated overnight with 20 ng/ml of PHA, harvested, washed, and checked for viability. The cells were then incubated at 6-10°C for 45-60 minutes with various concentrations of test antibodies, washed and incubated at 6-10°C for 45-60 minutes with FITC-labeled goat anti-human IgG. The cells were washed and fixed with BD Cytofix, refrigerated overnight and analyzed on a MACSQuant flow cytometer. The percentage of PD-1 positive cells at each antibody concentration was plotted vs log of the antibody concentration and EC_{50} values were generated in Prism.

Affinity measurements

PD-1 mAbs

Anti-PD-1 mAbs were tested for binding affinity to huPD1-ECD and cynoPD-1-ECD. Affinity measurements using Surface Plasmon Resonance (SPR) were performed using a ProteOn XPR36 system. A biosensor surface was prepared by coupling a mixture of anti-IgG Fc modified alginate polymer layer surface of a GLC chip using the manufacturer instructions for amine-coupling chemistry. Test mAbs were captured and their interactions with analytes (huPD1-ECD or cynoPD1-ECD) were monitored in PBS-based buffer at 25°C. The collected data were processed and fitted to a Langmuir 1:1 binding model. The result for each mAb was reported in the format of k_{on} (On-rate), k_{off} (Off-rate) and K_D (equilibrium dissociation constant).

TIM-3 ligand inhibition assay

TIM-3/galectin-9 competition ELISAs were done by binding 1 µg/ml recombinant human Fc-TIM-3 chimera (R&D Systems-cat#: 2365-TM-05) in PBS per well of a 96-well White Maxisorp plate (Nunc). The plates were washed and blocked with StartingBlock T20 (Pierce) and inhibitor at a 10 µg/ml concentration was added to the wells. Without washing, 7.5 µg/ml galectin-9 at was added to the wells and incubated for 30 min. Anti-galectin-9-biotin antibody polyclonal antibody (R&D Systems) at 0.5 µg/mL was then added and incubated for 30 minutes. The plates were washed and neutravidin-HRP-conjugated (Pierce) was added and the plates incubated for an additional 45 minutes. The plates were washed and POD Chemiluminescence substrate (Roche) was added immediately prior to reading plates and the luminescence was read on a luminometer.

Generation of antigens used in the study

Cloning, expression and purification of the antigens was done using standard methods. Various protein fragments were expressed as hexahistidine tag or Fc fusion proteins. The amino acid sequences of the used proteins without the tag sequences are shown in **SEQ ID NOs: 1-9, 138 and 89.**

Full length human PD1 (huPD1); **SEQ ID NO: 1**

PGWFLDSPDRPWNPPTFSPALLVVTEGDNATFTCSFSNTSESFVLNWYRMSPSNQT
DKLAAFPEDRSQPGQDCRFRVTQLPNGRDFHMSVVRARRNDSGTYLCGAISLAPK
AQIKESLRAELRVTERRAEVPTAHPSPSPRPAGQFQTLVVGVVGGLLGSLVLLVW

VLAVICSRAARGTIGARRTGQPLKEDPSAVPVFSVDYGELDFQWREKTPEPPVPCV
PEQTEYATIVFPSGMGTSSPARRGSADGPRSAQPLRPE DGHCSWPL

Extracellular domain of human PD1 (huPD1-ECD); **SEQ ID NO: 2**
PGWFLDSPDRPWNPPTFSPALLVVTEGDNATFTCSFSNTSESFVLNWYRMSPSNQT
DKLAAFPEDRSQPGQDCRFRVTQLPNGRDFHMSVVRARRNDSGTYLCGAISLAPK
AQIKESLRAELRVTERRAEVPTAHPSPSPRPAGQFQTL

Macaca fascicularis (cynomolgous, herein referred to as cyno) PD1 (cPD1); **SEQ ID NO: 3**)
PGWFLESPDRPWNAPTFSPALLLVTEGDNATFTCSFSNASESFVLNWYRMSPSNQ
TDKLAAFPEDRSQPGQDCRFRVTRLPNGRDFHMSVVRARRNDSGTYLCGAISLAP
KAQIKESLRAELRVTERRAEVPTAHPSPSPRPAGQFQALVVGVVGGLLGSLVLLV
WVLAVICSRAAQGTIEARRTGQPLKEDPSAVPVFSVDYGELDFQWREKTPEPPAP
CVPEQTEYATIVFPSGLGTSSPARRGSADGPRSPRPLRPEDGHCSWPL

Extracellular domain of cyno PD1 (cPD1-ECD); **SEQ ID NO: 4**
PGWFLESPDRPWNAPTFSPALLLVTEGDNATFTCSFSNASESFVLNWYRMSPSNQ
TDKLAAFPEDRSQPGQDCRFRVTRLPNGRDFHMSVVRARRNDSGTYLCGAISLAP
KAQIKESLRAELRVTERRAEVPTAHPSPSPRPAGQFQAL

Full length human PD-L1 (huPD-L1); **SEQ ID NO: 5**
FTVTVPKDLYVVEYGSNMTIECKFPVEKQLDLAALIVYWEMEDKNIIQFVHGEED
LKVQHSSYRQRARLLKDQLSLGNAALQITDVKLQDAGVYRCMISYGGADYKRIT
VKVNAPYNKINQRILVVDPVTSEHELTCQAEGYPKAEVIWTSSDHQVLSGKTTTT
NSKREEKLFNVTSTLRINTTTNEIFYCTFRRLDPEENHTAELVIPELPLAHPPNER

Extracellular domain of human PD-L1 (huPDL1-ECD) **SEQ ID NO: 6**
FTVTVPKDLYVVEYGSNMTIECKFPVEKQLDLAALIVYWEMEDKNIIQFVHGEED
LKVQHSSYRQRARLLKDQLSLGNAALQITDVKLQDAGVYRCMISYGGADYKRIT
VKVNAPYNKINQRILVVDPVTSEHELTCQAEGYPKAEVIWTSSDHQVLSGKTTTT
NSKREEKLFNVTSTLRINTTTNEIFYCTFRRLDPEENHTAELVIPELPLAHPPNERT

Extracellular domain of cynomolgus PD-L1 (cynoPDL1-ECD) **SEQ ID NO: 7**

AFTVTVPKDLYVVEYGSNMTIECKFPVEKQLDLTSLIVYWEMEDKNIIQFVHGEE
DLKVQHSNYRQRAQLLKDQLSLGNAALRITDVKLQDAGVYRCMISYGGADYKRI
TVKVNAPYNKINQRILVVDPVTSEHELTCQAEGYPKAEVIWTSSDHQVLSGKTTT
TNSKREEKLLNVTSTLRINTTANEIFYCIFRRLDPEENHTAELVIPELPLALPPNERT

Extracellular domain of human PD-L2 (huPDL2-ECD) **SEQ ID NO: 8**
LFTVTVPKELYIIEHGSNVTLECNFDTGSHVNLGAITASLQKVENDTSPHRERATLL
EEQLPLGKASFHIPQVQVRDEGQYQCIIIYGVAWDYKYLTLKVKASYRKINTHILK
VPETDEVELTCQATGYPLAEVSWPNVSVPANTSHSRTPEGLYQVTSVLRLKPPPG
RNFSCVFWNTHVRELTLASIDLQSQMEPRTHPT

Extracellular domain of mouse PD1 (musPD1-ECD) **SEQ ID NO: 9**
LEVPNGPWRSLTFYPAWLTVSEGANATFTCSLSNWSEDLMLNWNRLSPSNQTEK
QAAFCNGLSQPVQDARFQIIQLPNRHDFHMNILDTRRNDSGIYLCGAISLHPKAKIE
ESPGAELVVTERILETSTRYPSPSPKPEGRFQ

Full length human TIM-3, **SEQ ID NO: 138**
SEVEYRAEVGQNAYLPCFYTPAAPGNLVPVCWGKGACPVFECGNVVLRTDERDV
NYWTSRYWLNGDFRKGDVSLTIENVTLADSGIYCCRIQIPGIMNDEKFNLKLVIKP
AKVTPAPTRQRDFTAAFPRMLTTRGHGPAETQTLGSLPDINLTQISTLANELRDSR
LANDLRDSGATIRIGIYIGAGICAGLALALIFGALIFKWYSHSKEKIQNLSLISLANL
PPSGLANAVAEGIRSEENIYTIEENVYEVEEPNEYYCYVSSRQQPSQPLGCRFAMP

Extracellular domain of human TIM-3 (huTIM-3-ECD) **SEQ ID NO: 89**
SEVEYRAEVGQNAYLPCFYTPAAPGNLVPVCWGKGACPVFECGNVVLRTDERDV
NYWTSRYWLNGDFRKGDVSLTIENVTLADSGIYCCRIQIPGIMNDEKFNLKLVIKP
AKVTPAPTRQRDFTAAFPRMLTTRGHGPAETQTLGSLPDINLTQISTLANELRDSR
LANDLRDSGATIR

Example 2. Selection of human anti-PD-1 antibodies from phage display libraries

PD-1 binding Fabs were selected from *de novo* pIX phage display libraries as described in Shi *et al.,* J Mol Biol 397:385-96, 2010, Int. Patent Publ. No. WO2009/085462 and U.S. Patent Publ. No. US2010/0021477. Briefly, the libraries were generated by diversifying human scaffolds where germline VH genes IGHV1-69*01, IGHV3-23*01, and IGHV5-51*01 were recombined with the human IGHJ-4 minigene *via*

the H3 loop, and human germline VL kappa genes O12 (IGKV1-39*01), L6 (IGKV3-11*01), A27 (IGKV3-20*01), and B3 (IGKV4-1*01) were recombined with the IGKJ-1 minigene to assemble complete VH and VL domains. The positions in the heavy and light chain variable regions around H1, H2, L1, L2 and L3 loops corresponding to positions identified to be frequently in contact with protein and peptide antigens were chosen for diversification. Sequence diversity at selected positions was limited to residues occurring at each position in the IGHV or IGLV germline gene families of the respective IGHV or IGLV genes. Diversity at the H3 loop was generated by utilizing short to mid-sized synthetic loops of lengths 7-14 amino acids. The amino acid distribution at H3 was designed to mimic the observed variation of amino acids in human antibodies. Library design is detailed in Shi *et al.,* (2010) *J Mol Biol* 397:385-96. The scaffolds utilized to generate libraries were named according to their human VH and VL germline gene origin. The three heavy chain libraries were combined with the four germline light chains or combined with the diversified light chain libraries to generate 12 unique VH:VL combinations. These libraries were later combined further based on library versions to generate additional libraries for panning experiments against PD-1.

The libraries were panned against huPD1-ECD, cynoPD1-ECD, musPD1-ECD, huPD1-Fc and/or musPD1-Fc. The recombinant proteins were biotinylated (bt) and captured on streptavidin magnetic beads (Dynal), then exposed to the *de novo* pIX Fab libraries at a final concentration of 100nM or 10nM. Non-specific phages were washed away in PBS-Tween and bound phages were recovered by infection of MC1061F' *E. coli* cells. Phages were amplified from these cells overnight and panning was repeated for a total of three or four rounds. Following the final round of biopanning, monoclonal Fab was screened for binding to huPD1-ECD, huPD1-Fc, musPD1-Fc and/or cynoPD1-Fc in two ELISA formats. In Format 1, Fab was captured on an ELISA plate by anti-Fd antibody and the various forms of btPD1's were added to captured Fab, followed by detection of bt-PD1's with Streptavidin:HRP. In Format 2, the various forms of btPD1's were captured on ELISA plates by Streptavidin and secreted Fab was added to the captured antigen, followed by detection of the Fab with GoatAntiFab'2HRP. Clones that demonstrated binding to the proteins were sequenced in the heavy and light chain variable regions.

Fabs from the human PD-1 or mouse PD-1 selections were then tested for cross-reactivity to cynoPD1-Fc secreted in mammalian cell supernatant. Fab was captured on an ELISA plate by anti-Fd antibody and the cynoPD1-Fc supernatant was added to the captured Fab, followed by detection of cynoPD1-Fc with GoatAntiHumanFc:HRP. Based

on binding characteristics to cynoPD1-Fc, select antibodies were chosen for further characterization.

Select Fabs were chosen for further characterization and were cloned as IgG2sigma/κ. IgG2sigma has abolished effector functions and has V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions when compared to the wild type IgG2. IgG2sigma is described in U.S. Patent No. 8,961,967. The antibodies were evaluated for their ability to block human PD-1 binding to cynomolgus PD-L1, affinity to human and cynomolgus PD-1 proteins, and their ability to bind to cells endogenously expressing human PD-1 (Jurkat cells). The antibodies were subsequently evaluated for their ability to block human PD-L1 and human PD-L2 binding to huPD1.

Based on the results, several antibodies were chosen for affinity maturation. Characteristics of select antibodies chosen for affinity maturation are shown in **Table 7.**

Table 7.

	Ligand inhibition; IC_{50} (μg/ml)			Jurkat binding; EC_{50} μg/ml	ProteOn SPR affinity		
mAb	cynoPD-L1	huPD-L1	huPD-L2		k_{on} (1/Ms)	k_{off} (1/s)	K_D (nM)
PD1B11	0.017-0.018	0.019	0.029	0.03-0.24	4.68E+05	8.96E-03	19.2
PD1B70	0.010-0.021	0.040	0.059	0.69-1.32	1.84E+05	3.04E-02	166
PD1B71	0.014-0.015	0.024	0.035	0.13-0.47	2.31E+05	2.77E-02	120
Hu: human							
Cyno: cynomolgus							

Example 3. Affinity-maturation of human anti-PD-1 antibodies

Antibodies PD1B70, PD1B71 and PD1B114 (close homolog to PD1B11), were affinity matured in Fab format using phage display libraries with diversity at select VL positions and at HCDR1 and HCDR2. The design of affinity-maturation libraries for each Fab is shown in **Table 8.** Residue numbering is according to PD1B114 VH SEQ ID NO: 41 in Table 8.

Table 8.

Diversification of PD1B114, PD1B70 and PD1B71 VH		
Position	Parent amino acid	Residues used for diversification
30	S	D, K, S
31	S	D, N, S, T
32	Y	A, D, S, Y
33	A	A, D, G, S, W, Y
35	S	H, N, S
50	G	A, E, G, N, R, T, W, Y
52	I	A, D, I, N, R, S
54	I	E, I, N, S, Y
55	F	E, F, Q, S, Y
57	T	D, N, R, S, T, Y
59	N	E, G, N, Q, R, Y
Diversification of PD1B114, PD1B70 and PD1B71 VL		
Position	Parent amino acid	Residues used for diversification
30	S	D, N, R, S
31	S	N, S, T
32	Y	D, N, R, S, Y
49	Y	E, H, K, Y
50	D	D, G, S, W, Y
53	N	D, N, S, T, Y
91	R	A, D, E, G, H, N, R, S, W, Y
92	S	A, D, E, G, H, N, R, S, W, Y
93	N	A, D, E, G, H, N, R, S, W, Y
94	W	A, D, E, G, H, N, R, S, W, Y
96	L	F, I, L, N, R, W, Y

The libraries were constructed and phage was generated. The VH and the VL

phage libraries were then used for phage panning against huPD1-ECD and cynoPD1-ECD biotinylated recombinant proteins. Following phage panning, soluble Fabs were screened for binding to both human and cyno PD-1. Select Fabs were cloned as IgG2sigma isotype and characterized for their Jurkat cell binding and cynomolgus PD-L1 ligand inhibition at concentrations 1 μg/ml and 10 μg/ml.

Table 9 shows the characterization results of the parental and affinity-matured antibodies.

Table 9.

mAb	Ligand inhibition at indicated concentration*		Jurkat Cell binding; EC_{50} (μg/ml)
	1 μg/ml	10 μg/ml	
PD1B11	5%	5%	0.05
PD1B114	8%	13%	0.47
PD1B149	7%	7%	0.08
PD1B160	4%	3%	0.08
PD1B162	7%	6%	0.05
PD1B164	6%	3%	0.06
PD1B183	5%	5%	0.08
PD1B184	4%	4%	0.08
PD1B185	8%	5%	0.09
PD1B187	7%	5%	0.09
PD1B192	5%	5%	0.06
PD1B70	6%	6%	0.69
PD1B175	6%	5%	0.09
PD1B71	6%	9%	0.13
PD1B177	7%	8%	0.05
*value indicates percentage ligand not blocked			

The affinity matured antibodies were assessed in affinity experiments as described above using ProteOn SPR analyses for binding to huPD1-ECD and cynoPD1-ECD. The binding characteristics of the mAbs to cyno PD-1 are shown in **Table 10** and to human PD-1 in **Table 11.** STDEV were calculated for 3 or more replicates generated for human and cyno proteins. If less than 3 replicates were calculated, RANGE was indicated. RANGE is defined as the low and high values for the replicates tested. For samples in the **Table 10** or **Table 11** without value indicated in RANGE or STDEV, only one experiment was performed. The best affinity matured variants had affinities for human and cyno PD-1 in the single digit nM range following ~4-20 fold gains in affinity compared to their parental mAbs.

Table 10.

Sample	antigen: cyno PD-1					
	k_{on}	STDEV. kon	k_{off}	STDEV. koff	K_D	STDEV. K_D
	(1/Ms)	or RANGE	(1/s)	or RANGE	(nM)	or RANGE
PD1B70	2.10 E+05	(1.99-2.25) E+05	2.58 E-02	(2.45-2.75) E-02	123	109-138
PD1B175	2.14 E+05	(1.98-2.30) E+05	6.40 E-03	(6.06-6.73) E-03	30	26-34
PD1B71	3.04 E+05	2.35 E+04	2.03 E-02	7.27 E-04	66.8	5.68
PD1B177	2.92 E+05	(2.80-3.04) E+05	1.89 E-03	(1.84-1.93) E-03	6.47	6.1-6.9
PD1B114	2.94 E+05	1.69 E+04	2.39 E-02	1.45 E-03	81.5	6.8
PD1B149	3.20 E+05	(3.04-3.36) E+05	3.57 E-03	(3.48-3.65) E-03	11.2	(10.9-11.4)
PD1B160	3.17 E+05	(3.16-3.17) E+05	1.66 E-03	(1.63-1.68) E-03	5.23	5.1-5.3
PD1B162	3.87 E+05	(3.84-3.89) E+05	9.79 E-04	(9.59-9.98) E-04	2.53	2.5-2.6

PD1B164	2.67 E+05	(2.67-2.67) E+05	2.87 E-04	(2.82-2.91) E-04	1.07	1.06-1.09
PD1B11	2.93 E+05	(2.85-3.01) E+05	9.17 E-03	(0.8-1.00) E-02	31.3	(27.7-35.1)
PD1B183	3.20 E+05	(3.04-3.37) E+05	8.39 E-03	(8.01-8.76) E-03	26.3	23.9-28.8
PD1B184	2.38 E+05	(2.08-2.68) E+05	2.74 E-03	(2.55-2.92) E-03	11.5	9.5-14.1
PD1B185	3.11 E+05	(2.80-3.43) E+05	9.47 E-03	(9.38-9.55) E-03	30.5	27.5-34.1
PD1B187	2.94 E+05	(2.20-3.70) E+05	1.57 E-03	(1.28-1.85) E-03	5.32	3.5-8.4
PD1B192	3.07 E+05	(2.90-3.24) E+05	5.04 E-03	(4.86-5.22) E-03	16.4	15.0-18.0

Table 11.

	Antigen: human PD-1		
Sample	k_{on}	k_{off}	K_D
	(1/Ms)	(1/s)	(nM)
PD1B70	4.15E+05	4.18E-02	101
PD1B175	4.22E+05	9.72E-03	23
PD1B71	5.48E+05	2.73E-02	49.9
PD1B177	5.15E+05	2.57E-03	5
PD1B114	5.17E+05	2.79E-02	54.1
PD1B149	5.32E+05	6.20E-03	~12*
PD1B160	5.40E+05	3.71E-03	6.87
PD1B162	6.49E+05	3.86E-03	5.95
PD1B164	4.48E+05	1.31E-03	2.92
PD1B11	5.16E+05	8.52E-03	~17*
PD1B183	5.27E+05	8.44E-03	16
PD1B184	4.45E+05	5.09E-03	11.4
PD1B185	5.85E+05	7.65E-03	13.1
PD1B187	5.35E+05	2.78E-03	5.2

PD1B192	5.41E+05	1.17E-02	~228
*Values did not pass the data acceptance criteria (chi2 > 20%) and were therefore considered approximations.			

Example 4. Combinatorial variant PD-1 mAb production

Following the analysis of the affinity results, combinatorial sequences were considered.

PD1B11 and PD1B114 have very similar sequences. Because PD1B11 had approximately a 3-fold tighter affinity to human PD-1 and a 2-fold tighter affinity to cyno PD-1 compared to PD1B114, antibodies having combinations of their various CDRs were made. The HCDR3 of PD1B11 was placed into PD1B164 and PD1B162 (affinity-matured variants of PD1B114), using site directed mutagenesis while the HCDR2 of PD1B164 (affinity matured variant of PD1B114) was placed into PD1B187 (affinity matured variant of PD1B11). The resulting heavy chains were paired with parental light chains resulting in new antibodies PD1B194, PD1B195 and PD1B196, respectively.

