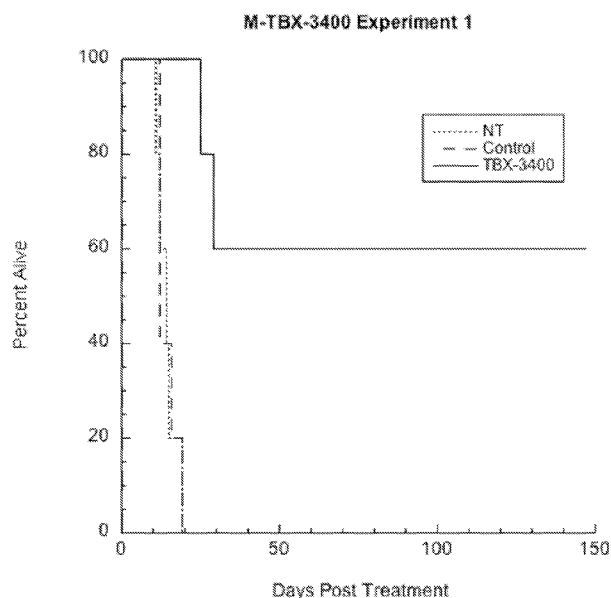




- (51) **International Patent Classification:**
C07K 14/47 (2006.01) *C12N 5/06* (2006.01)
- (21) **International Application Number:**
PCT/US2018/044740
- (22) **International Filing Date:**
01 August 2018 (01.08.2018)
- (25) **Filing Language:** English
- (26) **Publication Language:** English
- (30) **Priority Data:**
62/540,901 03 August 2017 (03.08.2017) US
- (71) **Applicant:** TAIGA BIOTECHNOLOGIES, INC.
[US/US]; 12635 E Montview Boulevard, Aurora, Colorado
80045 (US).
- (72) **Inventors:** REFAELI, Yosef; 1115 Steele Street, Denver,
Colorado 80206 (US). TURNER, Brian C.; 655 S. Glen-
coe Street, Denver, Colorado 80246 (US). BIRD, Grego-
ry Alan; 5952 S. Sheridan Way, Littleton, Colorado 80123
(US).
- (74) **Agent:** EWING, James F. et al.; FOLEY & LARDNER
LLP, 3000 K Street, N.W., Suite 600, Washington, District
of Columbia 20007-5109 (US).
- (81) **Designated States** (*unless otherwise indicated, for every
kind of national protection available*): AE, AG, AL, AM,
AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ,
CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO,
DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN,
HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP,
KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME,
MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ,
OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA,
SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN,
TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.
- (84) **Designated States** (*unless otherwise indicated, for every
kind of regional protection available*): ARIPO (BW, GH,
GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ,
UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ,
TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK,
EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV,
MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM,

(54) **Title:** METHODS AND COMPOSITIONS FOR THE TREATMENT OF CANCER

FIG. 1



(57) **Abstract:** Provided herein are methods for adoptive cell transfer for the treatment of cancer, comprising administering a therapeutically effective amount of immune cells having antitumor activity to a subject, wherein the immune cells are contacted with a protein transduction domain (PTD)-MYC fusion polypeptide prior to administration to the subject. In some embodiments, the PTD-MYC fusion polypeptide comprises: (i) an HIV TAT protein transduction domain; and (ii) a MYC polypeptide sequence.



TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW,
KM, ML, MR, NE, SN, TD, TG).

Declarations under Rule 4.17:

- *as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))*

Published:

- *with international search report (Art. 21(3))*
- *with sequence listing part of description (Rule 5.2(a))*

METHODS AND COMPOSITIONS FOR THE TREATMENT OF CANCER

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application claims the benefit of priority to U.S. Provisional Patent Application No. 62/540,901, filed August 3, 2017, the entire contents of which are incorporated herein by reference.

SEQUENCE LISTING

[0002] The instant application contains a Sequence Listing which has been submitted electronically in ASCII format and is hereby incorporated by reference in its entirety. Said ASCII copy, created on July 24, 2018, is named 106417-0333_SL.txt and is 22,015 bytes in size.

BACKGROUND OF THE INVENTION

[0003] Adoptive cell transfer (ACT) is a form of immunotherapy that involves the transfer of immune cells with antitumor activity into patients. ACT typically involves isolation of lymphocytes with antitumor activity from a patient, culturing the lymphocytes *in vitro* to expand the population, and then infusing the lymphocytes into the cancer-bearing host. Lymphocytes used for adoptive transfer can either be derived from the stroma of resected tumors (e.g., tumor infiltrating lymphocytes), from the lymphatics or lymph nodes, or from the blood. In some cases, the isolated lymphocytes are genetically engineered to express antitumor T cell receptors (TCRs) or chimeric antigen receptors (CARs). The lymphocytes used for infusion can be isolated from a donor (allogeneic ACT), or from the cancer-bearing host (autologous ACT).

SUMMARY OF THE INVENTION

[0004] Provided herein, in certain embodiments, are methods for adoptive cell transfer for the treatment of cancer. In some embodiments, provided are methods for the treatment of a cancer in a subject comprising administering a therapeutically effective amount of immune cells having antitumor activity to the subject, wherein the immune cells are contacted with a protein

transduction domain (PTD)-MYC fusion polypeptide prior to administration to the subject. In some embodiments, the immune cells comprise one or more lymphocytes. In some embodiments, the one or more lymphocytes comprise T cells and/or B cells. In some embodiments, the one or more lymphocytes comprise tumor-infiltrating lymphocytes. In some embodiments, the cancer is a metastatic cancer. In some embodiments, the cancer is a carcinoma, adenoma, adenocarcinoma, blastoma, sarcoma, or lymphoma. In some embodiments, the cancer is a basal cell carcinoma, biliary tract cancer, bladder cancer, breast cancer, cervical cancer, choriocarcinoma, CNS cancer, colon cancer, colorectal cancer, connective tissue cancer, cancer of the digestive system, endometrial cancer, esophageal cancer, eye cancer, gastric cancer, glial cell tumor, head and neck cancer, hepatoma, hepatic carcinoma, Hodgkin's lymphoma, Non-Hodgkin's lymphoma, intra-epithelial neoplasm, kidney cancer, larynx cancer, liver cancer, small-cell lung cancer, non-small cell lung cancer, melanoma, myeloma, neuroblastoma, oral cavity cancer, ovarian cancer, pancreatic cancer, prostate cancer, rectal cancer, renal cancer, cancer of the respiratory system, retinoblastoma, rhabdomyosarcoma, salivary gland carcinoma, squamous cell cancer, stomach cancer, testicular cancer, thyroid cancer, uterine cancer, cancer of the urinary system, or a vulvar cancer. In some embodiments, the immune cells are obtained from a donor subject having solid tumor. In some embodiments, the solid tumor is a metastatic tumor. In some embodiments, the immune cells are obtained from a donor subject having a melanoma or a colon cancer. In some embodiments, donor subject and the subject receiving the immune cells are the same (i.e., autologous ACT). In some embodiments, donor subject and the subject receiving the immune cells are different (i.e., allogeneic ACT).

