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(71) Applicant: BRISTOL-MYERS SQUIBB COMPANY
[US/US]; Route 206 & Province Line Road, Princeton, NJ
08543 (US).

(72) Inventors: DOYLE, Sean; 9657 26th Ave. NW, Seattle,
WA 98117 (US). SELBY, Mark, J.; 700 Bay Road, Red-
wood City, CA 94063 (US). CHADWICK, Eric; 2503
153rd Ave. Se, Bellevue, WA 98007 (US).

(74) Agent: REMILLARD, Jane, E. et al.; Nelson Mullins Ri-
ley & Scarborough LLP, One Post Office Square, Boston,
MA 02109-2127 (US).

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(54) Title: COMBINATION OF TIM-4 ANTAGONIST AND METHODS OF USE

(57) Abstract: Provided are methods and compositions for treating cancer using an effective amount of a PD-1 antagonist (e.g., an antibody) in combination with a TIM-4 antagonist (e.g., an antibody).

COMBINATION OF TIM-4 ANTAGONIST AND PD-1 ANTAGONIST AND METHODS OF USE

CROSS-REFERENCE TO RELATED APPLICATIONS

5 This application claims priority to U.S. Provisional Application No. 62/359073, filed July 6, 2016. The content of the aforementioned application is hereby incorporated by reference.

BACKGROUND

10 The National Cancer Institute has estimated that in the United States alone, 1 in 3 people will be struck with cancer during their lifetime. Moreover, approximately 50% to 60% of people contracting cancer will eventually succumb to the disease. Once tumor cells escape from the primary site they passage through the lymphatic and/or circulatory system and ultimately a few establish at distant sites to give metastases, and 95% of deaths from 15 solid tumors in the developed world are due to metastasis.

The widespread occurrence of cancer tumors underscores the need for improved anticancer regimens. However, despite advances in multimodal therapy, increases in overall survival in cancer patients have been limited. Accordingly, it is an object of the present invention to provide improved methods for treating subjects with such tumors.

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SUMMARY OF THE INVENTION

The present inventors have discovered for the first time that co-administration of a TIM-4 antagonist (*e.g.*, an antibody) and an anti-PD-1 antagonist (*e.g.*, an antibody) effectively inhibits tumor growth *in vivo*, even synergistically. Accordingly, it is an object of 25 the present invention to provide improved methods for treating subjects with cancer. Specifically, it is an object of the invention to provide efficacious combination treatment regimens wherein a TIM-4 antagonist is combined with an anti-PD-1 antagonist for the treatment of cancer.

In one aspect, the present invention provides a method for the treatment of cancer in a 30 subject by co-administering an effective amount of a PD-1 antagonist and a TIM-4 antagonist.

Suitable TIM-4 antagonists for use in the methods of the invention, include, without limitation, ligands, antibodies (e.g., monoclonal antibodies and bispecific antibodies) and multivalent agents. The TIM-4 antagonist can be a non-activating ligand or ligand binding partner (e.g., a small molecule, engineered PS emulator or soluble TIM-1, or other binding protein). In one embodiment, the TIM-4 antagonist is an anti-TIM4 antibody. In one embodiment, the TIM-4 antagonist is an antibody which binds to an epitope within the IgV domain of TIM-4. In another embodiment, the TIM-4 antagonist is an antibody such as 9F4, RMT4-53, RMT4-54, F31-563 or 21T112.

An exemplary anti-TIM-4 antibody contains heavy and light chains comprising the heavy and light chain CDRs of 9F4, RMT4-53, RMT4-54, F31-563 or 21T112, and optionally comprises a framework region with at least about 90% amino acid sequence identity with the framework region of the corresponding antibody. Anti-TIM-4 antibodies may also comprise heavy and light chain variable domains that are at least about 90%, 95% or 99% identical with that of antibody 9F4, RMT4-53, RMT4-54, F31-563 or 21T112. In certain embodiments, the anti-TIM-4 antagonist antibody competes for binding with, and/or binds to the same epitope on TIM-4 as, 9F4, RMT4-53, RMT4-54, F31-563 or 21T112.

Suitable PD-1 antagonists for use in the methods of the invention, include, without limitation, ligands, antibodies (e.g., monoclonal antibodies and bispecific antibodies), and multivalent agents. In one embodiment, the PD-1 antagonist is an anti-PD-1 or anti-PD-L1 antibody. In another embodiment, the PD-1 antagonist is an antibody, such as MK-3475 or CT-011. In one embodiment, the PD-1 antagonist is a fusion protein, e.g., an Fc fusion protein, such as AMP-244.

An exemplary anti-PD-1 antibody is 5C4 (referred to as 5C4 in WO 2006/121168; also known as MDX-1106, ONO-4538, and nivolumab) comprising heavy and light chains having the sequences shown in SEQ ID NOs: 11 and 12, respectively, or antigen binding fragments and variants thereof. In other embodiments, the antibody comprises the heavy and light chain CDRs or variable regions (VRs) of 5C4. Accordingly, in one embodiment, the antibody comprises the CDR1, CDR2, and CDR3 domains of the VH region of 5C4 having the sequence shown in SEQ ID NO: 13, and the CDR1, CDR2 and CDR3 domains of the VL region of 5C4 having the sequence shown in SEQ ID NO:15. In another embodiment, the antibody comprises the heavy chain CDR1, CDR2 and CDR3 domains having the sequences set forth in SEQ ID NOs: 17, 18, and 19, respectively, and the light chain CDR1, CDR2 and CDR3 domains having the sequences set forth in SEQ ID NOs: 20, 21, and 22, respectively.

In another embodiment, the antibody comprises VH and/or VL regions having the amino acid sequences set forth in SEQ ID NO: 13 and/or SEQ ID NO: 15, respectively. In another embodiment, the antibody comprises the heavy chain variable (VH) and/or light chain variable (VL) regions encoded by the nucleic acid sequences set forth in SEQ ID NO: 14 and/or SEQ ID NO: 16, respectively. In another embodiment, the antibody competes for binding with, and/or binds to the same epitope on PD-1 as, the above-mentioned antibodies. In another embodiment, the antibody has at least about 90% variable region amino acid sequence identity with the above-mentioned antibodies (e.g., at least about 90% , 95% or 99% variable region identity with SEQ ID NO: 13 or SEQ ID NO: 15).

10 In one embodiment, the PD-1 antagonist is an anti-PD-L1 antibody, such as MEDI4736 (also known as Anti-B7-H1) or MPDL3280A (also known as RG7446). An exemplary anti-PD-L1 antibody is 12A4 (referred to as 12A4 in WO 2007/005874 and US Patent No. 7,943,743). In one embodiment, the antibody comprises the heavy and light chain CDRs or VRs of 12A4. Accordingly, in one embodiment, the antibody comprises the CDR1, 15 CDR2, and CDR3 domains of the VH region of 12A4 having the sequence shown in SEQ ID NO: 1, and the CDR1, CDR2 and CDR3 domains of the VL region of 12A4 having the sequence shown in SEQ ID NO: 3. In another embodiment, the antibody comprises the heavy chain CDR1, CDR2 and CDR3 domains having the sequences set forth in SEQ ID NOs: 5, 6, and 7, respectively, and the light chain CDR1, CDR2 and CDR3 domains having the sequences set forth in SEQ ID NOs: 8, 9, and 10, respectively. In another embodiment, 20 the antibody comprises VH and/or VL regions having the amino acid sequences set forth in SEQ ID NO: 1 and/or SEQ ID NO: 3, respectively. In another embodiment, the antibody comprises the heavy chain variable (VH) and/or light chain variable (VL) regions encoded by the nucleic acid sequences set forth in SEQ ID NO: 2 and/or SEQ ID NO: 4, respectively. In another embodiment, the antibody competes for binding with, and/or binds to the same epitope on PD-L1 as, the above-mentioned antibodies. In another embodiment, the antibody has at least about 90% variable region amino acid sequence identity with the above-mentioned antibodies (e.g., at least about 90% , 95% or 99% variable region identity with SEQ ID NO: 1 or SEQ ID NO: 3).

30 In one embodiment, the invention provides a method of treating cancer in a subject, the method comprising administering to the subject an effective amount of a PD-1 antagonist and a TIM-4 antagonist, wherein

(a) the PD-1 antagonist is an anti-PD-1 antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 13, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 15; and

5 (b) the TIM-4 antagonist is an antibody.

In another embodiment, the invention provides a method of treating cancer in a subject, the method comprising administering to the subject an effective amount of a PD-1 antagonist and a TIM-4 antagonist, wherein

10 (a) the PD-1 antagonist is an anti-PD-L1 antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 1, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 3; and

15 (b) the TIM-4 antagonist is an antibody.

The efficacy of the treatment methods provided herein can be assessed using any suitable means. In one embodiment, the treatment produces at least one therapeutic effect selected from the group consisting of reduction in size of a tumor, reduction in number of metastatic lesions over time, complete response, partial response, and stable disease. In another embodiment, administration of a PD-1 antagonist and a TIM-4 antagonist results in at least a 1, 1.25, 1.50, 1.75, 2, 2.25, 2.50, 2.75, 3, 3.25, 3.5, 3.75, or 4-fold reduction in tumor volume, e.g., relative to 20 treatment with the PD-1 antagonist or TIM-4 antagonist alone, or relative to tumor volume before initiation of treatment. In another embodiment, administration of the PD-1 antagonist and TIM-4 antagonist results in at least a 1-fold, 2-fold, or more preferably a 3-fold reduction in tumor volume, e.g., relative to treatment with the PD-1 antagonist or TIM-4 antagonist alone, or relative to tumor volume before initiation of treatment. In a further embodiment, administration 25 of a PD-1 antagonist and a TIM-4 antagonist results in tumor growth inhibition of at least 50%, 60%, 70% or 80%, e.g., relative to treatment with the PD-1 antagonist or TIM-4 antagonist alone, or relative to tumor volume before initiation of treatment. In certain embodiments, tumor volume is reduced by 50%, 60%, 70%, 80%, 90% or more, e.g., relative to tumor size before initiation of the treatment.

30 The PD-1 antagonist and TIM-4 antagonist can be administered accordingly to a suitable dosage, route (e.g., intravenous, intraperitoneal, intramuscular, intrathecal or subcutaneous). The antagonist and agonist can also be administered according to any suitable schedule. For example, the antagonist and agonist can be simultaneously administered in a

single formulation. Alternatively, the antagonist and agonist can be formulated for separate administration, wherein they are administered concurrently or sequentially. In one embodiment, the PD-1 antagonist is administered prior to administration of the TIM-4 antagonist. In another embodiment, the TIM-4 antagonist is administered prior to 5 administration of the PD-1 antagonist. In a further embodiment, the TIM-4 antagonist and the PD-1 antagonist are administered simultaneously.

In one embodiment, the cancer is a cancer selected from the group consisting of carcinoma, sarcoma, blastoma, lymphoma and leukemia. In one embodiment, the cancer is selected from the group consisting of small-cell lung cancer, non-small cell lung cancer, 10 gastrointestinal cancer, colorectal cancer, stomach cancer, colon carcinoma, and glioblastoma. In another embodiment, the cancer is selected from the group consisting of chronic myeloid leukemia, acute lymphoblastic leukemia, Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), squamous cell carcinoma, small-cell lung cancer, non-small cell lung cancer, glioma, gastrointestinal cancer, renal cancer, ovarian cancer, liver 15 cancer, colorectal cancer, endometrial cancer, kidney cancer, prostate cancer, thyroid cancer, neuroblastoma, pancreatic cancer, glioblastoma multiforme, cervical cancer, stomach cancer, bladder cancer, hepatoma, breast cancer, colon carcinoma, and head and neck cancer, gastric cancer, germ cell tumor, pediatric sarcoma, sinonasal natural killer, multiple myeloma, acute myelogenous leukemia (AML), and chronic lymphocytic leukemia (CML).

20 Additional agents and therapies can be administered in combination with the agonists and antagonists described herein. In one embodiment, the methods comprise administration of an additional therapeutic agent (e.g., a cytoxin or chemotherapeutic agent.

Also provided herein are compositions comprising a PD-1 antagonist and a TIM-4 antagonist. In one embodiment, the antagonist is a ligand, antibody (e.g., monoclonal 25 antibody or bispecific antibody) or multivalent agent. In another embodiment, the PD-1 antagonist is an anti-PD-1 antibody comprising the heavy and light chain CDRs or VRs of 5C4. In another embodiment, the PD-1 antagonist is an anti-PD-L1 antibody comprising the heavy and light chain CDRs or VRs of 12A4.

Further provided are kits for treating a cancer in a subject, the kit comprising:
30 (a) a dose of a PD-1 antagonist;
(b) a dose of a TIM-4 antagonist; and

(c) instructions for using the PD-1 antagonist and TIM-4 antagonist in the methods described herein. In one embodiment, the TIM-4 antagonist is an antibody. In another embodiment, the PD-1 antagonist is an antibody. In a particular embodiment, the PD-1 antagonist is an anti-PD-1 antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 13, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 15. In another particular embodiment, the PD-1 antagonist is an anti-PD-L1 antibody comprising antibody comprises the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 1, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 3.

Other features and advantages of the instant disclosure will be apparent from the following detailed description and examples, which should not be construed as limiting.

15 BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 depicts an alignment of the amino acid sequences of human TIM-4 isoforms 1 and 2.

Figure 2 depicts the alignment of the amino acid sequences of cynomolgous monkey, mouse and human TIM-4 orthologs.

Figure 3A-E are graphs depicting the myeloid cell populations in spleen and tumors from mice after administration of a control, a TIM-4 antagonist antibody, a PD-1 antagonist antibody, or a combination of an anti-PD-1 antibody and an anti-TIM-4 antibody.

Figure 4A-F are graphs depicting the CD8+ cell populations in spleen and tumors from mice after administration of a control, an TIM-4 antagonist antibody, an anti-PD-1 antibody, or a combination of an anti-PD-1 antibody and an anti-TIM-4 antibody.

Figure 5A-F are graphs depicting the CD4+ populations in spleen and tumors from mice after administration of a control, an TIM-4 antagonist antibody, an anti-PD-1 antibody, or a combination of an anti-PD-1 antibody and an anti-TIM-4 antibody.

Figure 6A-D are graphs depicting tumor volume (mm³) in individual mice after administration of a control, an TIM-4 antagonist antibody, an anti-PD-1 antibody, or a combination of an anti-PD-1 antibody and an anti-TIM-4 antibody.

Figure 7A-B are graphs depicting the mean and median tumor volume (mm³) in mice after administration of a control, an TIM-4 antagonist antibody, an anti-PD-1 antibody, or a combination of an anti-PD-1 antibody and an anti-TIM-4 antibody.

5 **Figure 8** is a graph depicting the tumor volumes at day 28 in mice (mm³) in mice after administration of a control, an TIM-4 antagonist antibody, an anti-PD-1 antibody, or a combination of an anti-PD-1 antibody and an anti-TIM-4 antibody.

10 **Figure 9 (A)** is a graph depicting the survival rates of mice after administration of a control, an TIM-4 antagonist antibody, an anti-PD-1 antibody, or a combination of an anti-PD-1 antibody and an anti-TIM-4 antibody in the CT26 animal model; and **(B)** is a table indicating the median survival rate of each treatment group.

Figure 10 is a graph depicting the percentage progression free survival rates of mice after administration of a control, an TIM-4 antagonist antibody, an anti-PD-1 antibody, or a combination of an anti-PD-1 antibody and an anti-TIM-4 antibody in the MC38 animal model.

15 **Figure 11** shows the mean tumor volume relative to the days post implant in the mice that were in the experiment shown in Figure 10.

Figure 12A-C shows the individual tumor volumes relative to the days post implant in the mice that were in the experiment shown in Figures 10 and 11.

20 DETAILED DESCRIPTION OF THE INVENTION

As described herein, the invention is based on the discovery that co-administration of a TIM-4 antagonist (*e.g.*, an antibody) and a PD-1 antagonist (*e.g.*, an antibody) effectively inhibits tumor growth *in vivo*, even synergistically. Accordingly, the present invention provides a method for the treatment of cancer in a subject which comprises administering to a 25 subject (*e.g.*, human) an effective amount of a PD-1 antagonist and a TIM-4 antagonist.

I. Definitions

In order that the present description may be more readily understood, certain terms are first defined. Additional definitions are set forth throughout the detailed description. Unless defined otherwise, all technical and scientific terms used herein have the same meaning as 30 commonly understood by one of ordinary skill in the art, and conventional methods of

immunology, protein chemistry, biochemistry, recombinant DNA techniques and pharmacology are employed.

As used herein, the singular forms "a", "an" and "the" include plural referents unless the context clearly dictates otherwise. The use of "or" or "and" means "and/or" unless stated 5 otherwise. Furthermore, use of the term "including" as well as other forms, such as "include", "includes", and "included", is not limiting.

The term "about" as used herein when referring to a measurable value such as an amount, a temporal duration, and the like, is meant to encompass variations of $\pm 20\%$ or $\pm 10\%$, more preferably $\pm 5\%$, even more preferably $\pm 1\%$, and still more preferably $\pm 0.1\%$ 10 from the specified value, as such variations are appropriate to perform the disclosed methods. Unless otherwise indicated, all numbers expressing quantities of ingredients, properties such as molecular weight, reaction conditions, etc., used herein are to be understood as being modified by the term "about".

As used herein, the term "subject" or "patient" are used interchangeably herein and 15 refer to a mammal such as a human, mouse, rat, hamster, guinea pig, rabbit, cat, dog, monkey, cow, horse, pig and the like (e.g., a patient having cancer).

A "solid tumor" includes, for example, sarcoma, melanoma, carcinoma, prostate carcinoma, lung carcinoma, colon carcinoma, or other solid tumor cancer.

The terms "cancer", "cancerous", or "malignant" refer to or describe the physiological 20 condition in mammals that is typically characterized by unregulated cell growth. As used herein, the term includes pre-malignant as well as malignant cancers. Examples of cancer include, for example, leukemia, lymphoma, blastoma, carcinoma and sarcoma. More particular examples of such cancers include chronic myeloid leukemia, acute lymphoblastic leukemia, Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), 25 squamous cell carcinoma, small-cell lung cancer, non-small cell lung cancer, glioma, gastrointestinal cancer, renal cancer, ovarian cancer, liver cancer, colorectal cancer, endometrial cancer, kidney cancer, prostate cancer, thyroid cancer, neuroblastoma, pancreatic cancer, glioblastoma multiforme, cervical cancer, stomach cancer, bladder cancer, hepatoma, breast cancer, colon carcinoma, and head and neck cancer, gastric cancer, germ cell tumor, 30 pediatric sarcoma, sinonasal natural killer, multiple myeloma, acute myelogenous leukemia (AML), and chronic lymphocytic leukemia (CML).

The term "carcinoma" as used herein refers to a type of cancer that develops from epithelial cells. Specifically, a carcinoma is a cancer that begins in a tissue that lines the inner or outer surfaces of the body, and that generally arises from cells originating in the endodermal or ectodermal germ layer during embryogenesis. Various subtypes of carcinomas include, adenocarcinoma (featuring microscopic glandular-related tissue 5 cytology, tissue architecture, and/or gland-related molecular products, e.g., mucin), squamous cell carcinoma (observable features and characteristics indicative of squamous differentiation, e.g., intercellular bridges, keratinization, squamous pearls), adenosquamous carcinoma (a mixed tumor containing both adenocarcinoma and squamous cell carcinoma, wherein each of 10 these cell types comprise at least 10% of the tumor volume), anaplastic or undifferentiated carcinoma (a heterogeneous group of high-grade carcinomas that feature cells lacking distinct histological or cytological evidence of any of the more specifically differentiated neoplasms), large cell carcinoma (composed of large, monotonous rounded or overtly 15 polygonal-shaped cells with abundant cytoplasm), small cell carcinoma (cells are usually round and are less than approximately 3 times the diameter of a resting lymphocyte and little evident cytoplasm).

As used herein, the term "immune cell" refers to cells that play a role in the immune response, including lymphocytes, such as B cells and T cells; natural killer cells; myeloid cells, such as monocytes, macrophages, eosinophils, mast cells, basophils, and granulocytes.

20 The term "immune response" as used herein refers to a cellular immune response, including T cell mediated and/or B cell mediated immune responses such as stimulation of T lymphocytes, macrophages, and/or natural killer cells.

The terms "myeloid-derived suppressor cell" or "MDSC" are used interchangeably 25 herein to refer to a heterogeneous group of immune cells from the myeloid lineage (a family of cells that originate from bone marrow stem cells), to which dendritic cells, macrophages and neutrophils also belong. MDSCs strongly expand in pathological situations such as chronic infections and cancer, as a result of an altered hematopoiesis. MDSCs are further classified into two subtypes, monocytic MDSCs and granulocytic MDSCs. MDSC suppressor function lies in their ability to inhibit T cell proliferation and activation. For example, under chronic 30 inflammatory conditions (viral and bacterial infections) or cancer, myeloid differentiation is skewed towards the expansion of MDSCs. These MDSCs infiltrate inflammation sites and tumors, where they inhibit immune responses by inhibiting T cells (e.g., CD8+ T cells) and

NK cells, for example. MDSCs also accelerate angiogenesis, tumor progression and metastasis through the expression of cytokines and factors such as TGF-beta.

The terms "damage-associated molecular patterns" or "DAMPs" are used interchangeably herein to refer to intracellular molecules released by injured tissues. DAMPs 5 are molecules that have a physiological role inside the cell, but acquire additional functions when exposed to the extracellular environment. For example, DAMPs alert the body about danger, stimulate an inflammatory response, and finally promote the regeneration process. Beside their passive release by dead cells, some DAMPs can be secreted or exposed by living 10 cells undergoing a life-threatening stress. DAMPs have been linked to inflammation and related disorders. DAMPs include, but are not limited to, histones, genomic DNA, HMGB1, IL1a, IL33, ATP, F-actin, cyclophilin A, HSPs, uric acid crystals, S100s, mitochondrial DNA, mitochondrial transcription factor A, alreticulin.

The term "efferocytosis" refers to the process by which dying/dead cells (e.g., 15 apoptotic or necrotic) are removed by phagocytic cells (e.g., neutrophils, monocytes, macrophages, mast cells and dendritic cells). During efferocytosis, phagocytes accumulate to the sites of apoptotic cells and the cell membrane of phagocytic cells engulfs the apoptotic cell, forming a large fluid-filled vesicle, the "effersome" containing the dead cell. The effect of efferocytosis is that dead cells are removed before their membrane integrity is breached and their contents leak into the surrounding tissue. Efferocytosis triggers specific downstream 20 intracellular signal transduction pathways, for example resulting in anti-inflammatory, anti-protease and growth-promoting effects. Conversely, impaired efferocytosis has been linked to autoimmune disease and tissue damage.