PD1B175 and PD1B177 both contained the parental light chain even though the antibodies were generated using diversified VL libraries during affinity maturation. In an attempt to increase antibody affinities, PD1B175 heavy chain was paired with PD1L185 or PD1L187 affinity matured light chains, and PD1B177 heavy chain was paired with PD1L86, PD1L168 or PD1L190 affinity matured light chains, resulting in antibodies PD1B197, PD1B198, PD1B199, PD1B200 and PD1B201. VH and VL pairing of the antibodies is shown in **Table 20 in Example 5.**

The HCDR, LCDR, VH and VL sequences of these antibodies are shown in **Tables 14, 15, 16, 17, 18, 19, 21 and 22 in Example 5.** The antibodies were cloned as IgG2sigma/κ mAbs and transiently expressed in HEK293 expi cells for affinity measurements.

Affinities of the resulting antibodies were determined as described above. **Table 12** shows the measured affinities of the combinatorial mAb variants to cyno PD-1 and **Table 13** shows the affinities to human PD-1. STDEV were calculated for 3 or more replicates generated for human and cyno proteins. If less than 3 replicates were calculated, RANGE is indicated. RANGE is defined as the low and high values for the replicates tested. For samples without RANGE or STDEV, only one experiment was performed

Table 12.

Sample	binding to cyno PD-1					
	k_{on}	STDEV. kon	k_{off}	STDEV. koff	K_D	STDEV. KD
	(1/Ms)	or RANGE	(1/s)	or RANGE	(nM)	or RANGE
PD1B70 (Parent)	2.50E+05	(2.25-2.74) E+05	2.22 E-02	(2.18-2.26) E-02	88.98	(79.6-100)
PD1B197	2.75E+05	1.27 E+04	1.26 E-03	4.04 E-05	4.6	0.3
PD1B198	3.72E+05	1.61 E+04	4.16 E-03	9.29 E-05	11.18	0.54
PD1B11 (Parent)	3.50E+05	(3.49-3.50) E+05	9.42 E-03	(9.38-9.46) E-03	26.95	(26.8-27.1)
PD1B194	3.22E+05	2.86 E+04	1.93 E-04	5.86E-06	0.6	0.06
PD1B195	4.32E+05	(4.30-4.34) E+05	4.08 E-04	(3.96-4.19) E-04	0.94	(0.91-0.97)
PD1B196	3.03E+05	6.66 E+03	1.76 E-04	9.85 E-06	0.58	0.03
PD1B71 (Parent)	3.77E+05	(3.37-4.17) E+05	1.96 E-02	(1.85-2.07) E-02	51.99	(44.4-61.4)
PD1B199	3.40E+05	7.94 E+03	1.77 E-04	1.55 E-05	0.52	0.05
PD1B200	3.80E+05	2.21 E+04	4.22 E-04	1.99 E-05	1.11	0.08
PD1B201	3.05E+05	1.80 E+04	2.93 E-04	2.35 E-05	0.96	0.1

Table 13.

Sample	binding to human PD-1					
	k_{on}	STDEV. kon	k_{off}	STDEV. koff	K_D	STDEV. KD
	(1/Ms)	or RANGE	(1/s)	or RANGE	(nM)	or RANGE
PD1B70 (Parent)	7.69 E+05	(7.37-8.00) E+05	3.49 E-02	(3.41-3.56) E-02	45.35	(42.6-43.8)
PD1B197	6.58 E+05	2.26 E+04	3.24 E-03	1.74 E-04	4.9	0.3
PD1B198	8.95 E+05	6.44 E+04	9.34 E-03	9.90 E-04	10.43	1.34
PD1B11 (Parent)	9.33 E+05	(8.84-9.82) E+05	9.05 E-03	(8.67-9.43) E-03	9.7	(9.6-9.81)
PD1B194	8.97 E+05	1.45 E+05	9.60 E-04	2.78 E-05	1.07	0.18
PD1B195	1.23 E+06	1.79 E+05	1.52 E-03	6.51 E-05	1.23	0.19
PD1B196	8.83 E+05	6.39E+04	3.66 E-04	2.01E-05	0.41	0.04
PD1B71 (Parent)	9.55 E+05	(9.33-9.76) E+05	2.25 E-02	(2.19-2.30) E-02	23.52	(22.4-24.7)
PD1B199	9.33 E+05	6.92 E+04	5.64 E-04	1.98 E-05	0.6	0.05
PD1B200	1.05 E+06	1.40 E+05	1.22 E-03	3.21 E-05	1.17	0.16
PD1B201	8.58 E+05	8.22 E+04	9.57 E-04	3.06 E-05	1.12	0.11

Example 5. Structural characterization of anti-PD1 antibodies derived from phage display libraries

The cDNA sequences and amino acid translations of the antibodies were obtained using standard techniques throughout the generation of the antibodies using various campaigns. After polypeptide sequence determination, some antibody cDNAs encoding

the variable regions or full length antibodies were codon optimized using standard methods for scale-up expression.

Table 14 shows the HCDR1 sequences of select PD-1 antibodies.

Table 15 shows the HCDR2 sequences of select PD-1 antibodies.

Table 16 shows the HCDR3 sequences of select PD-1 antibodies.

Table 17 shows the LCDR1 sequences of select PD-1 antibodies.

Table 18 shows the LCDR2 sequences of select PD-1 antibodies.

Table 19 shows the LCDR3 sequences of select PD-1 antibodies.

Table 20 shows the VH and the VL pairing of select PD-1 antibodies.

Table 21 shows the VH sequences of select PD-1 antibodies.

Table 22 shows the VL sequences of select PD-1 antibodies.

Table 14.

Antibody	HCDR1					
	Sequence					SEQ ID NO:
PD1B114	S	Y	A	I	S	10
PD1B149	S	Y	A	I	S	10
PD1B160	S	Y	A	I	S	10
PD1B162	S	Y	A	I	S	10
PD1B164	S	Y	A	I	S	10
PD1B11	S	Y	A	I	S	10
PD1B183	S	Y	A	I	S	10
PD1B184	S	Y	A	I	S	10
PD1B185	S	Y	A	I	S	10
PD1B187	S	Y	A	I	S	10
PD1B192	S	Y	A	I	S	10
PD1B71	S	Y	A	I	S	10
PD1B177	D	Y	V	I	S	11
PD1B70	S	Y	A	I	S	10
PD1B175	S	Y	V	I	H	12
PD1B194	S	Y	A	I	S	10
PD1B195	S	Y	A	I	S	10
PD1B196	S	Y	A	I	S	10

PD1B197	S	Y	V	I	H	12
PD1B198	S	Y	V	I	H	12
PD1B199	D	Y	V	I	S	11
PD1B200	D	Y	V	I	S	11
PD1B201	D	Y	V	I	S	11

Table 15.

Antibody	HCDR2																	
	Sequence																	SEQ ID NO:
PD1B114	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B149	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B160	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B162	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B164	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B11	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B183	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B184	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B185	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B187	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B192	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B71	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B177	G	I	I	P	I	Y	G	T	A	N	Y	A	Q	K	F	Q	G	15
PD1B70	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B175	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B194	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B195	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B196	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B197	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B198	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B199	G	I	I	P	I	Y	G	T	A	N	Y	A	Q	K	F	Q	G	15

PD1B200	G	I	I	P	I	Y	G	T	A	N	Y	A	Q	K	F	Q	G	15
PD1B201	G	I	I	P	I	Y	G	T	A	N	Y	A	Q	K	F	Q	G	15

Table 16.

Antibody	HCDR3														
	Sequence														SEQ ID NO:
PD1B114	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B149	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B160	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B162	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B164	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B11	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B183	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B184	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B185	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B187	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B192	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B71	G	T	L	D	R	T	G	H	L	D	Y				18
PD1B177	G	T	L	D	R	T	G	H	L	D	Y				18
PD1B70	G	Y	V	R	A	T	G	M	L	D	Y				19
PD1B175	G	Y	V	R	A	T	G	M	L	D	Y				19
PD1B194	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B195	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B196	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B197	G	Y	V	R	A	T	G	M	L	D	Y				19
PD1B198	G	Y	V	R	A	T	G	M	L	D	Y				19
PD1B199	G	T	L	D	R	T	G	H	L	D	Y				18
PD1B200	G	T	L	D	R	T	G	H	L	D	Y				18
PD1B201	G	T	L	D	R	T	G	H	L	D	Y				18

Table 17.

Antibody	LCDR1											
	Sequence											SEQ ID NO:
PD1B114	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B149	R	A	S	Q	S	V	R	N	Y	L	A	21
PD1B160	R	A	S	Q	S	V	D	S	Y	L	A	22
PD1B162	R	A	S	Q	S	V	D	S	Y	L	A	22
PD1B164	R	A	S	Q	S	V	R	S	Y	L	A	23
PD1B11	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B183	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B184	R	A	S	Q	S	V	R	N	Y	L	A	21
PD1B185	R	A	S	Q	S	V	R	N	Y	L	A	21
PD1B187	R	A	S	Q	S	V	R	S	Y	L	A	23
PD1B192	R	A	S	Q	S	V	D	S	Y	L	A	22
PD1B71	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B177	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B70	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B175	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B194	R	A	S	Q	S	V	R	S	Y	L	A	23
PD1B195	R	A	S	Q	S	V	D	S	Y	L	A	22
PD1B196	R	A	S	Q	S	V	R	S	Y	L	A	23
PD1B197	R	A	S	Q	S	V	S	N	Y	L	A	24
PD1B198	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B199	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B200	R	A	S	Q	S	V	D	N	Y	L	A	25
PD1B201	R	A	S	Q	S	V	S	N	Y	L	A	24

Table 18.

Antibody	LCDR2	
	Sequence	SEQ

								SEQ ID NO:
PD1B114	D	A	S	N	R	A	T	26
PD1B149	D	A	S	N	R	A	T	26
PD1B160	D	A	S	D	R	A	T	27
PD1B162	D	A	S	N	R	A	T	26
PD1B164	D	A	S	Y	R	A	T	28
PD1B11	D	A	S	N	R	A	T	26
PD1B183	D	A	S	N	R	A	T	26
PD1B184	D	A	S	N	R	A	T	26
PD1B185	D	A	S	D	R	A	T	27
PD1B187	D	A	S	N	R	A	T	26
PD1B192	D	A	S	N	R	A	T	26
PD1B71	D	A	S	N	R	A	T	26
PD1B177	D	A	S	N	R	A	T	26
PD1B70	D	A	S	N	R	A	T	26
PD1B175	D	A	S	N	R	A	T	26
PD1B194	D	A	S	Y	R	A	T	28
PD1B195	D	A	S	N	R	A	T	26
PD1B196	D	A	S	N	R	A	T	26
PD1B197	D	A	S	N	R	A	T	26
PD1B198	D	A	S	S	R	A	T	29
PD1B199	D	A	S	T	R	A	T	30
PD1B200	D	A	S	N	R	A	T	26
PD1B201	D	A	S	N	R	A	T	26

Table 19.

Antibody	LCDR3	
	Sequence	SEQ ID NO:

PD1B114	Q	Q	R	S	N	W	P	L	T	31
PD1B149	Q	Q	R	N	Y	W	P	L	T	32
PD1B160	Q	Q	R	G	N	W	P	L	T	33
PD1B162	Q	Q	R	E	Y	W	P	L	T	34
PD1B164	Q	Q	R	D	Y	W	P	L	T	35
PD1B11	Q	Q	R	S	N	W	P	L	T	31
PD1B183	Q	Q	R	G	Y	W	P	L	T	36
PD1B184	Q	Q	R	N	Y	W	P	L	T	32
PD1B185	Q	Q	R	W	N	W	P	L	T	37
PD1B187	Q	Q	R	N	Y	W	P	L	T	32
PD1B192	Q	Q	R	N	Y	W	P	L	T	32
PD1B71	Q	Q	R	S	N	W	P	L	T	31
PD1B177	Q	Q	R	S	N	W	P	L	T	31
PD1B70	Q	Q	R	S	N	W	P	L	T	31
PD1B175	Q	Q	R	S	N	W	P	L	T	31
PD1B194	Q	Q	R	D	Y	W	P	L	T	35
PD1B195	Q	Q	R	E	Y	W	P	L	T	34
PD1B196	Q	Q	R	N	Y	W	P	L	T	32
PD1B197	Q	Q	R	A	Y	W	P	L	T	38
PD1B198	Q	Q	R	A	E	W	P	L	T	39
PD1B199	Q	Q	R	N	Y	W	P	L	T	32
PD1B200	Q	Q	R	S	A	W	P	L	T	40
PD1B201	Q	Q	R	N	Y	W	P	L	T	32

Table 20.

Antibody	VH peptide ID	VH SEQ ID NO:	VL peptide ID	VL SEQ ID NO:
PD1B114	PD1H24	41	PH9L3	49
PD1B149	PD1H24	41	PD1L128	50

PD1B160	PD1H131	42	PD1L101	51
PD1B162	PD1H131	42	PD1L67	52
PD1B164	PD1H131	42	PD1L71	53
PD1B11	PD1H3	43	PH9L3	49
PD1B183	PD1H3	43	PD1L109	54
PD1B184	PD1H3	43	PD1L128	50
PD1B185	PD1H3	43	PD1L132	55
PD1B187	PD1H3	43	PD1L148	56
PD1B192	PD1H3	43	PD1L133	57
PD1B71	PD1H108	44	PH9L3	49
PD1B177	PD1H164	45	PH9L3	49
PD1B70	PD1H107	46	PH9L3	49
PD1B175	PD1H163	47	PH9L3	49
PD1B194	PD1H170	48	PD1L71	53
PD1B195	PD1H170	48	PD1L67	52
PD1B196	PD1H170	48	PD1L148	56
PD1B197	PD1H163	47	PD1L185	58
PD1B198	PD1H163	47	PD1L187	59
PD1B199	PD1H164	45	PD1L86	60
PD1B200	PD1H164	45	PD1L168	61
PD1B201	PD1H164	45	PD1L190	62

Table 21.

VH peptide ID	VH SEQ ID NO:	VH sequence
PD1H24	41	QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQ APGQGLEWMGGIIPIFGTANYAQKFQGRVTITADESTSTA YMELSSLRSEDTAVYYCARPGLAAAYDTGNLDYWGQGT LVTVSS
PD1H131	42	QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQ APGQGLEWMGGIIPIFDTANYAQKFQGRVTITADESTSTA YMELSSLRSEDTAVYYCARPGLAAAYDTGNLDYWGQGT

		LVTVSS
PD1H3	43	QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIFGTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSLDYWGQGTLVTVSS
PD1H108	44	QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIFGTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARGTLDRTGHLDYWGQGTLVTVSS
PD1H164	45	QVQLVQSGAEVKKPGSSVKVSCKASGGTFSDYVISWVRQAPGQGLEWMGGIIPIYGTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARGTLDRTGHLDYWGQGTLVTVSS
PD1H107	46	QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIFGTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARGYVRATGMLDYWGQGTLVTVSS
PD1H163	47	QVQLVQSGAEVKKPGSSVKVSCKASGGTFKSYVIHWVRQAPGQGLEWMGGIIPIFGTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARGYVRATGMLDYWGQGTLVTVSS
PD1H170	48	QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIFDTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSLDYWGQGTLVTVSS

Table 22.

VL peptide ID	VL SEQ ID NO:	VL sequence
PH9L3	49	EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA

		VYYCQQRSNWPLTFGQGTKVEIK
PD1L128	50	EIVLTQSPATLSLSPGERATLSCRASQSVRNYLAWYQQKPG QAPRLLIHDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRNYWPLTFGQGTKVEIK
PD1L101	51	EIVLTQSPATLSLSPGERATLSCRASQSVDSYLAWYQQKPG QAPRLLIKDASDRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRGNWPLTFGQGTKVEIK
PD1L67	52	EIVLTQSPATLSLSPGERATLSCRASQSVDSYLAWYQQKPG QAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQREYWPLTFGQGTKVEIK
PD1L71	53	EIVLTQSPATLSLSPGERATLSCRASQSVRSYLAWYQQKPG QAPRLLIYDASYRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRDYWPLTFGQGTKVEIK
PD1L109	54	EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPG QAPRLLIKDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRGYWPLTFGQGTKVEIK
PD1L132	55	EIVLTQSPATLSLSPGERATLSCRASQSVRNYLAWYQQKPG QAPRLLIYDASDRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRWNWPLTFGQGTKVEIK
PD1L148	56	EIVLTQSPATLSLSPGERATLSCRASQSVRSYLAWYQQKPG QAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRNYWPLTFGQGTKVEIK
PD1L133	57	EIVLTQSPATLSLSPGERATLSCRASQSVDSYLAWYQQKPG QAPRLLIHDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRNYWPLTFGQGTKVEIK
PD1L185	58	EIVLTQSPATLSLSPGERATLSCRASQSVSNYLAWYQQKPG QAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRAYWPLTFGQGTKVEIK
PD1L187	59	EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPG QAPRLLIEDASSRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRAEWPLTFGQGTKVEIK
PD1L86	60	EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPG QAPRLLIHDASTRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRNYWPLTFGQGTKVEIK

PD1L168	61	EIVLTQSPATLSLSPGERATLSCRASQSVDNYLAWYQQKPG QAPRLLIHDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRSAWPLTFGQGTKVEIK
PD1L190	62	EIVLTQSPATLSLSPGERATLSCRASQSVSNYLAWYQQKPG QAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFA VYYCQQRNYWPLTFGQGTKVEIK

All anti-PD-1 antibodies were identified to have VH1-69 (**SEQ ID NO: 170**) and IGKV3-11 (L6) (**SEQ ID NO: 171**) frameworks.

SEQ ID NO: 170
QVQLVQSGAEVKKPGSSVKVSCKASGGTFS SYAIS WVRQAPGQGLEWMG GIIPIFGTANYAQKFQG RVTITADESTSTAYMELSSLRSEDTAVYYCAR

SEQ ID NO: 171
EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRAT GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRSNWP

Example 6. Generation and characterization of PD-1 antibodies in mice

BALB/c were immunized intraperitoneally with huPD1-ECD and assessed for specific IgG titers. Once sufficient titers were obtained, splenocytes were isolated and fused with FO cells. The resulting hybridomas were plated in 96 well plates and cultured for 10 days. Antigen specific clones were identified by standard capture ELISA for binding to huPD1-ECD. Human PD-1-specific hybridomas were further tested for their affinity to human and cyno PD-1, binding to Jurkat cells and cyno PD-L1 inhibition. Based on the results, clone PD1B28 was selected for humanization using framework adaptation.

Framework adaptation process was done as essentially described in U.S. Patent Publ. No. 2009/0118127 and Fransson *et al.,* (2010) *J Mol Biol* 398:214-231. Briefly, the heavy and light chain sequences were compared with the human germline sequences (only the "01" alleles as of Oct 01, 2007) using BLAST search against the IMGT database (Kaas, *et al.,* (2004) *Nucl Acids Res* 32, D208-D210; Lefranc *et al.,* (2005) *Nucl Acid Res* 33, D593-D597). From this set of human germline genes, redundant genes (100% identical at amino acid level) and those with unpaired cysteine residues were removed. The remaining closest matching human germline genes in both the framework and CDR

regions were chosen as the acceptor human frameworks. Several VL and VH germline human frameworks were selected based upon overall sequence homology and CDR lengths as well as CDR similarity. FR-4 was selected based on sequence similarity of the IGHJ/IGJK germline genes. Then, the CDRs of PD1B28 were transferred into the selected acceptor human frameworks to generate the HFA variants, except in the region corresponding to the HCDR1 of V_H. For this region a combination of CDR and HV, or a shorter HCDR2 (referred to as Kabat-7, see U.S. Patent Publ. No. 2009/0118127) were transferred from the non-human antibody into the human FRs because the remaining HCDR2 residues have not been found in contact in antigen-antibody complexes of known structures (Almagro, (2004) *J Mol Recognit.* 17:132). Backumtations were introduced into certain residue positions in the humanized antibodies. PD1B131 backmutations: VH: V37I_Q39L_W47S_R98S, VL: Y49K. PD1B132: VH W47S_R98S, VL: Y49K (residue numbering according to Chothia). Select antibodies were expressed as IgG2sigma/κ. The resulting antibodies were characterized for their binding to recombinant PD-1 and PD-1 expressed on cells (Jurkat cells), and their ligand inhibition (cyno PD-L1 and human PD-L1). Characteristics of select humanized antibodies are shown in **Table 23.** The VH and the VL sequences of the generated antibodies are shown in **Table 24 and Table 25**, respectively.