[0005] In some embodiments, the PTD-MYC fusion polypeptide comprises: (i) an HIV TAT protein transduction domain; and (ii) a MYC polypeptide sequence. In some embodiments, the PTD-MYC fusion polypeptide translocates to the nucleus of the immune cell. In some embodiments, the PTD-MYC fusion polypeptide exhibits a biological activity of MYC, such as the activation of MYC target genes. In some embodiments, the fusion peptide comprises SEQ ID NO: 1.

[0006] Described herein, in certain embodiments are compositions comprising (a) a MYC fusion peptide, comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence;

and (b) one or more primary immune cells isolated from a donor subject that has a tumor, wherein the one or more primary immune cells are reactive against a tumor-specific antigen. In some embodiments, the MYC fusion peptide translocates to the nucleus of the one or more primary immune cells. In some embodiments, the MYC fusion peptide exhibits a biological activity of MYC. In some embodiments, the MYC fusion peptide further comprises one or more molecules that link the protein transduction domain and the MYC polypeptide. In some embodiments, the MYC fusion peptide comprises a MYC fusion peptide with the following general structure:

[0007] protein transduction domain-X-MYC sequence,

[0008] wherein -X- is molecule that links the protein transduction domain and the MYC sequence. In some embodiments, the protein transduction domain sequence is a TAT protein transduction domain sequence. In some embodiments, the TAT protein transduction domain sequence is selected from the group consisting of TAT[48-57] and TAT[57-48]. In some embodiments, the MYC fusion peptide comprises SEQ ID NO: 1. In some embodiments, the MYC fusion peptide is acetylated. In some embodiments, the one or more immune cells have antitumor activity against the tumor cells. In some embodiments, the one or more immune cells comprises one or more lymphocytes. In some embodiments, the one or more lymphocytes comprises a T cell, a B cell, an NK cell, or any combination thereof. In some embodiments, the T cell is selected from the group consisting of naïve T cells, CD4+ T cells, CD8+ T cells, memory T cells, activated T cells, anergic T cells, tolerant T cells, chimeric B cells, and antigen-specific T cells. In some embodiments, the B cells are selected from the group consisting of naïve B cells, plasma B cells, activated B cells, memory B cells, anergic B cells, tolerant B cells, chimeric B cells, and antigen-specific B cells. In some embodiments, the one or more lymphocytes is a tumor-infiltrating lymphocyte, T-cell receptor modified lymphocyte, or a chimeric antigen receptor modified lymphocyte. In some embodiments, the tumor-infiltrating lymphocyte has a CD8+CD25+ signature. In some embodiments, the lymphocyte has a CD4+CD25+ signature. In some embodiments, the one or more immune cells comprises a detectable moiety.

[0009] Described herein, in certain embodiments are methods for treating a tumor in a subject, comprising administering one or more modified immune cells to the subject in need

thereof, wherein the one or more modified immune cells comprise a MYC fusion peptide comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence and are reactive to a tumor-specific antigen. In some embodiments, the one or more modified immune cells are derived from primary immune cells isolated from the subject. In some embodiments, the one or more modified immune cells are derived from primary immune cells isolated from a separate donor subject having the same type of tumor. In some embodiments, the one or more modified immune cells are prepared by contacting the primary immune cells *in vitro* with the MYC fusion peptide following isolation. In some embodiments, the methods further comprise expanding the primary immune cells *in vitro* prior to contacting with the MYC fusion peptide. In some embodiments, the methods further comprise expanding the primary immune cells following contacting with the MYC fusion peptide. In some embodiments, the cells are expanded using an anti-CD3 antibody. In some embodiments, the cells are expanded using an irradiated allogenic feeder cells. In some embodiments, the cells are expanded in the presence of an exogenous cytokine. In some embodiments, the cytokine is interleukin-2. In some embodiments, the MYC fusion peptide translocates to the nucleus of the immune cell. In some embodiments, the MYC fusion peptide exhibits a biological activity of MYC. In some embodiments, the MYC fusion peptide further comprises one or more molecules that link the protein transduction domain and the MYC polypeptide. In some embodiments, the MYC fusion peptide comprises a MYC fusion peptide with the following general structure:

[0010] protein transduction domain-X-MYC sequence,

[0011] wherein -X- is molecule that links the protein transduction domain and the MYC sequence. In some embodiments, the protein transduction domain sequence is a TAT protein transduction domain sequence. In some embodiments, the TAT protein transduction domain sequence is selected from the group consisting of TAT[48-57] and TAT[57-48]. In some embodiments, the MYC fusion peptide comprises SEQ ID NO: 1. In some embodiments, the MYC fusion peptide is acetylated. In some embodiments, the one or more modified immune cells have antitumor activity against the tumor cells in the subject. In some embodiments, the one or more modified immune cells have antitumor activity against tumor cells in the subject. In some embodiments, the cancer cells are solid tumor cells. In some embodiments, the solid tumor

is a metastatic tumor. In some embodiments, the cancer cells are melanoma or colon tumor cells. In some embodiments, the cancer cells are from a basal cell carcinoma, biliary tract cancer, bladder cancer, breast cancer, cervical cancer, choriocarcinoma, CNS cancer, colon cancer, colorectal cancer, connective tissue cancer, cancer of the digestive system, endometrial cancer, esophageal cancer, eye cancer, gastric cancer, glial cell tumor, head and neck cancer, hepatoma, hepatic carcinoma, Hodgkin's lymphoma, Non-Hodgkin's lymphoma, intra-epithelial neoplasm, kidney cancer, larynx cancer, liver cancer, small-cell lung cancer, non-small cell lung cancer, melanoma, myeloma, neuroblastoma, oral cavity cancer, ovarian cancer, pancreatic cancer, prostate cancer, rectal cancer, renal cancer, cancer of the respiratory system, retinoblastoma, rhabdomyosarcoma, salivary gland carcinoma, squamous cell cancer, stomach cancer, testicular cancer, thyroid cancer, uterine cancer, cancer of the urinary system, or a vulvar cancer. In some embodiments, the immune cells are obtained from a donor subject having solid tumor. In some embodiments, the one or more modified immune cells comprise one or more anergic immune cells. In some embodiments, the one or more immune cells comprises one or more lymphocytes. In some embodiments, the one or more lymphocytes comprises a T cell, a B cell, an NK, or any combination thereof. In some embodiments, the T cell is selected from the group consisting of naïve T cells, CD4⁺ T cells, CD8⁺ T cells, memory T cells, activated T cells, anergic T cells, tolerant T cells, chimeric B cells, and antigen-specific T cells. In some embodiments, the B cells are selected from the group consisting of naïve B cells, plasma B cells, activated B cells, memory B cells, anergic B cells, tolerant B cells, chimeric B cells, and antigen-specific B cells. In some embodiments, the one or more lymphocytes is a tumor-infiltrating lymphocyte, T-cell receptor modified lymphocyte, or a chimeric antigen receptor modified lymphocyte. In some embodiments, the lymphocyte has a CD8⁺CD28⁻CD152⁻ signature. In some embodiments, the lymphocyte has a CD8⁺CD25⁺ signature. In some embodiments, the lymphocyte has a CD4⁺CD25⁺ signature. In some embodiments, the methods further comprise isolating the primary immune cells from the donor subject. In some embodiments, the donor subject has cancer. In some embodiments, the one or more modified immune cells are administered intravenously, intraperitoneally, subcutaneously, intramuscularly, or intratumorally. In some embodiments, the methods further comprise lymphodepleting the subject prior to administration