"Autophagy" or "autophagocytosis" is the natural, destructive mechanism that 25 disassembles, through a regulated process, unnecessary or dysfunctional cellular components. Autophagy allows the orderly degradation and recycling of cellular components. During this process, targeted cytoplasmic constituents are isolated from the rest of the cell within a double-membraned vesicle known as an autophagosome. The autophagosome then fuses with a lysosome and the contents are degraded and recycled. There are three different forms of 30 autophagy that are commonly described, namely macroautophagy, microautophagy and chaperone-mediated autophagy. In the context of disease, autophagy has been seen as an adaptive response to stress which promotes survival, whereas in other cases it appears to promote cell death and morbidity.

The terms "treat," "treating," and "treatment," as used herein, refer to any type of intervention or process performed on, or administering an active agent or combination of active agents to the subject with the objective of reversing, alleviating, ameliorating, inhibiting, or slowing down or preventing the progression, development, severity or 5 recurrence of a symptom, complication, condition or biochemical indicia associated with a disease.

As used herein, "effective treatment" or "positive therapeutic response" refer to a treatment producing a beneficial effect, e.g., amelioration of at least one symptom of a disease or disorder, e.g., cancer. A beneficial effect can take the form of an improvement 10 over baseline, i.e., an improvement over a measurement or observation made prior to initiation of therapy according to the method. For example, a beneficial effect can take the form of slowing, stabilizing, stopping or reversing the progression of a cancer in a subject at any clinical stage, as evidenced by a decrease or elimination of a clinical or diagnostic symptom of the disease, or of a marker of cancer. Effective treatment may, for example, 15 decrease in tumor size, decrease the presence of circulating tumor cells, reduce or prevent metastases of a tumor, slow or arrest tumor growth and/or prevent or delay tumor recurrence or relapse.

The term "effective amount" or "therapeutically effective amount" refer to an amount of an agent or combination of agents that provides the desired biological, therapeutic, and/or 20 prophylactic result. That result can be reduction, amelioration, palliation, lessening, delaying, and/or alleviation of one or more of the signs, symptoms, or causes of a disease, or any other desired alteration of a biological system. In reference to solid tumors, an effective amount comprises an amount sufficient to cause a tumor to shrink and/or to decrease the growth rate of the tumor (such as to suppress tumor growth) or to prevent or delay other 25 unwanted cell proliferation. In some embodiments, an effective amount is an amount sufficient to delay tumor development. In some embodiments, an effective amount is an amount sufficient to prevent or delay tumor recurrence. An effective amount can be administered in one or more administrations. The effective amount of the drug or composition may: (i) reduce the number of cancer cells; (ii) reduce tumor size; (iii) inhibit, 30 retard, slow to some extent and may stop cancer cell infiltration into peripheral organs; (iv) inhibit (i.e., slow to some extent and may stop tumor metastasis; (v) inhibit tumor growth; (vi) prevent or delay occurrence and/or recurrence of tumor; and/or (vii) relieve to some extent one or more of the symptoms associated with the cancer. In one example, an

“effective amount” is the amount of a PD-1 antagonist (e.g., an antibody) and TIM-4 antagonist antibody (e.g., an antibody), in combination, to effect a significant decrease in cancer or slowing of progression of cancer, such as an advanced solid tumor. An effective amount of the combination therapy is administered according to the methods described herein 5 in an "effective regimen" which refers to a combination of the PD-1 antagonist and the TIM-4 antagonist, wherein the order of administration and dosage frequency is adequate to effect treatment.

"Optimal biologic dose (OBD)" is defined as the minimum dose of an agent or combination of agents that gives the most optimal and lasting *in vivo* response without 10 clinically unacceptable toxicity. Toxicity and therapeutic efficacy can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g., for determining the LD50 (the dose lethal to 50% of the population) and the ED50 (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD50/ED50.

15 As used herein, the terms “synergy”, “therapeutic synergy”, and “synergistic effect” refer to a phenomenon where treatment of patients with a combination of therapeutic agents (e.g., PD-1 antagonist in combination with TIM-4 antagonist) manifests a therapeutically superior outcome to the outcome achieved by each individual constituent of the combination used when used alone (see, e.g., T. H. Corbett *et al.*, 1982, *Cancer Treatment Reports*, 66, 20 1187). In this context a therapeutically superior outcome includes one or more of the following (a) an increase in therapeutic response that is greater than the sum of the separate effects of each agent alone at the same dose as in the combination; (b) a decrease in the dose of one or more agents in the combination without a decrease in therapeutic efficacy; (c) a decrease in the incidence of adverse events while receiving a therapeutic benefit that is equal 25 to or greater than the monotherapy of each agent at the same dose as in the combination, (d) a reduction in dose-limiting toxicities while receiving a therapeutic benefit that is greater than the monotherapy of each agent; (e) a delay or minimization of the induction of drug resistance. In xenograft models, a combination, used at its maximum tolerated dose, in which each of the constituents will be present at a dose generally not exceeding its individual 30 maximum tolerated dose, manifests therapeutic synergy when decrease in tumor growth achieved by administration of the combination is greater than the value of the decrease in tumor growth of the best constituent when the constituent is administered alone. Synergism of a drug combination may be determined, for example, according to the combination index

(CI) theorem of Chou-Talalay (Chou *et al.*, *Adv. Enzyme Regul.* 1984; 22:27-55; Chou, *Cancer Res.* 2010;70(2):440-446).

"Free of cancer" or "disease free" or NED (No Evidence of Disease) means that the patient has demonstrated a clinical response induced by treatment with the current standard of care therapies. By "clinical response," it is meant that the clinical signs, radiological signs, and symptoms of cancer have been significantly diminished or have disappeared entirely based on clinical diagnostics, although cancerous cells can still exist in the body. Thus, it is contemplated that clinical response encompasses partial and complete response. The presence of residual cancer cells can be enumerated by assays such as CTC (Circulating Tumor Cells) and can be predictive of recurrence.

"Relapse" or "recurrence" or "resurgence" are used interchangeably herein, and refer to the radiographic diagnosis of return, or signs and symptoms of return of cancer after a period of improvement or response.

As used herein, the term "antagonist" refers to a molecule which blocks (e.g., reduces or prevents) a biological activity. The term "inhibit" or "inhibition" means to reduce by a measurable amount.

As used herein, the term "ligand" refers to a molecule that forms a complex with a biomolecule (e.g., a receptor) to serve a biological purpose. In a narrower sense, is a signal triggering molecule, binding to a site on a target protein. The binding occurs by intermolecular forces, such as ionic bonds, hydrogen bonds and van der Waals forces. The docking (association) is usually reversible (dissociation). Actual irreversible covalent binding between a ligand and its target molecule is rare in biological systems. Ligand binding to a receptor (receptor protein) alters its chemical conformation (three dimensional shape). The conformational state of a receptor protein determines its functional state.

The "level" of a protein refers to the amount of protein in a sample as determined using any method known in the art for measuring protein levels, including electrophoresis, capillary electrophoresis, high performance liquid chromatography (HPLC), thin layer chromatography (TLC), hyperdiffusion chromatography, fluid or gel precipitation reactions, absorption spectroscopy, colorimetric assays, spectrophotometric assays, flow cytometry, immunodiffusion, solution phase assay, immunoelectrophoresis, Western blotting, radioimmunoassay (RIA), enzyme-linked immunosorbent assays (ELISAs), immunofluorescent assays and electrochemiluminescence immunoassays.

The term "sample" refers to a collection of fluids, cells or tissues isolated from a subject. Biological fluids are typically liquids at physiological temperatures and may include naturally occurring fluids present in, withdrawn from, expressed or otherwise extracted from a subject or biological source. Examples of biological fluids include blood, serum, serosal fluids, plasma, lymph, urine, cerebrospinal fluid, saliva, ocular fluids, cystic fluid, tear drops, feces, sputum, mucosal secretions, vaginal secretions, gynecological fluids, 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 and the like.

The term "control sample", as used herein, refers to any clinically relevant control sample, including, for example, a sample from a healthy subject or a sample made at an earlier time point from the subject to be assessed. For example, the control sample can be a sample taken from the subject prior to onset of cancer, at an earlier stage of disease, or before the administration of treatment or of a portion of treatment.

As used herein, the term "antibody" includes whole antibodies and any antigen binding fragment (*i.e.*, "antigen-binding fragments" (also known as "antigen-binding portions")). Whole antibodies are glycoproteins comprising at least two heavy (H) chains and two light (L) chains inter-connected by disulfide bonds. Each heavy chain is comprised of a heavy chain variable region (abbreviated herein as V_H) and a heavy chain constant region. The heavy chain constant region is comprised of three domains, C_{H1} , C_{H2} and C_{H3} . Each light chain is comprised of a light chain variable region (abbreviated herein as V_L) and a light chain constant region. The light chain constant region is comprised of one domain, C_L . The V_H and V_L regions can be further subdivided into regions of hypervariability, termed complementarity determining regions (CDR), interspersed with regions that are more conserved, termed framework regions (FR). Each V_H and V_L is composed of three CDRs and four FRs, arranged from amino-terminus to carboxy-terminus in the following order: FR1, CDR1, FR2, CDR2, FR3, CDR3, FR4. The variable regions of the heavy and light chains contain a binding domain that interacts with an antigen. The constant regions of the antibodies can mediate the binding of the immunoglobulin to host tissues or factors, including various cells of the immune system (*e.g.*, effector cells) and the first component (Clq) of the classical complement system. The term "antibody" also encompasses chimeric antibodies, humanized antibodies, fully human antibodies, as well as multimeric forms of antibodies, such as minibodies, bis-scFv, diabodies, triabodies, tetrabodies and chemically conjugated Fab' multimers.

The term "antibody fragment" (also referred to as "antigen-binding fragment" or "antigen-binding portion"), as used herein, refers to one or more fragments of an antibody that retain the ability to specifically bind to an antigen. It has been shown that the antigen-binding function of an antibody can be performed by fragments of a full-length antibody.

5 Examples of binding fragments encompassed within the term "antigen-binding fragment" of an antibody include (i) a Fab fragment, a monovalent fragment consisting of the V_L, V_H, C_L and C_{H1} domains; (ii) a F(ab')₂ fragment, a bivalent fragment is essentially a Fab with part of the hinge region (see, FUNDAMENTAL IMMUNOLOGY (Paul ed., 3.sup.rd ed. 1993); (iv) a Fd fragment consisting of the V_H and C_{H1} domains; (v) a Fv fragment consisting of the V_L 10 and V_H domains of a single arm of an antibody, (vi) a dAb fragment (Ward *et al.*, (1989) *Nature* 341:544-546), which consists of a V_H domain; (vii) an isolated complementarity determining region (CDR); and (viii) a nanobody (also known as a single-domain antibody (sdAb)), which is a heavy chain variable region containing a single variable domain and two constant domains. Single domain antibodies include V_HH fragments (single-domain 15 antibodies engineered from heavy-chain antibodies found in camelids, as well as VNAR fragments (single-domain antibodies obtained from heavy-chain antibodies (IgNAR, 'immunoglobulin new antigen receptor') of cartilaginous fishes).

"Antigen binding scaffolds" are proteins that bind specifically to a target (or antigen) or epitope, such as proteins comprising an Ig fold or an Ig-like fold. Antibodies or antigen 20 binding fragments thereof are also antigen binding scaffolds. Antigen binding scaffolds can be monovalent, multivalent, e.g., bivalent, trivalent, tetravalent, or bind 5, 6 or more epitopes. Multivalent antigen binding scaffolds can be monospecific or multispecific, i.e., binding to 25 multiple (at least 2, 3, 4 or 5) epitopes that are different from one another. For example, a multivalent monospecific antigen binding scaffold is a protein that binds to at least 2, 3, 4 or 5 identical epitopes, and may be a protein comprising at least 2, 3, 4 or 5 identical antigen binding portions. For example, TIM-4 binding scaffolds may comprise 2-10, e.g., 2-6, 2-5, 2-4 or 2-3 TIM-4 binding portions, which may be the same or different from one another.

A multivalent antibody includes antibodies comprising at least 2, 3, 4, 5, 6, 7, 8, 9, 10 30 or more antigen binding portions of antibodies, which antigen binding portions may comprise a portion of a heavy chain and a portion of a light chain. An antigen binding portion may be on a single polypeptide or comprise more than one polypeptide. For example, a multivalent antibody may comprise from 2-10 antigen binding portions, which may be the same or different from each other. A multivalent antibody may be monospecific or multispecific. A

multispecific antibody may be bispecific, trispecific, tetraspecific or bind to 5 or more different epitopes.

Furthermore, although the two domains of the Fv fragment, V_L and V_H , are coded for by separate genes, they can be joined, using recombinant methods, by a synthetic linker that 5 enables them to be made as a single protein chain in which the V_L and V_H regions pair to form monovalent molecules (known as single chain Fv (scFv); *see e.g.*, Bird *et al.* (1988) *Science* 242:423-426; and Huston *et al.* (1988) *Proc. Natl. Acad. Sci. USA* 85:5879-5883). Such single chain antibodies are also intended to be encompassed within the term "antigen-binding fragment" of an antibody. These antibody fragments are obtained using conventional 10 techniques known to those with skill in the art, and the fragments are screened for utility in the same manner as are intact antibodies.

As used herein, an antigen binding scaffold that "specifically binds" to an antigen or epitope thereof is an antigen binding scaffold that binds to the antigen or epitope thereof with a K_D of 10^{-7} M, 5×10^{-8} M, 10^{-8} M, 5×10^{-9} M, 10^{-9} M, 5×10^{-10} M, 10^{-10} M, 5×10^{-11} M, 10^{-11} M, 5×10^{-12} M, 10^{-12} M or less. For example, an antigen binding scaffold that specifically binds to TIM-4 is an antigen binding scaffold that binds to TIM-4 with a K_D of 10^{-7} M, 5×10^{-8} M, 10^{-8} M, 5×10^{-9} M, 10^{-9} M, 5×10^{-10} M, 10^{-10} M, 5×10^{-11} M, 10^{-11} M, 5×10^{-12} M, 10^{-12} M or less. For example, an antibody that "specifically binds to human PD-1" or "specifically binds to human PD-L1" is intended to refer to an antibody that binds to human 15 PD-1 or PD-L1, respectively, with a K_D of 10^{-7} M, 5×10^{-8} M, 10^{-8} M, 5×10^{-9} M, 10^{-9} M, 5×10^{-10} M, 10^{-10} M, 5×10^{-11} M, 10^{-11} M, 5×10^{-12} M, 10^{-12} M or less. An antigen binding scaffold that comprises 2 or more regions binding to an antigen or epitope may bind 20 specifically to the antigen or epitope even it has a lower affinity of binding to the antigen or epitope than the ranges provided above, as it will bind to the antigen or epitope with increased avidity.

A "bispecific" or "bifunctional antibody" is an artificial hybrid antibody having two different heavy/light chain pairs and two different binding sites. Bispecific antibodies can be produced by a variety of methods including fusion of hybridomas or linking of Fab' fragments. *See, e.g.*, Songsivilai & Lachmann, *Clin. Exp. Immunol.* 79:315-321 (1990); 25 Kostelny *et al.*, *J. Immunol.* 148, 1547-1553 (1992).

The term "monoclonal antibody" as used herein, refers to refers to an antibody from a population of substantially homogeneous antibodies that display a single binding specificity

and affinity for a particular epitope, except for possible variants that may arise during production of the monoclonal antibody, such variants generally being present in minor amounts. Accordingly, the term "human monoclonal antibody" or "monoclonal antibody composition" refers to an antibody which has variable and optional constant regions derived from human germline immunoglobulin sequences. The modifier "monoclonal" indicates the character of the antibody as being obtained from a substantially homogeneous population of antibodies, and is not to be construed as requiring production of the antibody by any particular method. For example, the monoclonal antibodies described herein may be made by a variety of techniques. In one embodiment, human monoclonal antibodies are produced by a hybridoma which includes a B cell obtained from a transgenic non-human animal, *e.g.*, a transgenic mouse, having a genome comprising a human heavy chain transgene and a light chain transgene fused to an immortalized cell.

The term "epitope" or "antigenic determinant" refers to a site on an antigen to which an immunoglobulin or antibody specifically binds. Epitopes can be formed both from contiguous amino acids or noncontiguous amino acids juxtaposed by tertiary folding of a protein. Epitopes formed from contiguous amino acids are typically retained on exposure to denaturing solvents, whereas epitopes formed by tertiary folding are typically lost on treatment with denaturing solvents. An epitope typically includes at least 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14 or 15 amino acids in a unique spatial conformation. Methods of determining spatial conformation of epitopes include techniques in the art and those described herein, for example, x-ray crystallography, HDX-MS and 2-dimensional nuclear magnetic resonance (see, *e.g.*, *Epitope Mapping Protocols in Methods in Molecular Biology*, Vol. 66, G. E. Morris, Ed. (1996)).

The term "epitope mapping" refers to the process of identification of the molecular determinants for antibody-antigen recognition.

The term "binds to the same epitope," with reference to two or more antibodies, means that the antibodies compete for binding to an antigen and bind to the same, overlapping, or encompassing continuous or discontinuous segments of amino acids. Those of skill in the art understand that the phrase "binds to the same epitope" does not necessarily mean that the antibodies bind to exactly the same amino acids. The precise amino acids to which the antibodies bind can differ. For example, a first antibody can bind to a segment of amino acids that is completely encompassed by the segment of amino acids bound by a second antibody. In another example, a first antibody binds one or more segments of amino

acids that significantly overlap the one or more segments bound by the second antibody. For the purposes herein, such antibodies are considered to "bind to the same epitope." Two antibodies "bind to the same epitope" as determined by a given method, e.g., HDX-MS, crystallography or target mutational analysis, requires that the same amino acids are

5 identified for both antibodies by one or more given methods.

Accordingly, also, encompassed by the present invention are antibodies that bind to an epitope which comprises all or a portion of an epitope recognized by the particular antibodies described herein (e.g., the same or an overlapping region or a region between or spanning the region) or that bind to the same epitope as determined by a given method, e.g.,

10 HDX-MS, crystallography or target mutational analysis.

Also encompassed by the present invention are antibodies that compete for binding with the antibodies described herein, and optionally bind the same epitope. Antibodies that compete for binding can be identified using routine techniques. Such techniques include, for example, an immunoassay, which shows the ability of one antibody to block the binding of a reference antibody to a common antigen. Numerous types of competitive binding assays are known, for example: Biacore, flow cytometry, solid phase direct or indirect radioimmunoassay (RIA), solid phase direct or indirect enzyme immunoassay (EIA),

15 sandwich competition assay (see Stahli *et al.*, *Methods in Enzymology* 9:242 (1983)); solid phase direct biotin-avidin EIA (see Kirkland *et al.*, *J. Immunol.* 137:3614 (1986)); solid phase direct labeled assay, solid phase direct labeled sandwich assay (see Harlow and Lane, *Antibodies: A Laboratory Manual*, Cold Spring Harbor Press (1988)); solid phase direct label RIA using I-125 label (see Morel *et al.*, *Mol. Immunol.* 25(1):7 (1988)); solid phase

20 direct biotin-avidin EIA (Cheung *et al.*, *Virology* 176:546 (1990)); and direct labeled RIA. (Moldenhauer *et al.*, *Scand. J. Immunol.* 32:77 (1990)). Typically, such an assay involves the use of purified antigen bound to a solid surface or cells bearing either of these, an unlabeled test immunoglobulin and a labeled reference immunoglobulin. Competitive inhibition is measured by determining the amount of label bound to the solid surface or cells in the

25 presence of the test immunoglobulin. Usually the test immunoglobulin is present in excess. Usually, when a competing antibody is present in excess, it will inhibit specific binding of a reference antibody to a common antigen by at least 50-55%, 55-60%, 60-65%, 65-70% 70-75% or more.

Epitope mapping methods also include x-ray analyses of crystals of antigen:antibody complexes which provides atomic resolution of the epitope. Other methods monitor the binding of the antibody to antigen fragments or mutated variations of the antigen where loss of binding due to a modification of an amino acid residue within the antigen sequence is often 5 considered an indication of an epitope component. In addition, computational combinatorial methods for epitope mapping can also be used. These methods rely on the ability of the antibody of interest to affinity isolate specific short peptides from combinatorial phage display or yeast display peptide libraries. The peptides are then regarded as leads for the definition of the epitope corresponding to the antibody used to screen the peptide library. For 10 epitope mapping, computational algorithms have also been developed which have been shown to map conformational discontinuous epitopes.

Chimeric molecules (or fusion molecules) comprising an antigen binding domain, or equivalent, fused to another polypeptide or molecule are also encompassed by the present invention. For example, the polypeptides may be fused or conjugated to an antibody Fc 15 region, or portion thereof (e.g., an Fc fusion protein). The antibody portion fused to a polypeptide may comprise the constant region, hinge region, CH1 domain, CH2 domain, and CH3 domain or any combination of whole domains or portions thereof. The polypeptides may also be fused or conjugated to the above antibody portions to form multimers. For example, Fc portions fused to the polypeptides of the present invention can form dimers 20 through disulfide bonding between the Fc portions. Higher multimeric forms can be made by fusing the polypeptides to portions of IgA and IgM. Methods for fusing or conjugating the polypeptides of the present invention to antibody portions are known in the art. See, e.g., U.S. Pat. Nos. 5,336,603, 5,622,929, 5,359,046, 5,349,053, 5,447,851, and 5,112,946; EP 307,434; EP 367,166; PCT Publication Nos. WO 96/04388 and WO 91/06570; Ashkenazi et 25 al., Proc. Natl. Acad. Sci. USA, 88:10535-10539 (1991); Zheng et al., J. Immunol., 154:5590-5600 (1995); and Vil et al., Proc. Natl. Acad. Sci. USA, 89:11337-11341 (1992).