Table 23.

mAb	Jurkat cell binding relative to PD1B28	Human PD-1 Affinity			PD-L1 Inhibition, IC_{50} (ng/ml)	
		kon (1/Ms)	koff (1/s)	K_D (pM)	Human PD-L1	Cyno PD-L1
PD1B28	100%	9.70 E+05	1.18 E-04	122	67	96
PD1B131	100%	8.27 E+05	1.05 E-04	127	79	96
PD1B132	100%	9.14 E+05	8.80 E-05	96	55	79

Table 24.

mAb	VH ID	VL ID	VH sequence	VH SEQ ID NO:
PD1B131	PD1H130	PD1L62	EVQLVESGGGLVQPGGSLRLSC AASGFAFSRYDMSWIRLAPGK GLESVAYISGGGANTYYLDNV KGRFTISRDNAKNSLYLQMNSL RAEDTAVYYCASPYLSYFDVW GQGTLVTVSS	63
PD1B132	PD1H129	PD1L62	EVQLVESGGGLVQPGGSLRLSC AASGFAFSRYDMSWVRQAPGK GLESVAYISGGGANTYYLDNV KGRFTISRDNAKNSLYLQMNSL RAEDTAVYYCASPYLSYFDVW GQGTLVTVSS	64

Table 25.

mAb	VH ID	VL ID	VL sequence	VL SEQ ID NO:
PD1B131	PD1H130	PD1L62	EIVMTQSPATLSVSPGERATLSC RASQSLSDYLHWYQQKPGQAP RLLIKSASQSISGIPARFSGSGSG TEFTLTISSLQSEDFAVYYCQNG HSFPYTFGQGTKLEIK	65
PD1B132	PD1H129	PD1L62	EIVMTQSPATLSVSPGERATLSC RASQSLSDYLHWYQQKPGQAP RLLIKSASQSISGIPARFSGSGSG TEFTLTISSLQSEDFAVYYCQNG HSFPYTFGQGTKLEIK	65

The CDR sequences of PD1B131 and PD1B132 are shown below:

HCDR1 (SEQ ID NO: 66)
RYDMS

HCDR2 (SEQ ID NO: 67)
YISGGGANTYYLDNVKG

HCDR3 (SEQ ID NO: 68)
PYLSYFDV

LCDR1 (SEQ ID NO: 69)
RASQSLSDYLH

LCDR2 (SEQ ID NO: 70)
SASQSIS

LCDR3 (SEQ ID NO: 71)
QNGHSFPYT

Example 7. Effect of isotype switching on anti-PD-1 antibody properties

Variable regions of antibodies PD1B196 and PD1B199 (of IgG2sigma/κ isotype) were cloned as IgG4 S228P isotypes and variable regions from antibody PD1B132 (of IgG2) into IgG2sigma isotype to assess possible differences in functionality and developability.

The antibodies were named PD1B244 (PD1B196 VH/VL on IgG4 S228P) PD1B245 (PD1B199 VH/VL on IgG4 S228P) AND PD1B243 (PD1B132 VH/VL on IgG2sigma).

Isotype switch had no consistent effect on the antibody properties however, for some of the antibodies, some change in EC_{50} values were seen in the CMV assay.

Exemplified below are heavy chain and light chain amino acid sequences of various antibodies. **Table 26** shows the summary of the VH, VL, heavy chain and light chain SEQ ID NOs: for select antibodies.

Table 26.

Antibody	VH peptide ID	VH SEQ ID NO:	VL peptide ID	VL SEQ ID NO:	HC SEQ ID NO	LC SEQ ID NO:
PD1B114	PD1H24	41	PH9L3	49	212	213
PD1B149	PD1H24	41	PD1L128	50	214	215
PD1B160	PD1H131	42	PD1L101	51	216	217
PD1B162	PD1H131	42	PD1L67	52	218	219
PD1B164	PD1H131	42	PD1L71	53	220	221
PD1B183	PD1H3	43	PD1L109	54	222	223
PD1B184	PD1H3	43	PD1L128	50	224	225
PD1B185	PD1H3	43	PD1L132	55	226	227
PD1B192	PD1H3	43	PD1L133	57	228	229
PD1B243	PD1H129	64	PD1L62	65	74	75
PD1B244	PD1H170	48	PD1L148	56	72	73
PD1B245	PD1H164	45	PD1L86	60	76	77

SEQ ID NO: 72 HC of PD1B244

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
DTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSL
DYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWN
SGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTKTYTCNVDHKPSNTKVDKR
VESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQ
FNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKG
LPSSIEKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESN
GQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQ
KSLSLSLGK

SEQ ID NO: 73 LC of PD1B244

EIVLTQSPATLSLSPGERATLSCRASQSVRSYLAWYQQKPGQAPRLLIYDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRNYWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 74 HC of PD1B243

EVQLVESGGGLVQPGGSLRLSCAASGFAFSRYDMSWVRQAPGKGLESVAYISGG
GANTYYLDNVKGRFTISRDNAKNSLYLQMNSLRAEDTAVYYCASPYLSYFDVWG
QGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALT
SGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTKTYTCNVDHKPSNTKVDKRVESKY
GPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWY
VDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSI
EKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPE
NNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQKSLSL
SLGK

SEQ ID NO: 75 LC of PD1B243

EIVMTQSPATLSVSPGERATLSCRASQSLSDYLHWYQQKPGQAPRLLIKSASQSISG
IPARFSGSGSGTEFTLTISSLQSEDFAVYYCQNGHSFPYTFGQGTKLEIKRTVAAPS
VFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSK
DSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 76 HC of PD1B245

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSDYVISWVRQAPGQGLEWMGGIIPIY
GTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARGTLDRTGHLDY
WGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSG
ALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTKTYTCNVDHKPSNTKVDKRVES
KYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFN
WYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLP
SSIEKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNG
QPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQK
SLSLSLGK

SEQ ID NO: 77 LC of PD1B245

EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLLIHDASTRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRNYWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 212 HC of PD1B114

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
GTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGN
LDYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSW
NSGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDK
TVERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEV
QFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNK
GLPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWES
NGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT
QKSLSLSPGK

SEQ ID NO: 213 LC of PD1B114

EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRSNWPLTFGQGTKVEIKRTVAAP
SVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDS
KDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO 214 HC of PD1B149

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
GTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGN
LDYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSW
NSGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDK
TVERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEV
QFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNK
GLPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWES
NGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT
QKSLSLSPGK

SEQ ID NO: 215 LC of PD1B149

EIVLTQSPATLSLSPGERATLSCRASQSVRNYLAWYQQKPGQAPRLLIHDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRNYWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 216 HC of PD1B160

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
DTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGN
LDYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSW
NSGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDK
TVERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEV
QFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNK
GLPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWES
NGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT
QKSLSLSPGK

SEQ ID NO: 217 LC of PD1B160

EIVLTQSPATLSLSPGERATLSCRASQSVDSYLAWYQQKPGQAPRLLIKDASDRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRGNWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 218 HC of PD1B162

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
DTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGN
LDYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSW
NSGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDK
TVERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEV
QFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNK
GLPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWES
NGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT
QKSLSLSPGK

SEQ ID NO: 219 LC of PD1B162

EIVLTQSPATLSLSPGERATLSCRASQSVDSYLAWYQQKPGQAPRLLIYDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQREYWPLTFGQGTKVEIKRTVAAP
SVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDS
KDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 220 HC of PD1B164

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
DTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGN
LDYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSW
NSGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDK
TVERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEV
QFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNK
GLPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWES
NGQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT
QKSLSLSPGK

SEQ ID NO: 221 LC of PD1B164

EIVLTQSPATLSLSPGERATLSCRASQSVRSYLAWYQQKPGQAPRLLIYDASYRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRDYWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 222 HC of PD1B183

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
GTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSL
DYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWN
SGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDKT
VERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQ
FNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKG
LPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESN
GQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQ
KSLSLSPGK

SEQ ID NO: 223 LC of PD1B183

EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLLIKDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRGYWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 224 HC of PD1B184

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
GTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSL
DYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWN
SGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDKT
VERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQ
FNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKG
LPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESN
GQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQ
KSLSLSPGK

SEQ ID NO: 225 LC of PD1B184

EIVLTQSPATLSLSPGERATLSCRASQSVRNYLAWYQQKPGQAPRLLIHDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRNYWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 226 HC of PD1B185

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
GTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSL
DYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWN
SGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDKT
VERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQ
FNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKG
LPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESN
GQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQ
KSLSLSPGK

SEQ ID NO: 227 LC of PD1B185

EIVLTQSPATLSLSPGERATLSCRASQSVRNYLAWYQQKPGQAPRLLIYDASDRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRWNWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 228 HC of PD1B192

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
GTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSL
DYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWN
SGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDKT
VERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQ
FNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKG
LPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESN
GQPENNYKTTPPMLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQ
KSLSLSPGK

SEQ ID NO: 229 LC or PD1B192

EIVLTQSPATLSLSPGERATLSCRASQSVDSYLAWYQQKPGQAPRLLIHDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRNYWPLTFGQGTKVEIKRTVAA
PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

Example 8. Characterization of PD-1 antibodies in cell-based assays

Select antibodies were characterized in MLR and CMV assays using protocols described in Example 1. The EC_{50} values for IFN-γ induction from MLR and CMV assays are shown in **Table 27.** In most cases, anti-PD-1 antibodies showed a dose-dependent increase in IFN-γ levels in both MLR and CMV assays.

Table 27.

Origin	mAb	MLR EC_{50}, nM	CMV EC_{50}, nM
Phage display	PD1B3	0.29	0.06
	PD1B91	0.05	0.03
	PD1B194	NT	NC
	PD1B195	NT	1.64
	PD1B196	0.14	0.31
	PD1B199	0.63	NC
	PD1B200	NT	3.81
	PD1B201	NT	2.60
	PD1B244	0.08	0.03

HFA	PD1B132	NT	0.07
	PD1B243	0.07	0.02
	NT: not tested		
	NC: no convergence		
	HFA: human framework adaptation		

In addition to IFN-γ, secreted levels of additional cytokines were also affected by PD-1 blockade in the two assays. Upon CMV stimulation, anti-PD-1 antibodies led to a dose-dependent induction of TNF-α and IL-4, whereas in the MLR assay they increased TNF-α and IL-2 levels.

Example 9. Generation of human anti-TIM-3 antibodies using phage display libraries

The *de novo* pIX Fab libraries described in Example 2 were panned against the extracellular domain of recombinant human TIM-3-Fc fusion protein (R&D Systems, #2365-TM; residues Ser22-Arg200 of full length TIM-3) (huTIM-3-Fc).

The recombinant protein was biotinylated (bt) and captured on streptavidin magnetic beads (Dynal), then exposed to the *de novo* pIX Fab libraries at a final concentration of 100nM. Non-specific phages were washed away in PBS-Tween and bound phages were recovered by infection of MC1061F' E. coli cells. Phages were amplified from these cells overnight and panning was repeated for a total of three rounds. Following the final round of biopanning, monoclonal Fab was screened for binding to biotinylated human TIM-3-Fc captured on ELISA plates by Streptavidin and secreted Fab was added to the captured antigen, followed by detection of the Fab with Goat Anti human kappa:HRP. Select antibodies were expressed and cloned on various IgG isotypes as indicated below, and characterized further.

Example 10. Generation of anti-TIM-3 antibodies in mice

Balb/c mice were immunized with recombinant human TIM-3-Fc fusion protein (R&D Systems, catalog #2365-TM) over the course of 18 days. Spleens were harvested, and a B cell enriched population was fused with FO mouse myeloma cells to generate mAb secreting hybridomas. The hybridoma supernatants were screened for binding by ELISA to TIM-3-Fc protein and an irrelevant human IgG1 Fc. TIM-3 specific supernatants were then assayed for the ability to bind to TIM-3 expressing THP-1 cells.

Select mAb HC and LC v-genes were cloned from the TIM-3 positive hybridomas using standard molecular biology techniques (RT-PCR followed by PCR fragment ligation into plasmid expression vectors). mAbs were expressed recombinantly, and the ELISA was repeated to confirm TIM-3 specific binding. Molecular models for murine antibody sequences to be human framework adapted were constructed using MOE (CCG, Montreal) and visually inspected. Potential problem positions that might influence antigen binding, VL/VH packing and/or core residues that might affect domain stabilities were identified. For both VL and VH, multiple human frameworks were proposed with or without back mutations to mouse framework sequences if problem positions were identified. The designed sequences were cloned into heavy and light chain plasmids and expressed in Expi293F cells. Expressed antibody in the culture supernatants were quantified and assessed for binding to HEK293 cells transfected with recombinant human TIM-3.

Example 11. Isotypes of anti-TIM-3 antibodies

The VH and VL of isolated anti-TIM-3 antibodies were cloned onto various heavy chain isotypes, optionally with various Fc substitutions, and allotypes with κ light chains during the course of antibody characterization to evaluate the effect, if any, of isotype switch on functionality or developability of the antibodies. The various isotypes used are shown in **Table 28.**

Table 28.

Isotype	Substitution when compared to wild type*	Purpose of substitution
IgG2sigma	V234A, G237A, P238S, H268A, V309L, A330S, P331S	Abolishing effector functions
IgG2sigma_K409R	V234A, G237A, P238S, H268A, V309L, A330S, P331S, K409R	Abolishing effector functions, improving heterodimer formation in bispecific antibody
IgG2sigma_F405L	V234A, G237A, P238S, H268A,	Abolishing effector functions, improving heterodimer formation in

	V309L, A330S, P331S, F405L	bispecific antibody
IgG4_PAA	S228P, F234A, L235A	Antibody stability, abolishing effector functions
IgG4_ PAA_F405L_R409K	S228P, F234A, L235A, F450L, R409K	Antibody stability, abolishing effector functions, improving heterodimer formation in bispecific antibody
IgG4_S228P	S228P	Antibody stability
IgG1	Wild type	
IgG1sigma	L234A, L235A, G237A, P238S, H268A, A330S, P331S	Abolishing effector functions
IgG1sigma_K409R	L234A, L235A, G237A, P238S, H268A, A330S, P331S, K409R	Abolishing effector functions, improving heterodimer formation in bispecific antibody
IgG1sigma_F405L	L234A, L235A, G237A, P238S, H268A, A330S, P331S, F405L	Abolishing effector functions, improving heterodimer formation in bispecific antibody
IgG1_AA	L234A, L235A	Abolishing effector functions
	*Residue numbering according to the EU Index	

The various allotypes used in the generated antibodies are shown in **Table 29.** Some of the antibodies had chimeric allotypes. Antibodies TM3B105 and TM3B403 for example differ by one amino acid substitution in a constant region at position 189. TM3B105 heavy and light chains SEQ ID NOs: 240 and 79, respectively; TM3B403 heavy and light chains SEQ ID NOs: 78 and 79, respectively. The two antibodies are expected to have the same characteristics.

Table 29.

Isotype/Allotype/Substitutions
IgG2sigma_G2m(n-)/(n)_K409R
IgG2sigma_G2m(n-)_K409R
IgG2sigma_G2m(n-)/(n)
IgG2sigma_F405L
IgG2_K409R
IgG2sigma_G2m(n-)
IgG2
IgG4_S228P
IgG4_S228P_F405L_R409K
IgG4_nG4m(a)_PAA_F405L_R409K
IgG4_PAA
IgG1sigma
IgG1_G1m(17)
IgG1_ G1m(17,1)_AA

In general, anti-TIM-3 antibodies with IgG2sigma Fc had greater activity in the CMV assay than anti-TIM-3 antibodies with huIgG4 Fc. In addition, antibodies with huIgG2 Fc demonstrated functionality that was intermediate between IgG2sigma and IgG4. Allotype had no effect on antibody activity.

Example 12. Structural characterization of anti-TIM-3 antibodies

The cDNA sequences and amino acid translations of the antibodies were obtained using standard techniques throughout the generation of the antibodies using various campaigns. After polypeptide sequence determination, some antibody cDNAs encoding the variable regions or full length antibodies were codon optimized using standard methods for scale-up expression. Antibodies TM3B103, TM3B105, M3B108, TM3B109 and TM3B113 were isolated from phage display libraries. Antibodies TM3B189, TM3B190, TM3B193, TM3B195 and TM3B196 were generated by immunizing mice.

Table 30 shows the HCDR1 sequences of select anti-TIM-3 antibodies.

Table 31 shows the HCDR2 sequences of select anti-TIM-3 antibodies.

Table 32 shows the HCDR3 sequences of select anti-TIM-3 antibodies.

Table 33 shows the LCDR1 sequences of select anti-TIM-3 antibodies.

Table 34 shows the LCDR2 sequences of select anti-TIM-3 antibodies.

Table 35 shows the LCDR3 sequences of select anti-TIM-3 antibodies.

Table 36 shows the VH sequences of select anti-TIM-3 antibodies.

Table 37 shows the VL sequences of select anti-TIM-3 antibodies.

Table 38 shows the frameworks of select anti-TIM-3 antibodies.

Table 30.

mAb name	HCDR1						
	Sequence						SEQ ID NO:
TM3B103	N	Y	W	M	S		90
TM3B105	S	Y	A	M	S		91
TM3B109	S	Y	A	M	S		91
TM3B108	G	Y	W	M	H		92
TM3B113	D	Y	W	M	S		93
TM3B189	S	Y	V	M	Y		94
TM3B190	S	D	Y	A	W	N	95
TM3B193	D	T	Y	L	H		96
TM3B195	S	Y	W	M	Q		97
TM3B196	S	Y	G	V	H		98
TM3B291	S	Y	W	M	Q		97

Table 31.

mAb	HCDR2																
	Sequence																SEQ ID NO:
TM3B103	A	I	S	G	S	G	G	S	T	Y	Y	A	D	S	V	K	G	99
TM3B105	A	I	S	G	S	G	G	S	T	Y	Y	A	D	S	V	K	G	99
TM3B109	A	I	S	G	S	G	G	S	T	Y	Y	A	D	S	V	K	G	99
TM3B108	A	I	S	Y	S	G	S	S	T	Y	Y	A	D	S	V	K	G	100
TM3B113	V	I	K	Y	S	G	G	S	K	Y	Y	A	D	S	V	K	G	101

TM3B189	Y	I	N	P	Y	N	D	G	T	K	Y	N	E	K	F	K	G	102
TM3B190	Y	I	N	Y	S	G	R	T	S	Y	N	P	S	L	K	S		103
TM3B193	R	I	D	P	T	N	G	N	I	K	Y	D	P	K	F	Q	G	104
TM3B195	A	I	Y	P	G	D	G	D	I	R	Y	T	Q	N	F	K	G	105
TM3B196	V	I	W	S	D	G	S	T	T	Y	N	S	A	L	K	S		106
TM3B291	A	I	Y	P	G	D	G	D	I	R	Y	T	Q	N	F	K	G	105

Table 32.

mAb	HCDR3															
	Sequence															SEQ ID NO:
TM3B103	D	H	W	D	P	N	F	L	D	Y						107
TM3B105	S	P	Y	A	P	L	D	Y								108
TM3B109	N	E	E	P	D	D	R	L	D	Y						109
TM3B108	G	T	N	W	L	D	Y									110
TM3B113	E	L	E	G	V	F	D	Y								111
TM3B189	D	D	Y	D	V	A	P	F	A	Y						112
TM3B190	G	G	N	F	D	Y										113
TM3B193	P	Y	Y	G	F	F	D	Y								114
TM3B195	W	E	K	S	T	T	V	V	Q	R	N	Y	F	D	Y	115
TM3B196	Q	A	N	Y	R	Y	D	S	A	M	D	Y				116
TM3B291	W	E	K	S	T	T	V	V	Q	R	N	Y	F	D	Y	115

Table 33.

mAb	LCDR1																	
	Sequence																	SEQ ID NO:
TM3B103	R	A	S	Q	S	V	S	S	S	Y	L	A						117
TM3B105	R	A	S	Q	S	V	N	D	Y	L	A							118
TM3B109	K	S	S	Q	S	V	L	A	S	S	N	N	K	N	Y	L	A	119
TM3B108	R	A	S	Q	S	V	S	S	S	Y	L	A						117

TM3B113	R	A	S	Q	S	V	S	N	S	T	L	A					120
TM3B189	R	A	S	E	S	L	D	S	Y	G	N	S	Y	I	H		121
TM3B190	Q	A	T	Q	D	I	V	K	N	L	N						122
TM3B193	K	A	S	Q	D	V	N	T	A	V	A						123
TM3B195	K	A	S	E	N	V	G	T	F	V	S						124
TM3B196	K	A	S	Q	S	V	D	Y	D	G	D	S	Y	M	N		125
TM3B291	K	A	S	E	N	V	G	T	F	V	S						124

Table 34.

mAb	LCDR2							
	Sequence							SEQ ID NO:
TM3B103	G	A	S	S	R	A	T	126
TM3B105	D	A	S	N	R	A	T	127
TM3B109	W	A	S	T	R	E	S	128
TM3B108	G	A	S	S	R	A	T	126
TM3B113	T	A	S	S	R	A	T	129
TM3B189	L	A	S	N	L	E	S	130
TM3B190	Y	V	T	E	L	A	E	131
TM3B193	S	A	T	Y	R	Y	T	132
TM3B195	G	A	S	N	R	Y	T	133
TM3B196	T	A	A	N	L	Q	S	134
TM3B291	G	A	S	N	R	Y	T	133

Table 35.