of the one or more modified immune cells. In some embodiments, the methods further comprise administering a cytokine to the subject. In some embodiments, the cytokine is administered prior to, during, or subsequent to administration of the one or more modified immune cells. In some embodiments, the cytokine is selected from a group consisting of interferon α , interferon β , interferon γ , complement C5a, IL-2, TNF α , CD40L, IL12, IL-23, IL15, IL17, CCL1, CCL11, CCL12, CCL13, CCL14-1, CCL14-2, CCL14-3, CCL15-1, CCL15-2, CCL16, CCL17, CCL18, CCL19, CCL19, CCL2, CCL20, CCL21, CCL22, CCL23-1, CCL23-2, CCL24, CCL25-1, CCL25-2, CCL26, CCL27, CCL28, CCL3, CCL3L1, CCL4, CCL4L1, CCL5, CCL6, CCL7, CCL8, CCL9, CCR10, CCR2, CCR5, CCR6, CCR7, CCR8, CCRL1, CCRL2, CX3CL1, CX3CR, CXCL1, CXCL10, CXCL11, CXCL12, CXCL13, CXCL14, CXCL15, CXCL16, CXCL2, CXCL3, CXCL4, CXCL5, CXCL6, CXCL7, CXCL8, CXCL9, CXCL9, CXCR1, CXCR2, CXCR4, CXCR5, CXCR6, CXCR7 and XCL2. In some embodiments, the tumor is metastatic. In some embodiments, the subject is a human or an animal. In some embodiments, the methods further comprise administering an additional cancer therapy. In some embodiments, the additional cancer therapy is selected from among chemotherapy, radiation therapy, immunotherapy, monoclonal antibodies, anti-cancer nucleic acids or proteins, anti-cancer viruses or microorganisms, and any combinations thereof. In some embodiments, the one or more modified immune cells comprises a detectable moiety.

[0012] Also described herein, in certain embodiments are methods for preparing modified immune cells for cancer therapy, comprising contacting one or more immune cells *in vitro* with a MYC fusion polypeptide, wherein the immune cells are from a donor that has been exposed to one or more tumor antigens and wherein the MYC fusion peptide comprises (i) a protein transduction domain; (ii) a MYC polypeptide sequence and are reactive to a tumor-specific antigen. In some embodiments, the one or more modified immune cells are derived from primary immune cells isolated from a subject having a tumor. In some embodiments, the methods further comprise expanding the primary immune cells *in vitro* prior to contacting with the MYC fusion peptide. In some embodiments, the methods further comprise expanding the primary immune cells following contacting with the MYC fusion peptide. In some embodiments, the cells are expanded using an anti-CD3 antibody. In some embodiments, the cells are expanded using an

irradiated allogenic feeder cells. In some embodiments, the cells are expanded in the presence of an exogenous cytokine. In some embodiments, the cytokine is interleukin-2. In some embodiments, the MYC fusion peptide translocates to the nucleus of the immune cell. In some embodiments, the MYC fusion peptide exhibits a biological activity of MYC. In some embodiments, the MYC fusion peptide further comprises one or more molecules that link the protein transduction domain and the MYC polypeptide. In some embodiments, the MYC fusion peptide comprises a MYC fusion peptide with the following general structure:

[0013] protein transduction domain-X-MYC sequence,

[0014] wherein -X- is molecule that links the protein transduction domain and the MYC sequence. In some embodiments, the protein transduction domain sequence is a TAT protein transduction domain sequence. In some embodiments, the TAT protein transduction domain sequence is selected from the group consisting of TAT[48-57] and TAT[57-48]. In some embodiments, the MYC fusion peptide comprises SEQ ID NO: 1. In some embodiments, the MYC fusion peptide is acetylated. In some embodiments, the one or more modified immune cells have antitumor activity. In some embodiments, the one or more modified immune cells have antitumor activity against tumor cells in the subject. In some embodiments, the one or more modified immune cells comprise one or more anergic immune cells. In some embodiments, the one or more immune cells comprises one or more lymphocytes. In some embodiments, the one or more lymphocytes comprises a T cell, a B cell, an NK, or any combination thereof. In some embodiments, the T cell is selected from the group consisting of naïve T cells, CD4⁺ T cells, CD8⁺ T cells, memory T cells, activated T cells, anergic T cells, tolerant T cells, chimeric B cells, and antigen-specific T cells. In some embodiments, the B cells are selected from the group consisting of naïve B cells, plasma B cells, activated B cells, memory B cells, anergic B cells, tolerant B cells, chimeric B cells, and antigen-specific B cells. In some embodiments, the one or more lymphocytes is a tumor-infiltrating lymphocyte, T-cell receptor modified lymphocyte, or a chimeric antigen receptor modified lymphocyte. In some embodiments, the lymphocyte has a CD8⁺CD28⁻CD152⁻ signature. In some embodiments, the lymphocyte has a CD8⁺CD25⁺ signature. In some embodiments, the lymphocyte has a CD4⁺CD25⁺ signature.

[0015] Also described herein, in certain embodiments, are compositions comprising: (a) one or more isolated primary immune cells that have been exposed to a tumor cell line; and (b) a MYC fusion peptide, comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence; wherein the one or more primary immune cells are reactive against a tumor-specific antigen.

[0016] Also described herein, in certain embodiments, are any of the aforementioned compositions for use in treating a cancer. Also described herein, in certain embodiments, are any of the aforementioned compositions for use in the manufacture of a medicament for treating a cancer.

[0017] Also described herein, in certain embodiments, are methods for increasing the efficacy of adoptive cell therapy or T-cell therapy in a subject comprising administering any of the aforementioned compositions.

[0018] Also described herein, in certain embodiments, are tumor-infiltrating lymphocytes comprising a MYC fusion peptide, comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence. In some embodiments, the tumor-infiltrating lymphocytes are derived from primary tumor-infiltrating lymphocytes isolated from a subject that has cancer.

[0019] Also described herein, in certain embodiments, are lymphocytes comprising a chimeric antigen receptor and a MYC fusion peptide, comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence. In some embodiments, the lymphocytes are derived from primary lymphocytes isolated from a subject that has cancer.

[0020] Also described herein, in certain embodiments, are methods for preparing a composition for adoptive cell therapy comprising contacting one or more primary immune cells with MYC fusion peptide, comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence, wherein one or more primary immune cells are isolated from a patient having a tumor, and wherein one or more primary immune cells are reactive to a tumor-specific antigen.

[0021] Also provided are kits comprising the MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells provided herein for use in treating a cancer. In some embodiments, the kit comprises one or more reagents for the detection of the administered MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells. In some

embodiments, the kit comprises cells for treatment with a MYC-fusion polypeptide provided herein, for example, hematopoietic stem cells, donor leukocytes, T cells, or NK cells. In some embodiments, the kit comprises associated instructions for using the MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells.