As used herein, the term “immunoconjugate” refers to an antibody linked to a therapeutic moiety, such as a cytotoxin, a drug or a radioisotope. When conjugated to a cytotoxin, these antibody conjugates are referred to as “immunotoxins.” A cytotoxin or 30 cytotoxic agent includes any agent that is detrimental to (e.g., kills) cells. Examples include taxol, cytochalasin B, gramicidin D, ethidium bromide, emetine, mitomycin, etoposide, tenoposide, vincristine, vinblastine, colchicin, doxorubicin, daunorubicin, dihydroxy anthracin dione, mitoxantrone, mithramycin, actinomycin D, 1-dehydrotestosterone,

glucocorticoids, procaine, tetracaine, lidocaine, propranolol, and puromycin and analogs or homologs thereof. Therapeutic agents include, but are not limited to, antimetabolites (e.g., methotrexate, 6-mercaptopurine, 6-thioguanine, cytarabine, 5-fluorouracil decarbazine), alkylating agents (e.g., mechlorethamine, thioepa chlorambucil, melphalan, carmustine 5 (BSNU) and lomustine (CCNU), cyclothosphamide, busulfan, dibromomannitol, streptozotocin, mitomycin C, and cis-dichlorodiamine platinum (II) (DDP) cisplatin), anthracyclines (e.g., daunorubicin (formerly daunomycin) and doxorubicin), antibiotics (e.g., dactinomycin (formerly actinomycin), bleomycin, mithramycin, and anthramycin (AMC)), and anti-mitotic agents (e.g., vincristine and vinblastine). Antibodies use in the present 10 invention can be conjugated to a radioisotope, e.g., radioactive iodine, to generate cytotoxic radiopharmaceuticals for treating cancer.

Immunoconjugates can be used to modify a given biological response, and the drug moiety is not to be construed as limited to classical chemical therapeutic agents. For example, the drug moiety may be a protein or polypeptide possessing a desired biological 15 activity. Such proteins may include, for example, an enzymatically active toxin, or active fragment thereof, such as abrin, ricin A, pseudomonas exotoxin, or diphtheria toxin; a protein such as tumor necrosis factor or interferon- γ ; or, biological response modifiers such as, for example, lymphokines, interleukin-1 ("IL-1"), interleukin-2 ("IL-2"), interleukin-6 ("IL-6"), granulocyte macrophage colony stimulating factor ("GM-CSF"), granulocyte colony 20 stimulating factor ("G-CSF"), or other growth factors.

Techniques for conjugating such therapeutic moiety to antibodies are well known, see, e.g., Arnon et al., "Monoclonal Antibodies For Immunotargeting Of Drugs In Cancer Therapy", in *Monoclonal Antibodies And Cancer Therapy*, Reisfeld *et al.* (eds.), pp. 243-56 (Alan R. Liss, Inc. 1985); Hellstrom et al., "Antibodies For Drug Delivery", in *Controlled 25 Drug Delivery* (2nd Ed.), Robinson et al. (eds.), pp. 623-53 (Marcel Dekker, Inc. 1987); Thorpe, "Antibody Carriers Of Cytotoxic Agents In Cancer Therapy: A Review", in *Monoclonal Antibodies '84: Biological And Clinical Applications*, Pinchera et al. (eds.), pp. 475-506 (1985); "Analysis, Results, And Future Prospective Of The Therapeutic Use Of 30 Radiolabeled Antibody In Cancer Therapy", in *Monoclonal Antibodies For Cancer Detection And Therapy*, Baldwin et al. (eds.), pp. 303-16 (Academic Press 1985), and Thorpe et al., "The Preparation And Cytotoxic Properties Of Antibody-Toxin Conjugates", *Immunol. Rev.*, 62:119-58 (1982).

As used herein, the term "multivalent" refers to a recombinant molecule that incorporates more than two biologically active segments. The protein fragments forming the multivalent molecule optionally may be linked through a polypeptide linker which attaches the constituent parts and permits each to function independently.

5 As used herein, "functional equivalents" refer to peptides or polypeptides (*e.g.*, antibodies or antigen binding portion thereof) which maintain a substantial amount of their original immunological activity. For example, certain amino acids may be substituted for other amino acids in a protein structure without appreciable loss of interactive binding capacity with structures such as, for example, antigen-binding regions of antibodies or 10 binding sites on substrate molecules. It is thus contemplated that various changes may be made in the amino acid sequences of the disclosed compositions without appreciable loss of their biological utility or activity. Amino acid substitutions may be based on the relative similarity of the amino acid side-chain substituents, for example, their hydrophobicity, hydrophilicity, charge, size, and the like. Exemplary substitutions which take various of the 15 foregoing characteristics into consideration are well known to those of skill in the art, and such immunologically functional equivalents are also encompassed within the present invention.

"Percent (%) amino acid sequence identity" herein is defined as the percentage of amino acid residues in a candidate sequence that are identical with the amino acid residues in 20 a selected sequence, after aligning the sequences and introducing gaps, if necessary, to achieve the maximum percent sequence identity, and not considering any conservative substitutions as part of the sequence identity. Alignment for purposes of determining percent amino acid sequence identity can be achieved in various ways that are within the skill in the art, for instance, using publicly available computer software such as BLAST, BLAST-2, 25 ALIGN, ALIGN-2 or Megalign (DNASTAR) software. Those skilled in the art can determine appropriate parameters for measuring alignment, including any algorithms needed to achieve maximal alignment over the full-length of the sequences being compared.

For purposes herein, the % amino acid sequence identity of a given amino acid sequence A to, with, or against a given amino acid sequence B (which can alternatively be 30 phrased as a given amino acid sequence A that has or comprises a certain % amino acid sequence identity to, with, or against a given amino acid sequence B) is calculated as follows: 100 times the fraction X/Y where X is the number of amino acid residues scored as identical matches by a sequence alignment program, such as BLAST, BLAST-2, ALIGN, ALIGN-2 or

Megalign (DNASTAR), in that program's alignment of A and B, and where Y is the total number of amino acid residues in B. It will be appreciated that where the length of amino acid sequence A is not equal to the length of amino acid sequence B, the % amino acid sequence identity of A to B will not equal the % amino acid sequence identity of B to A.

5 **II. PD-1 Antagonists**

As used herein, the terms “Protein PD-1,” “PD-1,” PD1,” “PDCD1” are used interchangeably with “Programmed Death 1,” “Programmed Cell Death 1.” The complete human PD-1 sequence can be found under GenBank Accession No. U64863 (SEQ ID NO:23).

10 As used herein, the terms “PD-L1”, “PDL1”, “PDCD1L1”, “PDCD1LG1”, “CD274”, “B7 homolog 1”, “B7-H1”, “B7-H”, and “B7H1” are used interchangeably with “Programmed Cell Death 1 Ligand 1.” The complete human PD-L1 amino acid sequence - isoform a precursor – can be found under GenBank Accession No. NP_054862.1 (SEQ ID NO:24). The complete human PD-L1 amino acid sequence –isoform b precursor - can be 15 found under GenBank Accession No. NP_001254635.1 (SEQ ID NO:25).

The protein Programmed Death 1 (PD-1) is an inhibitory member of the CD28 family of receptors, that also includes CD28, CTLA-4, ICOS and BTLA. PD-1 is expressed on activated B cells, T cells, and myeloid cells (Agata *et al.*, *supra*; Okazaki *et al.* (2002) *Curr. Opin. Immunol.* 14: 391779-82; Bennett *et al.* (2003) *J Immunol* 170:711-8). The initial 20 members of the family, CD28 and ICOS, were discovered by functional effects on augmenting T cell proliferation following the addition of monoclonal antibodies (Hutloff *et al.* (1999) *Nature* 397:263-266; Hansen *et al.* (1980) *Immunogenetics* 10:247-260). PD-1 was discovered through screening for differential expression in apoptotic cells (Ishida *et al.* (1992) *EMBO J* 11:3887-95). The other members of the family, CTLA-4 and BTLA, were 25 discovered through screening for differential expression in cytotoxic T lymphocytes and TH1 cells, respectively. CD28, ICOS and CTLA-4 all have an unpaired cysteine residue allowing for homodimerization. In contrast, PD-1 is suggested to exist as a monomer, lacking the unpaired cysteine residue characteristic in other CD28 family members.

The PD-1 protein is a 55 kDa type I transmembrane protein that is part of the Ig gene 30 superfamily (Agata *et al.* (1996) *Int Immunol* 8:765-72). PD-1 contains a membrane proximal immunoreceptor tyrosine inhibitory motif (ITIM) and a membrane distal tyrosine-based switch motif (ITSM) (Thomas, M.L. (1995) *J Exp Med* 181:1953-6; Vivier, E and

Daeron, M (1997) *Immunol Today* 18:286-91). Although structurally similar to CTLA-4, PD-1 lacks the MYPPPY motif (SEQ ID NO: 27) that is critical for B7-1 and B7-2 binding.

Consistent with PD-1 being an inhibitory member of the CD28 family, PD-1 deficient animals develop various autoimmune phenotypes, including autoimmune cardiomyopathy and a lupus-like syndrome with arthritis and nephritis (Nishimura *et al.* (1999) *Immunity* 11:141-51; Nishimura *et al.* (2001) *Science* 291:319-22). Additionally, PD-1 has been found to play a role in autoimmune encephalomyelitis, systemic lupus erythematosus, graft-versus-host disease (GVHD), type I diabetes, and rheumatoid arthritis (Salama *et al.* (2003) *J Exp Med* 198:71-78; Prokunina and Alarcon-Riquelme (2004) *Hum Mol Genet* 13:R143; Nielsen *et al.* (2004) *Lupus* 13:510). In a murine B cell tumor line, the ITSM of PD-1 was shown to be essential to block BCR-mediated Ca²⁺-flux and tyrosine phosphorylation of downstream effector molecules (Okazaki *et al.* (2001) *PNAS* 98:13866-71).

Two ligands for PD-1 have been identified, PD-L1 and PD-L2, that have been shown to downregulate T cell activation upon binding to PD-1 (Freeman *et al.* (2000) *J Exp Med* 192:1027-34; Latchman *et al.* (2001) *Nat Immunol* 2:261-8; Carter *et al.* (2002) *Eur J Immunol* 32:634-43). Both PD-L1 and PD-L2 are B7 homologs that bind to PD-1, but do not bind to other CD28 family members. PD-L1 is abundant in a variety of human cancers (Dong *et al.* (2002) *Nat. Med.* 8:787-9). The interaction between PD-1 and PD-L1 results in a decrease in tumor infiltrating lymphocytes, a decrease in T-cell receptor mediated proliferation, and 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 can be reversed by inhibiting the local interaction of PD-1 with PD-L1, and the effect is additive when the interaction of PD-1 with PD-L2 is blocked as well (Iwai *et al.* (2002) *Proc. Nat'l. Acad. Sci. USA* 99:12293-7; Brown *et al.* (2003) *J. Immunol.* 170:1257-66).

The methods of the present invention involve the use of a PD-1 antagonist (*e.g.*, an antibody) in combination with a TIM-4 antagonist (*e.g.*, an antibody), for treating cancer. Accordingly, PD-1 antagonists of the invention bind to ligands of PD-1 and interfere with, reduce, or inhibit the binding of one or more ligands to the PD-1 receptor, or bind directly to the PD-1 receptor, without engaging in signal transduction through the PD-1 receptor. In one embodiment, the PD-1 antagonist binds directly to PD-1 and blocks PD-1 inhibitory signal transduction. In another embodiment the PD-1 antagonist binds to one or more ligands of PD-1 (*e.g.*, PD-L1 and PD-L2) and reduces or inhibits the ligand(s) from triggering inhibitory

signal transduction through the PD-1. In one embodiment, the PD-1 antagonist binds directly to PD-L1, inhibiting or preventing PD-L1 from binding to PD-1, thereby blocking PD-1 inhibitory signal transduction.

PD-1 antagonists used in the methods and compositions of the present invention 5 include PD-1 binding scaffold proteins and include, but are not limited to, PD-1 ligands, antibodies and multivalent agents. In a particular embodiment, the antagonist is a fusion protein, such as AMP-224. In another embodiment, the antagonist is an anti-PD-1 antibody (“PD-1 antibody”). Anti-human-PD-1 antibodies (or VH and/or VL domains derived therefrom) suitable for use in the invention can be generated using methods well known in 10 the art. Alternatively, art recognized anti-PD-1 antibodies can be used. For example, antibodies nivolumab (OPDIVOTM), MK-3475 (pembrolizumab (KEYTRUDATM), PDR001 or CT-011 can be used. Additionally, monoclonal antibodies 5C4, 17D8, 2D3, 4H1, 4A11, 15 7D3, and 5F4, described in WO 2006/121168, the teachings of which are hereby incorporated by reference, can be used. Antibodies that compete with any of these art-recognized antibodies for binding to PD-1 also can be used.

An exemplary anti-PD-1 antibody is 5C4 comprising heavy and light chains having the sequences shown in SEQ ID NOS: 11 and 12, respectively, or antigen binding fragments and variants thereof. In other embodiments, the antibody comprises the heavy and light chain CDRs or variable regions of 5C4. Accordingly, in one embodiment, the antibody comprises 20 the CDR1, CDR2, and CDR3 domains of the VH of 5C4 having the sequence set forth in SEQ ID NO: 13, and the CDR1, CDR2 and CDR3 domains of the VL of 5C4 having the sequences set forth in SEQ ID NO: 15. In another embodiment, the antibody comprises CDR1, CDR2 and CDR3 domains having the sequences set forth in SEQ ID NOS: 17, 18, and 25 19, respectively, and CDR1, CDR2 and CDR3 domains having the sequences set forth in SEQ ID NOS: 20, 21, and 22, respectively. In another embodiment, the antibody comprises VH and/or VL regions having the amino acid sequences set forth in SEQ ID NO: 13 and/or SEQ ID NO: 15, respectively. In another embodiment, the antibody comprises the heavy chain variable (VH) and/or light chain variable (VL) regions encoded by the nucleic acid sequences set forth in SEQ ID NO: 14 and/or SEQ ID NO: 16, respectively. In another 30 embodiment, the antibody competes for binding with and/or binds to the same epitope on PD-1 as the above-mentioned antibodies. In another embodiment, the antibody has at least about 90% variable region amino acid sequence identity with the above-mentioned antibodies (e.g.,

at least about 90%, 95% or 99% variable region identity with SEQ ID NO: 13 or SEQ ID NO: 15).

In certain embodiments, the PD1 antibodies exhibit one or more desirable functional properties, such as high affinity binding to PD-1, e.g., binding to human PD-1 with a K_D of 5 10^{-7} M or less; lack of significant cross-reactivity to other CD28 family members, e.g., CD28, CTLA-4 and ICOS; the ability to stimulate T cell proliferation in a mixed lymphocyte reaction (MLR) assay; the ability to increase IFN- γ and/or IL-2 secretion in an MLR; the ability to inhibit binding of one or more PD-1 ligands (e.g., PD-L1 and/or PD-L2) to PD-1; the ability to stimulate antigen-specific memory responses; the ability to stimulate antibody 10 responses and/or the ability to inhibit growth of tumor cells *in vivo*.

In another embodiment, the PD-1 antagonist is an anti-PD-L1 antibody. Anti-human-PD-L1 antibodies (or VH and/or VL domains derived therefrom) suitable for use in the invention can be generated using methods well known in the art. Alternatively, art-recognized anti-PD-L1 antibodies can be used. For example, MEDI4736 (also known as 15 Anti-B7-H1; durvalumab (IMFINZITM)), MPDL3280A (atezolizumab (TECENTRIQTM) also known as RG7446), and avelumab (BAVENCIOTM) and can be used. Additionally, monoclonal antibodies 12A4, 3G10, 10A5, 5F8, 10H10, 1B12, 7H1, 11E6, 12B7, and 13G4 described in WO 2007/005874 and US Patent No. 7,943,743, the teachings of which are hereby incorporated by reference, can be used. Antibodies that compete with any of these 20 art-recognized antibodies for binding to PD-L1 also can be used.

An exemplary anti-PD-L1 antibody is 12A4 (WO 2007/005874 and US Patent No. 7,943,743). In one embodiment, the antibody comprises the heavy and light chain CDRs or VRs of 12A4. Accordingly, in one embodiment, the antibody comprises the CDR1, CDR2, and CDR3 domains of the VH region of 12A4 having the sequence shown in SEQ ID NO: 1 25 and the CDR1, CDR2 and CDR3 domains of the VL region of 12A4 having the sequence shown in SEQ ID NO: 3. In another embodiment, the antibody comprises the heavy chain CDR1, CDR2 and CDR3 domains having the sequences set forth in SEQ ID NOS: 5, 6, and 7, respectively, and the light chain CDR1, CDR2 and CDR3 domains having the sequences set forth in SEQ ID NOS: 8, 9, and 10, respectively. In another embodiment, the antibody 30 comprises VH and/or VL regions having the amino acid sequences set forth in SEQ ID NO: 1 and/or SEQ ID NO: 3, respectively. In another embodiment, the antibody comprises the heavy chain variable (VH) and/or light chain variable (VL) regions encoded by the nucleic acid sequences set forth in SEQ ID NO: 2 and/or SEQ ID NO: 4, respectively. In another

embodiment, the antibody competes for binding with, and/or binds to the same epitope on PD-L1 as, the above-mentioned antibodies. In another embodiment, the antibody has at least about 90% variable region amino acid sequence identity with the above-mentioned antibodies (e.g., at least about 90%, 95% or 99% variable region identity with SEQ ID NO: 1 or SEQ 5 ID NO: 3).

Anti-PD-1 or anti-PD-L1 antibodies may bind to PD-1 or PD-L1, respectively, with a K_D of 10^{-7} M, 5×10^{-8} M, 10^{-8} M, 5×10^{-9} M, 10^{-9} M, 5×10^{-10} M, 10^{-10} M or less.

In certain embodiments, an anti-PD-1 or anti-PD-L1 antibody is an IgG antibody, such as an IgG1, IgG2 or IgG4 antibody. In specific embodiments, an anti-PD-1 or PD-11 10 antibody has an effectorless constant region. Anti-PD-1 or PD-L1 antibodies may be IgG4 antibodies, e.g., IgG4 antibodies having an S228P mutation.

III. TIM-4 antagonists

As used herein, the terms "TIM-4" and "TIMD-4", also known as "T-cell immunoglobulin and mucin domain-containing protein 4", "T-cell immunoglobulin mucin 15 receptor 4," "T-cell membrane protein 4" and "SMUCKLER" are used interchangeably. The complete amino acid and nucleotide sequences of isoform-1 of human TIM-4 can be found under GenBank Accession Nos. NP_612388.2 and NM_138379.2, respectively (SEQ ID 20 NOS: 26-27). The complete amino acid and nucleotide sequences of isoform-2 of human TIM-4 can be found under GenBank Accession Nos. NP_001140198.1 and NM_001146726.1, respectively (SEQ ID NOS: 28-29) (Jones *et al.*, Int Arch Allergy Immunol. 2006;141(4):331-6). An alignment of the two human isoforms is depicted in

Figure 1.

The amino acid sequences of TIM-4 isoforms from cynomolgus monkey can be found under GenBank Accession Nos. XP_005558436 and EHH54702, and the amino acid 25 sequences for murine TIM-4 isoforms can be found under GenBank Accession Nos. NP_848874 and NP_599009. An alignment of the amino acid sequences of mouse, cynomolgus monkey and human TIM-4 polypeptides is provided in **Figure 2**, and the percent identity of the full-length and IgV regions of human, cynomolgus and murine orthologs are set forth in **Table 1**.

Table 1. TIM-4 Orthologs

	hTIM4-FL	hTIM4 IgV
Cyno	87%	93%
Mouse	49%	64%

5

The protein "TIM-4" is type I membrane protein that is a member of T-cell immunoglobulin and mucin domain-containing (TIM) family. The human TIM family contains three members (TIM-1, TIM-3 and TIM-4) on human chromosome 5q33.2, located in a chromosomal region that has been linked with asthma, allergy and autoimmunity. TIM 10 proteins are type I cell-surface glycoproteins with common structural features including an N-terminal immunoglobulin (IgV)-like domain with highly conserved cysteine residues, a threonine-rich mucin domain with O-linked and N-linked glycosylations, a single transmembrane domain, and a cytoplasmic region. While the cytoplasmic domain of human TIM-1 and TIM-3 contain tyrosine phosphorylation motif(s), TIM-4 lacks a phosphotyrosine 15 motif.

TIM-4 is expressed on myeloid cells, including dendritic cells (DCs) and macrophages from spleen, lymph nodes, or peritoneal cavity. TIM-4 has been demonstrated to bind to TIM-1, MerTK, Integrin $\alpha\beta 3$ and phosphatidyl serine via the IgV domain. MerTK and Integrin $\alpha\beta 3$ may play a role in certain TIM4 mediated biological activities 20 (Nishi et al. (2014) Mol Cell Biol. 34(8):1512-20 and Toda et al. (2012) Mol Cell Biol. 32(1):118-25). TIM-4 differentially regulates T cell homeostasis by inhibiting naïve T cells during the induction phase of an immune response and enhancing T cell responses at the effector phase (Rodriguez-Manzanet *et al.*, *supra*). For example, TIM-4 binds to TIM-1, which is present on activated T cells, and co-stimulates T cell proliferation.

25 TIM-4 binds to the phosphatidylserine (PS) through the FG-CC' binding cleft in the N-terminal immunoglobulin variable (IgV) domain. TIM-4 is upregulated following the release of DAMPs, and enhances efferocytosis of apoptotic cells by macrophages (Kobayashi, 2007; Albacker, et al., J. Immunol. 2010; 185:6839-6849; Mizui et al. Int. Immunol. 2008; 20:695-708). In contrast, TIM-4 knockout mice have defects in lysosomal 30 degradation, antigen presentation and cross-priming (Miyanishi 2012, Rodrigues-Manzaneta, 2010).

TIM-4 has been demonstrated to be expressed on macrophages and DCs in the tumor microenvironment (TME), and may modulate the interaction between myeloid cells and

antigen-specific cytotoxic T cells within tumors. Bahgadadi *et al.* (*supra*) reported that apoptotic tumor cells were ingested by TIM-4+ bone-marrow derived macrophages (BMDMs), suggesting that TIM-4 contributes to immune tolerance by promoting excessive lysosomal degradation of ingested tumor cells, leading to impaired tumor antigen 5 presentation by TIM-4+ tumor-associated macrophages (TAMs). Inhibition of this function appears to promote antitumor activity following monotherapy with an anti-TIM-4 blocking antibody and demonstrates synergistic antitumor activity when combined with an anti-TIM-3 blocking antibody in the context of the vaccine therapy model (Baghadadi 2013).

The term "antagonist" as used with reference to TIM-4 refers to any molecule that 10 partially or fully inhibits one or more biological activities of TIM-4, *in vitro*, *in situ*, or *in vivo*. Examples of such biological activities include binding of TIM-4 to PS, efferocytosis of tumor cells, and suppression of tumor antigen presentation, as well as those further reported in the literature. TIM-4 antagonists may function in a direct or indirect manner. For example, the TIM-4 antagonist may function to partially or fully inhibit one or more 15 biological activities of TIM-4, *in vitro*, *in situ*, or *in vivo* as a result of blocking direct binding to TIM-4 to PS and/or blocking the interaction of TIM-4 to other PS receptors and reducing cellular ability to engage in efferocytosis of tumor cells. The antagonist may also function by interfering with TIM-4 interaction with TIM-1 thereby preventing direct T cell regulatory activity. The TIM-4 antagonist may also function indirectly to partially or fully 20 inhibit one or more biological activities of TIM-4, *in vitro*, *in situ*, or *in vivo* as a result of, e.g., inhibiting another molecule which then blocks TIM-4 activation or expression. It is contemplated that an antagonist may act as a molecule which functions indirectly to block, inhibit or decrease TIM-4 expression or activity.