mAb	LCDR3									
	Sequence									SEQ ID NO:
TM3B103	Q	Q	Y	G	S	S	P	L	T	135

TM3B105	Q	Q	G	G	H	A	P	I	T	136
TM3B109	Q	Q	Y	Y	S	T	P	L	T	137
TM3B108	Q	Q	Y	G	S	S	P	L	T	135
TM3B113	Q	Q	S	Y	T	S	P	W	T	139
TM3B189	Q	Q	N	N	E	D	P	F	T	140
TM3B190	L	Q	F	Y	E	F	P	L	T	141
TM3B193	Q	Q	H	Y	S	T	P	Y	T	142
TM3B195	G	Q	S	Y	S	Y	P	T		143
TM3B196	Q	Q	S	N	E	D	P	F	T	144
TM3B291	G	Q	S	Y	S	Y	P	T		143

Table 36.

mAb name	VH name	VH sequence	SEQ ID NO:
TM3B103	TM3H21	EVQLLESGGGLVQPGGSLRLSCAASGFTFSN YWMSWVRQAPGKGLEWVSAISGSGGSTYY ADSVKGRFTISRDNSKNTLYLQMNSLRAED TAVYYCAKDHWDPNFLDYWGQGTLVTVSS	145
TM3B105	TM3H24	EVQLLESGGGLVQPGGSLRLSCAASGFTFSS YAMSWVRQAPGKGLEWVSAISGSGGSTYY ADSVKGRFTISRDNSKNTLYLQMNSLRAED TAVYYCAKSPYAPLDYWGQGTLVTVSS	146
TM3B108	TM3H30	EVQLLESGGGLVQPGGSLRLSCAASGFTFSG YWMHWVRQAPGKGLEWVSAISYSGSSTYY ADSVKGRFTISRDNSKNTLYLQMNSLRAED TAVYYCAKGTNWLDYWGQGTLVTVSS	147
TM3B109	TM3H31	EVQLLESGGGLVQPGGSLRLSCAASGFTFSS YAMSWVRQAPGKGLEWVSAISGSGGSTYY ADSVKGRFTISRDNSKNTLYLQMNSLRAED TAVYYCAKNEEPDDRLDYWGQGTLVTVSS	148
TM3B113	TM3H65	EVQLLESGGGLVQPGGSLRLSCAASGFTFSD	149

		YWMSWVRQAPGKGLEWVSVIKYSGGSKYY ADSVKGRFTISRDNSKNTLYLQMNSLRAED TAVYYCAKELEGVFDYWGQGTLVTVSS	
TM3B189	TM3H141	EVQLQQSGPELLKPGASVKMSCKASGYTFT SYVMYWVKQKPGQGLEWIGYINPYNDGTK YNEKFKGKATLTSDKSSSTAYMELSRLTSED SAVYYCTRDDYDVAPFAYWGQGTLVTVSA	150
TM3B190	TM3H96	DVQLQESGPGLVKPSQSLSLTCTVTGYSITS DYAWNWIRQFPGNKLEWMGYINYSGRTSY NPSLKSRISITRDTSKNQFFLQLNSVTTEDTA TYYCTSGGNFDYWGQGTTLTVSS	151
TM3B193	TM3H99	EVQLQQSGAELVKPGASVKLSCTASGFHIKD TYLHWVKQRPEQGLEWIGRIDPTNGNIKYD PKFQGKATITSDTSSNTAYLQLSSLTSEDTAV YYCARPYYGFFDYWGQGTTLTVSS	152
TM3B195	TM3H144	EVQLQQSGAELARPGASVKLSCKASGYTFT SYWMQWVKQRPGQGLEWIGAIYPGDGDIR YTQNFKGKATLTADKSSSTAYMQLSSLASE DSAVYYCARWEKSTTVVQRNYFDYWGQGT TLTVSS CORRECT?	153
TM3B196	TM3H102	QVQLKESGPGLVAPSQSLSITCTISGFSLTSY GVHWVRQPPGKGLEWLVVIWSDGSTTYNS ALKSRLSISKDNSKSQVFLKMNSLQTDDTA MYYCARQANYRYDSAMDYWGQGTSVTVS S	154
TM3B291	TM3H162	EVQLVQSGAEVKKPGESLKISCKGSGYSFTS YWMQWVRQMPGKGLEWMGAIYPGDGDIR YTQNFKGQVTISADKSISTAYLQWSSLKASD TAMYYCARWEKSTTVVQRNYFDYWGQGT TVTVSS	172

Table 37.

mAb name	VL name	VL sequence	SEQ ID

			NO:
TM3B103	PH9L1	EIVLTQSPGTLSLSPGERATLSCRASQSVSSS YLAWYQQKPGQAPRLLIYGASSRATGIPDRF SGSGSGTDFTLTISRLEPEDFAVYYCQQYGS SPLTFGQGTKVEIK	155
TM3B105	TM3L33	EIVLTQSPATLSLSPGERATLSCRASQSVNDY LAWYQQKPGQAPRLLIYDASNRATGIPARFS GSGSGTDFTLTISSLEPEDFAVYYCQQGGHA PITFGQGTKVEIK	156
TM3B108	PH9L1	EIVLTQSPGTLSLSPGERATLSCRASQSVSSS YLAWYQQKPGQAPRLLIYGASSRATGIPDRF SGSGSGTDFTLTISRLEPEDFAVYYCQQYGS SPLTFGQGTKVEIK	155
TM3B109	PYYL6	DIVMTQSPDSLAVSLGERATINCKSSQSVLA SSNNKNYLAWYQQKPGQPPKLLIYWASTRE SGVPDRFSGSGSGTDFTLTISSLQAEDVAVY YCQQYYSTPLTFGQGTKVEIK	157
TM3B113	TM3L12	EIVLTQSPGTLSLSPGERATLSCRASQSVSNS TLAWYQQKPGQAPRLLIYTASSRATGIPDRF SGSGSGTDFTLTISRLEPEDFAVYYCQQSYTS PWTFGQGTKVEIK	158
TM3B189	TM3L61	DIVLTQSPASLAVSLGQRATISCRASESLDSY GNSYIHWYQQKPGQPPKLLIYLASNLESGVP ARFSGSGSKTDFTLTIDPVEADDPATYYCQQ NNEDPFTFGSGTKLEIK	159
TM3B190	TM3L62	DIVMTQSPSSMSASLGDRITITCQATQDIVKN LNWYQQKPGKPPSFLIHYVTELAEGVPSRFS GSGSGSDYSLTISNLESEDFADYYCLQFYEFP LTFGAGTKLELK	160
TM3B193	TM3L52	DIVMTQSHKFMSTSVGDRVSITCKASQDVN TAVAWYQQKPGQSPKLLIYSATYRYTGVPD RFTGSGSGTDFTFTISSVQAEDLAVYYCQQH YSTPYTFGSGTKLEIK	161
TM3B195	TM3L67	DVQMIQSPKSMSMSVGERVTLSCKASENVG	162

		TFVSWYQQKPDQSPKLLIYGASNRYTGVPD RFTGSGSATDFTLTISSVQAEDLADYHCGQS YSYPTFGSGTKLEM	
TM3B196	TM3L64	DIQMTQSPASLAVSLGQRATISCKASQSVDY DGDSYMNWYQQKPGQPPKLLIYTAANLQS GIPARFSGSGSGTDFTLNIHPVEEEDAATYYC QQSNEDPFTFGSGTKLEIK	163
TM3B291	TM3L85	DIQMTQSPSSLSASVGDRVTITCKASENVGT FVSWYQQKPGKAPKLLIYGASNRYTGVPSR FSGSGSGTDFTLTISSLQPEDFATYYCGQSYS YPTFGQGTKLEIK	173

Table 38.

mAb name	VH name	VH framework		VL name	VL framework	
		Name	SEQ ID NO:		Name	SEQ ID NO:
TM3B103	TM3H21	IGHV3-23	174	PH9L1	IGKV3-20	180
TM3B105	TM3H24	IGHV3-23	174	TM3L33	IGKV3-11	171
TM3B108	TM3H30	IGHV3-23	174	PH9L1	IGKV3-20	180
TM3B109	TM3H31	IGHV3-23	174	PYYL6	IGKV4-1	181
TM3B113	TM3H65	IGHV3-23	174	TM3L12	IGKV3-20	180
TM3B189	TM3H141	IGHV1-02	175	TM3L61	IGKV4-1	181
TM3B190	TM3H96	IGHV4-30	176	TM3L62	IGKV1-39	182
TM3B193	TM3H99	IGHV1-03	177	TM3L52	IGKV1-33	183
TM3B195	TM3H144	IGHV1-03	177	TM3L67	IGKV1-39	182
TM3B196	TM3H102	IGHV2-26	178	TM3L64	IGKV4-1	181
TMB291	TM3H162	IGHV5-51	179	TM3L85	IGKV1-39	182

IGHV3-23 SEQ ID NO: 174
EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMSWVRQAPGKGLEWVS
AISGSGGSTYYADSVKG RFTISRDNSKNTLYLQMNSLRAEDTAVYYCAK

IGHV1-02 SEQ ID NO: 175
QVQLVQSGAEVKKPGASVKVSCKASGYTFT GYYMH WVRQAPGQGLEWMG
RINPNSGGTNYAQKFQG RVTSTRDTSISTAYMELSRLRSDDTVVYYCAR

IGHV4-30 SEQ ID NO: 176
QVQLQESGPGLVKPSQTLSLTCTVSGGSISSGDYYWSWIRQPPGKGLEWIGYIYYS
GSTYYNPSLKSRVTISVDTSKNQFSLKLSSVTAADTAVYYCAR

IGHV1-03 SEQ ID NO: 177
QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYAMHWVRQAPGQRLEWMG
WINAGNGNTKYSQKFQGRVTITRDTSASTAYMELSSLRSEDTAVYYCAR

IGHV2-26 SEQ ID NO: 178
QVTLKESGPVLVKPTETLTLTCTVSGFSLSNARMGVSWIRQPPGKALEWLA
HIFSNDEKSYSTSLKSRLTISKDTSKSQVVLTMTNMDPVDTATYYCARI

IGHV5-51 SEQ ID NO: 179
EVQLVQSGAEVKKPGESLKISCKGSGYSFTSYWIGWVRQMPGKGLEWMGIIYPG
DSDTRYSPSFQGQVTISADKSISTAYLQWSSLKASDTAMYYCAR

IGKV3-20 SEQ ID NO: 180
EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIY
GASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSP

IGKV3-11 SEQ ID NO: 171
EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRSNWP

IGKV4-1SEQ ID NO: 181
EIVLTQSPGTLSLSPGERATLSCRASQSVSSSYLAWYQQKPGQAPRLLIY
GASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSP

IGKV1-39 SEQ ID NO: 182
DIQMTQSPSSLSASVGDRVTITCRASQSISSYLNWYQQKPGKAPKLLIY
AASSLQSGVPSRFSGSGSGTDFTLTISSLQPEDFATYYCQQSYSTPI

GKV1-33 SEQ ID NO: 183
DIQMTQSPSSLSASVGDRVTITCQASQDISNYLNWYQQKPGKAPKLLIY
DASNLETGVPSRFSGSGSGTDFTFTISSLQPEDIATYYCQQYDNLP

Example 13. Characterization of anti-TIM-3 antibodies

Select antibodies were characterized for their binding to human or cyno cells, and their ability to block ligand galectin 9 binding. **Table 39** shows the characteristics of select antibodies in these assays. The cell binding data represents the calculated EC_{50} values of the antibodies binding to cells transfected with the indicated TIM-3 recombinant protein expressed in µg/ml units. The galectin-9 inhibition represents the maximal level of inhibition of galectin-9 binding to human TIM-3 seen with the indicated antibodies. The tested antibodies were tested as IgG2sigma isotypes.

Epitope mapping assays were performed by coating recombinant huTIM-3-Fc protein on MSD plates. Plates were blocked and washed, followed by the addition of the mixture of the MSD-tag-labeled anti-TIM-3 mAbs incubated with increasing concentrations of unlabeled anti-TIM-3 mABs. After incubation with gentle shaking at room temperature, plates were washed and analyzed with a SECTOR Imager 6000. Antibodies that competed with each other for binding to human TIM-3 were considered to bind to similar epitopes. Positive inhibition was noted if >75% of the binding was inhibited. Partial inhibition was 40-75% inhibition. < 40% inhibition was denoted as negative.

Table 39.

mAb	Cell binding EC_{50}, µg/ml		Galectin 9 Inhibition, % inhibition	Epitope Bin
	Human cells	Cyno cells		
TM3B103	0.71	0.09	71.2	1
TM3B105	0.46	0.03	69.8	1

TM3B107			74.8	2
TM3B108	0.42	0.03	64.2	1
TM3B109			77.0	1
TM3B113			75.6	2
TM3B189	0.74	0.19	76.4	3
TM3B190	0.35	0.08	60.7	1
TM3B193			47.4	3
TM3B219	0.60	0.10	38.0	3
TM3B196			57.0	4

Example 14. Development of a functional *in vitro* assay to characterize anti-TIM-3 antibodies

Functional assessment of inhibitory receptors such as PD-1 can be done using T cells from normal donor that are stimulated by allogeneic dendritic cells or specific antigens, such as Tetanus toxoid or CMV. In this setting, changes in T cell function with antibody treatment can be detected by measuring supernatant cytokine levels or markers of T cell activation. Effects of anti-TIM-3 antibodies can be very variable in these types of assays, with little overall change in the state of activation or functionality of bulk T cell (non-antigen-specific). On the other hand, using tetramer approaches to follow single T cell sub-populations/clones in these assays does not provide the resolution needed to detect functional effects of anti-TIM-3 antibodies, due to the low frequency and heterogeneous functional profile of these T cell clones. In addition, this approach necessitates the prior identification of the epitopes recognized by CMV-specific T cells in each donor.

CD137 was recently described as a surrogate marker for activated antigen-specific T cells (Wolf *et al.*, (2007) *Blood* 110(1):201-210; Klinger *et al.*, (2013) *PLoS One* 8(9): e74231). In our assays, using CD137 enabled the identification of antigen specific $CD8^+$ and $CD4^+$ T cells that expand in response to CMV antigen stimulation and allowed the detection of the functional effects of anti-TIM-3 antibodies. In addition to CD137 expression, cytokine secretion by MSD was also evaluated in these assays.

The activity of select anti-TIM-3 antibodies was tested in CMV pp65-stimulated PBMCs. In these assays, anti-TIM-3 antibodies augmented T cell activation, as evidenced by increased CD137 expression on both $CD8^+$ and $CD4^+$ T cells. In addition, selected anti-TIM-3 antibodies also enhanced secretion of IFN-γ and TNF-α in this assay.

Table 40 shows the results of the CMV assay where enhanced surface expression of CD137 was evaluated on CD8+ or CD4+ cells for select TIM-3 antibodies. The table shows the p values generated using the Two-tailed T-test (unequal variance).

Table 40.

	$CD8^+ CD137^+$, p values			$CD4^+ CD137^+$, p values		
	Mean	Std Dev	n	Mean	Std Dev	n
TM3B103	0.043	0.025	5	0.071	0.112	3
TM3B105	0.029	0.036	6	0.01	0.017	3
TM3B107	0.182	0.188	5	0.157	0.125	3
TM3B108	0.022	0.018	5	0.01	0.01	3
TM3B109	0.035	0.041	5	0.017	0.015	3
TM3B113	0.082	0.064	6	0.05	0.026	3
TM3B189	0.027	0.026	6	0.007	0.011	3
TM3B190	0.078	0.159	6	0.004	0.005	3
TM3B193	0.467	0.252	3	0.1	NA	1
TM3B195	0.035	0.043	7	0.01	0.01	3
TM3B196	0.328	0.183	6	0.733	0.058	3
TM3B197	0.473	0.303	4	0.3	NA	1

Example 15. Generation of bispecific PD-1/TIM-3 antibodies

Select monospecific PD-1 and TIM-3 antibodies were expressed as IgG1/κ, IgG2/κ or IgG4/κ. Substitutions were made at positions 405 and 409 (EU numbering) in the monospecific antibodies to promote subsequent *in vitro* arm exchange and formation of the bispecific antibodies. The IgG1 and IgG2 anti-PD-1 and anti-TIM-3 antibodies were engineered to have a F405L and a K409R substitution, respectively, to promote arm exchange and generation the bispecific antibodies. On IgG4, the 409 WT position is R, hence the IgG4 anti-PD-1 antibody was not engineered and the IgG4 anti-TIM-3 antibody was engineered to have F405L and R409K substitutions. In addition to position 405 and 409 substitutions, the IgG4 mAbs were engineered to have S228P substitution and the IgG2 antibodies were optionally engineered to include IgG2sigma substitution (V234A, G237A, P238S, H268A, V309L, A330S and P331S).

The monospecific antibodies were expressed and purified using standard methods using a Protein A column (HiTrap MabSelect SuRe column). After elution, the pools were dialyzed into D-PBS, pH 7.2

Bispecific PD-1/TIM-3 antibodies were generated by combining a monospecific PD-1 mAb and a monospecific TIM-3 mAb in *in vitro* Fab arm exchange as described in Int. Patent Publ. No. WO2011/131746. Briefly, at about 1-20 mg/ml at a molar ratio of 1:1 of each antibody in PBS, pH 7-7.4 and 75 mM 2-mercaptoethanolamine (2-MEA) was mixed together and incubated at 25-37ºC for 2-6 h, followed by removal of the 2-MEA via dialysis, diafiltration, tangential flow filtration and/or spinned cell filtration using standard methods.

The bispecific antibodies were further purified after the *in vitro* Fab-arm exchange using hydrophobic interaction chromatography to minimize residual parental PD-1 and TIM-3 antibodies using standard methods.

Select monospecific anti-PD-1 antibodies and anti-TIM-3 antibodies were combined in matrix in *in vitro* Fab arm exchange to generate bispecific antibodies. **Table 41, Table 42 and Table 43** show the VH, the VL, the HC and the LC sequences of the generated bispecific antibodies and their isotypes. The G2 antibody allotypes were G2m(n)/(n-) or G2m(n-).

In some experiments, control antibodies were used that were monovalent for either PD-1 or TIM-3 with the second arm being inert binding to gp120. The gp120 binding arm had a VH of SEQ ID NO: 184 and the VL of SEQ ID NO: 185. **Table 44** shows the generated control antibodies.