BRIEF DESCRIPTION OF THE DRAWINGS

[0022] FIG. 1 illustrates results for survival of melanoma tumor-bearing mice following infusion of lymphocytes from tumor-bearing donor mice treated with TAT-MYC for 1 hour. Mice were treated with TAT-MYC lymphocytes, lymph cells treated with a control protein or left untreated. Day of death recorded with day of treatment as Day 0.

[0023] FIG. 2 illustrates results for survival of melanoma tumor-bearing mice following infusion of lymphocytes from tumor-bearing donor mice treated with TAT-MYC (repeat of experiment shown in Fig. 1). Mice were treated with TAT-MYC lymphocytes, lymph cells treated with a control protein or left untreated. Day of death recorded with day of treatment as Day 0.

[0024] FIG. 3 illustrates results for survival of melanoma tumor-bearing mice following infusion of different amounts of lymphocytes from tumor-bearing donor mice treated with TAT-MYC. Mice were treated with TAT-MYC lymphocytes, lymph cells treated with a control protein or left untreated. Day of death recorded with day of treatment as Day 0.

[0025] FIG. 4 illustrates results for survival of melanoma tumor-bearing mice following infusion of different amounts of lymphocytes from tumor-bearing donor mice treated with TAT-MYC. Mice were treated with TAT-MYC lymphocytes, lymph cells treated with a control protein or left untreated. Day of death recorded with day of treatment as Day 0.

[0026] FIG. 5 illustrates results for survival of colon tumor-bearing mice following infusion of lymphocytes from tumor-bearing donor mice treated with TAT-MYC. Mice were treated with TAT-MYC lymphocytes, lymph cells treated with a control protein or left untreated. Day of death recorded with day of treatment as Day 0.

[0027] FIG. 6 illustrates results for survival of colon tumor-bearing mice following infusion of different amounts of lymphocytes from tumor-bearing donor mice treated with TAT-MYC.

Mice were treated with TAT-MYC lymphocytes or left untreated. Day of death recorded with day of treatment as Day 0.

DETAILED DESCRIPTION OF THE INVENTION

[0028] The present disclosure is not to be limited in terms of the particular embodiments described in this application, which are intended as single illustrations of individual aspects of the disclosure. All the various embodiments of the present disclosure will not be described herein. Many modifications and variations of the disclosure can be made without departing from its spirit and scope, as will be apparent to those skilled in the art. Functionally equivalent methods and apparatuses within the scope of the disclosure, in addition to those enumerated herein, will be apparent to those skilled in the art from the foregoing descriptions. Such modifications and variations are intended to fall within the scope of the appended claims. The present disclosure is to be limited only by the terms of the appended claims, along with the full scope of equivalents to which such claims are entitled.

[0029] It is to be understood that the present disclosure is not limited to particular uses, methods, reagents, compounds, compositions or biological systems, which can, of course, vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments only, and is not intended to be limiting.

[0030] In addition, where features or aspects of the disclosure are described in terms of Markush groups, those skilled in the art will recognize that the disclosure is also thereby described in terms of any individual member or subgroup of members of the Markush group.

[0031] As will be understood by one skilled in the art, for any and all purposes, particularly in terms of providing a written description, all ranges disclosed herein also encompass any and all possible subranges and combinations of subranges thereof. Any listed range can be easily recognized as sufficiently describing and enabling the same range being broken down into at least equal halves, thirds, quarters, fifths, tenths, etc. As a non-limiting example, each range discussed herein can be readily broken down into a lower third, middle third and upper third, etc. As will also be understood by one skilled in the art all language such as “up to,” “at least,”

“greater than,” “less than,” and the like, include the number recited and refer to ranges which can be subsequently broken down into subranges as discussed above. Finally, as will be understood by one skilled in the art, a range includes each individual member. Thus, for example, a group having 1-3 cells refers to groups having 1, 2, or 3 cells. Similarly, a group having 1-5 cells refers to groups having 1, 2, 3, 4, or 5 cells, and so forth.

[0032] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this disclosure belongs.

I. Definitions

[0033] The terminology used herein is for the purpose of describing particular embodiments only and is not intended to be limiting of the disclosure. As used herein, the singular forms “a”, “an” and “the” are intended to include the plural forms as well, unless the context clearly indicates otherwise.

[0034] As used herein, the term “about” means that a value can vary +/- 20%, +/- 15%, +/- 10% or +/- 5% and remain within the scope of the present disclosure. For example, “a concentration of about 200 IU/mL” encompasses a concentration between 160 IU/mL and 240 IU/mL.

[0035] As used herein, the term “administration” of an agent to a subject includes any route of introducing or delivering the agent to a subject to perform its intended function.

Administration can be carried out by any suitable route, including intravenously, intramuscularly, intraperitoneally, or subcutaneously. Administration includes self-administration and the administration by another.

[0036] The term “amino acid” refers to naturally occurring and non-naturally occurring amino acids, as well as amino acid analogs and amino acid mimetics that function in a manner similar to the naturally occurring amino acids. Naturally encoded amino acids are the 20 common amino acids (alanine, arginine, asparagine, aspartic acid, cysteine, glutamine, glutamic acid, glycine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, proline, serine, threonine, tryptophan, tyrosine, and valine) and pyrrolysine and selenocysteine. Amino acid analogs refers to agents that have the same basic chemical structure as a naturally occurring

amino acid, *i.e.*, an α carbon that is bound to a hydrogen, a carboxyl group, an amino group, and an R group, such as, homoserine, norleucine, methionine sulfoxide, methionine methyl sulfonium. Such analogs have modified R groups (such as, norleucine) or modified peptide backbones, but retain the same basic chemical structure as a naturally occurring amino acid. In some embodiments, amino acids forming a polypeptide are in the D form. In some embodiments, the amino acids forming a polypeptide are in the L form. In some embodiments, a first plurality of amino acids forming a polypeptide are in the D form and a second plurality are in the L form.

[0037] Amino acids are referred to herein by either their commonly known three letter symbols or by the one-letter symbols recommended by the IUPAC-IUB Biochemical Nomenclature Commission. Nucleotides, likewise, are referred to by their commonly accepted single-letter code.

[0038] The terms “polypeptide,” “peptide,” and “protein” are used interchangeably herein to refer to a polymer of amino acid residues. The terms apply to naturally occurring amino acid polymers as well as amino acid polymers in which one or more amino acid residues is a non-naturally occurring amino acid, *e.g.*, an amino acid analog. The terms encompass amino acid chains of any length, including full length proteins, wherein the amino acid residues are linked by covalent peptide bonds.

[0039] As used herein, a “control” is an alternative sample used in an experiment for comparison purpose. A control can be “positive” or “negative.” For example, where the purpose of the experiment is to determine a correlation of the efficacy of a therapeutic agent for the treatment for a particular type of disease, a positive control (a composition known to exhibit the desired therapeutic effect) and a negative control (a subject or a sample that does not receive the therapy or receives a placebo) are typically employed.