A TIM-4 antagonist may be any molecule that directly or indirectly inhibits or 25 decreases the activity of TIM-4 and reduces tumor growth, whether on its own or in combination with another treatment, such as a PD-1 antagonist. Exemplary TIM-4 antagonists include TIM-4 binding scaffolds, such as anti-TIM-4 antibodies ("TIM-4 antibodies"), e.g., chimeric, humanized or fully human antibodies, an antigen binding portion thereof, or molecules that are based on or derived from any of these. TIM-4 antagonists may 30 also be non-antibody proteins. For example, TIM-4 antagonist also include modified TIM-4 ligands or binding proteins, e.g., TIM-1 and molecules that are derived from or based on PS. In addition, TIM-4-regulated biology may be interfered with by fusion proteins such as TIM-4: Ig.

A TIM-4 antagonist may be monovalent or multivalent. In certain embodiments, a TIM-4 antagonist is bivalent, trivalent, tetravalent, or binds to 5, 6, 7, 8, 9, 10 or more TIM-4 epitopes, which may be the same or different TIM-4 epitopes. In certain embodiments, a TIM-4 antagonist is a multivalent monospecific TIM-4 binding scaffold, e.g., a protein

5 comprising a TIM-4 binding scaffold that comprises at least 2, 3, 4, 5, 6, 7, 8, 9, 10 or more regions that specifically bind to the same TIM-4 epitope, which binding regions may be composed of the same or a different amino acid sequence. For example, a TIM-4 antagonist may be a TIM-4 binding scaffold comprising 2, 3, 4, 5, 6, 7, 8, 9, 10 or more repeats of the same TIM-4 binding region, e.g., the N-terminal IgV region or portion thereof containing the

10 FG-CC' binding cleft of the IgV region.

In certain embodiments, a TIM-4 antagonist binds specifically to TIM-4, but does not bind significantly or specifically to other members of the TIM family, such as TIM-1 or TIM-

3. In other embodiments, a TIM-4 antagonist binds specifically to TIM-4 and TIM-1.

In another embodiment, the TIM-4 antagonist is an antibody, e.g., an antibody that

15 binds to human TIM-4 with a K_D of 10^{-7} M, 5×10^{-8} M, 10^{-8} M, 5×10^{-9} M, 10^{-9} M, 5×10^{-10} M, 10^{-10} M or less, wherein the antibody inhibits tumor growth, and/or increases tumor antigen specific CTLs. Antibodies binding to human TIM-4 are known in the art. Any of these antibodies may be used in combination with a PD-1 antagonist, provided that their combination results in inhibition of tumor growth or reduction in tumor size, e.g., in a subject

20 having cancer. Exemplary antibodies include those which bind specifically to the N-terminal IgV domain of TIM-4 (e.g., the FG-CC" binding cleft). Anti-TIM-4 antibodies include those which bind human TIM-4, e.g., 9F4 (BioLegend). Anti-mouse anti-TIM4 antibodies include RMT4-53 (BioXCell, GeneTex), RMT4-54 (BioLegend) F31-5G3 and 21H12 (BioLegend, BD Biosciences, respectively). Variants of these antibodies, such as antibodies comprising

25 the CDRs of these antibodies, or antibodies that compete for binding to human or murine TIM-4 with one of these antibodies or antibodies that bind to the same or similar epitope on TIM-4 as one of these antibodies may be used in combinations with a PD-1 antagonist.

In certain embodiments, an anti-TIM4 antibody for use in the methods described herein binds to a region on human TIM-4 that is not the IgV domain, and can be, e.g., the

30 stem domain or any other region in the extracellular domain of TIM-4, provided that the antibody antagonizes the biological activity of TIM-4, and has at least an additive effect with a PD-1 antagonist when used for treating cancer (relative to either agent alone).

In another embodiment, the TIM-4 antagonist is a multivalent agent, such as a multimer (e.g., a polypeptide construct with trimerizing domain and a polypeptide that binds TIM-4).

Agents, which compete for binding to TIM-4 with any of the exemplary agents listed 5 herein, and which inhibit tumor growth or reduces tumor size may also be used. Antibodies having VH and VL chains comprising an amino acid sequence that is at least 90%, 95%, 98% or 99% identical to those of any of the anti-TIM-4 antibodies listed herein may be used.

Administration of a TIM-4 antagonist may be monitored by imaging with a phosphatidylserine (PS)-binding agent (e.g. annexin V peptides, TIM-4 binding fragments), 10 where the PS binding agent may be labeled for imaging, e.g. PET, SPECT, fluorescence, etc. The binding agent is brought into contact with the target tumor cells, and the presence of bound agent is indicative of PS being present and able to interact with TIM-4.

For example, Annexin V is a ubiquitous intracellular protein in humans that has a 15 nanomolar affinity for the membrane-bound constitutive anionic phospholipid phosphatidylserine (PS), which is selectively expressed on the surface of apoptotic or physiologically stressed cells. As such, radiolabeled forms of annexin V have been used in both animal models and human Phase I and Phase II trials for utilizing the tracer as an early surrogate marker of therapeutic efficacy (e.g., Blankenberg et al. Proc. Am. J. Thoracic. Soc. 2009; 6:469-476). For example, the *in vivo* monitoring of PS within the tumor 20 microenvironment may be performed with anti-annexin V linked to a label radiotracer (for PET or SPECT), or with a TIM-4 fusion protein linked to a label or radiotracer. Labels could include q-dots for near IR imaging, or other more conventional labels. pK/Kd studies may be performed with radiolabeled antibodies.

IV. Compositions

25 In one aspect, the present invention provides composition comprising a PD-1 antagonist and a TIM-4 antagonist (e.g., formulated together in a single composition or separately formulated). In one embodiment, the PD-1 antagonist is nivolumab, pembrolizumab, durvalumab, atezolizumab, avelumab or PDR001. In one embodiment, the composition comprises a PD-1 antagonist and a TIM-4 antagonist, wherein (a) the PD-1 30 antagonist is an anti-PD-1 antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 13, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in

SEQ ID NO: 15; and (b) the TIM-4 antagonist is an antibody. In another embodiment, the composition comprises a PD-1 antagonist and a TIM-4 antagonist, wherein (a) the PD-1 antagonist is an anti-PD-L1 antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 1, and the CDR1, 5 CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 3 and (b) the TIM-4 antagonist is an antibody.

Pharmaceutical compositions suitable for administration to human patients are typically formulated for parenteral administration, e.g., in a liquid carrier, or suitable for reconstitution into liquid solution or suspension for intravenous administration.

10 In general, such compositions typically comprise a pharmaceutically acceptable carrier. As used herein, the term "pharmaceutically acceptable" means approved by a government regulatory agency or listed in the U.S. Pharmacopeia or another generally recognized pharmacopeia for use in animals, particularly in humans. The term "carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the compound is administered. Such 15 pharmaceutical carriers can be sterile 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, glycerol polyethylene glycol ricinoleate, and the like. Water or aqueous solution saline and aqueous dextrose and glycerol solutions may be employed as carriers, particularly for injectable solutions. Liquid compositions for parenteral administration can be formulated 20 for administration by injection or continuous infusion. Routes of administration by injection or infusion include intravenous, intraperitoneal, intramuscular, intrathecal and subcutaneous.

For oral use, the pharmaceutical compositions of the present invention, may be administered, for example, in the form of tablets or capsules, powders, dispersible granules, or cachets, or as aqueous solutions or suspensions. In the case of tablets for oral use, carriers 25 which are commonly used include lactose, corn starch, magnesium carbonate, talc, and sugar, and lubricating agents such as magnesium stearate are commonly added. For oral administration in capsule form, useful carriers include lactose, corn starch, magnesium carbonate, talc, and sugar. When aqueous suspensions are used for oral administration, emulsifying and/or suspending agents are commonly added.

30 In addition, sweetening and/or flavoring agents may be added to the oral compositions. For intramuscular, intraperitoneal, subcutaneous and intravenous use, sterile solutions of the active ingredient(s) are usually employed, and the pH of the solutions should

be suitably adjusted and buffered. For intravenous use, the total concentration of the solute(s) should be controlled in order to render the preparation isotonic.

For preparing suppositories according to the invention, a low melting wax such as a mixture of fatty acid glycerides or cocoa butter is first melted, and the active ingredient is 5 dispersed homogeneously in the wax, for example by stirring. The molten homogeneous mixture is then poured into conveniently sized molds and allowed to cool and thereby solidify.

Liquid preparations include solutions, suspensions and emulsions. Such preparations are exemplified by water or water/propylene glycol solutions for parenteral injection. Liquid 10 preparations may also include solutions for intranasal administration.

Aerosol preparations suitable for inhalation may include solutions and solids in powder form, which may be in combination with a pharmaceutically acceptable carrier, such as an inert compressed gas.

Also included are solid preparations which are intended for conversion, shortly before 15 use, to liquid preparations for either oral or parenteral administration. Such liquid forms include solutions, suspensions and emulsions.

V. Patient Populations

Provided herein are effective methods for treating cancer in a patient, e.g., using a combination of a TIM-4 antagonist and PD-1 antagonist. In one embodiment, the patient 20 suffers from a cancer selected from the group consisting of carcinoma, sarcoma, blastoma, leukemia and lymphoma. In another embodiment, the patient suffers from a cancer selected from the group consisting of non-small cell lung cancer, small cell lung cancer, gastrointestinal cancer, colorectal cancer, stomach cancer, colon carcinoma and glioblastoma. In another embodiment, the patient suffers from a cancer selected from the 25 group consisting of chronic myeloid leukemia, acute lymphoblastic leukemia, Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), squamous cell carcinoma, small-cell lung cancer, non-small cell lung cancer, glioma, gastrointestinal cancer, renal cancer, ovarian cancer, liver cancer, endometrial cancer, kidney cancer, prostate cancer, thyroid cancer, neuroblastoma, pancreatic cancer, glioblastoma multiforme, cervical cancer, 30 stomach cancer, bladder cancer, hepatoma, breast cancer, colon carcinoma, and head and neck cancer, gastric cancer, germ cell tumor, pediatric sarcoma, sinonasal natural killer,

multiple myeloma, acute myelogenous leukemia (AML), and chronic lymphocytic leukemia (CML).

VI. Additional Agents/Therapies

The combinations of the present invention (e.g., PD-1 antagonist in combination with 5 TIM-4 antagonist) may also be used in conjunction with other well known therapies that are selected for their particular usefulness against the cancer that is being treated. Combinations of the instant invention may alternatively be used sequentially with known pharmaceutically acceptable agent(s) when inappropriate.

For example, the PD-1 antagonists and TIM-4 antagonist described herein can further 10 be used in combination (e.g., simultaneously or separately) with an additional treatment, such as irradiation, chemotherapy (e.g., using camptothecin (CPT-11), 5-fluorouracil (5-FU), cisplatin, doxorubicin, irinotecan, paclitaxel, gemcitabine, cisplatin, paclitaxel, doxorubicin, 5-fu, or camptothecin + apo2l/TRAIL (a 6X combo)), one or more proteasome inhibitors (e.g., bortezomib or MG132), one or more Bcl-2 inhibitors (e.g., BH3I-2' (bcl-xl inhibitor), 15 AT-101 (R-(-)-gossypol derivative), ABT-263 (small molecule), GX-15-070 (obatoclax), or MCL-1 (myeloid leukemia cell differentiation protein-1) antagonists), iAP (inhibitor of apoptosis protein) antagonists (e.g., smac7, smac4, small molecule smac mimetic, synthetic smac peptides (see Fulda *et al.*, *Nat Med* 2002;8:808-15), ISIS23722 (LY2181308), or AEG- 35156 (GEM-640)), HDAC (histone deacetylase) inhibitors, anti-CD20 antibodies (e.g., 20 rituximab), angiogenesis inhibitors (e.g., bevacizumab), anti-angiogenic agents targeting VEGF and VEGFR, synthetic triterpenoids (see Hyer *et al.*, *Cancer Research* 2005;65:4799- 808), c-FLIP (cellular FLICE-inhibitory protein) modulators (e.g., natural and synthetic ligands of PPAR γ (peroxisome proliferator-activated receptor γ), 5809354 or 5569100), kinase inhibitors (e.g., Sorafenib), and/or genotoxic drugs.

25 The PD-1 antagonists and TIM-4 antagonists described herein can further be used in combination with one or more anti-proliferative cytotoxic agents. Classes of compounds that may be used as anti-proliferative cytotoxic agents include, but are not limited to, the following:

Alkylating agents (including, without limitation, nitrogen mustards, ethylenimine 30 derivatives, alkyl sulfonates, nitrosoureas and triazenes): Uracil mustard, Chlormethine, Cyclophosphamide (CYTOXANTM) fosfamide, Melphalan, Chlorambucil, Pipobroman,

Triethylenemelamine, Triethylenethiophosphoramine, Busulfan, Carmustine, Lomustine, Streptozocin, Dacarbazine, and Temozolomide.

Antimetabolites (including, without limitation, folic acid antagonists, pyrimidine analogs, purine analogs and adenosine deaminase inhibitors): Methotrexate, 5-Fluorouracil,

5 Floxuridine, Cytarabine, 6-Mercaptopurine, 6-Thioguanine, Fludarabine phosphate, Pentostatine, and Gemcitabine.

Suitable anti-proliferative agents for use in the methods of the invention, include, without limitation, taxanes, paclitaxel (paclitaxel is commercially available as

TAXOL.RTM.), docetaxel, discodermolide (DDM), dictyostatin (DCT), Peloruside A, 10 epothilones, epothilone A, epothilone B, epothilone C, epothilone D, epothilone E, epothilone F, furanoepothilone D, desoxyepothilone B1, [17]-dehydrodesoxyepothilone B, [18]dehydrodesoxyepothilones B, C12,13-cyclopropyl-epothilone A, C6-C8 bridged epothilone A, trans-9,10-dehydroepothilone D, cis-9,10-dehydroepothilone D, 16-desmethyllepothilone B, epothilone B10, discoderomolide, patupilone (EPO-906), KOS-862, 15 KOS-1584, ZK-EPO, ABJ-789, XAA296A (Discodermolide), TZT-1027 (soblidotin), ILX-651 (tasidotin hydrochloride), Halichondrin B, Eribulin mesylate (E-7389), Hemiasterlin (HTI-286), E-7974, Cyrtophycins, LY-355703, Maytansinoid immunoconjugates (DM-1), MKC-1, ABT-751, T1-38067, T-900607, SB-715992 (ispinesib), SB-743921, MK-0731, STA-5312, eleutherobin, 17beta-acetoxy-2-ethoxy-6-oxo-B-homo-estra-1,3,5(10)-trien-3-ol, 20 cyclostreptin, isolaulimalide, laulimalide, 4-epi-7-dehydroxy-14,16-didemethyl-(+)-discodermolides, and cryptothilone 1, in addition to other microtubuline stabilizing agents known in the art.

In cases where it is desirable to render aberrantly proliferative cells quiescent in conjunction with or prior to treatment with the chemotherapeutic methods of the invention,

25 hormones and steroids (including synthetic analogs), such as 17a-Ethinylestradiol, Diethylstilbestrol, Testosterone, Prednisone, Fluoxymesterone, Dromostanolone propionate, Testolactone, Megestrolacetate, Methylprednisolone, Methyl-testosterone, Prednisolone, Triamcinolone, Chlorotrianisene, Hydroxyprogesterone, Aminoglutethimide, Estramustine, Medroxyprogesteroneacetate, Leuprolide, Flutamide, Toremifene, ZOLADEX™, can also be 30 administered to the patient. When employing the methods or compositions of the present invention, other agents used in the modulation of tumor growth or metastasis in a clinical setting, such as antimimetics, can also be administered as desired.

Methods for the safe and effective administration of chemotherapeutic agents are known to those skilled in the art. In addition, their administration is described in the standard literature. For example, the administration of many of the chemotherapeutic agents is described in the Physicians' Desk Reference (PDR), e.g., 1996 edition (Medical Economics Company, Montvale, N.J. 07645-1742, USA); the disclosure of which is incorporated herein by reference thereto.

A PD-1 antagonist and a TIM4 antagonist may also be combined with one or more immunotherapy agent that stimulates the immune system, e.g., an agent that binds to human CTLA-4, LAG-3, GITR, OX40, IDO, CSF-1R etc.

The chemotherapeutic agent(s) and/or radiation therapy can be administered according to therapeutic protocols well known in the art. It will be apparent to those skilled in the art that the administration of the chemotherapeutic agent(s) and/or radiation therapy can be varied depending on the disease being treated and the known effects of the chemotherapeutic agent(s) and/or radiation therapy on that disease. Also, in accordance with the knowledge of the skilled clinician, the therapeutic protocols (e.g., dosage amounts and times of administration) can be varied in view of the observed effects of the administered therapeutic agents on the patient, and in view of the observed responses of the disease to the administered therapeutic agents.

VII. Treatment Protocols

Suitable treatment protocols for treating cancer in a patient include, for example, administering to the patient an effective amount of a PD-1 antagonist (e.g., antibody) and a TIM-4 antagonist (e.g., antibody).

As used herein, adjunctive or combined administration (co-administration) includes simultaneous administration of the two antagonists in the same or different dosage form, or separate administration of the two antagonist (e.g., sequential administration). Thus, the PD-1 antagonist (e.g., antibody) and TIM-4 antagonist (e.g., antibody) can be simultaneously administered in a single formulation. Alternatively, the PD-1 antagonist and TIM-4 antagonist can be formulated for separate administration and are administered concurrently or sequentially.

For example, the PD-1 antagonist can be administered first followed by (e.g., immediately followed by) the administration of the TIM-4 antagonist, or vice versa. In one embodiment, the PD-1 antagonist is administered prior to administration of the TIM-4

antagonist, e.g., the PD-1 antagonist is infused into the patient first, followed from 10 minutes to 3 hours later by an infusion of the TIM-4 antagonist. In one embodiment, the TIM-4 antagonist is administered prior to administration of the PD-1 antagonist, e.g., the TIM-4 antagonist is infused into the patient first, followed from 10 minutes to 3 hours later by an infusion of the PD-1 antagonist. Such concurrent or sequential administration preferably results in both antagonists being simultaneously present in treated patients. In another embodiment, the TIM-4 antagonist and the PD-1 antagonist are administered simultaneously.

In one embodiment, a subject is administered a single dose of a TIM-4 antagonist and a single dose of the PD-1 antagonist, e.g., an anti-PD-1 or anti-PD-L1 antibody. In certain embodiments, multiple (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10 or more) doses of a TIM-4 antagonist and multiple (e.g., 2, 3, 4, 5, 6, 7, 8, 9, 10 or more) doses of a PD-1 antagonist are administered to a subject in need of treatment. Administration of the TIM-4 antagonist and the PD-1 antagonist may be on the same day, or alternatively, the TIM-4 antagonist may be administered 1 or more days before or after the PD-1 antagonist.

In one embodiment, administrations of a TIM-4 antagonist and a PD-1 antagonist may be done weekly or monthly, in which regimen, they may be administered on the same day (e.g., simultaneously), or one after the other (e.g., one or more minutes, hours or days before or after one another). In one embodiment, the TIM-4 antagonist and PD-1 antagonist are administered every three days.

In one embodiment, the dose of the PD-1 antagonist and/or TIM-4 antagonist is varied over time. For example, the PD-1 antagonist and/or TIM-4 antagonist may be initially administered at a high dose and may be lowered over time. In another embodiment, the PD-1 antagonist and/or TIM-4 antagonist is initially administered at a low dose and increased over time.

In another embodiment, the amount of the PD-1 antagonist and/or TIM-4 antagonist administered is constant for each dose. In another embodiment, the amount of the PD-1 antagonist and/or TIM-4 antagonist varies with each dose. For example, the maintenance (or follow-on) dose of the antagonist can be higher or the same as the loading dose which is first administered. In another embodiment, the maintenance dose of the antagonist can be lower or the same as the loading dose. A clinician may utilize preferred dosages as warranted by the condition of the patient being treated. The dose may depend upon a number of factors,

including stage of disease, etc. The specific dose that should be administered based upon the presence of one or more of such factors is within the skill of the artisan. Generally, treatment is initiated with smaller dosages which are less than the optimum dose of the compound. Thereafter, the dosage is increased by small amounts until the optimum effect under the 5 circumstances is reached. For convenience, the total daily dosage may be divided and administered in portions during the day if desired. Intermittent therapy (e.g., one week out of three weeks or three out of four weeks) may also be used.

In one embodiment, the TIM-4 antagonist (e.g., antibody) is administered at a dose of 0.1, 0.3, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 mg/kg body weight. In another embodiment, the PD-1 10 antagonist (e.g., antibody) is administered at a dose of 0.5, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 mg/kg body weight. Generally, 200 μ g/mouse is approximately 10 mg/kg and 100 μ g/mouse is approximately 5 mg/kg. Therefore, based on the experiments described herein, one or more doses of 1-20 mg/kg body weight, 1-10 mg/kg body weight, 5-20 mg/kg body weight or 5-10 mg/kg body weight of a TIM-4 antagonist and PD-1 antagonist may be administered to a 15 subject. In certain embodiments, a dose of 0.3 mg/kg to 10 mg/kg body weight of a TIM-4 antagonist is used and a dose of at least 1 mg/kg, e.g., 1-10 mg/kg body weight of a PD-1 antagonist is used.

VIII. Outcomes

Patients, e.g., humans, treated according to the methods disclosed herein preferably 20 experience improvement in at least one sign of cancer. In one embodiment, improvement is measured by a reduction in the quantity and/or size of measurable tumor lesions. In another embodiment, lesions can be measured on chest x-rays or CT or MRI films. In another embodiment, cytology or histology can be used to evaluate responsiveness to a therapy.

In one embodiment, the patient treated exhibits a reduction in size of a tumor, 25 reduction in number of metastatic lesions over time, complete response, partial response, and stable disease. In another embodiment, the patient treated experiences tumor shrinkage and/or decrease in growth rate, i.e., suppression of tumor growth. In another embodiment, unwanted cell proliferation is reduced or inhibited. In yet another embodiment, one or more of the following can occur: the number of cancer cells can be reduced; tumor size can be reduced; 30 cancer cell infiltration into peripheral organs can be inhibited, retarded, slowed, or stopped; tumor metastasis can be slowed or inhibited; tumor growth can be inhibited; recurrence of

tumor can be prevented or delayed; one or more of the symptoms associated with cancer can be relieved to some extent.