SEQ ID NO: 184 VH of gp120 binding mAb
QVQLVQSGAEVKKPGASVKVSCQASGYRFSNFVIHWVRQAPGQRFEWMGWINP
YNGNKEFSAKFQDRVTFTADTSANTAYMELRSLRSADTAVYYCARVGPYSWDDS
PQDNYYMDVWGKGTTVIVSS

SEQ ID NO: 185 VL of gp120 binding mAb
EIVLTQSPGTLSLSPGERATFSCRSSHSIRSRRVAWYQHKPGQAPRLVIHGVSNRAS
GISDRFSGSGSGTDFTLTITRVEPEDFALYYCQVYGASSYTFGQGTKLERK

Table 41.

mAb	PD-1 binding arm				
	VH1	VH1 SEQ ID NO:	VL1	VL1 SEQ ID NO:	Isotype
PTBB14	PD1H170	48	PD1L148	56	IgG2sigma
PTBB15	PD1H170	48	PD1L148	56	IgG2sigma
PTBB16	PD1H129	64	PD1L62	65	IgG2sigma
PTBB17	PD1H129	64	PD1L62	65	IgG2sigma
PTBB24	PD1H170	48	PD1L148	56	IgG2sigma
PTBB30	PD1H170	48	PD1L148	56	IgG2sigma
PTBB27	PD1H170	48	PD1L148	56	IgG2
PTBB28	PD1H170	48	PD1L148	56	IgG2
PTBB18	PD1H129	64	PD1L62	65	IgG4_S228P
PTBB20	PD1H170	48	PD1L148	56	IgG4_S228P
PTBB21	PD1H170	48	PD1L148	56	IgG4_S228P

Table 42.

mAb	TIM-3 binding arm				
	VH2	VH2 SEQ ID NO:	VL2	VL2 SEQ ID NO:	Isotype
PTBB14	TM3H144	153	TM3L67	162	IgG2sigma
PTBB15	TM3H24	146	TM3L33	156	IgG2sigma
PTBB16	TM3H144	153	TM3L67	162	IgG2sigma
PTBB17	TM3H24	146	TM3L33	156	IgG2sigma
PTBB24	TM3H162	172	TM3L85	173	IgG2sigma
PTBB30	TM3H24	146	TM3L33	156	IgG2sigma
PTBB27	TM3H162	172	TM3L85	173	IgG2
PTBB28	TM3H24	146	TM3L33	156	IgG2
PTBB18	TM3H24	146	TM3L33	156	IgG4_S228
PTBB20	TM3H24	146	TM3L33	156	IgG4_S228
PTBB21	TM3H162	172	TM3L85	173	IgG4_S228

Table 43.

mAb	SEQ ID NO:			
	PD-1 binding arm		TIM-3 binding arm	
	HC1	LC1	HC2	LC2

PTBB14	186	188	190	193
PTBB15	186	188	191	194
PTBB16	187	189	190	193
PTBB17	187	189	191	194
PTBB24	186	188	192	195
PTBB30	186	188	248	194
PTBB27	241	188	244	195
PTBB28	241	188	245	194
PTBB18	242	189	246	194
PTBB20	243	188	246	194
PTBB21	243	188	247	195

SEQ ID NO: 186

QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
DTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSL
DYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWN
SGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDKT
VERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQ
FNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKG
LPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESN
GQPENNYKTTPPMLDSDGSFLLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQ
KSLSLSPGK

SEQ ID NO: 187

EVQLVESGGGLVQPGGSLRLSCAASGFAFSRYDMSWVRQAPGKGLESVAYISGG
GANTYYLDNVKGRFTISRDNAKNSLYLQMNSLRAEDTAVYYCASPYLSYFDVWG
QGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALT
SGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDKTVERKC
CVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQFNWYV
DGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIE
KTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPEN
NYKTTPPMLDSDGSFLLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSL
SPGK

SEQ ID NO: 188

EIVLTQSPATLSLSPGERATLSCRASQSVRSYLAWYQQKPGQAPRLLIYDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQRNYWPLTFGQGTKVEIKRTVAA

PSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQD
SKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 189
EIVMTQSPATLSVSPGERATLSCRASQSLSDYLHWYQQKPGQAPRLLIKSASQSISG
IPARFSGSGSGTEFTLTISSLQSEDFAVYYCQNGHSFPYTFGQGTKLEIKRTVAAPS
VFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSK
DSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 190
EVQLQQSGAELARPGASVKLSCKASGYTFTSYWMQWVKQRPGQGLEWIGAIYPG
DGDIRYTQNFKGKATLTADKSSSTAYMQLSSLASEDSAVYYCARWEKSTTVVQR
NYFDYWGQGTTLTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVS
WNSGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVD
KTVERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPE
VQFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSN
KGLPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWE
SNGQPENNYKTTPPMLDSDGSFFLYSRLTVDKSRWQQGNVFSCSVMHEALHNHY
TQKSLSLSPGK

SEQ ID NO: 191
EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMSWVRQAPGKGLEWVSAISGSG
GSTYYADSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCAKSPYAPLDYWGQ
GTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTS
GVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKVDKTVERKCC
VECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQFNWYVD
GVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKT
ISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNY
KTTPPMLDSDGSFFLYSRLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG
K

SEQ ID NO: 192
EVQLVQSGAEVKKPGESLKISCKGSGYSFTSYWMQWVRQMPGKGLEWMGAIYP
GDGDIRYTQNFKGQVTISADKSISTAYLQWSSLKASDTAMYYCARWEKSTTVVQ
RNYFDYWGQGTTVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTV

SWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVTSSNFGTQTYTCNVDHKPSNTKV
DKTVERKCCVECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDP
EVQFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVS
NKGLPSSIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVE
WESNGQPENNYKTTPPMLDSDGSFFLYSRLTVDKSRWQQGNVFSCSVMHEALHN
HYTQKSLSLSPGK

SEQ ID NO: 193
DVQMIQSPKSMSMSVGERVTLSCKASENVGTFVSWYQQKPDQSPKLLIYGASNR
YTGVPDRFTGSGSATDFTLTISSVQAEDLADYHCGQSYSYPTFGSGTKLEMKRTV
AAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTE
QDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 194
EIVLTQSPATLSLSPGERATLSCRASQSVNDYLAWYQQKPGQAPRLLIYDASNRAT
GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQQGGHAPITFGQGTKVEIKRTVAAP
SVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDS
KDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 195
DIQMTQSPSSLSASVGDRVTITCKASENVGTFVSWYQQKPGKAPKLLIYGASNRY
TGVPSRFSGSGSGTDFTLTISSLQPEDFATYYCGQSYSYPTFGQGTKLEIKRTVAAP
SVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDS
KDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

SEQ ID NO: 241
QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
DTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSL
DYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWN
SGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKT
VERKCCVECPPCPAPPVAGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVQ
FNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVVHQDWLNGKEYKCKVSNKG
LPAPIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESN
GQPENNYKTTPPMLDSDGSFLLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQ
KSLSLSPGK

SEQ ID NO: 242
EVQLVESGGGLVQPGGSLRLSCAASGFAFSRYDMSWVRQAPGKGLESVAYISGG
GANTYYLDNVKGRFTISRDNAKNSLYLQMNSLRAEDTAVYYCASPYLSYFDVWG
QGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALT
SGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTKTYTCNVDHKPSNTKVDKRVESKY
GPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWY
VDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSI
EKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPE
NNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQKSLSL
SLGK

SEQ ID NO: 243
QVQLVQSGAEVKKPGSSVKVSCKASGGTFSSYAISWVRQAPGQGLEWMGGIIPIF
DTANYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYYCARPGLAAAYDTGSL
DYWGQGTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWN
SGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTKTYTCNVDHKPSNTKVDKR
VESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQ
FNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKG
LPSSIEKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESN
GQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQ
KSLSLSLGK

SEQ ID NO: 244
EVQLVQSGAEVKKPGESLKISCKGSGYSFTSYWMQWVRQMPGKGLEWMGAIYP
GDGDIRYTQNFKGQVTISADKSISTAYLQWSSLKASDTAMYYCARWEKSTTVVQ
RNYFDYWGQGTTVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTV
SWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKV
DKTVERKCCVECPPCPAPPVAGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDP
EVQFNWYVDGVEVHNAKTKPREEQFNSTFRVVSVLTVVHQDWLNGKEYKCKVS
NKGLPAPIEKTISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVE
WESNGQPENNYKTTPPMLDSDGSFFLYSRLTVDKSRWQQGNVFSCSVMHEALHN
HYTQKSLSLSPGK

SEQ ID NO: 245

EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMSWVRQAPGKGLEWVSAISGSG
GSTYYADSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCAKSPYAPLDYWGQ
GTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTS
GVHTFPAVLQSSGLYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKTVERKCC
VECPPCPAPPVAGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVQFNWYVD
GVEVHNAKTKPREEQFNSTFRVVSVLTVVHQDWLNGKEYKCKVSNKGLPAPIEK
TISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENN
YKTTPPMLDSDGSFFLYSRLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSP
GK

SEQ ID NO: 246
EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMSWVRQAPGKGLEWVSAISGSG
GSTYYADSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCAKSPYAPLDYWGQ
GTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTS
GVHTFPAVLQSSGLYSLSSVVTVPSSSLGTKTYTCNVDHKPSNTKVDKRVESKYG
PPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWYV
DGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIE
KTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPE
NNYKTTPPVLDSDGSFLLYSKLTVDKSRWQEGNVFSCSVMHEALHNHYTQKSLS
LSLGK

SEQ ID NO: 247
EVQLVQSGAEVKKPGESLKISCKGSGYSFTSYWMQWVRQMPGKGLEWMGAIYP
GDGDIRYTQNFKGQVTISADKSISTAYLQWSSLKASDTAMYYCARWEKSTTVVQ
RNYFDYWGQGTTVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTV
SWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTKTYTCNVDHKPSNTKV
DKRVESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDP
EVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVS
NKGLPSSIEKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVE
WESNGQPENNYKTTPPVLDSDGSFLLYSKLTVDKSRWQEGNVFSCSVMHEALHN
HYTQKSLSLSLGK

SEQ ID NO: 248
EVQLLESGGGLVQPGGSLRLSCAASGFTFSSYAMSWVRQAPGKGLEWVSAISGSG
GSTYYADSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVYYCAKSPYAPLDYWGQ

GTLVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTS
GVHTFPAVLQSSGLYSLSSVVTVPSSNFGTQTYTCNVDHKPSNTKVDKTVERKCC
VECPPCPAPPAAASSVFLFPPKPKDTLMISRTPEVTCVVVDVSAEDPEVQFNWYVD
GVEVHNAKTKPREEQFNSTFRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKT
ISKTKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNY
KTTPPMLDSDGSFFLYSRLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG
K

Table 44.

Control mAb	Arm 1 VH/VL with F405L substitution	Arm 2 VH/VL with K409R substitution	Isotype
TM3B342	gp120	TM3B195	IgG2sigma
TM3B343	gp120	TM3B299	IgG2sigma
B23B74	gp120	B23B32	IgG2sigma
PTBB23	gp120	TM3B291	IgG2sigma
PD1B355	PD1B246	gp120	IgG2sigma
PD1B356	PD1B248	gp120	IgG2sigma

Example 16. Characterization of bispecific PD-1/TIM-3 antibodies

The generated antagonistic bispecific antibodies were tested in the CMV assay for their ability to enhance antigen-specific T cell responses. Functionality was measured by assessing CD137 expression on both $CD4^+$ and CD^+ T cells and by IFN-γ and TNF-α levels in the culture supernatants as described in Example 14. **Table 45 and Table 46** summarize the activity of bispecific PD-1/TIM-3 antibodies in this assay for the different readouts. As shown in this table, select bispecific molecules led to significant increases in CD137 expression on $CD4^+$ and $CD8^+$ T cells and in levels of secreted IFN-γ and TNF-α.

Overall, the PD-1/TIM-3 bispecifics with huIgG2sigma Fc had the most robust activity, followed by those molecules with huIgG2 and then huIgG4.

Table 45.

mAb		Statistical Significance			
		$CD4^+CD137^+$		$CD8^+CD137^+$	
Isotype	name	Avg p value	St Dev	Avg p value	St Dev
IgG2sigma	PTBB14	0.1144	0.1591	0.0002	0.0001
IgG2sigma	PTBB15	0.0467	0.0988	0.0001	0.0000
IgG2sigma	PTBB16	0.0017	0.0023	0.0001	0.0000
IgG2sigma	PTBB17	0.4148	0.5051	0.0001	0.0001
IgG2sigma	PTBB24	0.0031	0.0051	0.0001	0.0000
IgG2	PTBB27	0.0009	0.0011	0.0001	0.0000
IgG2	PTBB28	0.0003	0.0002	0.0001	0.0000
IgG4	PTBB18*	0.0353		0.0071	
IgG4	PTBB20	0.6025	0.1710	0.0004	0.0004
IgG4	PTBB21	0.1071	0.1372	0.0059	0.0081
*one p value reported					

Table 46.

mAb		Statistical significance			
		IFN-γ		TNF-α	
Isotype	name	Avg p value	St Dev	Avg p value	St Dev
IgG2sigma	PTBB14	0.0001	0.0000	0.0112	0.0157
IgG2sigma	PTBB15	0.0001	0.0000	0.0005	0.0008
IgG2sigma	PTBB16	0.0001	0.0000	0.0012	0.0016
IgG2sigma	PTBB17	0.0001	0.0000	0.0001	0.0000
IgG2sigma	PTBB24	0.0001	0.0001	0.0008	0.0008
IgG2	PTBB27	0.0026	0.0030	0.3406	0.4757
IgG2	PTBB28	0.0001	0.0000	0.1437	0.1229

IgG4	PTBB18	0.0001	#DIV/0!	0.0008	#DIV/0!
IgG4	PTBB20	0.0544	0.0768	0.1754	0.2140
IgG4	PTBB21	0.0174	0.0245	0.2685	0.1103
*one p value reported					

Example 17. Anti-PD1 antibodies upregulate TIM-3 expression on tumors

Effect of anti-PD-1 antibody treatment in expression of TIM-3 on tumors were evaluated in CT26 or MC38 colon carcinoma mouse model.

Balb/c mice were implanted subcutaneously with $1x10^6$ CT26 colon carcinoma tumors. Seven days after tumor cell implant, tumors were measured and mice were randomized by tumor size. Treatment with PBS or 10mg/kg anti-mouse PD-1 antibodies (clone RMP1-14, BioXCell) began on day 7 after tumor cell implant and continued biweekly for the remainder of the study. To analyze T cell expression of TIM-3, tumors were harvested at day 22 and dissociated using GentleMACS (Miltenyi). Staining for flow cytometry was carried out with Live/Dead and markers for CD3, CD4, CD8 and TIM-3. Flow cytometry was performed on a LSR Fortessa (BD). Data was analyzed using the Flow Jo software.

Wild-type C57Bl/6 female mice were implanted subcutaneously with $5x10^5$ MC-38 colon carcinoma cells suspended in PBS. Tumors were measured and mice were randomized by tumor size (50-100mm^3). Treatment with PBS or 10mg/kg anti-mouse PD-1 (clone RMP1-14, BioXCell) began after randomization and continued biweekly for the remainder of the study. To profile tumor infiltrating T cells, tumors were harvested and dissociated using GentleMACS (Miltenyi) 12, 15, 19, or 22 days after implant.

Staining for flow cytometry was carried out with Live/Dead and markers for CD45, Thy1, CD3, CD4, CD8, TIM-3, CD137, OX40, GITR, TIGIT. Flow cytometry data was collected on a LSR Fortessa (BD). Data was analyzed using the FlowJo software (v9.9.4) and visualized with GraphPad Prism. Statistics were generated by GraphPad Prism.

Analysis of TIM-3 expression on CD8+ T cells isolated from CT26 tumors at day 22 revealed an increase of TIM-3 expression in the PD-1 treated samples, compared to PBS control. **Figure 1A** shows the mean fluorescent intensity of TIM-3 expression in the two treatment groups.

TIM-3 expression was also increased in MC-38 tumors in the anti-PD-1 mAb treated samples when compared to PBS control. **Figure 1B** shows the geometric mean fluorescent intensity of TIM-3 expression in the $CD8^+$ TIL population. **Figure 1C** shows the percentage (%) relative frequency of TIM-3^+ $CD8^+$ cells of total $CD8^+$ TILs.

These data show that TIM-3 is upregulated in response to anti-PD-1 treatment, supporting the rational for targeting TIM-3 in PD-1 treated subjects.

CD137, OX40 and GITR expression was also analyzed on CD8+ T cells infiltrating MC38 tumors isolated from mice treated with anti-mouse PD-1 antibodies. These results showed that both the frequency and level (gMFI) of TNF family costimulatory receptors CD137, OX40 and GITR expression was increased following PD-1 blockade. **Figure 2A** and **Figure 2B** show the gMFI and relative frequency of CD137 expression on CD8 TILs, respectively. **Figure 3A** and **Figure 3B** show the gMFI and relative frequency of OX40 expression on CD8 TILs, respectively, and **Figure 4A** and **Figure 4B** show the gMFI and relative expression of GITR on CD8 TILs, respectively.

These data support the rational for targeting CD137, OX40 and/or GITR in PD-1 treated subjects.

Example 18. Activity of anti-TIM-3 antibodies following PD-1 blockade

The activity of anti-TIM-3 antibodies was also tested following anti-PD-1 antibody blockade in the CMV assay. In these experiments, PBMCs from one normal donor (CMV-sera positive) were incubated with pp65 peptide pools and anti-PD-1 antibodies for 5 days. On day 5, supernatants were harvested and cells were re-stimulated with pp65 peptide pool in the presence of either anti-TIM-3 or anti-PD-1 antibody. IFN-γ levels in the supernatant were measured 24 hours later. Treatment with anti-TIM-3 antibodies after 5 days of anti-PD-1 blockade resulted in a significant increase of IFN-γ levels. This effect was significant ($p=0.0183$) compared to continued anti-PD-1 treatment. In the experiment, anti-TIM-3 antibody TM3B403 and anti-PD-1 antibody PD1B244 were used. **Figure 5** shows the increased IFN-γ levels in the CMV assay, where PBMCs were treated with anti-TIM-3 antibody TM3B105 following 5 days of treatment with anti-PD-1 PD1B244. Values represent average of six biological replicates used for each condition.

Example 19. Epitope mapping of anti-TIM-3 antibodies

Solution hydrogen/deuterium exchange-mass spectrometry (HDX-MS) was performed to identify the binding epitopes of TMB403 and TMB291. For the

experiments, the VH and the VL of TM3B403 and TM3B291 were cloned as IgG1 Fabs with a hexahistidine tag in the C-terminus. The Fabs, were generated from transient transfections of HEK293 Expi cells in suspension shake flasks. TIM-3 IgG1 Fc Chimera, Ser22-Arg200 (Accession # Q8TDQ0), produced in Mouse myeloma cell line (NS0 derived) from R&D Systems (Catalog # 2365-TM) was used.

For H/D exchange, the procedures used to analyze the Fab perturbation were similar to those described previously (Hamuro *et al.,* Biomolecular Techniques 14: 171-182, 2003; Horn *et al.,* Biochemistry 45: 8488-8498, 2006) with some modifications. Briefly, deglycosylated human TIM-3/Fc fusion protein or deglycosylated human TIM-3-Fc plus Fab mixture was incubated with deuterium oxide labeling buffer at 0°C for various times up to 2 hours. Deuterium exchange was quenched by adding guanidine hydrochloride and the quenched sample was subjected to on-column pepsin digestion and LC-MS analysis. The mass spectra were recorded in MS only mode. For the calculation of deuterium incorporation, the mass spectra for a given peptide were combined across the extracted ion chromatogram peak and the weighted average m/z was calculated. The mass increase from the mass of the native peptide (0 min) to the weighted averaged mass corresponds to the level of deuterium incorporation. About 98.4% of the protein could be mapped to specific peptides.

The deuterium levels at the identified peptides were monitored from the mass shift on LC-MS. The selected deuterium buildup curves, which show significant difference in deuterium levels and/or slopes, over exchange time for the peptides were plotted. Deglycosylated human Tim-3/Fc fusion protein showed significant reduction in deuterium uptakes upon binding to TM3B403 at sequences $_{32}$**WGKGACPVFECGNVVL**$_{47}$, (SEQ ID NO: 261) and upon binding to TM3B291 at sequences $_{90}$**RIQIPGIMNDEKF**$_{102}$.(SEQ ID NO: 262). These regions with significant reduction in deuterium uptakes upon binding to Fabs can thus be regarded as main epitopes of the mAbs.

A segment, $_{50}$DERDVNY$_{56}$, (SEQ ID NO: 263) demonstrated modest reduction in deuterium exchange upon binding to TM3B403 or TM3B291. This region may be also considered as a potential epitope for both antibodies.

The major binding epitopes for TM3B403 or TM3B291 are different. However, they may share the similar modest protection region, $_{50}$**DERDVNY**$_{56}$, (SEQ ID NO: 263) based on the HDX mapping results. To help assess if this region contributes to common binding epitope region for both Fab molecules, competition ELISA was performed. Recombinant human Tim-3/Fc protein was directly coated on plates which were then blocked and washed. A mixture of Ruthenium (Ru)-labeled TM3B291 Fab which was

pre-incubated with different concentrations of unlabeled TM3B105 or TM3B291. Plates were incubated, washed and MSD Read Buffer T was dispensed into each well followed by reading with a SECTOR Imager 6000 (Meso Scale Discovery, Gaithersburg, MD).

The competition analysis demonstrated that that TM3B403 competed for binding to TIM-3 with TM3B291. This result could indicate that the modestly protected region, DERDVNY (SEQ ID NO: 263) is part of the epitope for both antibodies or that the antibodies may be sterically blocking each other's binding due to the close proximity of their epitopes.

Example 20. TIM-3 blockade increases TIGIT expression on $CD8^+$ TILs

Effect of anti-TIM-3 antibody treatment on expression of TIGIT in tumors was evaluated in CT26 and MC38 colon carcinoma mouse models. The studies were conducted as described in Example 17 except that 10 mg/ml anti-TIM-3 antibody RMT3-23 (Bioxcell) was used.

TIGIT expression on CD8+ TILs (**Figure 19A, Figure 20A**) and relative frequency of TIGIT+ TILs (**Figure 19B, Figure 20B**) were elevated in both CT26 (Figure 19A, Figure 19B) and MC38 (Figure 20A, Figure 20B) tumor models following TIM-3 blockage.

Example 21. TIM-3 expression is increased after ex vivo PD-1 blockade in melanoma patient PBMC

PBMCs from treatment naïve melanoma patients were stimulated with melanoma antigen peptide pools (NY-ESO, gp100, MART-1) in the presence of anti-PD-1 or anti-TIM-3 function blocking antibodies. Expression of TIM-3 was evaluated on peptide-restimulated cells on day 6. Results showed significant increases in the frequency of TIM-3+ CD8+ T cells in the anti-PD-1 treated samples compared to controls or TIM-3 treated PBMCs (**Figure 21**).

On day 0, frozen PBMCs from treatment naïve melanoma patients were rapidly thawed in a 37 °C water bath. Cells were thawed, washed and counted in complete RPMI media (RPMI + 10 % FBS + 1% sodium pyruvate + 1 % NEAA + 1 % pen/strep). Cells were plates at 200,000 cells per well in a 96 well, U-bottom plate in the presence or absence of anti-PD-1 or anti-TIM-3 function blocking antibodies (PD1B244 and TM3B403, respectively) and 1μg/mL of melanoma antigen peptide pools (NY-ESO, gp100, MART-1) for 6 days at 37C. Cells were restimulated with the peptide pool at day 6

and analyzed by flow cytometry for expression of PD-1 and TIM-3 as well as T cell activation and proliferation markers.