[0040] As used herein, the term “effective amount” or “therapeutically effective amount” refers to a quantity of an agent sufficient to achieve a desired therapeutic effect. In the context of therapeutic applications, the amount of a therapeutic peptide administered to the subject can depend on the type and severity of the infection and on the characteristics of the individual, such as general health, age, sex, body weight and tolerance to drugs. It can also depend on the degree,

severity and type of disease. The skilled artisan will be able to determine appropriate dosages depending on these and other factors.

[0041] As used herein, the term “expression” refers to the process by which polynucleotides are transcribed into mRNA and/or the process by which the transcribed mRNA is subsequently being translated into peptides, polypeptides, or proteins. If the polynucleotide is derived from genomic DNA, expression can include splicing of the mRNA in a eukaryotic cell. The expression level of a gene can be determined by measuring the amount of mRNA or protein in a cell or tissue sample. In one aspect, the expression level of a gene from one sample can be directly compared to the expression level of that gene from a control or reference sample. In another aspect, the expression level of a gene from one sample can be directly compared to the expression level of that gene from the same sample following administration of the compositions disclosed herein. The term “expression” also refers to one or more of the following events: (1) production of an RNA template from a DNA sequence (*e.g.*, by transcription) within a cell; (2) processing of an RNA transcript (*e.g.*, by splicing, editing, 5' cap formation, and/or 3' end formation) within a cell; (3) translation of an RNA sequence into a polypeptide or protein within a cell; (4) post-translational modification of a polypeptide or protein within a cell; (5) presentation of a polypeptide or protein on the cell surface; and (6) secretion or presentation or release of a polypeptide or protein from a cell.

[0042] The term “linker” refers to synthetic sequences (*e.g.*, amino acid sequences) that connect or link two sequences, *e.g.*, that link two polypeptide domains. In some embodiments, the linker contains 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 of amino acid sequences.

[0043] The terms “lyophilized,” “lyophilization” and the like as used herein refer to a process by which the material (*e.g.*, nanoparticles) to be dried is first frozen and then the ice or frozen solvent is removed by sublimation in a vacuum environment. An excipient can be included in pre-lyophilized formulations to enhance stability of the lyophilized product upon storage. The lyophilized sample can further contain additional excipients.

[0044] As used herein the term immune cell refers to any cell that plays a role in the immune response. Immune cells are of hematopoietic origin, and include lymphocytes, such as B cells

and T cells; natural killer cells; myeloid cells, such as monocytes, macrophages, dendritic cells, eosinophils, neutrophils, mast cells, basophils, and granulocytes.

[0045] The term “lymphocyte” refers to all immature, mature, undifferentiated and differentiated white lymphocyte populations including tissue specific and specialized varieties. It encompasses, by way of non-limiting example, B cells, T cells, NKT cells, and NK cells. In some embodiments, lymphocytes include all B cell lineages including pre-B cells, progenitor B cells, early pro-B cells, late pro-B cells, large pre-B cells, small pre-B cells, immature B cells, mature B cells, plasma B cells, memory B cells, B-1 cells, B-2 cells and anergic AN1/T3 cell populations.

[0046] As used herein, the term T-cell includes naïve T cells, CD4⁺ T cells, CD8⁺ T cells, memory T cells, activated T cells, anergic T cells, tolerant T cells, chimeric B cells, and antigen-specific T cells.

[0047] The term “B cell” or “B cells” refers to, by way of non-limiting example, a pre-B cell, progenitor B cell, early pro-B cell, late pro-B cell, large pre-B cell, small pre-B cell, immature B cell, mature B cell, naïve B cells, plasma B cells, activated B cells, anergic B cells, tolerant B cells, chimeric B cells, antigen-specific B cells, memory B cell, B-1 cell, B-2 cells and anergic AN1/T3 cell populations. In some embodiments, the term B cell includes a B cell that expresses an immunoglobulin heavy chain and/or light chain on its cells surface. In some embodiments, the term B cell includes a B cell that expresses and secretes an immunoglobulin heavy chain and/or light chain. In some embodiments, the term B cell includes a cell that binds an antigen on its cell-surface. In some embodiments disclosed herein, B cells or AN1/T3 cells are utilized in the processes described. In certain embodiments, such cells are optionally substituted with any animal cell suitable for expressing, capable of expressing (*e.g.*, inducible expression), or capable of being differentiated into a cell suitable for expressing an antibody including, *e.g.*, a hematopoietic stem cell, a naïve B cell, a B cell, a pre-B cell, a progenitor B cell, an early Pro-B cell, a late pro-B cell, a large pre-B cell, a small pre-B cell, an immature B cell, a mature B cell, a plasma B cell, a memory B cell, a B-1 cell, a B-2 cell, an anergic B cell, or an anergic AN1/T3 cell.

[0048] As used herein "adoptive cell therapeutic composition" refers to any composition comprising cells suitable for adoptive cell transfer. In exemplary embodiments, the adoptive cell therapeutic composition comprises a cell type selected from a group consisting of a tumor infiltrating lymphocyte (TIL), TCR (i.e. heterologous T-cell receptor) modified lymphocytes and CAR (i.e. chimeric antigen receptor) modified lymphocytes. In another embodiment, the adoptive cell therapeutic composition comprises a cell type selected from a group consisting of T-cells, CD8⁺ cells, CD4⁺ cells, NK-cells, delta-gamma T-cells, regulatory T-cells and peripheral blood mononuclear cells. In another embodiment, TILs, T-cells, CD8⁺ cells, CD4⁺ cells, NK-cells, delta-gamma T-cells, regulatory T-cells or peripheral blood mononuclear cells form the adoptive cell therapeutic composition. In one embodiment, the adoptive cell therapeutic composition comprises T cells.

[0049] As used herein "tumor-infiltrating lymphocytes" or TILs refer to white blood cells that have left the bloodstream and migrated into a tumor.

[0050] The terms "MYC" and "MYC gene" are synonyms. They refer to a nucleic acid sequence that encodes a MYC polypeptide. A MYC gene comprises a nucleotide sequence of at least 120 nucleotides that is at least 60% to 100% identical or homologous, *e.g.*, at least 60, 65%, 70%, 75%, 80%, 85%, 86%, 87%, 88%, 90%, 91%, 92%, 94%, 95%, 96%, 97%, 98%, or any other percent from about 70% to about 100% identical to sequences of NCBI Accession Number NM—002467. In some embodiments, the MYC gene is a proto-oncogene. In certain instances, a MYC gene is found on chromosome 8, at 8q24.21. In certain instances, a MYC gene begins at 128,816,862 bp from pter and ends at 128,822,856 bp from pter. In certain instances, a MYC gene is about 6 kb. In certain instances, a MYC gene encodes at least eight separate mRNA sequences—5 alternatively spliced variants and 3 unspliced variants.