5 In another embodiment, the methods of treatment produce a comparable clinical benefit rate (CBR = CR (complete response), PR (partial response) or SD (stable disease) ≥ 6 months) better than that achieved by a PD-1 (e.g., antibody) or TIM-4 antagonist (e.g., antibody) alone. In other embodiments, the improvement of clinical benefit rate is about 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80% or more, e.g., compared to treatment with a PD-1 antagonist or TIM-4 antagonist alone or relative to tumor growth on the first day of treatment or immediately before initiation of treatment.

10 10 In another embodiment, administration of a PD-1 antagonist and a TIM-4 antagonist results in at least a three-fold reduction (e.g., a 3.5-fold reduction) in tumor volume, e.g., relative to treatment with the PD-1 antagonist or the TIM-4 antagonist alone or relative to tumor growth on the first day of treatment or immediately before initiation of treatment.

15 15 In a further embodiment, administration of a PD-1 antagonist and a TIM-4 antagonist results in tumor growth inhibition of at least 80%, e.g., relative to treatment with the PD-1 antagonist or TIM-4 antagonist alone or relative to tumor growth on the first day of treatment or immediately before initiation of treatment.

20 20 In certain embodiments, administration of a PD-1 antagonist and a TIM-4 antagonist reduces tumor mass by at least 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 95%, 99% relative to the tumor mass prior to initiation of the treatment or on the first day of treatment. In some embodiment, the tumor mass is no longer detectable following treatment as described herein. In some embodiments, a subject is in partial or full remission. In certain embodiments, a subject has an increased overall survival, median survival rate, and/or progression free survival.

25 **IX. Kits and Unit Dosage Forms**

Also provided herein are kits which include a pharmaceutical composition containing (a) a PD-1 antagonist and (b) a TIM-4 antagonist and a pharmaceutically-acceptable carrier, in a therapeutically effective amount adapted for use in the preceding methods. In one embodiment, the PD-1 antagonist is an antibody (e.g., 5C4 or 12A4, respectively). In another 30 embodiment, the TIM-4 antagonist is an antibody. The kits optionally also can include instructions, e.g., comprising administration schedules, to allow a practitioner (e.g., a

physician, nurse, or patient) to administer the composition contained therein to a patient having cancer. The kit also can include a syringe.

Optionally, the kits include multiple packages of the single-dose pharmaceutical compositions each containing an effective amount of the PD-1 antagonist and the TIM-4 antagonist for a single administration in accordance with the methods provided above. Instruments or devices necessary for administering the pharmaceutical composition(s) also may be included in the kits. For instance, a kit may provide one or more pre-filled syringes containing an amount of the PD-1 antagonist and the TIM-4 antagonist.

In one embodiment, the present invention provides a kit for treating cancer in a patient, the kit comprising:

- (a) a dose of a PD-1 antagonist;
- (b) a dose of a TIM-4 antagonist; and
- (c) instructions for using the PD-1 antagonist and TIM-4 antagonist in the methods described herein.

In one embodiment, the present invention provides a kit for treating cancer in a patient, the kit comprising:

- (a) one or more doses of a PD-1 antagonist;
- (b) one or more doses of a TIM-4 antagonist; and
- (c) instructions for using the PD-1 antagonist and TIM-4 antagonist in the methods described herein.

In certain embodiments, the TIM-4 antagonist is an antibody. In certain embodiments, the PD-1 antagonist is an antibody. In particular embodiments, the PD-1 antagonist is an anti-PD-1 antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 13, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 15. In another particular embodiment, the PD-1 antagonist is an anti-PD-L1 antibody comprising antibody comprises the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 1, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 3.

The following examples are merely illustrative and should not be construed as limiting the scope of this disclosure in any way as many variations and equivalents will become apparent to those skilled in the art upon reading the present disclosure.

The contents of all references, Genbank entries, patents and published patent 5 applications cited throughout this application are expressly incorporated herein by reference.

EXAMPLES

Materials and Methods

Animals

10 Ten to eleven-week-old female C57/BL6 mice (Harlan) were used in the studies. Mice received food and water *ad libitum* and were maintained in a controlled environment according to Association for Assessment and Accreditation of Laboratory Animal Care (AAALAC) International regulations. All animal studies have been approved by the appropriate ethics committee and have therefore been performed in accordance with the 15 ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments.

Antibodies

Anti-mouse PD-1 mAb (anti-mPD-1 mAb) clone 4H2, mouse IgG1 isotype was produced and purified by Bristol-Myers Squibb (Biologics Discovery, CA). The anti-TIM-4 mAb, RMT4-53 has been previously described (Yeung *et al.*, *J. Immunol.* 2009; 191:4447-20 4455). Briefly, Sprague Dawley rats were immunized with a TIM-4-Ig fusion protein containing the extracellular domain of murine TIM-4 (aa 1-288) to the Fc portion of mouse IgG2a, and fusing LN cells with P3U1 myeloma cells. RMT4-53 reacts with TIM-4/NRK cells but not with parental NRK or other transfected TIM family (TIM-1 B6/NRK, TIM-1 BALB/NRK, TIM-2/NRK, TIM-3 B6/NRK, TIM-3 BALB/NRK) cells. Antibodies were 25 certified to have <0.5 EU/mg endotoxin levels, >95% purity and <5% high molecular weight species. Stock solutions of anti-mPD-1 mAb (clone 4H2; mouse IgG1) and anti-mTIM-4 antibody (clone RM 4-53; rat IgG2a), were kept at 4°C prior to use. Dosing solutions of anti-mPD-1 mAb and anti-mTIM-4 mAb were prepared in sterile phosphate buffered saline (pH 7.0) and maintained at 4°C.

Cell Lines

CT-26 is an undifferentiated colon carcinoma cell line with a fibroblast morphology (ATCC). Mice inoculated subcutaneously develop lethal tumors at 80% frequency with 10^3 cells and at 100% with 10^4 cells. Pulmonary metastases develop when mice are inoculated, 5 intravenously, with 10^4 cells (Wang et al. *J. Immunol.* 1995; 154:4685-4692). The MC38 cell line was derived from C57BL6 murine colon adenocarcinoma cells (CD44+ALDH1+) with an epithelial morphology.

Expression of TIM-4

Expression of TIM-4 on mouse peritoneal macrophages and TAMs was confirmed by 10 flow cytometry.

Immunohistochemistry

Immunohistochemical analysis using Rabbit anti-TIM4 (Atlas Antibodies, HPA015625) demonstrated expression of TIM-4 on tissue resident macrophages present in human tonsil, lung and liver. Expression was demonstrated on fixed tissue was sectioned and 15 applied to Superfrost Plus glass slides at a thickness of 5 microns. Sections were allowed to dry overnight at room temperature. Tissues were then deparaffinized in xylene followed by hydration to distilled water in graded series of ethanol. Antigen Retrieval was performed using HIER with Biocare Medical EDTA, pH 8.2 (1X) at approximately 115°C for 1 minute and then cooled for 20 minutes at RT. Sections were rinsed in TNT Buffer (TBS + 0.5% 20 Tween) and isolated using a PAP pen. Blocking was performed with Biocare Medical Background Sniper for 30 minutes at RT, followed by 1X TNT Buffer rinse, incubation in Dako Peroxidase Block for 10 minutes at RT and 2X TNT Buffer rinses. Primary Antibody staining was performed using a 1/1000 dilution in Biocare Medical Renaissance antibody diluent and background reducer for 60 minutes at RT and 3X TNT Buffer rinses. Detection 25 antibody was applied using appropriate number of drops to cover section with Biocare Medical MACH3 Rabbit polymer-HRP and incubating for 30 minutes at RT followed by and 3X TNT Buffer rinses. Samples were incubated with Biocare Medical Betazoid DAB and reaction stopped by placing slides in water). Sections were counterstained in Hematoxylin, Leica for 30 seconds, washed in tap water, incubated in Bluing Solution for 45 seconds to 1 30 minute, washed 2X in water, dehydrated in graded series of alcohol, and Xylene and then mounted.

Example 1: Inhibition of Efferocytosis by anti-TIM-4 Antibody

Thymocytes were isolated from BALB/c mice and cultured with or without dexamethasone for 4 hours to induce apoptosis. Cells were then incubated with pHrodo dye for 5-10 minutes according to manufacturer's instructions (Thermofisher Scientific). Briefly, 5 2×10^7 cells were washed in media without serum 1 x and resuspended in 1 mL of Diluent C (Catalog Number G8278). Cells were then stained by adding 1 mL of 2x dye solution (by adding 4 mL of the PKH26 ethanolic dye solution (Catalog Number P9691) to 1 mL of Diluent C) and mixing sample by pipetting. Incubation was for 5 minutes with periodic mixing. Staining reaction was stopped by adding an equal volume of serum and incubating 1 10 minute. Cells were then washed 1 x in media without serum and 2x in PBS. 3×10^6 pHrodo-labeled cells were injected i.p. into BALB/c mice and peritoneal macrophages were collected by lavage by injecting 3 - 5 mL of ice cold PBS+ 2mM EDTA into the peritoneal cavity of euthanized mice, palpating the abdomen and then collecting the solution via needle aspiration. Experimental animals were injected with RMT4-53 anti-TIM4 antibody 1 hour 15 prior to injection of the pHrodo-labeled thymocytes T.

The uptake of apoptotic cells by peritoneal macrophages was measured using flow cytometry. Briefly, peritoneal macrophages were washed 2x in PBS, resuspended in 100 μ L viability dye (eBioscience) and incubated on ice 15 minutes. Cells were washed 1 x in PBS and incubated in FcBlock (BioLegend) for 15 minutes on ice. Cells were then stained with 20 antibodies to CD45 (BD), Tim-4 (BioLegend), CD 11b (BioLegend), CD 206 (BioLegend), and F4/80+ (BioLegend), for 30 minutes on ice. Cells were then washed 1 x and resuspended in Perm/Fix. Samples were then collected on a LSR Fortessa X20 (BD). To identify cells that engaged in efferocytosis, CD45 +/CD11b+ myeloid cells from the live cell population were gated on. Out of these cells, macrophages that engaged in efferocytosis were defined as 25 F4/80+, OPKH26+. The results are set forth in Table 2.

Table 2.

	% Apoptotic Cells
Control	25-40
Control + Dex	50-70
Experimental + anti-TIM4	10-15
Experimental + Dex and TIM-4	22-42

The results demonstrate an increase in efferocytosis of apoptotic cells (50-70%) relative to non-apoptotic cells (25-40%). The addition of anti-TIM-4 blocking anybody greatly reduced uptake of both viable and apoptotic cells. This demonstrates that TIM-4 is an important mediator of cellular uptake.

5

Example 2: Inhibition of Tumor Growth *In Vivo* in the CT26 model by Combination Treatment with Anti-TIM-4 Antibody and Anti-PD-1 Antibody

An experiment was conducted in a murine tumor model to test the hypothesis that the combination of anti-TIM-4 and anti-PD-1 would potentiate anti-tumor efficacy. SC CT-26 10 mice, a colon adenocarcinoma tumor model (TGM-1438), were evaluated for tumor growth after treatment with the anti-TIM4 antibody, RTM-453, alone or in combination with an anti-PD1 antibody, IgG1 D265A.

Female BALB/c mice (Harlan; approximately 8-9 wk old) were subcutaneously implanted with 10^6 CT-26 cells on day 0. Sixty mice were evaluated by body weight and 15 tumor measurements after 2X/wk dosing according to the following schedule:

Table 3: Dosing Schedule:

Group	N	Treatment	Dosing Regimen
1	15	mIgG1 isotype control mAb	0.45 mg, IP, d. 7, 10, 13
2	15	mPD-1 IgG1 D265A mAb + Control mAb	0.2 + 0.25* mg, IP, d. 7, 10, 13
4	15	mTIM-4 rat IgG2a mAb + Control mAb	0.25 + 0.2* mg, IP, d. 7, 10, 13
6	15	mTIM-4 + mPD-1 mAbs	0.25 mg + 0.2 mg, IP, d. 7, 10, 13

On the days of dosing, the two antibodies were combined and a total of $100\mu\text{L}$ of combined antibody was injected into the mice. Spleens and tumors from five mice per group 20 were collected and processed for flow cytometry analysis of tumor infiltrating lymphocytes (TILs), and spleen immune cells on day 16 according to the following protocol:

Briefly, cells were resuspended in PBS and aliquoted into plates (spleens $2 \times 10^6/\text{well}$, all tumor cells/well) and then were washed 2x in PBS, backspace. Cells were then resuspended in $100 \mu\text{L}$ viability dye (eBioscience) and incubated on ice 15 minutes. Cells 25 were washed 1 x in PBS and incubated in $50 \mu\text{L}$ FcBlock (BioLegend) for 15 minutes on ice

and then stained with 50 μ L of antibody stain mix as described below for 30 minutes. Cells were then washed 1 x with FACS buffer and resuspended in Perm/Fix. Samples were then collected on a LSR Fortessa X20 (BD).

T cell Panel

Fluor	Specificity	Clone	Antibody	Cat#	Company	Ab diln
AlexaFluor 488 (FITC)	CD107a	1D4B	Alexa Fluor® 488 anti-mouse CD107a (LAMP-1) Antibody	121608	BioLegend	1:100
PE	CD115	AFS98	PE anti-mouse CD115 (CSF-1R) Antibody	135506	BioLegend	1:100
PerCp-Cy5.5	CD49b	DX5	PerCp/Cy5.5 anti-mouse CD49b (pan-NK cells) antibody	108916	BioLegend	1:100
BUV 395	CD45	30-F11	BUV395 rat anti-mouse CD45 antibody	564279	eBioscience	1:100
BV 421	CD8	53-6.7	Brilliant Violet 421™ anti-mouse CD8a antibody	100738	BioLegend	1:200
BV 510	CD4	RM4-5	Brilliant Violet 510™ anti-mouse CD4 antibody	100553	BioLegend	1:200
BV 605	Thy1.2	53-2.1	Brilliant Violet 605™ anti-mouse CD90.2 (Thy-1.2) antibody	140317	BioLegend	1:200
BV 711	CD62L	MEL-14	Brilliant Violet 711™ anti-mouse CD62L antibody	104445	BioLegend	1:200
BV 785	CD44	IM7	Brilliant Violet 785™ anti-mouse CD44 antibody	103041	BioLegend	1:200
APC	FoxP3	FJK-16s	Anti-Mouse/Rat Foxp3 APC	17-5773-80B	eBioscience	1:100
PE-Cy7	Ki-67	Sola15	Anti-Mouse/Rat Ki-67 PE Cyanine7	25-5698-82	eBioscience	1:100
APC eFluor 780 (APC-Cy7)	Fixable Viability Dye	n/a	eFluor® 780 Fixable Viability Dye	65-0865-14	eBioscience	1:1000
Fc Block	block	93	TruStain FcX anti-mouse CD16/32	101320	BioLegend	

5

Ki-67+ to assess proliferating cells;

CD44/CD62L analysis to determine subset T cells into naïve, activated, and memory T cell subsets;

FoxP3 to assess population of Tregs (FoxP3+CD4+ T cells);

10

CD8+CD107a+ to assess degranulating CD8+ T cells (Antigen/Tumor-specific CTL)

Myeloid Panel

Fluor	Specificity	Clone	Antibody	Cat#	Company	
AlexaFluor 488 (FITC)	CD11c	N418	FITC anti-mouse CD11c antibody	117306	BioLegend	1:100
PE	CD115	AFS98	PE anti-mouse CD115 (CSF-1R) Antibody	135506	BioLegend	1:100
PerCp-Cy5.5	CD49b	DX5	PerCp/Cy5.5 anti-mouse CD49b (pan-NK cells) antibody	108916	BioLegend	1:100
BUV 395	CD45	30-F11	BUV395 rat anti-mouse CD45 antibody	564279	eBioscience	1:100
BV 421	Ly6C	HK1.4	Brilliant Violet 421™ anti-mouse Ly6C antibody	128032	BioLegend	1:200
BV 605	Thy 1.2 (CD90.2)	53-2.1	Brilliant Violet 605™ anti-mouse Thy1.2 antibody	140317	BioLegend	1:200
BV 711	Gr-1	RB6-8C5	Brilliant Violet 711™ anti-mouse Gr-1 antibody	104731	BioLegend	1:200
BV 785	F4-80	BM8	Brilliant Violet 785™ anti-mouse F4/80 antibody	123141	BioLegend	1:200
APC	Ly6G	1A8	APC anti-mouse Ly-6G antibody	127614	BioLegend	1:200
AF-700	CD11b	M1/70	Alexa Fluor 700 anti-mouse/human CD11b antibody	101222	BioLegend	1:100
PE-Cy7	MHC II	M5/114.15.2	PE-Cy7 anti-mouse I-A/I-E antibody	107630	eBioscience	1:100
APC eFluor 780 (APC-Cy7)	Fixable Viability Dye	n/a	eFluor® 780 Fixable Viability Dye	65-0865-14	eBioscience	1:1000
Fc Block	block	93	TruStain FcX anti-mouse CD16/32	101320	BioLegend	

Myeloid Cell subsetting for TAMs, MDSCs, NK cells, B cells

15

CD80, CD86 staining for activated APCs

The myeloid populations in spleen and tumors are shown in **Figure 3**. The CD8+ T cell populations in spleen and tumor infiltrating lymphocytes are shown in **Figure 4**. The CD4+ cell populations in spleen and tumor infiltrating lymphocytes are shown in **Figure 5**.

The data demonstrate a shift in myeloid cell populations, and an increase in activated T cells in tumor infiltrating lymphocytes.

The remaining ten mice per group were monitored for tumor growth. Tumor size and body weights were measured twice weekly. Tumor size (measured as mm³) was calculated by 5 multiplying the tumor length by the square of the tumor width divided by 2. Treatments were initiated when subcutaneous tumors reached a median size of 200 mm³ (established model).

Anti-TIM-4 antibody administered in combination with anti-PD-1 antibody provided enhanced anti-tumor activity above the activity of either agent alone. The combination demonstrated decreased tumor growth rates (**Figures 6-8**), and an increased overall survival 10 rate (**Figure 9**). In sum, the combination of the TIM-4 mAb and the PD-1 mAb resulted in synergistic activity compared to the activity elicited by single agents alone. Therefore, results from this study demonstrate that a combination regimen of anti-mTIM-4 mAb and PD-1 mAb is well-tolerated and result in marked antitumor activity.

15 **Example 3: Inhibition of Tumor Growth *In Vivo* in the MC38 model by Combination Treatment with Anti-TIM-4 Antibody and Anti-PD-1 Antibody**

MC38 mice, a colon adenocarcinoma tumor model, were evaluated for tumor growth after treatment with the anti-TIM4 antibody, RTM-453, alone or in combination with an anti-PD1 antibody, IgG1 D265A.

20 Female C57/BL6 mice (Harlan; approximately 8-9 wk old) were subcutaneously implanted with 10⁶ MC38 cells on day 0. Mice were dosed beginning on Day 6. Three doses at 200 μ g/injection were administered IP every 4th day. When the two antibodies were administered to a mouse, the antibodies were first combined, and administered together into the mouse. Tumors were measured using calipers and tumor volumes calculated using the 25 formula (L²xW)/2. Progression Free Survival defined as # of days for tumor to reach 4x initial tumor volume.

The results are provided in Figures 10, 11 and 12, and Table X. Figure 10, which provides the percentage of progression free survival relative to the days post implant, show that administration of the combination of the anti-PD-1 and anti-TIM4 antibodies results in a 30 higher percentage of progression free survival relative to each antibody separately and relative to the isotype control. Table X shows that the combination treatment provided more

complete regressions relative to each antibody separately and relative to the isotype control. Figure 11, which shows the MC38 mean tumor volume relative to the days post implant, indicates that the combination treatment reduced the mean tumor volume more relative to each antibody separately and relative to the isotype control. Figure 12A-C shows the MC38 individual tumor volumes, and confirms the results shown in the Figures 10, 11 and Table X.

Table 5: Number of mice with complete regressions of tumors

Treatment	Complete Regressions
Isotype Control	0/10
Anti-PD1 Alone	2/10
anti-TIM4 (RMT4-53 mIgG1 D265A) Alone	0/10
Anti-PD1 + anti-TIM4 (RMT4-53 mIgG1 D265A)	4/10

Thus, administration of a combination of an anti-TIM4 antibody with an anti-PD-1 antibody in the CT26 and MC39 animal models resulted in a stronger anti-tumor effect relative to each antibody alone.