Example 22. Anti-TIM-3 antibodies increase the frequency of activated NK cells in IL-2 stimulated PBMCs

The effects of anti-TIM-3 antibody TM3B403 on the frequency of activated NK cells was determined in assays where human PBMCs were stimulated with IL-2 (20U). Frequency of CD69 and CD25, markers of NK cell activation, were evaluated by flow cytometry 48 hours post-treatment at a range of mAb concentrations. TM3B403 increased the frequency of activated NK cells when the activation was assessed by percentage of CD69 positive cells (**Figure 22A**) or percentage of CD25 positive cells (**Figure 22B**).

WHAT IS CLAIMED

1) An isolated antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising a heavy chain complementarity determining region 1 (HCDR1), a HCDR2 and a HCDR3 of SEQ ID NOs: 82, 83 and 84, respectively and a light chain complementarity determining region 1 (LCDR1), a LCDR2 and a LCDR3 of SEQ ID NOs: 86, 87 and 88, respectively.
2) The antibody or the antigen-binding portion thereof of claim 1, wherein the antibody has one, two, three, four or five of the following properties:
    a) enhances an activation of antigen specific $CD4^+$ or $CD8^+$ T cells in a dose dependent manner, wherein the activation is measured using a cytomegalovirus antigen recall assay (CMV assay) as described in Example 1;
    b) binds human PD-1 with an equilibrium dissociation constant (K_D) of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;
    c) binds human PD-1 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C;
    d) binds cynomolgus PD-1 of SEQ ID NO: 3 with the K_D of less than about 100 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C; or
    e) binds cynomolgus PD-1 of SEQ ID NO: 3 with the K_D of less than about 1 nM, wherein the K_D is measured using ProteOn XPR36 system at +25°C.
3) The antibody or the antigen-binding portion thereof of claim 1 or 2, comprising the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs:
    a) 10, 13, 16, 20, 26 and 31, respectively;
    b) 10, 13, 16, 21, 26 and 32, respectively;
    c) 10, 14, 16, 22, 27 and 33, respectively;
    d) 10, 14, 16, 22, 26 and 34, respectively;
    e) 10, 14, 16, 23, 28 and 35, respectively;
    f) 10, 13, 17, 20, 26 and 31, respectively;
    g) 10, 13, 17, 20, 26 and 36, respectively;
    h) 10, 13, 17, 21, 26 and 32, respectively;
    i) 10, 13, 17, 21, 27 and 37, respectively;
    j) 10, 13, 17, 23, 26 and 32, respectively;
    k) 10, 14, 17, 23, 28 and 35, respectively;

l) 10, 14, 17, 22, 26 and 34, respectively; or

m) 10, 14, 17, 23, 26 and 32, respectively.

4) The antibody or the antigen-binding portion thereof of any of the claims 1-3, comprising

a) a heavy chain variable region (VH) of SEQ ID NOs: 41, 42, 43 or 48;

b) a light chain variable region (VL) of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55 or 56; or

c) the VH of SEQ ID NOs: 41, 42, 43 or 48 and the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55 or 56.

5) The antibody or the antigen-binding portion thereof of any of the claims 1-4, comprising

a) the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 49;

b) the VH of SEQ ID NO: 41 and the VL of SEQ ID NO: 50;

c) the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 51;

d) the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 52;

e) the VH of SEQ ID NO: 42 and the VL of SEQ ID NO: 53;

f) the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 49;

g) the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 54;

h) the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 50;

i) the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 55;

j) the VH of SEQ ID NO: 43 and the VL of SEQ ID NO: 56;

k) the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 53;

l) the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 52; or

m) the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56.

6) The antibody or the antigen-binding portion thereof of any of the claims 1-5, wherein the antibody is human or humanized.

7) The antibody or the antigen-binding portion thereof of any of the claims 1-6, wherein the antibody is

a) an IgG1 isotype, optionally comprising one, two, three, four, five, six, seven, eight, nine or ten substitutions in an Fc region;

b) an IgG2 isotype, optionally comprising one, two, three, four, five, six, seven, eight, nine or ten substitutions in the Fc region;

c) an IgG3 isotype, optionally comprising one, two, three, four, five, six, seven, eight, nine or ten substitutions in the Fc region;

d) an IgG4 isotype, optionally comprising one, two, three, four, five, six, seven, eight, nine or ten substitutions in the Fc region;
e) an IgG1 isotype comprising L234A, L235A, G237A, P238S, H268A, A330S and P331S substitutions;
f) an IgG2 isotype comprising V234A, G237A, P238S, H268A, V309L, A330S and P331S substitutions;
g) an IgG4 isotype comprising F234A, L235A, G237A, P238S and Q268A substitutions;
h) an IgG1 isotype comprising L234A, L235A or L234A and L235A substitutions;
i) an IgG4 isotype comprising F234A, L235A or F234A and L235A substitutions;
j) an IgG2 isotype comprising a V234A substitution;
k) an IgG4 isotype comprising a S228P substitution; or
l) an IgG4 isotype comprising S228P, F234A and L235A substitutions, wherein residue numbering is according to the EU Index.

8) The antibody or the antigen-binding portion thereof of any of the claims 1-7, comprising
a) the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 10, 14, 17, 23, 26 and 32, respectively;
b) the VH of SEQ ID NO: 48 and the VL of SEQ ID NO: 56; and/or
c) a heavy chain (HC) of SEQ ID NO: 72 and a light chain (LC) of SEQ ID NO: 73.

9) The antibody or the antigen-binding portion thereof of any of the claims 1-8, wherein the antibody is a bispecific antibody, optionally binding PD-L1 (SEQ ID NO: 5), PD-L2 (SEQ ID NO: 8), LAG-3 (SEQ ID NO: 293), TIM-3 (SEQ ID NO: 138), CEACAM-1 (SEQ ID NO: 296), CEACAM-5 (SEQ ID NO: 307), OX-40 (SEQ ID NO: 279), GITR (SEQ ID NO: 271), CD27 (SEQ ID NO: 280), VISTA (SEQ ID NO: 286), CD137 (SEQ ID NO: 281), TIGIT (SEQ ID NO: 301) or CTLA-4 (SEQ ID NO: 292).

10) A pharmaceutical composition comprising the antibody or the antigen-binding portion thereof of any of the claims 1-9 and a pharmaceutically accepted carrier.

11) A polynucleotide
a) encoding the VH of SEQ ID NOs: 41, 42, 43 or 48;
b) encoding the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55 or 56;
c) encoding the VH of SEQ ID NOs: 41, 42, 43 or 48 and the VL of SEQ ID NOs: 49, 50, 51, 52, 53, 54, 55 or 56; or
d) comprising the polynucleotide sequence of SEQ ID NOs: 196 or 197.

12) A vector comprising the polynucleotide of claim 11.

13) A host cell comprising the vector of claim 12.

14) A method of producing an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising culturing the host cell of claim 13 in conditions that the antibody of the antigen-binding portion thereof is expressed, and recovering the antibody or the antigen-binding portion thereof produced by the host cell.

15) An isolated antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising

a) the HCDR1, the HCDR2, the HCDR3, the LCDR1, the LCDR2 and the LCDR3 of SEQ ID NOs: 66, 67, 68, 69, 70 and 71, respectively;

b) the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65; and/or

c) the HC of SEQ ID NO: 74 and the LC of SEQ ID NO: 75.

16) A pharmaceutical composition comprising the antibody or the antigen-binding portion thereof of claim 15 and a pharmaceutically accepted carrier.

17) A polynucleotide

a) encoding the VH of SEQ ID NO: 64;

b) encoding the VL of SEQ ID NO: 65;

c) encoding the VH of SEQ ID NO: 64 and the VL of SEQ ID NO: 65; or

d) comprising the polynucleotide sequence of SEQ ID NOs: 198 or 199.

18) A vector comprising the polynucleotide of claim 17.

19) A host cell comprising the vector of claim 18.

20) A method of producing an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof, comprising culturing the host cell of claim 19 in conditions that the antibody or the antigen-binding portion thereof is expressed, and recovering the antibody or the antigen-binding portion thereof produced by the host cell.

21) A method of treating a cancer in a subject, comprising administering a therapeutically effective amount of the isolated antibody or the antigen-binding portion thereof of any of the claims 1-9 or the pharmaceutical composition of claim 10 to the subject in need thereof for a time sufficient to treat the cancer.

22) A method of treating a cancer in a subject, comprising administering a therapeutically effective amount of the isolated antibody or the antigen-binding portion thereof of claim 15 or the pharmaceutical composition of claim 16 to the subject in need thereof for a time sufficient to treat the cancer.

23) The method of claim 21 or 22, wherein the cancer is a solid tumor or a hematological malignancy.

24) The method of any of the claims 21-23, wherein the solid tumor is a melanoma, a lung cancer, a squamous non-small cell lung cancer (NSCLC), a non-squamous NSCLC, a colorectal cancer, a prostate cancer, a castration-resistant prostate cancer, a stomach cancer, an ovarian cancer, a gastric cancer, a liver cancer, a pancreatic cancer, a thyroid cancer, a squamous cell carcinoma of the head and neck, carcinomas of the esophagus or gastrointestinal tract, a breast cancer, a fallopian tube cancer, a brain cancer, an urethral cancer, a genitourinary cancer, an endometriosis, a cervical cancer or a metastatic lesion of the cancer.

25) The method of any of the claims 21-23, wherein the hematological malignancy is a lymphoma, a myeloma or a leukemia.

26) A method of enhancing an immune response in a subject, comprising administering a therapeutically effective amount of the antibody or the antigen-binding portion thereof of any of the claims 1-9 or 15 or a pharmaceutical composition of claim 10 or 16 to the subject in need thereof for a time sufficient to enhance the immune response.

27) The method of claim 26, wherein the subject has a cancer or a viral infection.

28) The method of any of the claims 21-27, wherein the antibody or the antigen-binding portion thereof is administered in combination with a second therapeutic agent.

29) The method of any of the claims 21-28, wherein the second therapeutic agent is

a) a standard of care drug for treatment of the solid tumor or the hematological malignancy;

b) an agonist of a T cell activating molecule;

c) an agonist of CD86 (SEQ ID NO: 264), CD80 (SEQ ID NO: 265), CD28 (SEQ ID NO: 266), ICOS (SEQ ID NO: 267), ICOS ligand (SEQ ID NO: 268), TMIGD2 (SEQ ID NO: 269), CD40 (SEQ ID NO: 270), GITR (SEQ ID NO: 271), 4-1BB ligand (SEQ ID NO: 271), OX40 ligand (SEQ ID NO: 272), CD70 (SEQ ID NO: 274), CD40L (SEQ ID NO: 275), TNFRSF25 (SEQ ID NO: 264), LIGHT (SEQ ID NO: 277), GITR ligand (SEQ ID NO: 278), OX-40 (SEQ ID NO: 279), CD27 (SEQ ID NO: 280), CD137 (SEQ ID NO: 281), NKG2D (SEQ ID NO: 282), CD48 (SEQ ID NO: 283), CD226 (SEQ ID NO: 284), or MICA (SEQ ID NO: 285);

d) an inhibitor of a T cell inhibitory molecule;

e) an inhibitor of PD-1 (SEQ ID NO: 1), PD-L1 (SEQ ID NO: 5), PD-L2 (SEQ ID NO: 8), VISTA (SEQ ID NO: 286), BTNL2 (SEQ ID NO: 287), B7-H3 (SEQ ID

NO: 288), B7-H4 (SEQ ID NO: 289), HVEM (SEQ ID NO: 290), HHLA2 (SEQ ID NO: 291), CTLA-4 (SEQ ID NO: 292), LAG-3 (SEQ ID NO: 293), TIM-3 (SEQ ID NO: 138), BTLA (SEQ ID NO: 294), CD160 (SEQ ID NO: 295), CEACAM-1 (SEQ ID NO: 296), LAIR1 (SEQ ID NO: 297), TGFβ (SEQ ID NO: 298), IL-10 (SEQ ID NO: 299), CD96 (SEQ ID NO: 300), TIGIT (SEQ ID NO: 301), NKG2A (SEQ ID NO: 302), CD112 (SEQ ID NO: 303), CD47 (SEQ ID NO: 304), SIRPA (SEQ ID NO: 305) or CD244 (SEQ ID NO: 306);

f) an antagonistic antibody specifically binding TIM-3;

g) an antagonistic antibody specifically binding TIM-3 comprising the VH and the VL of

i) SEQ ID NOs: 145 and 155, respectively;

ii) SEQ ID NOs: 146 and 156, respectively;

iii) SEQ ID NOs: 148 and 157, respectively;

iv) SEQ ID NOs: 147 and 155, respectively;

v) SEQ ID NOs: 149 and 158, respectively;

vi) SEQ ID NOs: 150 and 159, respectively;

vii) SEQ ID NOs:151 and 160, respectively;

viii) SEQ ID NOs: 152 and 161, respectively;

ix) SEQ ID NOs: 153 and 162, respectively;

x) SEQ ID NOs: 154 and 163, respectively; or

xi) SEQ ID NOs: 172 and 173, respectively;

h) a fibroblast growth factor receptor (FGFR) inhibitor;

i) a vaccine;

j) an agonistic antibody specifically binding GITR;

k) an agonistic antibody specifically binding OX40;

l) an agonistic antibody specifically binding OX40, comprising the VH and the VL of SEQ ID NOs: 309 and 310, respectively;

m) an agonistic antibody specifically binding OX40, comprising the VH and the VL of SEQ ID NOs: 311 and 312, respectively;

n) an agonistic antibody specifically binding CD137;

o) radiation therapy; or

p) surgery.

30) The method of any of the claims 21-29, wherein the antibody or the antigen-binding portion thereof and the second therapeutic agent are administered simultaneously, sequentially or separately.

31) An anti-idiotypic antibody binding to the antibody or the antigen-binding portion thereof of any of the claims 1-9 or 15.

32) A kit comprising the antibody of any of the claims 1-9 or 15.

Figure 1A.

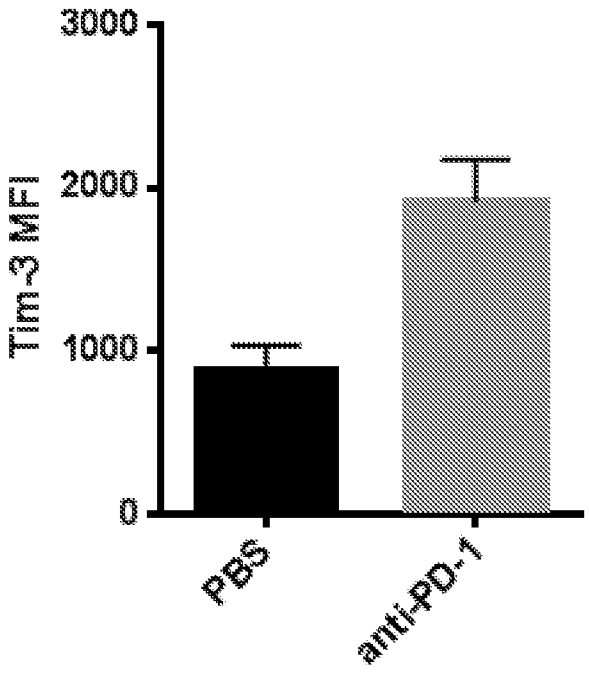


Figure 1B.

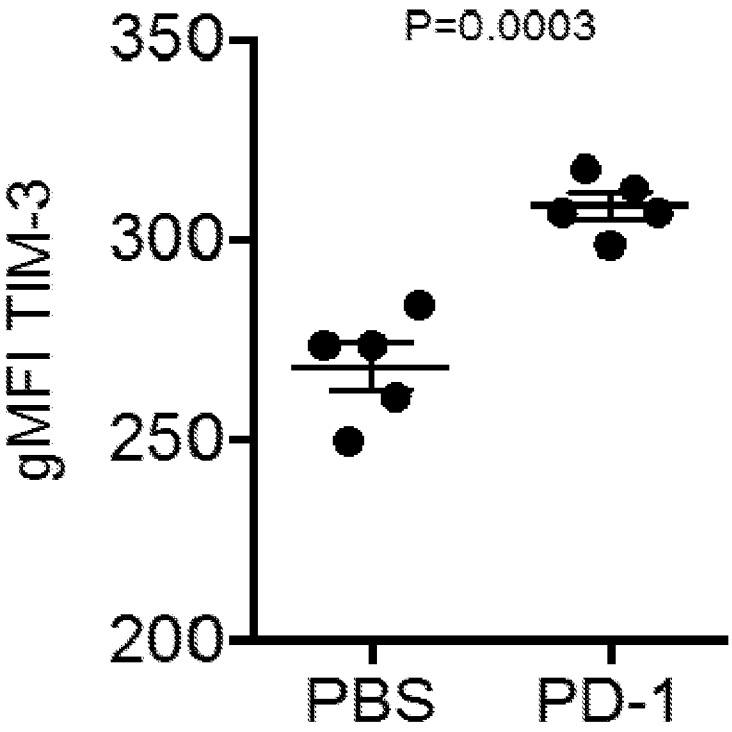


Figure 1C.

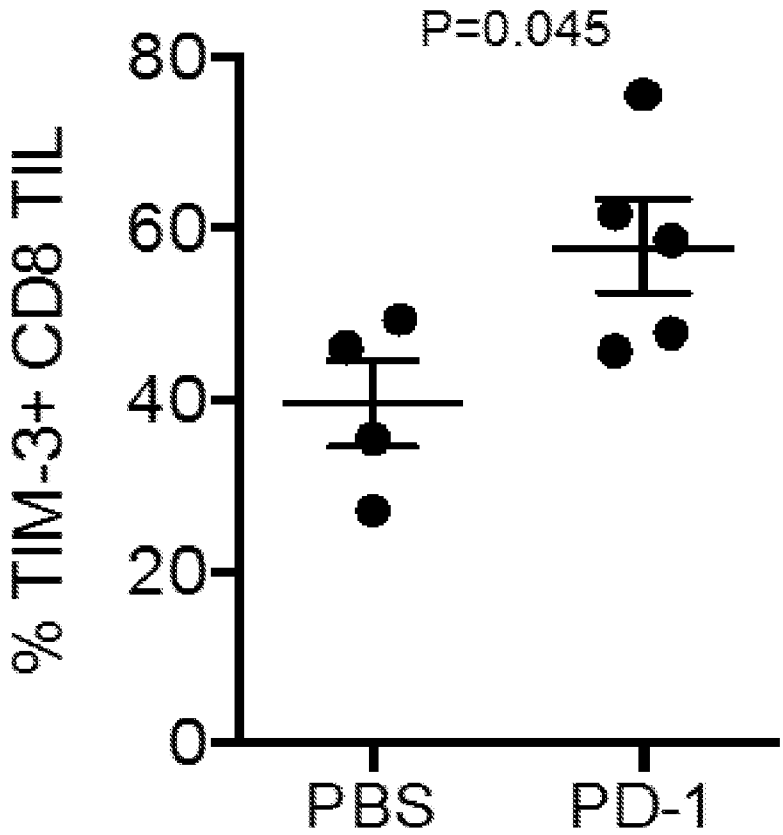


Figure 2A.

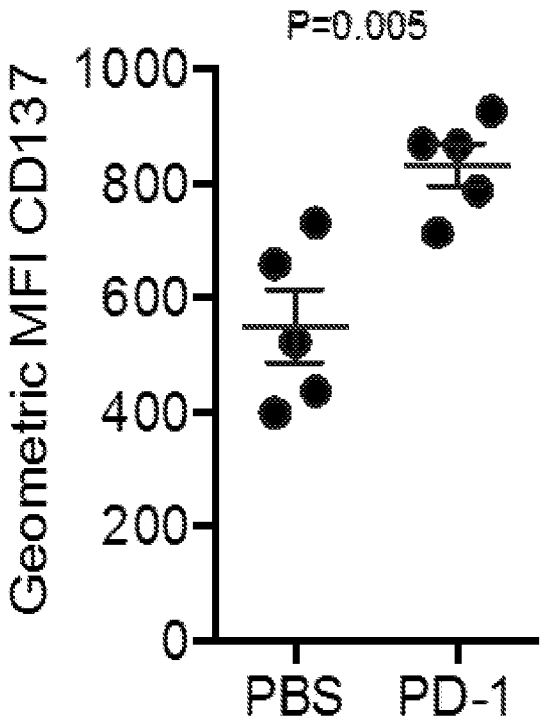


Figure 2B.

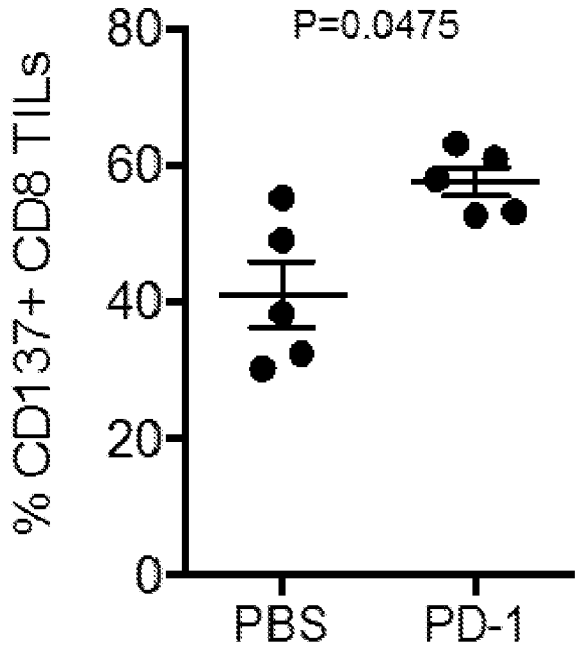


Figure 3A.