[0051] The terms "MYC protein," "MYC polypeptide," and "MYC sequence" are synonyms and refer to the polymer of amino acid residues disclosed in NCBI Accession Number UniProtKB/Swiss-Prot:P01106.1 (MYC isoform 1) or NP_002458.2 (UniProtKB/Swiss-Prot:P01106.2; MYC isoform 2), and functional homologs, analogs or fragments thereof. The sequence of or UniProtKB/Swiss-Prot:P01106.1 is:

MPLNVSFTNRNYDL DYDSVQPYFYCDEEENFYQQQQQSELQPPAPSEDIWKKFELLPTP
 PLSPSRRSGLCSPSYVAVTPFSLRGDNDGGGGSFSTADQLEMVTELLGGDMVNQSFICDP
 DDETFIKNIIIQDCMWSGFSAAAKLVSEKLASYQAARKDSGSPNPARGHSVCSTSSLYLQ
 DLSAAASECIDPSVVFYPYPLNDSSSPKSCASQDSSAFSPSSDSSLSTESSPQGSPEPLVLHE
 ETPPTTSSDSEEEQEDEEEIDVVSVEKRQAPGKRSESGSPSAGGHKPPHSPLVLKRCHVS
 THQHNYAAPPSTRKDYPAAKRVKLDSVRVLRQISNNRKCTSPRSSDTEENVKRRTHNVL
 ERQRRNELKRSFFALRDQIPELENNEKAPKVVLKKATAYILSVQAEEQKLISEEDLLRKR
 REQLKHKLEQLRNSCA (SEQ ID NO: 2)

The sequence of NP_002458.2 (UniProtKB/Swiss-Prot:P01106.2) is:

MDFFRVVENQQPPATMPLNVSFTNRNYDL DYDSVQPYFYCDEEENFYQQQQQSELQPP
 APSEDIWKKFELLPTPPLSPSRRSGLCSPSYVAVTPFSLRGDNDGGGGSFSTADQLEMVT
 ELLGGDMVNQSFICDPDDETFIKNIIIQDCMWSGFSAAAKLVSEKLASYQAARKDSGSPN
 PARGHSVCSTSSLYLQDLSAAASECIDPSVVFYPYPLNDSSSPKSCASQDSSAFSPSSDSSL
 STESSPQGSPEPLVLHEETPPTTSSDSEEEQEDEEEIDVVSVEKRQAPGKRSESGSPSAGG
 HSKPPHSPLVLKRCHVSTHQHNYAAPPSTRKDYPAAKRVKLDSVRVLRQISNNRKCTSP
 RSSDTEENVKRRTHNVLERQRRNELKRSFFALRDQIPELENNEKAPKVVLKKATAYILS
 VQAEEQKLISEEDLLRKRREQLKHKLEQLRNSCA (SEQ ID NO: 11)

[0052] In some embodiments, the MYC polypeptide is a complete MYC polypeptide sequence. In some embodiments, the MYC polypeptide is a partial MYC polypeptide sequence. In some embodiments, the MYC polypeptide comprises at least 400 consecutive amino acids of SEQ ID NO: 2 OR 11. In some embodiments, the MYC polypeptide comprises at least 400 consecutive amino acids of SEQ ID NO: 2 OR 11 and retains at least one MYC activity. In some embodiments, the MYC polypeptide comprises at least 400, at least 410, at least 420, at least 430, or at least 450 consecutive amino acids of SEQ ID NO: 2 OR 11. In some embodiments, the MYC polypeptide comprises at least 400, at least 410, at least 420, at least 430, or at least 450 consecutive amino acids of SEQ ID NO: 2 OR 11 and retains at least one MYC activity. In some embodiments, the MYC polypeptide is c-MYC. In some embodiments, the MYC polypeptide sequence comprises the sequence shown below:

MDFFRVVENQQPPATMPLNVSTNRNYDL DYDSVQPYFYCDEEENFYQQQQQSELQPP
 APSEDIWKKFELLPTPPLSPSRRSGLCSPSYVAVTPFSLRGDNDGGGGSFSTADQLEMVT
 ELLGGDMVNVQSFICDPDDETFIKNIIQDCMWSGFSAAAKLVSEKLASYQAARKDSGSPN
 PARGHSVCSTSSLYLQDL SAAASECIDPSVVFYPYPLNDSSSPKSCASQDSSAFSPSSDSL
 STESSPQGSPEPLVLHEETPPTTSSDSEEEQEDEEEIDVVSVEKRQAPGKRSESGSPSAGG
 HSKPPHSPLVLKRCHVSTHQHNYAAPPSTRKDYPAAKRVKLDSVRVLRQISNNRKCTSP
 RSSDTEENVKRRTHNVLERQRRNELKRSFFALRDQIPELENNEKAPKVVLKKATAYILS
 VQAEEQKLISEEDLLRKRRLEQLKHKLEQLR (SEQ ID NO: 3).

[0053] In some embodiments, the MYC polypeptide sequence comprises the sequence shown below:

PLNVSTNRNYDL DYDSVQPYFYCDEEENFYQQQQQSELQPPAPSEDIWKKFELLPTPPL
 SPSRRSGLCSPSYVAVTPFSLRGDNDGGGGSFSTADQLEMVTELLGGDMVNVQSFICDPD
 DETFIKNIIQDCMWSGFSAAAKLVSEKLASYQAARKDSGSPNPARGHSVCSTSSLYLQD
 LSAAASECIDPSVVFYPYPLNDSSSPKSCASQDSSAFSPSSDSLSTESSPQGSPEPLVLHEE
 TPPTTSSDSEEEQEDEEEIDVVSVEKRQAPGKRSESGSPSAGGHHSKPPHSPLVLKRCHVST
 HQHNYAAPPSTRKDYPAAKRVKLDSVRVLRQISNNRKCTSPRSSDTEENVKRRTHNVLE
 RQRRNELKRSFFALRDQIPELENNEKAPKVVLKKATAYILSVQAEEQKLISEEDLLRKRR
 EQLKHKLEQLR (SEQ ID NO: 4).

[0054] In some embodiments, a MYC polypeptide comprises an amino acid sequence that is at least 40% to 100% identical, *e.g.*, at least 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 90%, 91%, 92%, 94%, 95%, 96%, 97%, 98%, 99%, or any other percent from about 40% to about 100% identical to the sequence of NCBI Accession Number NP002458.2 or UniProtKB/Swiss-Prot Accession Number P01106.1. In some embodiments, MYC polypeptide refers to a polymer of 439 amino acids, a MYC polypeptide that has not undergone any post-translational modifications. In some embodiments, MYC polypeptide refers to a polymer of 439 amino acids that has undergone post-translational modifications. In some embodiments, the MYC polypeptide is 48,804 kDa. In some embodiments, the MYC polypeptide contains a basic Helix-Loop-Helix Leucine Zipper (bHLH/LZ) domain. In some embodiments, the bHLH/LZ domain comprises the sequence of:

ELKRSFFALRDQIPELENNEKAPKVVLKKATAYILSVQAEEQKLISEEDLLRKRREQLKH KLEQLR (SEQ ID NO: 5). In some embodiments, the MYC polypeptide is a transcription factor (*e.g.*, Transcription Factor 64). In some embodiments, the MYC polypeptide contains an E-box DNA binding domain. In some embodiments, the MYC polypeptide binds to a sequence comprising CACGTG. In some embodiments, the MYC polypeptide promotes one or more of cell survival and/or proliferation. In some embodiments, a MYC polypeptide includes one or more of those described above, and includes one or more post-translational modifications (*e.g.*, acetylation). In some embodiments, the MYC polypeptides comprise one or more additional amino acid residues at the N-terminus or C-terminus of the polypeptide. In some embodiments, the MYC polypeptides are fusion proteins. In some embodiments, the MYC polypeptides are linked to one or more additional peptides at the N-terminus or C-terminus of the polypeptide.