SUMMARY OF THE SEQUENCE LISTING

SEQ ID NO:	SEQUENCE
1	Heavy Chain Variable Region (VH) Amino Acid Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) QVQLVQSGAEVKPGSSVKVSCKTSGDTFSTYAI SWVRQAPGQGLEWMGGII PIFGKAHYAQKFQGRVTITADESTSTAYMELSSLRSEDTAVYFCARKFHFVS GSPFGMDVWGQGTTVTVSS
2	Heavy Chain Variable Region (VH) Nucleotide Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) cag gtc cag ctg gtg cag tct ggg gct gag gtg aag aag cct ggg tcc tcg gtg aag gtc tcc tgc aag act tct gga gac acc ttc agc acc tat gct atc agc tgg gtg cga cag gcc cct gga caa ggg ctt gag tgg atg gga ggg atc atc cct ata ttt ggt aaa gca cac tac gca cag aag ttc cag ggc aga gtc acg att acc gcg gac gaa tcc acg agc aca gcc tac atg gag ctg agc agc ctg aga tct gag gac acg gcc gtg tat ttt tgt gcg aga aag ttt cac ttt gtt tcg ggg agc ccc ttc ggt atg gac gtc tgg ggc caa ggg acc acg gtc acc gtc tcc
3	Light Chain Variable Region (VL) Amino Acid Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLLIY DASNRATGIPARFSGSGSGTDFLTISLEPEDFAVYYCQQRSNWPTFG QGTKVEIK
4	Light Chain Variable Region (VL) Nucleotide Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) gaa att gtg ttg aca cag tct cca gcc acc ctg tct ttg tct cca ggg gaa aga gcc acc ctc tcc tgc agg gcc agt cag agt gtt agc agc tac tta gcc tgg tac caa cag aaa cct ggc cag gct ccc agg ctc ctc atc tat gat gca tcc aac agg gcc act ggc atc cca gcc agg ttc agt ggc agt ggg tct ggg aca gac ttc act ctc acc atc agc agc cta gag cct gaa gat ttt gca gtt tat tac tgt cag cag cgt agc aac tgg ccg acg ttc ggc caa ggg acc aag gtg gaa atc aaa
5	Heavy Chain CDR1 Amino Acid Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) TYAIS
6	Heavy Chain CDR2 Amino Acid Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) GIIPIFGKAHYAQKFQ

SEQ ID NO:	SEQUENCE
7	Heavy Chain CDR3 Amino Acid Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) KFHFVSGSPFGMDV
8	Light Chain CDR1 Amino Acid Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) RASQSVSSYLA
9	Light Chain CDR2 Amino Acid Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) DASN RAT
10	Light Chain CDR3 Amino Acid Sequence Anti-PD-L1 mAb (12A4; 12A4 in WO 2007/005874 and US Patent No. 7,943,743) QQRSNWPT
11	Heavy Chain Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (variable region underlined; constant region bold) <u>QVQLVESGGVVQPGRLRLDCKASGITFSNSGMHWVRQAPGKGL</u> <u>EWVAVIWYDGSKRYYADSVKGRFTISRDNSKNTLFLQMNSLRAEDT</u> <u>AVYYCATNDDYWQGTLVTVSS</u> ASTKGPSVFPLAPCSRSTSESTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAPLQSSGLYSISSLVVT VPSSSLGTTKTYTCNVDHKPNTKVDKRVESKYGPPCPCCPAPEFLG GPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGV EVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKG LPSSIEKTISAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPS DIAVEWESNGQPENNYKTPPVLDSDGSFFLYSRLTVDKSRWQEGNV FSCSVVMHEALHNHYTQKSLSLSLGK
12	Light Chain Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (variable region underlined; constant region bold) <u>EIVLTQSPATLSLSPGERATLSCRASQSVSSYLA</u> WYQQKPGQAPRLLI <u>YDASNRATGIPARFSGSGSGTDFTLT</u> ISSLEPEDFAVYYCQQSSNWPR TFGQGTKEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREA KVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTTLSKADYEKHK VYACEVTHQGLSSPVTKSFNRGEC
13	Heavy Chain Variable Region (VH) Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:4 from WO 2006/121168) QVQLVESGGVVQPGRLRLDCKASGITFSNSGMHWVRQAPGKGLEWVAV I WYDGSKRYYADSVKGRFTISRDNSKNTLFLQMNSLRAEDT AVYYCATND DYWGQGT LVTVSS
14	Heavy Chain Variable Region (VH) Nucleotide Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168)

SEQ ID NO:	SEQUENCE
	(SEQ ID NO:60 from WO 2006/121168) cag gtg cag ctg gtg gag tct ggg gga ggc gtg gtc cag cct ggg agg tcc ctg aga ctc gac tgt aaa gcg tct gga atc acc ttc agt aac tct ggc atg cac tgg gtc cgc cag gct cca ggc aag ggg ctg gag tgg gtg gca gtt att tgg tat gat gga agt aaa aga tac tat gca gac tcc gtg aag ggc cga ttc acc atc tcc aga gac aat tcc aag aac acg ctg ttt ctg caa atg aac agc ctg aga gcc gag gac acg gct gtg tat tac tgt gcg aca aac gac gac tac tgg ggc cag gga acc ctg gtc acc gtc tcc tca
15	Light Chain Variable Region (VL) Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:11 from WO 2006/121168) EIVLTQSPATLSLSPGERATLSCRASQSVSSYLA WYQQKPGQAPRLLIYD ASNRATGIPARFSGSGSGTDFLT TISSLEPEDFAVYYCQQSSNW PRTFGQ GTKVEIK
16	Light Chain Variable Region (VL) Nucleotide Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:67 from WO 2006/121168) gaa att gtg ttg aca cag tct cca gcc acc ctg tct ttg tct cca ggg gaa aga gcc acc ctc tcc tgc agg gcc agt cag agt gtt agt agt tac tta gcc tgg tac caa cag aaa cct ggc cag gct ccc agg ctc ctc atc tat gat gca tcc aac agg gcc act ggc atc cca gcc agg ttc agt ggc agt ggg tct ggg aca gac ttc act ctc acc atc agc agc cta gag cct gaa gat ttt gca gtt tat tac tgt cag cag agt agc aac tgg cct cgg acg ttc ggc caa ggg acc aag gtg gaa atc aaa
17	Heavy Chain CDR1 Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:18 from WO 2006/121168) NSGMH
18	Heavy Chain CDR2 Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:25 from WO 2006/121168) VIWYDGSKRYYADSVKG
19	Heavy Chain CDR3 Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:32 from WO 2006/121168) NDDY
20	Light Chain CDR1 Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:39 from WO 2006/121168) RASQSVSSYLA

SEQ ID NO:	SEQUENCE
21	Light Chain CDR2 Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:46 from WO 2006/121168) DASN RAT
22	Light Chain CDR3 Amino Acid Sequence Anti-PD-1 mAb (5C4 in WO 2006/121168) (SEQ ID NO:53 from WO 2006/121168) QQSSNWPRT
23	Complete PD-1 sequence (GenBank Accession No.: U64863) agttccctt ccgctcacct ccgcctgagc agtggagaag gcggcactct ggtggggctg ctccaggcat gcagatccca caggccctt ggccagtcgt ctggcggtg ctacaactgg gctggggcc aggatggttc ttagactccc cagacaggcc ctggAACCCC cccacccct tcccagccct gctcgtgggt accgaagggg acaacgcccac cttcacctgc agcttctcca acacatcgg a gacttcgtg ctaaactggt accgcatgag ccccagcaac cagacggaca agctggccgc cttcccccgg gaccgcagcc agcccgccca ggactgcccgc ttccgtgtca cacaactgcc caacgggcgt gacttccaca tgagcgtggt cagggcccg cgcaatgaca gccgcaccta cctctgtggg gccatctccc tggcccccgg ggcgcagatc aaagagagcc tgcgggcaga gctcagggtt acagagagaa gggcagaatg gcccacagcc caccggcagcc cctcaccagg gccagccggc cagttccaaa ccctgggtggt tgggtctgtg ggccgcctgc tgggcagccctt ggtgctgtca gtctgggtcc tggccgtcat ctgctcccg gccgcacag ggacaatagg agccaggcgc accggccagc ccctgaaggg ggaccctca gccgtgcctg tggctctgtt ggactatggg gagctggatt tccagtggcg agagaagacc cccggagccccc ccgtgcctgt tggcccttag cagacggat atgcaccat tggctttctt agcggatgg gcacccatc ccccgcccg aggggcttag ccgacggccc tggagtgcc cagccactga ggcctgagga tggacactgc tcttggccccc tctgaccggc ttccttggcc accagtgttc tgcagaccct ccaccatgg cccgggttag cgcatttctt caggagaagc aggccagggtg caggccattt caggccgtcc aggggcttag ctgcctgggg ggcgcacgggg ctccagccct cacctgcacc aggcacagcc ccaccacagg actcatgtct caatgcccac agtgagccca ggcagcagggt gtcaccgtcc cctacaggga gggccagatg cagtcactgc ttcagggtc gccagcacag agctgcctgc gtccagctcc ctgaatctt gctgctgtc ctgctgtc tgctgctgtcc tgcggcccg ggctgaaggc gccgtggccc tgcctgacgc cccggagcc cctgcctgaa ctggggggctt gggtggagat ggccttggag cagccaagggt gcccctggca gtggcatccc gaaacggccc ggacgcaggg cccaaagactg ggcacaggag tgggagggtac atggggctgg ggactccccc ggagttatct gtcctctgca ggcttagaga agttttaggg aaggtcagaa gagctctgg ctgtgggtgg cagggcagga aaccctccc acctttacac atgcccaggc agcaccttag gccctttgtt gggcaggggaa gctgaggcg taagcggca ggcagagctg gaggccttcc aggcaccca gcaacttggc ctctgcgc cgcatttcc cccagcccttcc cacaccactc gggagaggga catcctacgg tcccaagggtc aggaggcg ggctgggtt gactcaggcc ctcccgatg gtggccaccc ggggtttggg agggcagaag tgcaggcacc tagggccccc catgtgccc ccctggagc tctccttggaa acccatttct gaaatttattt aaaggggtt gccggctcc caccaggggc tgggtggaa ggtacaggcg ttccccccgg gcttagtacc cccgcgtggc ctatccactc ctcacatcca cacactgcac cccacttcttcc gggcaggggc caccacatc caggcggccca gcaaggcactt gagtggtgg gacaaggat ccccttccc tgggttcta ttatattata attataatta aatatgagag catgtt
24	Human PD-L1 amino acid sequence - isoform a precursor (GenBank Accession No. NP_054862.1) MRIFAVFIFM TYWHLLNAFT VTVPKDLYVV EYGSNMTIEC KFPVEKQLDL AALIVYWEME DKNIIQFVHG EEDLKVKQHSS YRQRARLLKD QLSLGNAALQ ITDVKLQDAG VYRCMISYGG

SEQ ID NO:	SEQUENCE
	ADYKRITVKV NAPYNKINQR ILVVDPVTSE HELTCQAEGY PKAEVIWTSS DHQVLSGKTT TTNSKREEKL FNVTSRLRIN TTTNEIFYCT FRRLDPEENH TAEVVIPELP LAHPPNERTH LVILGAILLC LGVALTFIFR LRKGRRMMDVK KCGIQDTNSK KQSDTHLEET
25	Human PD-L1 amino acid sequence - isoform b precursor (GenBank Accession No. NP_001254635.1) MRIFAVFIFM TYWHLLNAPY NKNQRILVV DPVTSEHELT CQAEGYPKAE VIWTSSDHQV LSGKTTTNS KREEKLFNVT STLRINTTN EIFYCTFRRRL DPEENHTAEL VIPELPLAHP PNERTHLVIL GAILLCLGVA LTFIFRLRKG RMMDVKKCGI QDTNSKKQSD THLEET
26	Human TIM-4 amino acid sequence - isoform 1 (GenBank Accession No. NP_612388.2) MSKEPLILWLMIEFWWLYLTPVTSETVVTEVLGHRVTLPCLYSSWSHNSNSMCWGKDQCPYSGCKEALIR TDGMRVTSRKSAYRLQGTIPRGDVSLTILNPSESDSGVYCCRIEVPGWFNDVKINVRLNLQRASSTTHR TATTTT RRTTTSPTRQMTTPAALPTTVVTPDLTTGTPLQMTTIAVFTTANTCLSLTPSTLPEEAT GLLTPEPSKEGPILTAESETVLPDSWSSVESTSADTVLLTSKESKVWDLPSTS HVSMWKTS DSVSSPQP GASDTAVPEQNKTGQMDGIPMSMKNEMPISQLMIIAPSLGFVLFALFVAFLRGKLMETYCSQKHT RLDYIGDSKNVLNDVQHGRED EDEGLFTL
27	Homo sapiens T-cell (TIMD4), transcript variant 1, mRNA (GenBank Accession No. NM_138379.2) ATAAGAGGTTGGGCTTGGATAGATAGACAGACTCCTGGGTCAACCGTCAAAATGTCAAAGAAC CTCTCATTCTCTGGCTGATGATTGAGTTGGTGGCTTACCTGACACCAGTCACTCAGAGACTGTTG GACGGAGGTTGGGTACCCGGTGACTTGCCTGTCTGACTCATCCTGGTCTCACAAACAGAACAGC ATGTGCTGGGGAAAGACCACTGCCCCTACTCCGGTTGCAAGGAGGGCCTCATCCGACTGATGGAATGA GGGTGACCTCAAGAAAGTCAGCAAATATAGACTTCAGGGACTATCCGAGAGGTATGTCCTTGAC CATCTAAACCCAGTGAAAGTGACAGCGGTGTACTGCTGCCATAGAAGTGCCTGGCTGGTTCAAC GATGTAAAGATAAACGTGCGCCTGAATCTACAGAGACGCTAACAAACACAGCACAGAACAGAACACCA CCACACGCAGAACACAACAAGCCCCACCACCCGACAATGACAACAACCCAGCTGCACCTCC AACAACAGTCGTGACCACACCCGATCTCACAACCGAACACCACCTCCAGATGACAACCATTGCCGTCTC ACAACAGCAAACACGTGCCTTCACTAACCCCAAGCACCCTCCGGAGGAAGCCACAGGTCTCTGACTC CCGAGCCTCTAAGGAAGGGCCCACCTCACTGCAGAATCAGAAACTGTCTCCCCAGTGATTCTGGAG TAGTGTGAGTCTACTTCTGCTGACACTGTCCTGCTGACATCCAAAGAGTCCAAAGTTGGATCTCCA TCAACATCCCACGTGTCATGTGAAAACGAGTGATTCTGTCTTCCTCAGCCTGGAGCATCTGATA CAGCAGTCCTGAGCAGAACAAAACAACAAAAACAGGACAGATGGATGGAATACCCATGTCAATGAAGAA TGAAATGCCCATCTCCAAACTACTGATGATCATGCCCTCCTGGATTGTGCTCTCGCATTGTT GTGGCGTTCTCTGAGAGGGAAACTCATGGAAACCTATTGTCGAGAACACACAAGGCTAGACTACA TTGGAGATAGAAAAATGTCCTCAATGACGTGCAGCATGGAAGGGAAAGACGAAGACGGCTTTACCT CTAACAAACGCAGTAGCATGTTAGATTGAGGATGGGGCATGACACTCCAGTGTCAAATAAGCTTAGTA GATTCTCTGTTCATAAAAAGACTCACTTATTCCATGGATGTCATTGATCCAGGCTTGCTTAGTTCA ATGAATGAAGGGTACTTAGAGACCACAACTCTCTGTCAAAA
	Human TIM-4 amino acid sequence isoform 2 (GenBank Accession No. NP_001140198.1) MSKEPLILWLMIEFWWLYLTPVTSETVVTEVLGHRVTLPCLYSSWSHNSNSMCWGKDQCPYSGCKEALIR TDGMRVTSRKSAYRLQGTIPRGDVSLTILNPSESDSGVYCCRIEVPGWFNDVKINVRLNLQRASSTTHR TATTTT RRTTTSPTRQMTTPAALPTTVVTPDLTTGTPLQMTTIAVFTTANTCLSLTPSTLPEEAT GLLTPEPSKEGPILTAESETVLPDSWSSVESTSADTVLLTSKASDTAVPEQNKTGQMDGIPMSMKN EMPISQLMIIAPSLGFVLFALFVAFLRGKLMETYCSQKHTLDYIGDSKNVLNDVQHGRED EDEGLFTL

SEQ ID NO:	SEQUENCE
	<p>Homo sapiens T-cell (TIMD4), transcript variant 2, mRNA (GenBank Accession No. NM_001146726.1)</p> <p>ATAAGAGGTTGGGCTTGGATAGATAGACAGACTCCTGGTCCGGTCAACCGTCAAAATGTCCAAAGAAC CTCTCATTCTCTGGCTGATGATTGAGTTGGTGGCTTACCTGACACCAGTCACCCAGACTGAGACTGTTGT GACGGAGGTTTGGGTACCCGGTGACTTGCCTGCTGTACTCATCCTGGTCTCACAAACAGCAACAGC ATGTGCTGGGGAAAGACCAGTGCCTACTCCGGTGCAAGGAGGCGCTCATCCGACTGATGGAATGA GGGTGACCTCAAGAAAGTCAGCAAATATAGACTTCAGGGGACTATCCGAGAGGTGATGTCCTTGAC CATCTTAAACCCAGTGAAAGTGACAGCGGTGTACTGCTGCCGATAGAAGTGCCTGGCTGGTTCAAC GATGTAAGATAAACGTGCGCTGAATCTACAGAGAGCCTAACAAACACAGCACAGAACAGCAACCACCA CCACACGCAGAACACAACAAGCCCCACCACCGACAAATGACAACAACCCAGCTGCACCTCC AACAAACAGTCGTGACCACACCGATCTCACAAACCGAACACCACCTCCAGATGACAACCATTGCCGTCTC ACAACAGCAAACACGTGCCTTCACTAACCCCAAGGCACCCCTCCGGAGGAAGGCCACAGGTCTGACTC CCGAGCCTCTAAGGAAGGGCCCCTCCTCACTGCAGAATCAGAAACTGTCCTCCCCAGTGATTCCCTGGAG TAGTGTGAGTCTACTTCTGCTGACACTGTCCTGCTGACATCCAAGCATCTGATACAGCAGTTCCCTGAG CAGAACAAAACAACAAAACAGGACAGATGGATGGAATACCCATGTCAATGAAGAATGAAATGCCCATCT CCCAACTACTGATGATCATGCCCTCCTGGATTGCTCTTCGCTATTGTTGTGGCTTCTCCT GAGAGGGAAACTCATGGAAACCTATTGTTCGCAGAAACACACAAGGCTAGACTACATTGGAGATAGTAAA AATGTCCTCAATGACGTGCACCATGGAAGGGAAAGACGAAGACGGCCTTTACCCCTAACAAACGCAGTA GCATGTTAGATTGAGGATGGGGCATGACACTCCAGTGTCAAATAAGTCTTAGTAGATTCCCTGTT ATAAAAAAAGACTCACTTATTCCATGGATGTCATTGATCCAGGCTTGCTTAGTTCATGAATGAAGGGTA CTTAGAGACCACAACCTCTGTCAAAAAA</p>

CLAIMS

What is claimed is:

1. A method of treating cancer in a subject, the method comprising administering to the subject an effective amount of a PD-1 antagonist and a TIM-4 antagonist.
- 5
2. The method of claim 1, wherein the TIM-4 antagonist is selected from the group consisting of a ligand, antibody, and multivalent agent.
- 10 3. The method of claim 2, wherein the TIM-4 antagonist is an antibody which binds to the IgV domain of TIM-4.
4. The method of claim 2, wherein the anti-TIM-4 antagonist binds to the FG-CC" cleft of the IgV domain.
- 15 5. The method of any one of claims 1-4, wherein the PD-1 antagonist is selected from the group consisting of a ligand, antibody and multivalent agent.
6. The method of any one of claims 1-5, wherein the PD-1 antagonist is an anti-PD-1 antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 13, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 15.
- 20
7. The method of any one of claims 1-6, wherein the PD-1 antagonist is an anti-PD-1 antibody comprising:
 - 25 (a) a heavy chain variable region CDR1 having the sequence set forth in SEQ ID NO: 17;
 - (b) a heavy chain variable region CDR2 having the sequence set forth in SEQ ID NO: 18;
 - 30 (c) a heavy chain variable region CDR3 having the sequence set forth in SEQ ID NO: 19;
 - (d) a light chain variable region CDR1 having the sequence set forth in SEQ ID NO: 20;

(e) a light chain variable region CDR2 having the sequence set forth in SEQ ID NO: 21; and

(f) a light chain variable region CDR3 having the sequence set forth in SEQ ID NO: 22.

5

8. The method of any one of claims 1-6, wherein the PD-1 antagonist is an anti-PD-1 antibody comprising heavy and light chain variable regions having the sequences set forth in SEQ ID NOs: 13 and 15, respectively.

10 9. The method of any one of claims 1-6, wherein the PD-1 antagonist is an anti-PD-1 antibody comprising heavy and light chains having the sequences as set forth in SEQ ID NOs: 11 and 12, respectively.

15 10. The method of any one of claims 1-6, wherein the PD-1 antagonist is an anti-PD-L1 antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 1, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 3.

20 11. The method of any one of claims 1-6, wherein the PD-1 antagonist is an anti-PD-L1 antibody comprising:

(a) a heavy chain variable region CDR1 having the sequence set forth in SEQ ID NO: 5

(b) a heavy chain variable region CDR2 having the sequence set forth in SEQ ID NO: 6;

25 (c) a heavy chain variable region CDR3 having the sequence set forth in SEQ ID NO: 7;

(d) a light chain variable region CDR1 having the sequence set forth in SEQ ID NO: 8;

30 (e) a light chain variable region CDR2 having the sequence set forth in SEQ ID NO: 9; and

(f) a light chain variable region CDR3 having the sequence set forth in SEQ ID NO: 10.

12. The method of any one of claims 1-6, wherein the PD-1 antagonist is an anti-PD-L1 antibody comprising heavy and light chain variable regions having the sequences set forth in SEQ ID NOs: 1 and 3, respectively.

5 13. The method of any one of claims 1-12, wherein administration of the PD-1 antagonist and TIM-4 antagonist reduces tumor size by at least 50%.

14. The method of any one of claims 1-12, wherein administration of the PD-1 antagonist and TIM-4 antagonist, reduces tumor size by at least 80%.

10

15. The method of any one of claims 1-12, wherein administration of the PD-1 antagonist and TIM-4 antagonist results in tumor growth inhibition of at least 80%.

15 16. The method of any one of claims 1-12, wherein the treatment produces at least one therapeutic effect chosen from a reduction in size of a tumor, reduction in number of metastatic lesions over time, complete response, partial response, and stable disease.

17. A method of treating cancer in a subject, the method comprising administering to the subject an effective amount of a PD-1 antagonist and a TIM-4 antagonist, wherein

20 (a) the PD-1 antagonist is an antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 13, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 15; and

(b) the TIM-4 antagonist is an antibody.

25

18. A method of treating cancer in a subject, the method comprising administering to the subject an effective amount of a PD-1 antagonist and a TIM-4 antagonist, wherein

30 (a) the PD-1 antagonist is an antibody comprising the CDR1, CDR2 and CDR3 domains in a heavy chain variable region having the sequence set forth in SEQ ID NO: 1, and the CDR1, CDR2 and CDR3 domains in a light chain variable region having the sequence set forth in SEQ ID NO: 3; and

(b) the TIM-4 antagonist is an antibody.