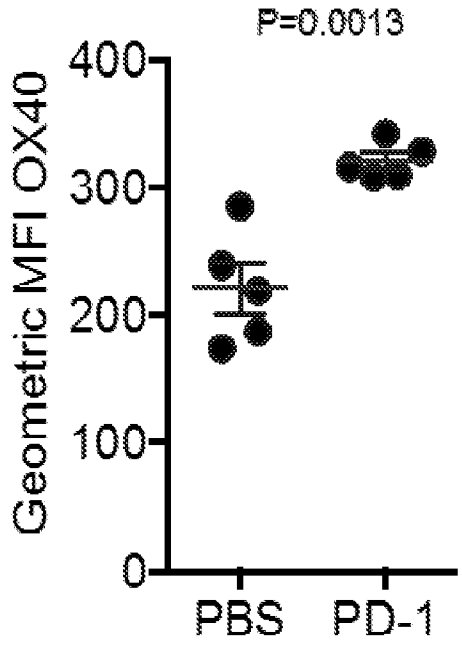


Figure 3B.

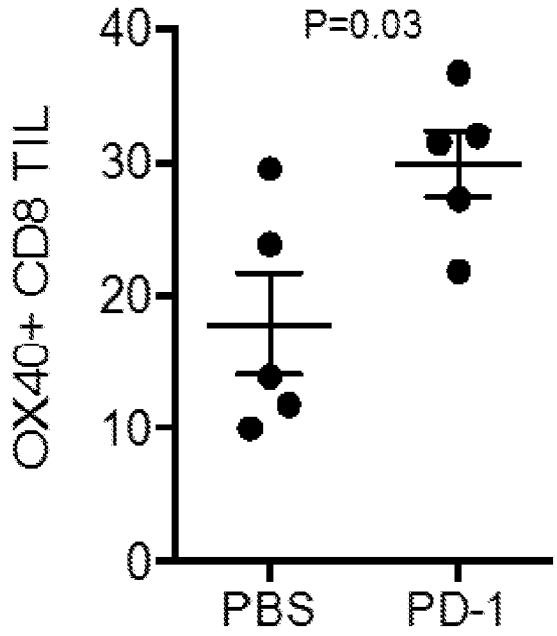


Figure 4A.

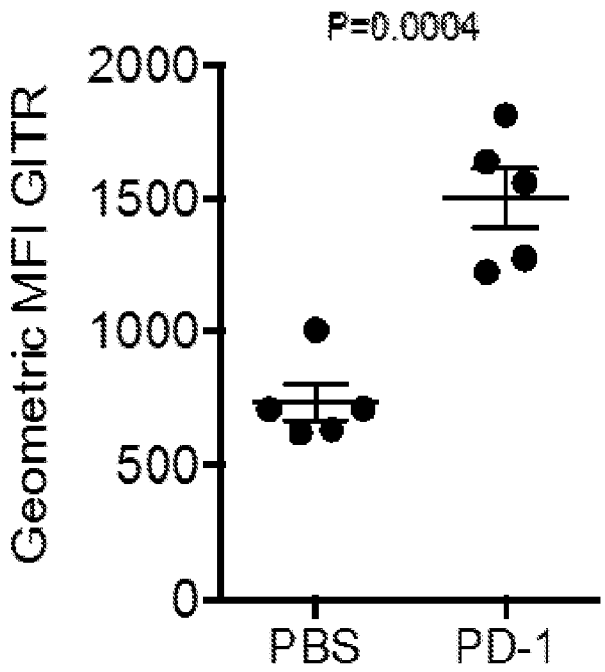


Figure 4B.

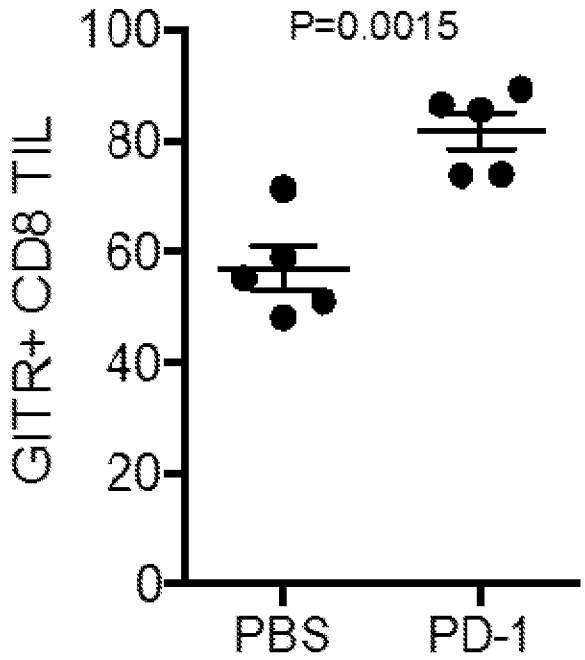


Figure 5.

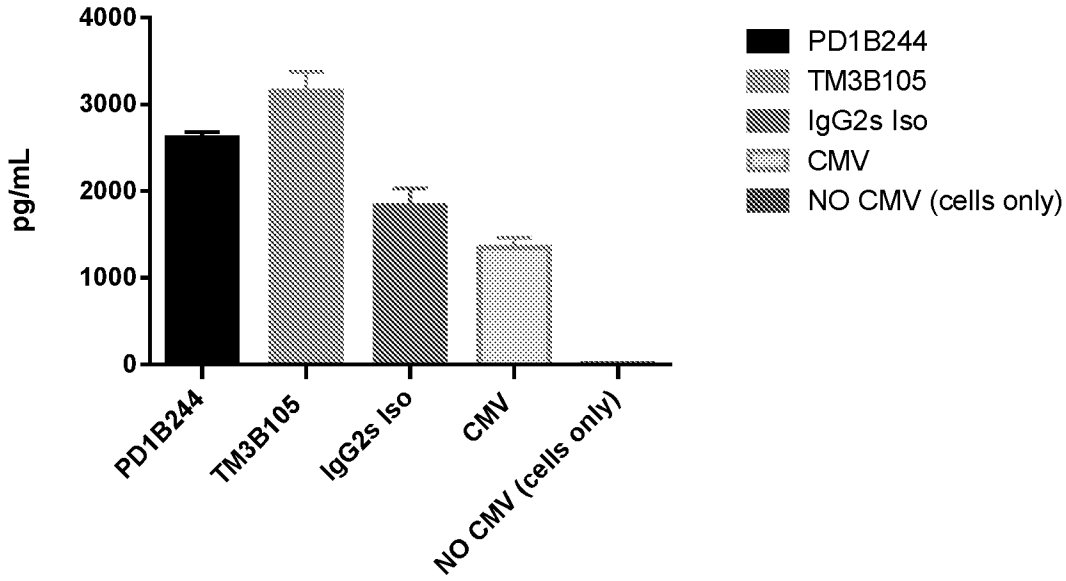

4000
3000
2000
1000
0
pg/mL
PD1B244
TM3B105
IgG2s Iso
CMV
NO CMV (cells only)
PD1B244
TM3B105
IgG2s Iso
CMV
NO CMV (cells only)

Figure 6.

Antibody	HCDR1					
	Sequence					SEQ ID NO:
PD1B114	S	Y	A	I	S	10
PD1B149	S	Y	A	I	S	10
PD1B160	S	Y	A	I	S	10
PD1B162	S	Y	A	I	S	10
PD1B164	S	Y	A	I	S	10
PD1B11	S	Y	A	I	S	10
PD1B183	S	Y	A	I	S	10
PD1B184	S	Y	A	I	S	10
PD1B185	S	Y	A	I	S	10
PD1B187	S	Y	A	I	S	10
PD1B71	S	Y	A	I	S	10
PD1B177	D	Y	V	I	S	11
PD1B70	S	Y	A	I	S	10
PD1B175	S	Y	V	I	H	12
PD1B194	S	Y	A	I	S	10
PD1B195	S	Y	A	I	S	10
PD1B196	S	Y	A	I	S	10
PD1B197	S	Y	V	I	H	12
PD1B198	S	Y	V	I	H	12
PD1B199	D	Y	V	I	S	11
PD1B200	D	Y	V	I	S	11
PD1B201	D	Y	V	I	S	11
HCDR1 genus	X_1	Y	X_2	I	X_3	82

PD-1 mAb HCDR1 genus sequence:

$X_1YX_2IX_3$ (SEQ ID NO: 82),

wherein

X_1 is S or D;

X_2 is V or A; and

X_3 is H or S.

Figure 7.

Antibody	HCDR2																	
	Sequence																	SEQ ID NO:
PD1B114	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B149	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B160	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B162	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B164	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B11	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B183	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B184	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B185	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B187	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B71	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B177	G	I	I	P	I	Y	G	T	A	N	Y	A	Q	K	F	Q	G	15
PD1B70	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B175	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B194	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B195	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B196	G	I	I	P	I	F	D	T	A	N	Y	A	Q	K	F	Q	G	14
PD1B197	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B198	G	I	I	P	I	F	G	T	A	N	Y	A	Q	K	F	Q	G	13
PD1B199	G	I	I	P	I	Y	G	T	A	N	Y	A	Q	K	F	Q	G	15
PD1B200	G	I	I	P	I	Y	G	T	A	N	Y	A	Q	K	F	Q	G	15
PD1B201	G	I	I	P	I	Y	G	T	A	N	Y	A	Q	K	F	Q	G	15
HCDR2 genus	G	I	I	P	I	X_4	X_5	T	A	N	Y	A	Q	K	F	Q	G	83

PD-1 mAb HCDR2 genus sequence

GIIPIX_4X_5TANYAQKFQG (SEQ ID NO: 83),

wherein

X_4 is Y or F; and

X_5 is G or D.

Figure 8.

Antibody	HCDR3														
	Sequence														SEQ ID NO:
PD1B114	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B149	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B160	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B162	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B164	P	G	L	A	A	A	Y	D	T	G	N	L	D	Y	16
PD1B11	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B183	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B184	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B185	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B187	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B194	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B195	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
PD1B196	P	G	L	A	A	A	Y	D	T	G	S	L	D	Y	17
HCDR3 genus 1	P	G	L	A	A	A	Y	D	T	G	X_6	L	D	Y	84

PD-1 mAb HCDR3 genus 1

PGLAAAYDTGX_6LDY (SEQ ID NO: 84),

wherein

X_6 is N or S.

Figure 9.

Antibody	HCDR3											
	Sequence											SEQ ID NO:
PD1B71	G	T	L	D	R	T	G	H	L	D	Y	18
PD1B177	G	T	L	D	R	T	G	H	L	D	Y	18
PD1B70	G	Y	V	R	A	T	G	M	L	D	Y	19
PD1B175	G	Y	V	R	A	T	G	M	L	D	Y	19
PD1B197	G	Y	V	R	A	T	G	M	L	D	Y	19
PD1B198	G	Y	V	R	A	T	G	M	L	D	Y	19
PD1B199	G	T	L	D	R	T	G	H	L	D	Y	18
PD1B200	G	T	L	D	R	T	G	H	L	D	Y	18
PD1B201	G	T	L	D	R	T	G	H	L	D	Y	18
HCDR3 genus 2	G	X_7	X_8	X_9	X_{10}	T	G	X_{11}	L	D	Y	85

PD-1 mAb HCDR3 genus 2

$GX_7X_8X_9X_{10}TGX_{11}LDY$ (SEQ ID NO; 85),

wherein

X_7 is T or Y;

X_8 is L or V;

X_9 is D or R;

X_{10} is R or A; and

X_{11} is H or M.

Figure 10.

Antibody	LCDR1											
	Sequence											SEQ ID NO:
PD1B114	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B149	R	A	S	Q	S	V	R	N	Y	L	A	21
PD1B160	R	A	S	Q	S	V	D	S	Y	L	A	22
PD1B162	R	A	S	Q	S	V	D	S	Y	L	A	22
PD1B164	R	A	S	Q	S	V	R	S	Y	L	A	23
PD1B11	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B183	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B184	R	A	S	Q	S	V	R	N	Y	L	A	21
PD1B185	R	A	S	Q	S	V	R	N	Y	L	A	21
PD1B187	R	A	S	Q	S	V	R	S	Y	L	A	23
PD1B71	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B177	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B70	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B175	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B194	R	A	S	Q	S	V	R	S	Y	L	A	23
PD1B195	R	A	S	Q	S	V	D	S	Y	L	A	22
PD1B196	R	A	S	Q	S	V	R	S	Y	L	A	23
PD1B197	R	A	S	Q	S	V	S	N	Y	L	A	24
PD1B198	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B199	R	A	S	Q	S	V	S	S	Y	L	A	20
PD1B200	R	A	S	Q	S	V	D	N	Y	L	A	25
PD1B201	R	A	S	Q	S	V	S	N	Y	L	A	24
LCDR1 genus	R	A	S	Q	S	V	X_{12}	X_{13}	Y	L	A	86

PD-1 mAb LCDR1 genus

RASQSV$X_{12}X_{13}$YLA (SEQ ID NO: 86),

wherein

X_{12} is S, R or D; and

X_{13} is S or N.

Figure 11.

Antibody	LCDR2							
	Sequence							SEQ ID NO:
PD1B114	D	A	S	N	R	A	T	26
PD1B149	D	A	S	N	R	A	T	26
PD1B160	D	A	S	D	R	A	T	27
PD1B162	D	A	S	N	R	A	T	26
PD1B164	D	A	S	Y	R	A	T	28
PD1B11	D	A	S	N	R	A	T	26
PD1B183	D	A	S	N	R	A	T	26
PD1B184	D	A	S	N	R	A	T	26
PD1B185	D	A	S	D	R	A	T	27
PD1B187	D	A	S	N	R	A	T	26
PD1B71	D	A	S	N	R	A	T	26
PD1B177	D	A	S	N	R	A	T	26
PD1B70	D	A	S	N	R	A	T	26
PD1B175	D	A	S	N	R	A	T	26
PD1B194	D	A	S	Y	R	A	T	28
PD1B195	D	A	S	N	R	A	T	26
PD1B196	D	A	S	N	R	A	T	26
PD1B197	D	A	S	N	R	A	T	26
PD1B198	D	A	S	S	R	A	T	29
PD1B199	D	A	S	T	R	A	T	30
PD1B200	D	A	S	N	R	A	T	26
PD1B201	D	A	S	N	R	A	T	26
LCDR2 genus	D	A	S	X_{14}	R	A	T	87

PD-1 mAb LCDR2 genus

$DASX_{14}RAT$ (SEQ ID NO: 87),

wherein

X_{14} is N, D, Y, S or T.

Figure 12.

Antibody	LCDR3									
	Sequence									SEQ ID NO:
PD1B114	Q	Q	R	S	N	W	P	L	T	31
PD1B149	Q	Q	R	N	Y	W	P	L	T	32
PD1B160	Q	Q	R	G	N	W	P	L	T	33
PD1B162	Q	Q	R	E	Y	W	P	L	T	34
PD1B164	Q	Q	R	D	Y	W	P	L	T	35
PD1B11	Q	Q	R	S	N	W	P	L	T	31
PD1B183	Q	Q	R	G	Y	W	P	L	T	36
PD1B184	Q	Q	R	N	Y	W	P	L	T	32
PD1B185	Q	Q	R	W	N	W	P	L	T	37
PD1B187	Q	Q	R	N	Y	W	P	L	T	32
PD1B71	Q	Q	R	S	N	W	P	L	T	31
PD1B177	Q	Q	R	S	N	W	P	L	T	31
PD1B70	Q	Q	R	S	N	W	P	L	T	31
PD1B175	Q	Q	R	S	N	W	P	L	T	31
PD1B194	Q	Q	R	D	Y	W	P	L	T	35
PD1B195	Q	Q	R	E	Y	W	P	L	T	34
PD1B196	Q	Q	R	N	Y	W	P	L	T	32
PD1B197	Q	Q	R	A	Y	W	P	L	T	38
PD1B198	Q	Q	R	A	E	W	P	L	T	39
PD1B199	Q	Q	R	N	Y	W	P	L	T	32
PD1B200	Q	Q	R	S	A	W	P	L	T	40
PD1B201	Q	Q	R	N	Y	W	P	L	T	32
LCDR3 genus	Q	Q	R	X_{15}	X_{16}	W	P	L	T	88

PD-1 mAb LCDR3 genus:

$QQRX_{15}X_{16}WPLT$ (SEQ ID NO: 88),

wherein

X_{15} is S, N, G, E, D, W, E or A; and

X_{16} is N, Y, E or A.

Figure 13.

mAb name	HCDR1						
	Sequence						SEQ ID NO:
TM3B103	N	Y	W	M	S		90
TM3B105	S	Y	A	M	S		91
TM3B109	S	Y	A	M	S		91
TM3B108	G	Y	W	M	H		92
TM3B113	D	Y	W	M	S		93
HCDR1 genus	X_{17}	Y	X_{18}	M	X_{19}		164

TIM3 mAb HCDR1 genus:

$X_{17}YX_{18}MX_{19}$ (SEQ ID NO: 164),

Wherein

X_{17} is N, S, G or D;

X_{18} is W or A; and

X_{19} is S or H.

Figure 14.

mAb	HCDR2																	
	Sequence																	SEQ ID NO:
TM3B103	A	I	S	G	S	G	G	S	T	Y	Y	A	D	S	V	K	G	99
TM3B105	A	I	S	G	S	G	G	S	T	Y	Y	A	D	S	V	K	G	99
TM3B109	A	I	S	G	S	G	G	S	T	Y	Y	A	D	S	V	K	G	99
TM3B108	A	I	S	Y	S	G	S	S	T	Y	Y	A	D	S	V	K	G	100
TM3B113	V	I	K	Y	S	G	G	S	K	Y	Y	A	D	S	V	K	G	101
HCDR2 genus	X_{20}	I	X_{21}	X_{22}	S	G	G	S	X_{23}	Y	Y	A	D	S	V	K	G	165

TIM-3 mAb HCDR2 genus

$X_{20}IX_{21}X_{22}SGGSX_{23}YYADSVKG$ (SEQ ID NO: 165),

wherein

X_{20} is A or V;

X_{21} is S or K;

X_{22} is G or Y; and

X_{23} is T or K.

Figure 15.

mAb	HCDR3										
	Sequence										SEQ ID NO:
TM3B103	D	H	W	D	P	N	F	L	D	Y	107
TM3B105	S	P	-	-	Y	A	P	L	D	Y	108
TM3B109	N	E	E	P	D	D	R	L	D	Y	109
TM3B108	G	T	N				W	L	D	Y	110
TM3B113	E	L	E			G	V	F	D	Y	111
HCDR3 genus	X_{24}	X_{25}	X_{26}	X_{27}	X_{28}	X_{29}	X_{30}	X_{31}	D	Y	166

$X_{24}X_{25}X_{26}X_{27}X_{28}X_{29}X_{30}X_{31}$DY (SEQ ID NO: 166),

wherein

X_{24} is D, S, N, G or E;

X_{25} is H, P, E, T or L;

X_{26} is W, E, N or deleted;

X_{27} is D, P or deleted;

X_{28} is P, Y, D or deleted;

X_{29} is N, A, D, G or deleted;

X_{30} is F, P, R, W or V; and

X_{31} is L or F.

Figue 16.

mAb	LCDR1																
	Sequence																SEQ ID NO:
TM3B103	R	A	S	Q	S	V	S	S	-					S	Y	L	A	117
TM3B105	R	A	S	Q	S	V	N	-						D	Y	L	A	118
TM3B109	K	S	S	Q	S	V	L	A	S	S	N	N	K	N	Y	L	A	119
TM3B108	R	A	S	Q	S	V	S	S						S	Y	L	A	117
TM3B113	R	A	S	Q	S	V	S	N						S	T	L	A	120
LCDR1 genus	X_{32}	X_{33}	S	Q	S	V	X_{34}	X_{35}	X_{36}	X_{37}	X_{38}	X_{39}	X_{40}	X_{41}	X_{42}	L	A	167

$X_{32}X_{33}SQSVX_{34}X_{35}X_{36}X_{37}X_{38}X_{39}X_{40}X_{41}X_{42}LA$ (SEQ ID NO: 167),

wherein

X_{32} is R or K;

X_{33} is A or S;

X_{34} is S, N or L;

X_{35} s S, A, N or deleted;

X_{36} is S or deleted;

X_{37} is S or deleted;

X_{38} is N or deleted;

X_{39} is N or deleted;

X_{40} is K or deleted;

X_{41} is S, D or N; and

X_{42} is Y or T.