[0055] Proteins suitable for use in the methods described herein also includes functional variants, including proteins having between 1 to 15 amino acid changes, *e.g.*, 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, or 15 amino acid substitutions, deletions, or additions, compared to the amino acid sequence of any protein described herein. In other embodiments, the altered amino acid sequence is at least 75% identical, *e.g.*, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to the amino acid sequence of any protein inhibitor described herein. Such sequence-variant proteins are suitable for the methods described herein as long as the altered amino acid sequence retains sufficient biological activity to be functional in the compositions and methods described herein. Where amino acid substitutions are made, the substitutions can be conservative amino acid substitutions. Among the common, naturally occurring amino acids, for example, a “conservative amino acid substitution” is illustrated by a substitution among amino acids within each of the following groups: (1) glycine, alanine, valine, leucine, and isoleucine, (2) phenylalanine, tyrosine, and tryptophan, (3) serine and threonine, (4) aspartate and glutamate, (5) glutamine and asparagine, and (6) lysine, arginine and histidine. The BLOSUM62 table is an amino acid substitution matrix derived from about 2,000 local multiple alignments of protein sequence segments, representing highly conserved regions of more than 500 groups of related proteins (Henikoff *et al.*, (1992), *Proc. Natl Acad. Sci. USA*, 89:10915-

10919). Accordingly, the BLOSUM62 substitution frequencies are used to define conservative amino acid substitutions that, in some embodiments, are introduced into the amino acid sequences described or disclosed herein. Although it is possible to design amino acid substitutions based solely upon chemical properties (as discussed above), the language “conservative amino acid substitution” preferably refers to a substitution represented by a BLOSUM62 value of greater than -1. For example, an amino acid substitution is conservative if the substitution is characterized by a BLOSUM62 value of 0, 1, 2, or 3. According to this system, preferred conservative amino acid substitutions are characterized by a BLOSUM62 value of at least 1 (*e.g.*, 1, 2 or 3), while more preferred conservative amino acid substitutions are characterized by a BLOSUM62 value of at least 2 (*e.g.*, 2 or 3).

[0056] The phrases “E-box sequence” and “enhancer box sequence” are used interchangeably herein and mean the nucleotide sequence CANNTG, wherein N is any nucleotide. In certain instances, the E-box sequence comprises CACGTG. In certain instances, the basic helix-loop-helix domain of a transcription factor encoded by MYC binds to the E-box sequence. In certain instances the E-box sequence is located upstream of a gene (*e.g.*, p21, Bcl-2, or ornithine decarboxylase). In certain instances, the MYC polypeptide contains an E-box DNA binding domain. In certain instances, the E-box DNA binding domain comprises the sequence of KRRTHNVLERQRRN (SEQ ID NO: 6). In certain instances, the binding of the transcription factor encoded by MYC to the E-box sequence, allows RNA polymerase to transcribe the gene downstream of the E-box sequence.

[0057] The term “MYC activity” or “MYC biological activity” or “biologically active MYC” includes one or more of enhancing or inducing cell survival, cell proliferation, and/or antibody production. By way of example and not by way of limitation, MYC activity includes enhancement of expansion of anti-CD3 and anti-CD28 activated T-cells and/or increased proliferation of long-term self-renewing hematopoietic stem cells. MYC activity also includes entry into the nucleus of a cell, binding to a nucleic acid sequence (*e.g.*, binding an E-box sequence), and/or inducing expression of MYC target genes.

[0058] The terms “patient,” “subject,” “individual,” and the like are used interchangeably herein, and refer to an animal, typically a mammal. In one embodiment, the patient, subject, or

individual is a mammal. In one embodiment, the patient, subject or individual is a human. In some embodiments the patient, subject or individual is an animal, such as, but not limited to, domesticated animals, such as equine, bovine, murine, ovine, canine, and feline.

[0059] The terms “protein transduction domain (PTD)” or “transporter peptide sequence” (also known as cell permeable proteins (CPP) or membrane translocating sequences (MTS)) are used interchangeably herein to refer to small peptides that are able to ferry much larger molecules into cells independent of classical endocytosis. In some embodiments, a nuclear localization signal can be found within the protein transduction domain, which mediates further translocation of the molecules into the cell nucleus.

[0060] The terms “treating” or “treatment” as used herein covers the treatment of a disease in a subject, such as a human, and includes: (i) inhibiting a disease, *i.e.*, arresting its development; (ii) relieving a disease, *i.e.*, causing regression of the disease; (iii) slowing progression of the disease; and/or (iv) inhibiting, relieving, or slowing progression of one or more symptoms of the disease. With respect to a cancer, “treating” or “treatment” also encompasses regression of a tumor, slowing tumor growth, inhibiting metastasis of a tumor, inhibiting relapse or recurrent cancer and/or maintaining remission.

[0061] It is also to be appreciated that the various modes of treatment or prevention of medical diseases and conditions as described are intended to mean “substantial,” which includes total but also less than total treatment or prevention, and wherein some biologically or medically relevant result is achieved. The treatment can be a continuous prolonged treatment for a chronic disease or a single, or few time administrations for the treatment of an acute condition.

[0062] The term “therapeutic” as used herein means a treatment and/or prophylaxis. A therapeutic effect is obtained by suppression, remission, or eradication of a disease state.

II. Overview

[0063] The present disclosure relates, in part, to the treatment of cancer in a subject by administering a composition comprising one or more immune cells having anti-tumor activity (e.g., immune cells that modulate a response against a tumor, such as tumor-infiltrating lymphocytes (TILs)), wherein the one or more immune cells are contacted with a PTD-MYC fusion polypeptide *in vitro* prior to administration to the subject. In some embodiments, the

immune cells are obtained from a donor subject that has a tumor. In some embodiments, the cells are autologous to the subject receiving treatment. In some embodiments, the tumor is a melanoma tumor.

[0064] The present disclosure is based, at least in part, on the discovery, that treating lymphocytes isolated from a donor subject having a melanoma tumor with a MYC fusion polypeptide containing a MYC polypeptide and a protein transduction domain (PTD), such as the HIV TAT protein transduction domain, and administering the treated lymphocytes to a subject bearing a melanoma tumor significantly increases the survival of the tumor-bearing subject. The examples provided herein demonstrate that immune cells extracted from the lymph nodes of a melanoma-bearing mouse had significantly increased therapeutic efficacy when the cells were treated with a TAT-MYC fusion protein *in vitro* prior to administration to a second melanoma-bearing mice. These data support that adoptive cell transfer using anti-tumor immune cells treated with a PTD-MYC fusion polypeptide can be employed in the treatment of cancers, such as melanoma.