19. The method of any one of claims 1-18, wherein the antagonists are formulated for intravenous administration.
20. The method of any one of claims 1-18, wherein the PD-1 antagonist is administered prior to administration of the TIM-4 antagonist.
21. The method of any one of claims 1-18, wherein the TIM-4 antagonist is administered prior to administration of the PD-1 antagonist.
- 10 22. The method of any one of claims 1-18, wherein the TIM-4 antagonist and the PD-1 antagonist are administered simultaneously.
23. The method of any one of claims 1-22, wherein the cancer is a cancer selected from the group consisting of leukemia, lymphoma, blastoma, carcinoma and sarcoma.
- 15 24. The method of any one of claims 1-22, wherein the cancer is selected from the group consisting of small-cell lung cancer, non-small cell lung cancer, gastrointestinal cancer, colorectal cancer, glioblastoma and colon carcinoma.
- 20 25. The method of any one of claims 1-24, which comprises administration of an additional therapeutic agent.
26. The method of claim 25, wherein the additional therapeutic agent is a cytotoxin or chemotherapeutic agent.
- 25 27. The method of any one of claims 1-26, wherein the subject is a human.
28. A composition comprising a PD-1 antagonist and a TIM-4 antagonist.
- 30 29. A method of treating cancer in a subject, comprising administering the subject the composition of claim 28.
30. A kit for treating a cancer in a subject, the kit comprising:
 - (a) a dose of a PD-1 antagonist;

- (b) a dose of a TIM-4 antagonist; and
- (c) instructions for using the PD-1 antagonist and TIM-4 antagonist in the method of any one of claims 1-29.

hum_TIM4_1_NP_612388	10	20	30	40	50
hum_TIM4_2_NP_001140198	MSKEPLILWIMIEFWWLYLTPVTSETVVTEVLGHHRVTLPCLYSSWSHNSN	MSKEPLILWIMIEFWWLYLTPVTSETVVTEVLGHHRVTLPCLYSSWSHNSN			
hum_TIM4_1_NP_612388	60	70	80	90	100
hum_TIM4_2_NP_001140198	SMCWGKDQCPYSGCKEALIRTDGMRVTSRKSAKYRLQGTIPRGDVSLTIL	SMCWGKDQCPYSGCKEALIRTDGMRVTSRKSAKYRLQGTIPRGDVSLTIL			
hum_TIM4_1_NP_612388	110	120	130	140	150
hum_TIM4_2_NP_001140198	NPSESSDSSGVYCCRIEVPGWENDVKINVRLNQLRASITTHRATTIRRRT	NPSESSDSSGVYCCRIEVPGWENDVKINVRLNQLRASITTHRATTIRRRT			
hum_TIM4_1_NP_612388	160	170	180	190	200
hum_TIM4_2_NP_001140198	TTSPTTTRQMTTTPAALPITVVTTPDLTTGTPLQMTTIAVFTTANTCSSL	TTSPTTTRQMTTTPAALPITVVTTPDLTTGTPLQMTTIAVFTTANTCSSL			
hum_TIM4_1_NP_612388	210	220	230	240	250
hum_TIM4_2_NP_001140198	TPSTLPEEATGLITPEPSKEGPILTAESETVLPDSWSSVESTSADTVILL	TPSTLPEEATGLITPEPSKEGPILTAESETVLPDSWSSVESTSADTVILL			
hum_TIM4_1_NP_612388	260	270	280	290	300
hum_TIM4_2_NP_001140198	TSKESKWDLPLPSTS HVSMWKTSDSVSSPQPGASDTAVPEQNKTTKTGQMD	TSK---ASDTAVPEQNKTTKTGQMD			
hum_TIM4_1_NP_612388	310	320	330	340	350
hum_TIM4_2_NP_001140198	GIPMSMKNEMPISQLLMIAPSIGFVLEFAFLVAFILRGKLMEYCSQKHT	GIPMSMKNEMPISQLLMIAPSIGFVLEFAFLVAFILRGKLMEYCSQKHT			
hum_TIM4_1_NP_612388	360	370			
hum_TIM4_2_NP_001140198	RLDXYIGDSKNVLNDVQHGREDEDGLEFTL	RLDXYIGDSKNVLNDVQHGREDEDGLEFTL			

■ Ig-like domain S/T rich domain (mucin) Signal/Transmembrane domain
 CYS (C-C) pairs are based on 4JH and 4QC structures

FIGURE 1

2/12

TIM4_cyno1_EHH54702	10	20	30	40	50	60	70
TIM4_cyno2_XP_005558436	MSKEPLILWLVMEFWWLILTPVTSETVVTIVLGHRTLPCLYSSWQNSNSMCWGKDKCPYSGCCKEALIRTDGTR	75					
hum_TIM4_INP_612388	MSKEPLILWLVMEFWWLILTPVTSETVVTIVLGHRTLPCLYSSWQNSNSMCWGKDKCPYSGCCKEALIRTDGTR	75					
TIM4_mus_NP_848874	MSKGLLLWLVTELWMLYLTPAASEDTIIGFLGQPVTLPCHYLSWSRNSMCWGKGSCKPNSKCNAELLRTDGTR	75					
TIM4_cyno1_EHH54702	80	90	100	120	130	140	150
TIM4_cyno2_XP_005558436	VTSRKSAKYRLLPGTIQRGNVSLTILNPREGDSGVYCCRIEVPGWENDVKINVRLNLQR	150					
hum_TIM4_INP_612388	VTSRKSAKYRLLPGTIQRGNVSLTILNPREGDSGVYCCRIEVPGWENDVKINVRLNLQR	142					
TIM4_mus_NP_848874	VTSRKSAKYRLLPGTIQRGNVSLTILNPSESDSGVYCCRIEVPGWENDVKINVRLNLQR	150					
TIM4_cyno1_EHH54702	160	170	180	190	200	210	220
TIM4_cyno2_XP_005558436	TTSPPPTPHVTTT-AALPTTVMTTPDLTTPLQTAVETTAN-TCPSPPTPSTLPEAATGLLPEPSKEGPI	223					
hum_TIM4_INP_612388	TTSPPPTPHVTTT-AALPTTVMTTPDLTTPLQTAVETTAN-TCPSPPTPSTLPEAATGLLPEPSKEGPI	215					
TIM4_mus_NP_848874	TTSPPPTPHVTTT-AALPTTVMTTPDLTTPLQTAVETTAN-TCPSPPTPSTLPEAATGLLPEPSKEGPI	223					
TIM4_cyno1_EHH54702	230	240	250	260	270	280	290
TIM4_cyno2_XP_005558436	LTAEESEPVLPSS-DSWSSTESTESP-ADTVVLLTRESKVKWDLPPSTSHPSSMWWKTSDSVSSPQPGASD	296					
hum_TIM4_INP_612388	LTAEESEPVLPSS-DSWNSTESTESP-ADTVVLLTRESKVKWDLPPSTSHPSSMWWKTSDSVSSPQPGASD	288					
TIM4_mus_NP_848874	LTAEESEPVLPSS-DSWSSTESTSADTVVLLTRESKVKWDLPPSTSHPSSMWWKTSDSVSSPQPGASD	297					
TIM4_cyno1_EHH54702	310	320	330	340	350	360	370
TIM4_cyno2_XP_005558436	QMDGMPMPMKNEMPISQQLMILAPSLGFVILLA-LIMAFLLRGKLMETNCLQKHTRLDCIGDSKVNLD	370					
hum_TIM4_INP_612388	QMDGMPMPMKNEMPISQQLMILAPSLGFVILLA-LIMAFLLRGKLMETNCLQKHTRLDCIGDSKVNLD	362					
TIM4_mus_NP_848874	QIN-----SRQTLIACCVGFVIMVILLIAFLIRGKVITGANCLQRHKREDNTEDSDSVINDMSHGRDD	371					
TIM4_cyno1_EHH54702	380						
TIM4_cyno2_XP_005558436	EDGLFTL 377						
hum_TIM4_INP_612388	EDGLFTL 369						
TIM4_mus_NP_848874	EDGLFTL 378						

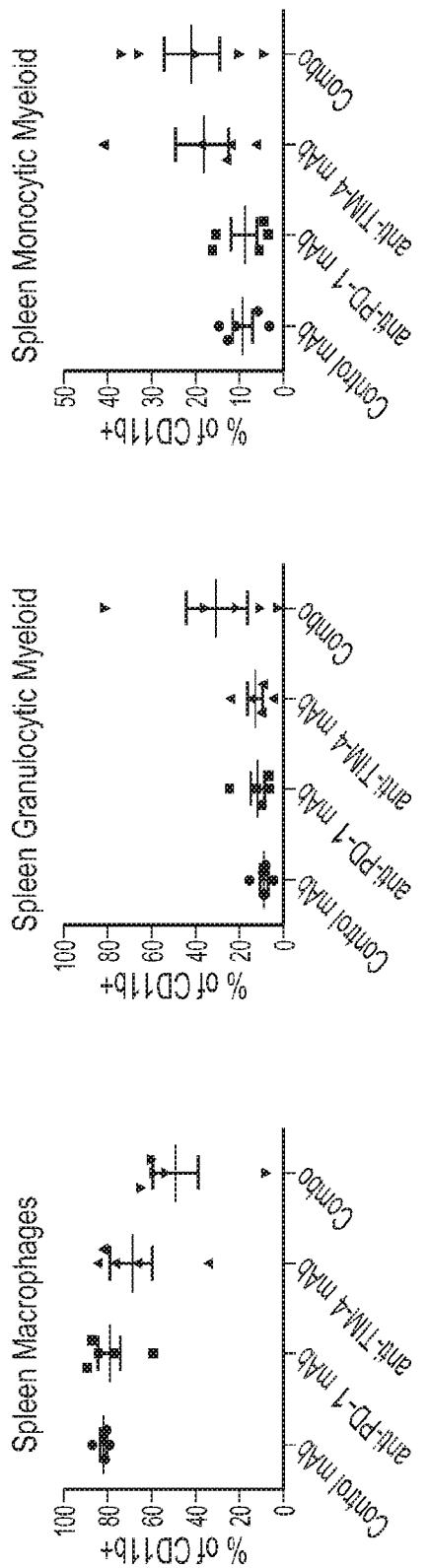


FIGURE 3A
Treatment Group

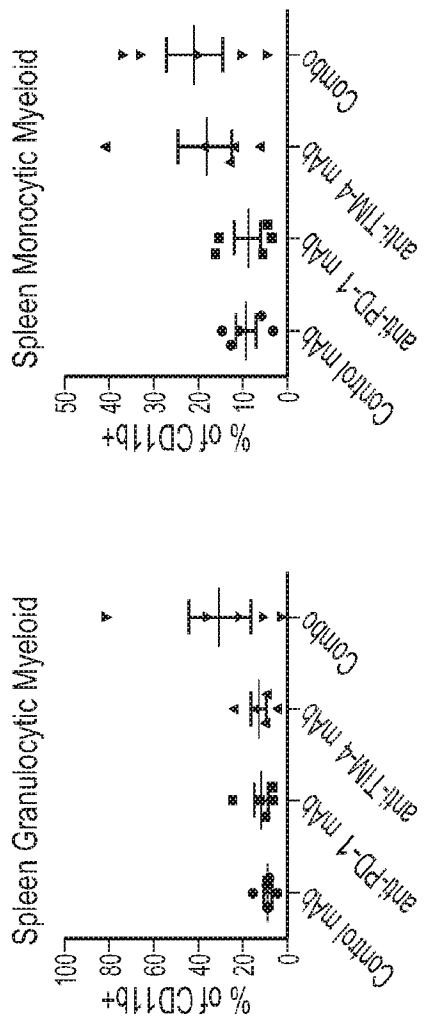


FIGURE 3B
Treatment Group

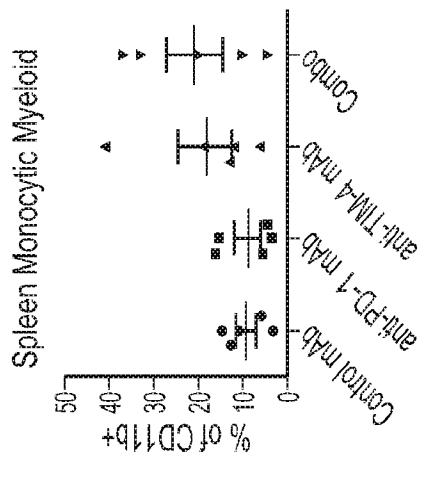


FIGURE 3C
Treatment Group

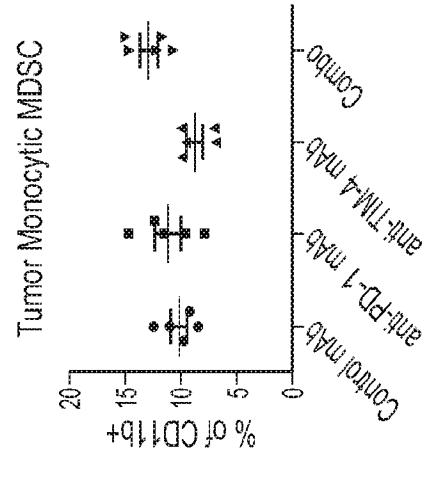


FIGURE 3D
Treatment Group

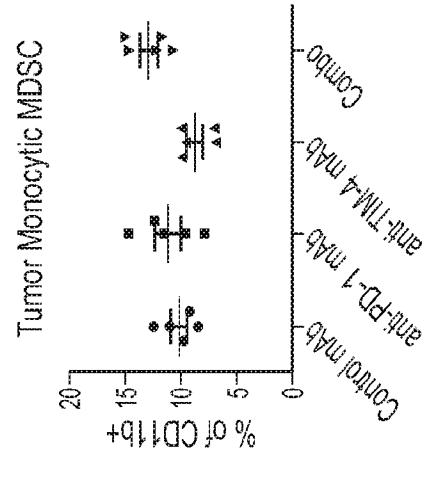


FIGURE 3E
Treatment Group

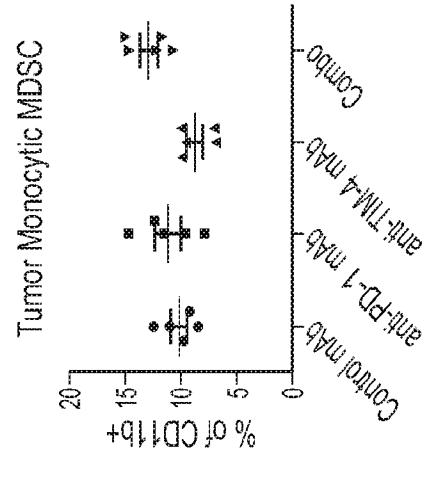
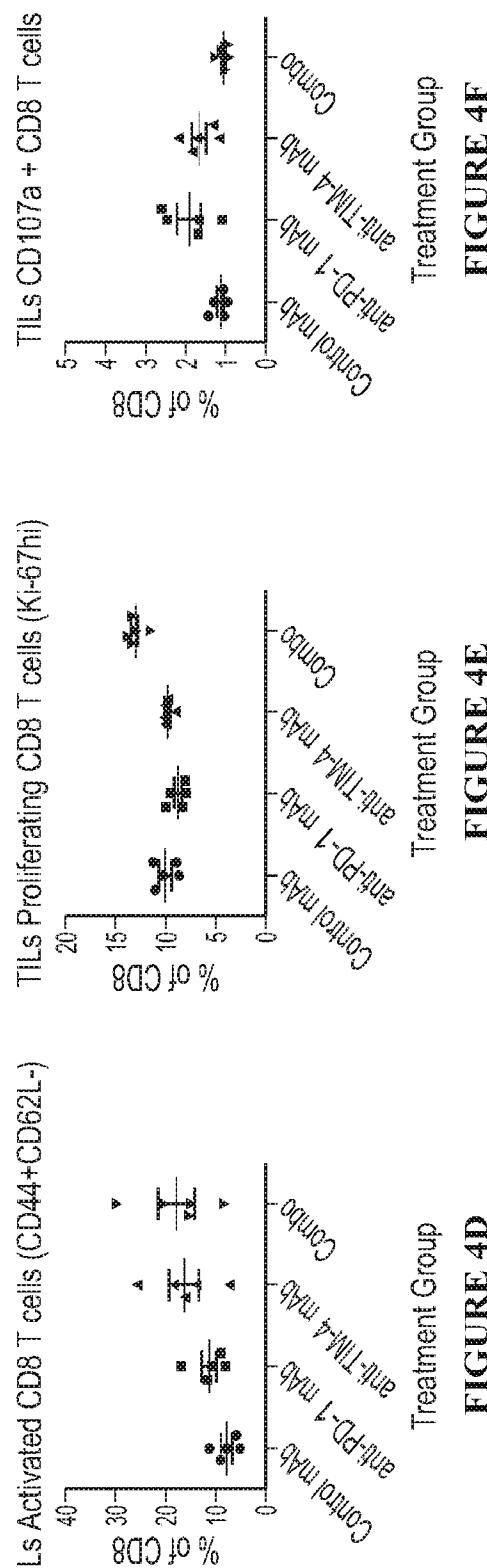
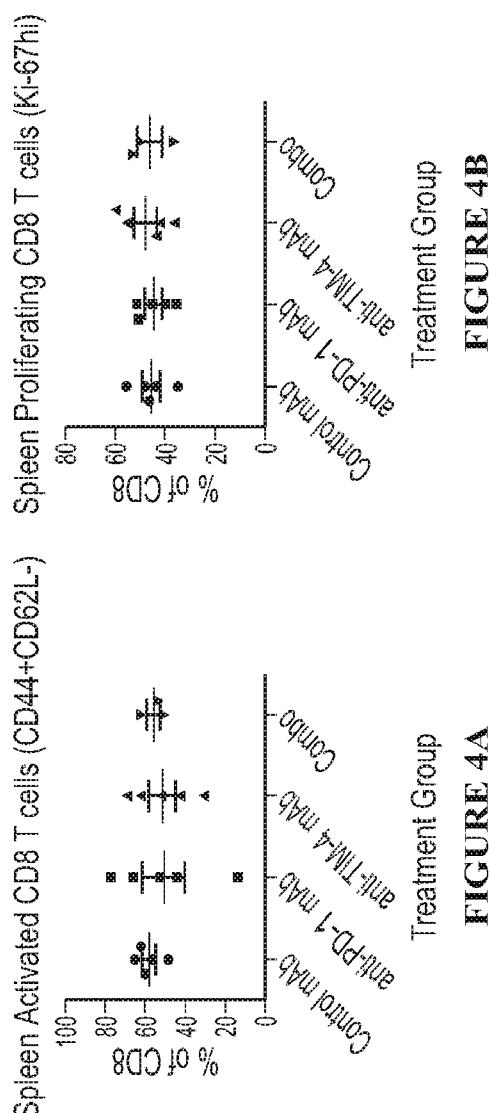
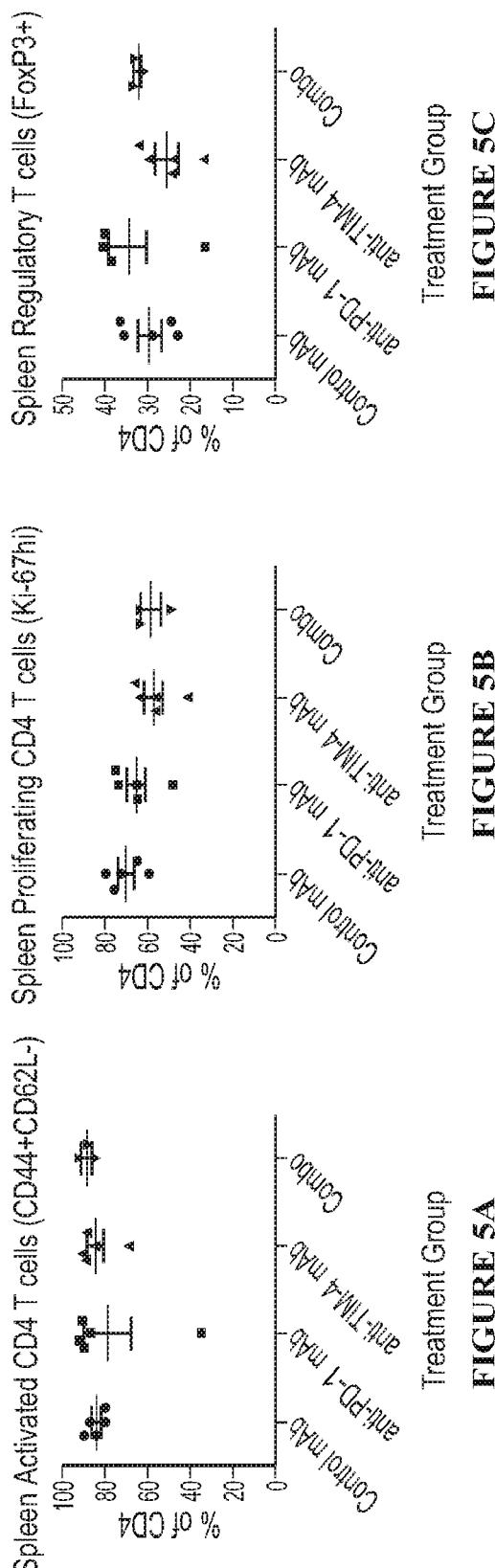
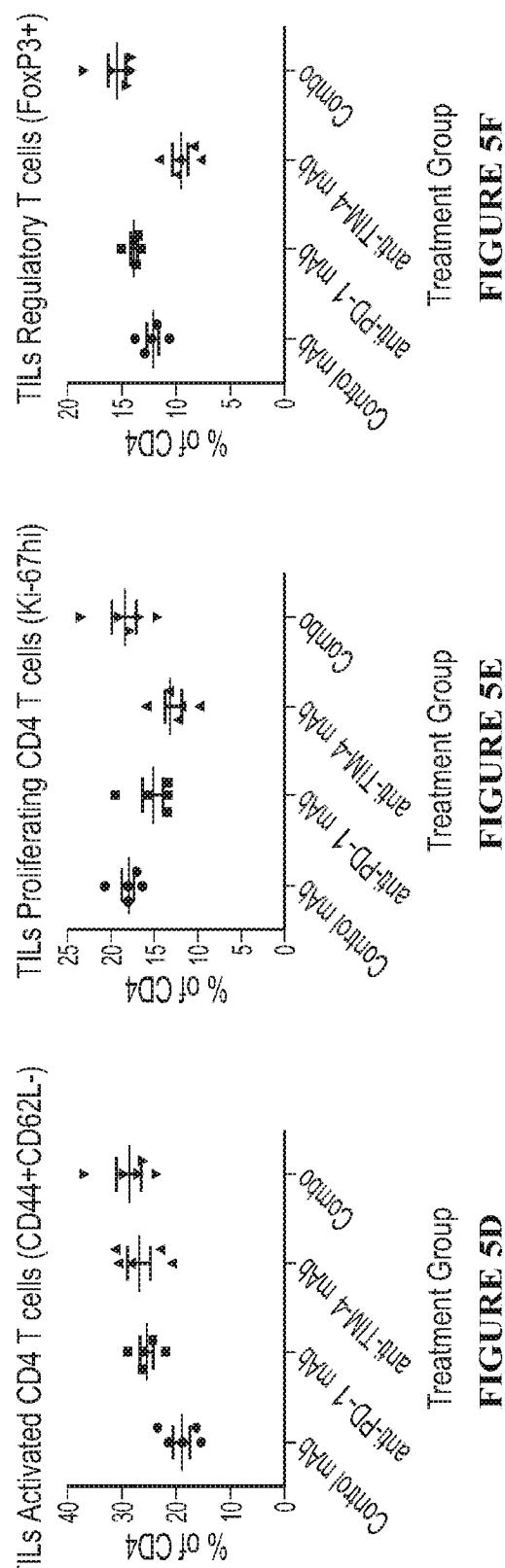