Figure 17.

mAb	LCDR2							
	Sequence							SEQ ID NO:
TM3B103	G	A	S	S	R	A	T	126
TM3B105	D	A	S	N	R	A	T	127
TM3B109	W	A	S	T	R	E	S	128
TM3B108	G	A	S	S	R	A	T	126
TM3B113	T	A	S	S	R	A	T	129
LCDR2 genus	X_{43}	A	S	X_{44}	R	X_{45}	X_{46}	168

TIM-3 mAb LCDR2 genus

$X_{43}ASX_{44}RX_{45}X_{46}$ (SEQ ID NO: 168),

wherein

X_{43} is G, D, W or T;

X_{44} is S, N or T;

X_{45} is A or E; and

X_{46} is T or S.

Figure 18.

mAb	LCDR3									
	Sequence									SEQ ID NO:
TM3B103	Q	Q	Y	G	S	S	P	L	T	135
TM3B105	Q	Q	G	G	H	A	P	I	T	136
TM3B109	Q	Q	Y	Y	S	T	P	L	T	137
TM3B108	Q	Q	Y	G	S	S	P	L	T	138
TM3B113	Q	Q	S	Y	T	S	P	W	T	139
LCDR3 genus	Q	Q	X_{47}	X_{48}	X_{49}	X_{50}	P	X_{51}	T	169

$QQX_{47}X_{48}X_{49}X_{50}PX_{51}T$ (SEQ ID NO: 169),

wherein

X_{47} is Y, G or S;

X_{48} is G or Y;

X_{49} is S, H or T;

X_{50} is S, A or T; and

X_{51} is L, I or W.

Figure 19A.

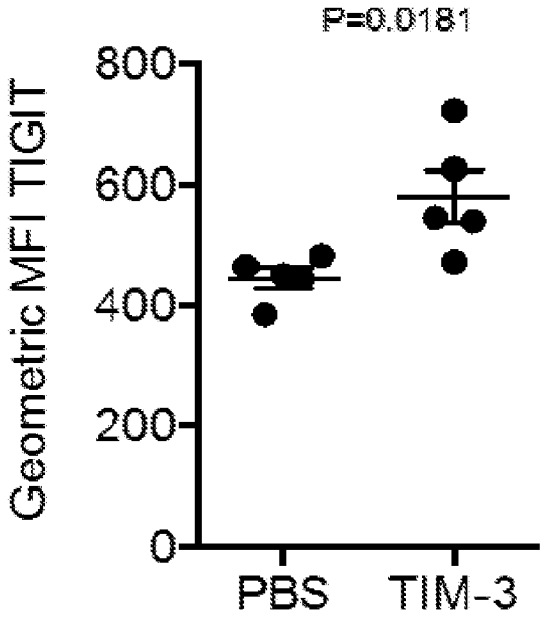


Figure 19B.

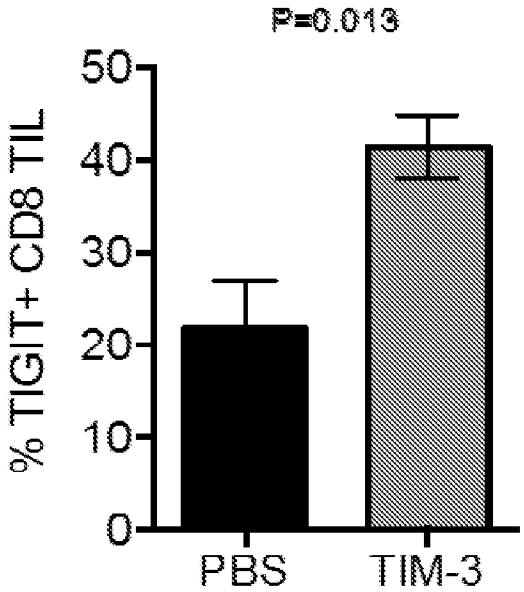


Figure 20A.

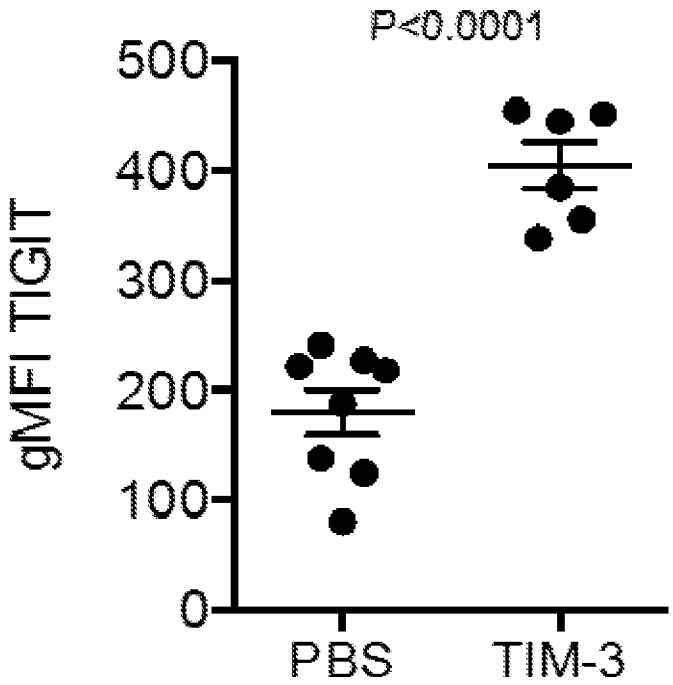


Figure 20B.

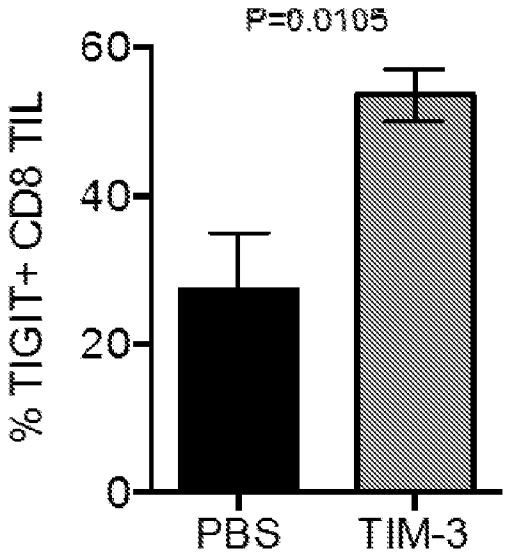


Figure 21.

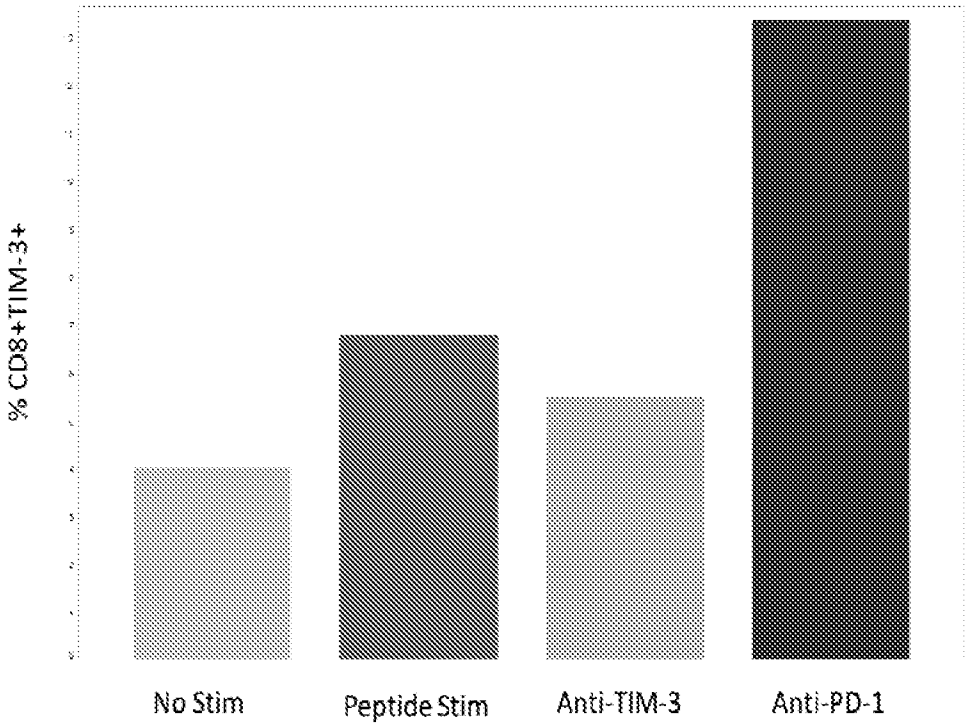


Figure 22A.

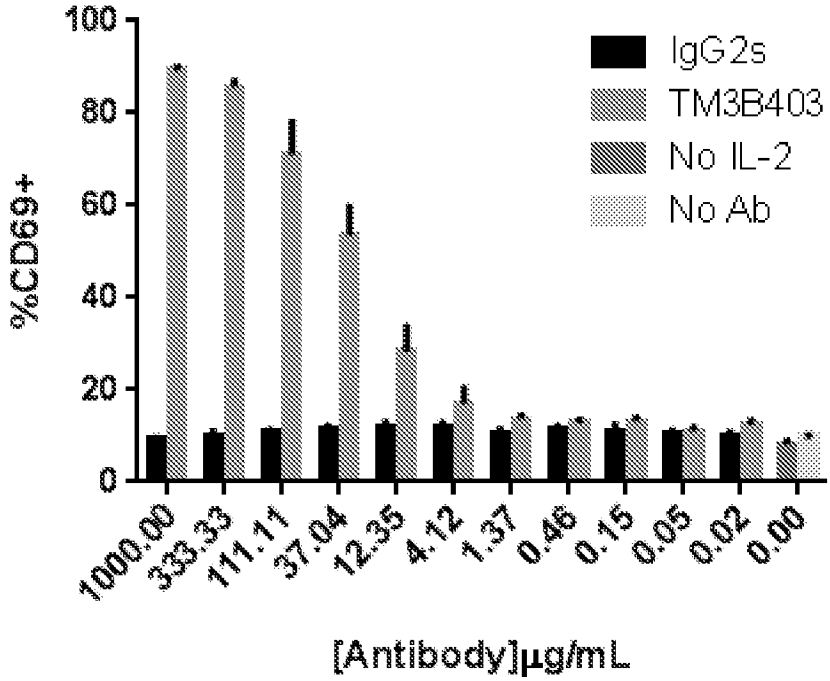
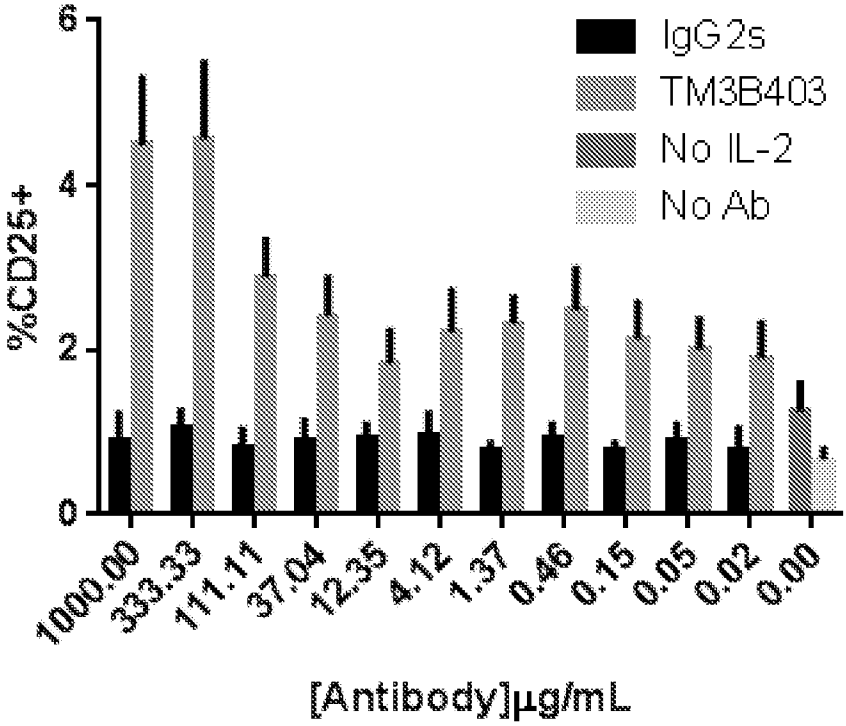

100
80
60
40
20
0
%CD69+
IgG2s
TM3B403
No IL-2
No Ab
1000.00
333.33
111.11
37.04
12.35
4.12
1.37
0.46
0.15
0.05
0.02
0.00
[Antibody]µg/mL

Figure 22B.


6
4
2
0
%CD25+
IgG2s
TM3B403
No IL-2
No Ab
1000.00
333.33
111.11
37.04
12.35
4.12
1.37
0.46
0.15
0.05
0.02
0.00
[Antibody]µg/mL

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 16/59833

A. CLASSIFICATION OF SUBJECT MATTER
IPC(8) - A61K 39/395, A61K 39/00 (2017.01)
CPC - C07K 2317/70, C07K 2317/75, C07K 16/28, C07K 2317/56

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
See Search History Document

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
See Search History Document

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
See Search History Document

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2015/0165025 A1 (MEDAREX L.L.C.) 18 June 2015 (18.06.2015) para [0006], [0007], [0132], [0133]	1-3, 11-20, 22
A	US 2003/0096977 A1 (KOIKE et al.) 22 May 2003 (22.05.2003) para [0351], SEQ ID NO: 40	1-3, 11-20, 22
A	US 2014/0112915 A1 (BARDROFF et al.) 24 April 2014 (24.04.2014) para [0022], SEQ ID NO: 4	1-3, 11-20, 22
A	US 2012/0017292 A1 (KOVALIC et al.) 19 January 2012 (19.01.2012) para [0004], SEQ ID NO: 47420	1-3, 11-20, 22
A	US 2003/0040044 A1 (HEAVNER et al.) 27 February 2003 (27.02.2003) claim 31, SEQ ID NO: 4	1-3, 11-20, 22
A	US 2003/0226155 A1 (Sadeghi et al) 04 December 2003 (04.12.2003) Table 3, SEQ ID NO: 74	1-3, 11-20, 22
A	US 2012/0108795 A1 (KEHOE et al.) 03 May 2012 (03.05.2012) para [0041], [0082], SEQ ID NOs: 5, 59	1-3, 11-20, 22

☒ Further documents are listed in the continuation of Box C. ☐

* Special categories of cited documents:
"A" document defining the general state of the art which is not considered to be of particular relevance
"E" earlier application or patent but published on or after the international filing date
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
"O" document referring to an oral disclosure, use, exhibition or other means
"P" document published prior to the international filing date but later than the priority date claimed
"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
28 March 2017	13 APR 2017

Name and mailing address of the ISA/US	Authorized officer:
Mail Stop PCT, Attn: ISA/US, Commissioner for Patents P.O. Box 1450, Alexandria, Virginia 22313-1450 Facsimile No. 571-273-8300	Lee W. Young PCT Helpdesk: 571-272-4300 PCT OSP: 571-272-7774

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 16/59833

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2012/0114652 A1 (ELVIN et al.) 10 May 2012 (10.05.2012) para [0077], SEQ ID NO: 114	1-3, 11-20, 22
A	US 2005/0009136 A1 (NIXON et al.) 13 January 2005 (13.01.2005) para [0303], SEQ ID NO:328	1-3, 11-20, 22
A	US 2009/0220485 A1 (TANHA) 03 September 2009 (03.09.2009) para [0022], SEQ ID NO: 35	1-3, 11-20, 22
A	US 2006/0222645 A1 (LEE et al.) 05 October 2006 (05.10.2006) para [0010], SEQ ID NO: 21	1-3, 11-20, 22
A	US 2014/0322218 A1 (XIAO et al.) 30 October 2014 (30.10.2014) para [0013], SEQ ID NO: 119	1-3, 11-20, 22
A	US 2005/0215770 A1 (Bell et al.) 29 September 2005 (29.09.2005) para [0044], SEQ ID NO: 133	1-3, 11-20, 22
A	US 2007/0048315 A1 (PRESTA) 01 March 2007 (01.03.2007) claim 7, SEQ ID NO: 69	1-3, 11-20, 22
A	US 2004/0133357 A1 (ZHONG et al.) 08 July 2004 (08.07.2004) para [0244], SEQ ID NO:167	1-3, 11-20, 22
A	WO 2012/061448 A1 (BOEHRINGER INGELHEIM INTERNATIONAL GMBH) 10 May 2012(10.05.2012) pg 3, ln 25, SEQ ID NO: 14	1-3, 11-20, 22
A	US 2010/0260754 A1 (CHEDID et al.) 14 October 2010 (14.10.2010) para [0023], SEQ ID NO: 11	1-3, 11-20, 22
A	US 2015/0183874 A1 (THE BOARD OF TRUSTEES OF THE LELAND STANFORD JUNIOR UNIVERSITY) 02 July 2015 (02.07.2015) para [0037]	1-3, 11-20, 22

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 16/59833

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☒ Claims Nos.: 4-10, 21, 23-32
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

---------- see extra sheet ----------

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.

3. ☒ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
1-3, 11-20 and 22 limited to SEQ ID NOs: 12, 15, 16, 20, 26, 31, 49, 10, 14, 17, 23, 26, 32, 66-71

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.

☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.

☒ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 16/59833

Continuation of Box No. III, Observations where unity of invention is lacking:

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Group I+: Claims 1-3, 11-20 and 22, directed to an isolated antagonistic antibody specifically binding PD-1, and a method of use thereof to treat cancer. The antibody will be searched to the extent that the HCDR1-3 and LCDR1-3 of SEQ ID NOs: 82-84 and 86-88, respectively, encompass sequences comprising the first species of each variable, as listed below:
SEQ ID NO: 82 (SYVIH) (X1=S, X2=V, X3=H) [corresponding to SEQ ID NO: 12];
SEQ ID NO: 83 (GIIPIYGTANYAQKFQG) (X4=Y, X5=G) [corresponding to SEQ ID NO: 15];
SEQ ID NO: 84 (PGLAAAYDTGNLDY) (X6=N) [corresponding to SEQ ID NO: 16];
SEQ ID NO: 86 (RASQSVSSYLA) (X12=S, X13=S) [corresponding to SEQ ID NO: 20];
SEQ ID NO: 87 (DASNRAT) (X14=N) [corresponding to SEQ ID NO: 26];
SEQ ID NO: 88 (QQRSNWPLT) (X15=S, X16=N) [corresponding to SEQ ID NO: 31];
VL SEQ ID NO: 49 comprises LCDR1-3 of SEQ ID NO: 20, 26, 31, respectively.
[NOTE, None of the listed VH in the Specification comprises SEQ ID NOs: 12, 15 and 16].

It is believed that claims 1, 2, 11-14, 22 encompass this first named invention, and thus these claims will be searched without fee to the extent that they encompass SEQ ID NOs: 82, 83, 84, 86, 87, 88, 12, 15, 16, 20, 26, 31, and 49. Additional antibodies will be searched upon the payment of additional fees. Applicants must specify the claims that encompass any additionally elected anti-PD-1 antibodies. Applicants must further indicate, if applicable, the claims which encompass the first named invention, if different than what was indicated above for this group. Failure to clearly identify how any paid additional invention fees are to be applied to the "+" group(s) will result in only the first claimed invention to be searched. An exemplary election would be an isolated antagonistic antibody specifically anti-PD-1 antibody comprising HCDR1-3 and LCDR1-3 of SEQ ID NOs: 66-68 and 69-71, respectively; VH SEQ ID NO: 64; VL SEQ ID NO: 65 (claims 1, 2, 15-19 and 22).

The inventions listed as Groups I+ do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

Special Technical Features

The technical feature of each of the inventions listed as Group I+ is the isolated antagonistic antibody specifically binding PD-1, recited therein. Each invention of Group I+ requires a specific antibody comprising CDRs of unique amino acid sequences, not required by any of the other inventions.

Common Technical Features

The inventions of Group I+ share the technical feature of an antagonistic antibody specifically binding PD-1 or an antigen-binding portion thereof; a polynucleotide encoding an antibody specifically binding PD-1; and use thereof to treat cancer. However, this shared technical feature does not represent a contribution over prior art, because the shared technical feature is anticipated by US 2015/0165025 A1 to Medarex, L.L.C. et al. (hereinafter 'Medarex') (para [0006] "isolated monoclonal antibodies, i[n] particular human monoclonal antibodies, that bind to PD-1 and that exhibit numerous desirable properties. These properties include, for example, high affinity binding to human PD-1 ... the invention provides a method of inhibiting growth of tumor cells in vivo using anti-PD-1 antibodies.", para [0007] "the invention pertains to an isolated monoclonal antibody, or as antigen-binding portion thereof"; para [0132] "Nucleic acid molecules encoding the antibodies, or antigen-binding portions thereof, of the invention are also encompassed by the invention, as well as expression vectors comprising such nucleic acids and host cells comprising such expression vectors."; para [0133], " the antibody of the invention enhances, stimulates or increases the immune response in the subject.").

As the technical features were known in the art at the time of the invention, this cannot be considered a special technical feature that would otherwise unify the inventions.

Group I+ therefore lack unity under PCT Rule 13 because they do not share the same or corresponding special technical feature.