FIGURE 3F
Treatment Group



**FIGURE 5C****FIGURE 5F**

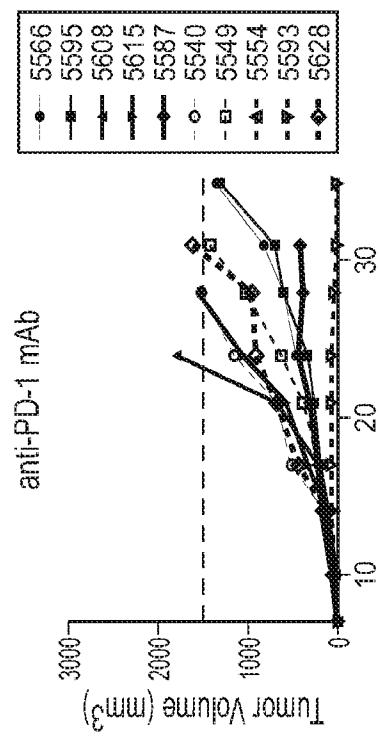


FIGURE 6B

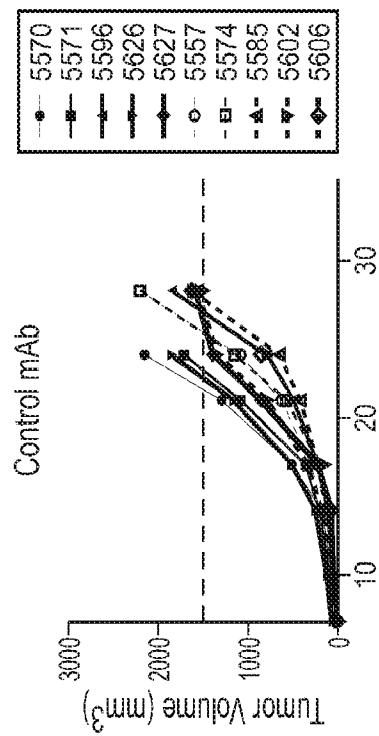


FIGURE 6A

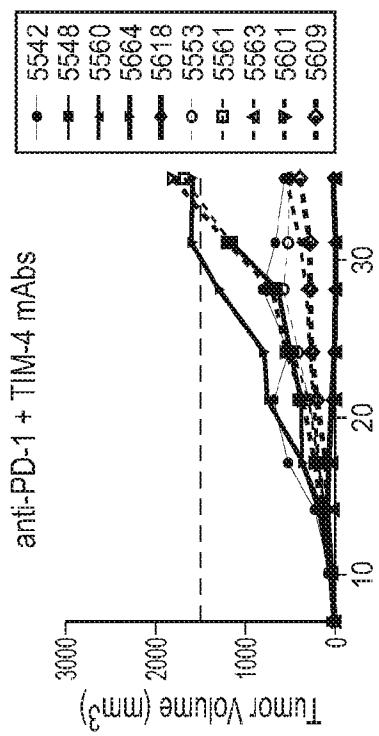


FIGURE 6D

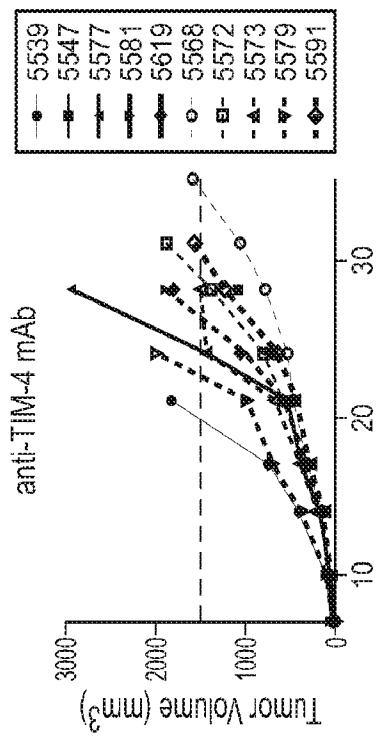


FIGURE 6C

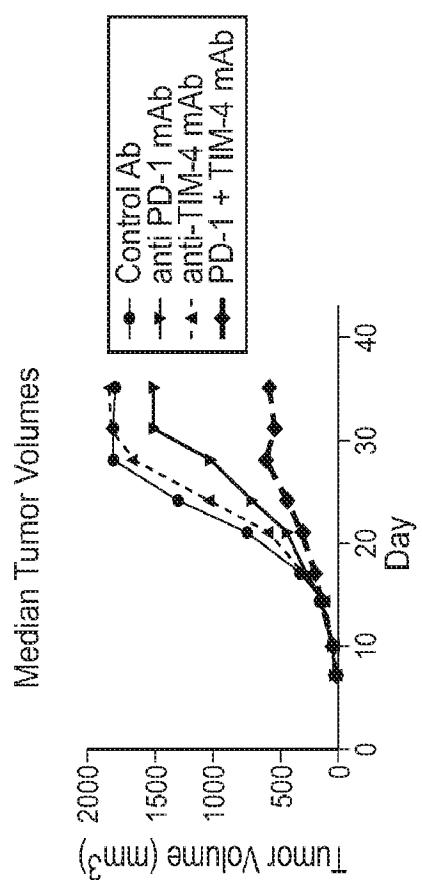


FIGURE 7B

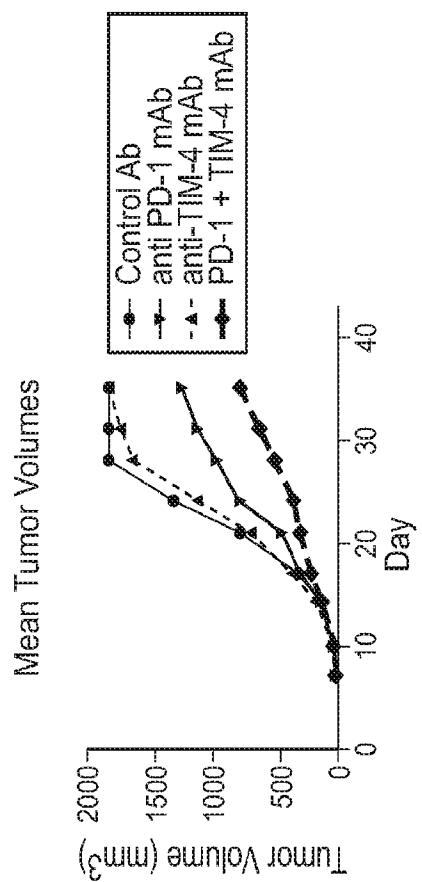


FIGURE 7A

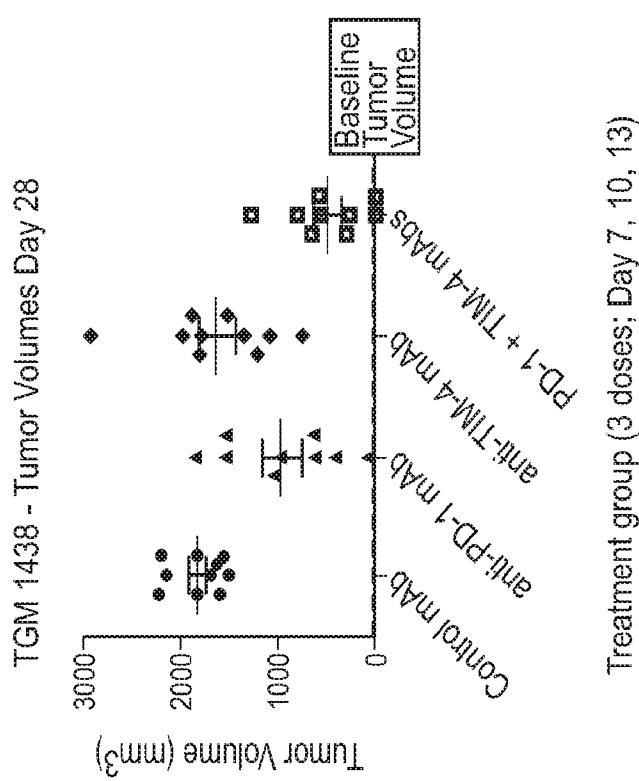


FIGURE 8

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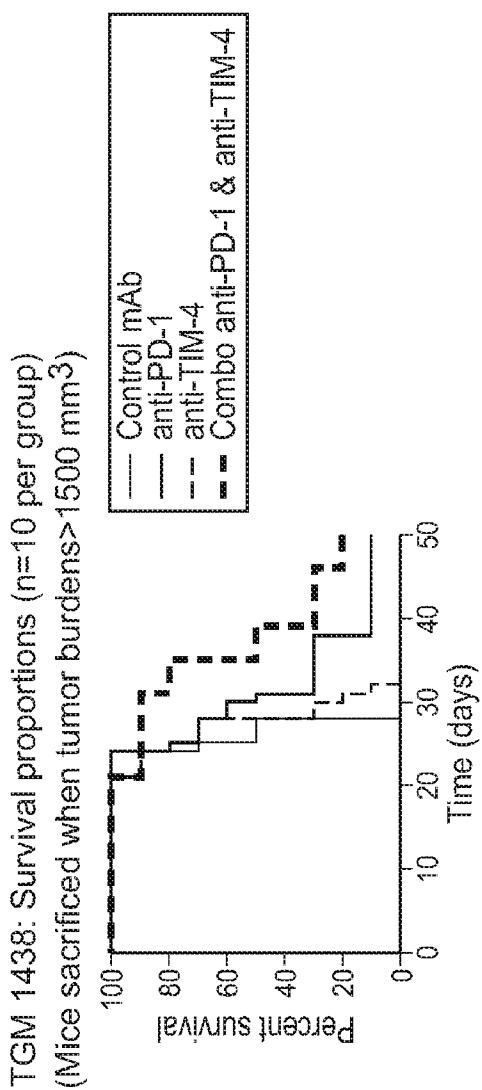


FIGURE 9A

	Control mAb	anti-PD-1	anti-TIM-4	Combo anti-PD-1 & anti-TIM-4
Median survival	26.5	30.5	28	37

FIGURE 9B

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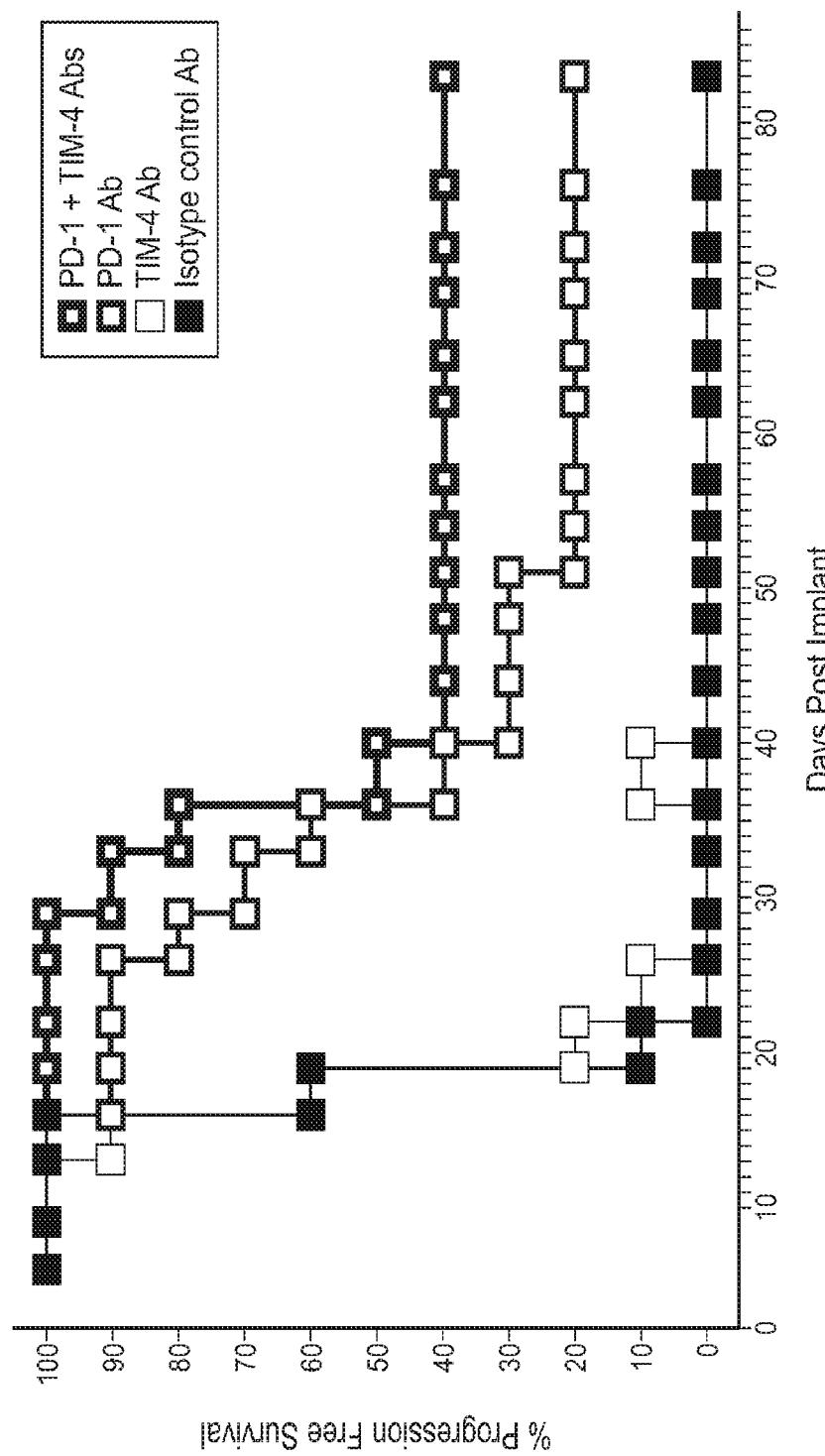


FIGURE 10

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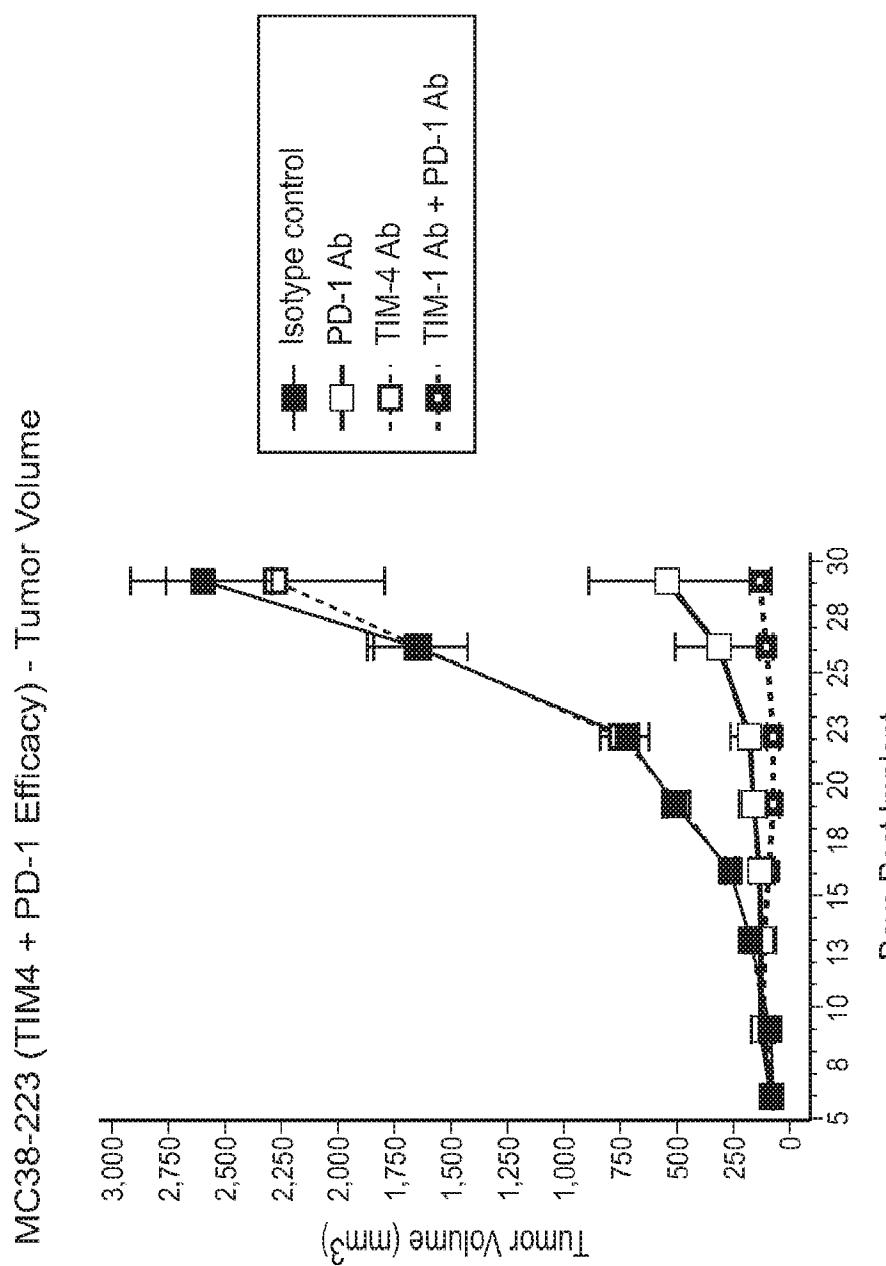


FIGURE 11

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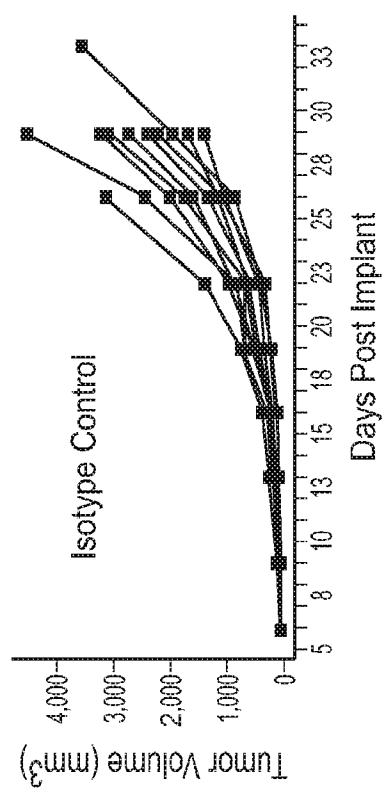


FIGURE 12A

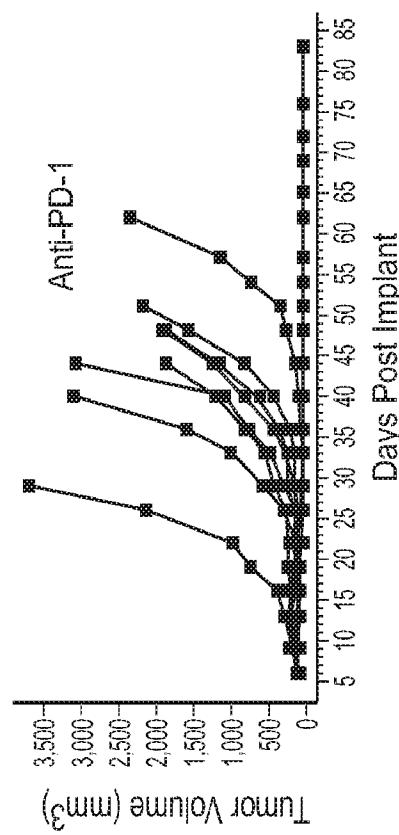


FIGURE 12B

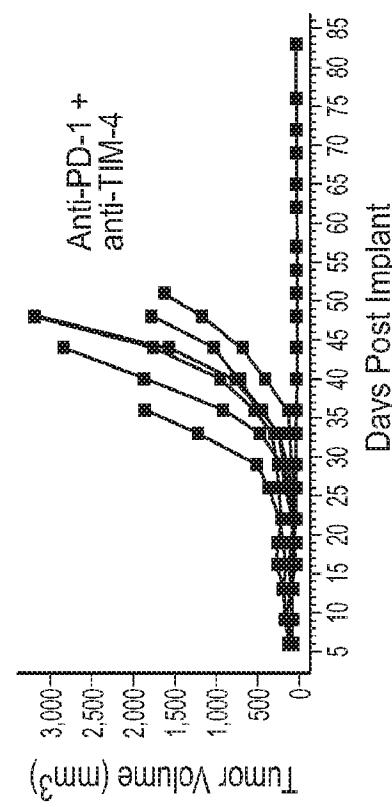


FIGURE 12C

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2017/040665

A. CLASSIFICATION OF SUBJECT MATTER
INV. C07K16/28
ADD. A61K39/00

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)
C07K A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, BIOSIS, CHEM ABS Data, EMBASE, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	<p>JENNIFER E. KIM ET AL: "Combination Therapy with Anti-PD-1, Anti-TIM-3, and Focal Radiation Results in Regression of Murine Gliomas", CLINICAL CANCER RESEARCH, vol. 23, no. 1, 29 June 2016 (2016-06-29), pages 124-136, XP055406617, US ISSN: 1078-0432, DOI: 10.1158/1078-0432.CCR-15-1535 see last paragraph of the Introduction part and abstract</p> <p style="text-align: center;">-----</p> <p style="text-align: center;">-/-</p>	1-30



Further documents are listed in the continuation of Box C.



See patent family annex.

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Date of the actual completion of the international search

Date of mailing of the international search report

29 September 2017

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Name and mailing address of the ISA/
European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040,
Fax: (+31-70) 340-3016

Authorized officer

Vadot, Pierre

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2017/040665

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	SHOHEI KOYAMA ET AL: "Adaptive resistance to therapeutic PD-1 blockade is associated with upregulation of alternative immune checkpoints", NATURE COMMUNICATIONS, vol. 7, 17 February 2016 (2016-02-17), page 10501, XP055309591, DOI: 10.1038/ncomms10501 see last paragraph of the Introduction part -----	1-30
A	K. SAKUISHI ET AL: "Targeting Tim-3 and PD-1 pathways to reverse T cell exhaustion and restore anti-tumor immunity", NATURE REVIEWS IMMUNOLOGY, vol. 6, no. 10, 27 September 2010 (2010-09-27), pages 715-2194, XP055151727, ISSN: 1474-1733, DOI: 10.1084/jem.20100643 see Introduction part -----	1-30
Y	MUHAMMAD BAGHDADI ET AL: "Combined blockade of TIM-3 and TIM-4 augments cancer vaccine efficacy against established melanomas", CANCER IMMUNOLOGY, IMMUNOTHERAPY, vol. 62, no. 4, 10 November 2012 (2012-11-10), pages 629-637, XP055181430, ISSN: 0340-7004, DOI: 10.1007/s00262-012-1371-9 see last paragraph of the Introduction part -----	1-30