[0065] In some embodiments, the method for the treatment of cancer in a subject comprises administering immune cells that have been contacted *in vitro* with a PTD-MYC fusion polypeptide. In some embodiments, the immune cells for use in the present methods are primed *in vivo* with tumor antigen. In some embodiments, the immune cells are from a donor having cancer. In some embodiments, the immune cells are from a donor having a solid tumor, such as a melanoma, carcinoma, adenoma, adenocarcinoma, blastoma, sarcoma, or lymphoma. In some embodiments, the immune cells are contacted *in vivo* with a tumor antigen. In some embodiments, the immune cells are from a donor that has been exposed to a one or more tumor antigens. In some embodiments, the immune cells are from a donor that has been exposed to an anti-tumor vaccine. In some embodiments, the immune cells are B cells, T cells, NK cells, or any combination thereof. In some embodiments, the immune cells are tumor infiltrating lymphocytes (TIL). In some embodiments, the immune cells are chimeric antigen receptor (CAR)-T cells.

[0066] In some embodiments, the method for the treatment of cancer in a subject comprises administering one or more modified immune cells to the subject in need thereof, wherein the one or more modified immune cells comprise a MYC fusion peptide comprising (i) a protein

transduction domain; (ii) a MYC polypeptide sequence and are reactive to a tumor-specific antigen.

[0067] In some embodiments, the method for the treatment of cancer in a subject comprises the steps of:

a) contacting immune cells *in vitro* with a MYC fusion polypeptide, wherein the immune cells are from a donor that has been exposed to one or more tumor antigens and the MYC fusion peptide comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence; and

b) administering the contacted immune cells to the cancer-bearing subject, whereby the cancer is treated.

[0068] In some embodiments, contacting the immune cells *in vitro* with a PTD-MYC fusion polypeptide is performed by culturing the immune cells in the presence of the MYC fusion polypeptide. In some embodiments, the immune cells are cultured in the presence of one or more cytokines and/or growth factors (e.g., interleukin-2 (IL-2), IL-4, IL-7, IL-9, and IL-15). In some embodiments, the immune cells are not expanded prior to administration. In some embodiments, the immune cells are expanded prior to administration. In some embodiments, the donor and subject for treatment are the same.

[0069] In some embodiments, the immune cells are tumor-infiltrating lymphocytes. In some embodiments, the tumor-infiltrating lymphocytes are autologous tumor-infiltrating lymphocytes. Accordingly, in some embodiments, the method for the treatment of cancer in a subject comprises administering lymphocytes that have been contacted *in vitro* with a PTD-MYC fusion polypeptide, wherein the immune cells are from lymphocytes are autologous tumor-infiltrating lymphocytes from the subject.

[0070] In some embodiments, the method for the treatment of cancer in a subject comprises the steps of:

a) contacting lymphocytes *in vitro* with a PTD-MYC fusion polypeptide, wherein the lymphocytes are autologous tumor-infiltrating lymphocytes from the subject, and

b) administering the contacted autologous tumor-infiltrating lymphocytes to the subject, whereby the cancer is treated.

Methods of Obtaining and Preparing Immune Cells For Transfer

[0071] Immune cells for use in the methods provided herein can be obtained using any suitable method known in the art. In some embodiments, the immune cells are primary immune cells. In some embodiments, the immune cells are lymphocytes, such as T and B cells. In some embodiments, the immune cells are natural killer (NK) cells. In some embodiments, the immune cells are a mixture of lymphocytes and NK cells. In some embodiments, the immune cells are peripheral blood mononuclear cells (PBMC). In some embodiments, the immune cells are T cells that have infiltrated a tumor (e.g., tumor infiltrating lymphocytes). In some embodiments, the T cells are removed during surgery of a tumor. For example, in some embodiments, the T cells are isolated after removal of tumor tissue by biopsy. In some embodiments, the immune cells are modified following isolation from a donor. In some embodiments, the immune cells are chimeric antigen receptor (CAR)-T cells.

[0072] In some embodiments, the T cells are isolated from sample containing a population of cells, such as a blood, lymph or tissue biopsy sample. T cells can be isolated from a population of cells by any means known in the art. In one embodiment, the method comprises obtaining a bulk population of T cells from a tumor sample by any suitable method known in the art. For example, a bulk population of T cells can be obtained from a tumor sample by dissociating the tumor sample into a cell suspension from which specific cell populations can be selected. Suitable methods of obtaining a bulk population of T cells can include, but are not limited to, any one or more of mechanically dissociating (e.g., mincing) the tumor, enzymatically dissociating (e.g., digesting) the tumor, and aspiration (e.g., as with a needle).

[0073] The bulk population of T cells obtained from a tumor sample can comprise any suitable type of T cell. Preferably, the bulk population of T cells obtained from a tumor sample comprises tumor infiltrating lymphocytes (TILs).

[0074] The tumor sample can be obtained from any mammal. Unless stated otherwise, as used herein, the term "mammal" refers to any mammal including, but not limited to, mammals of the order Logomorpha, such as rabbits; the order Carnivora, including Felines (cats) and Canines (dogs); the order Artiodactyla, including Bovines (cows) and Swines (pigs); or of the order Perssodactyla, including Equines (horses). The mammals can be non-human primates, e.g., of the order Primates, Ceboids, or Simoids (monkeys) or of the order Anthropoids (humans and apes).

In some embodiments, the mammal can be a mammal of the order Rodentia, such as mice and hamsters. Preferably, the mammal is a non-human primate or a human. An exemplary mammal is a human. In some embodiments, the subject to receive the immune cells is also the donor of the tumor sample (i.e., autologous ACT)

[0075] T cells can be obtained from a number of sources, including peripheral blood mononuclear cells, bone marrow, lymph node tissue, spleen tissue, and tumors. In certain embodiments, T cells can be obtained from a unit of blood collected from a subject using any number of techniques known to the skilled artisan, such as Ficoll separation. In one embodiment, cells from the circulating blood of an individual are obtained by apheresis or leukopheresis. The apheresis product typically contains lymphocytes, including T cells, monocytes, granulocytes, B cells, other nucleated white blood cells, red blood cells, and platelets. In one embodiment, the cells collected by apheresis can be washed to remove the plasma fraction and to place the cells in an appropriate buffer or media for subsequent processing steps. In one embodiment of the invention, the cells are washed with phosphate buffered saline (PBS). In an alternative embodiment, the wash solution lacks calcium and can lack magnesium or can lack many if not all divalent cations. Initial activation steps in the absence of calcium lead to magnified activation. As those of ordinary skill in the art would readily appreciate, a washing step can be accomplished by methods known to those in the art, such as by using a semi-automated “flow-through” centrifuge (for example, the Cobe 2991 cell processor) according to the manufacturer's instructions. After washing, the cells can be resuspended in a variety of biocompatible buffers, such as, for example, Ca-free, Mg-free PBS. Alternatively, the undesirable components of the apheresis sample can be removed and the cells directly resuspended in culture media.

[0076] In another embodiment, T cells are isolated from peripheral blood lymphocytes by lysing the red blood cells and depleting the monocytes, for example, by centrifugation through a PERCOLL™ gradient. A specific subpopulation of T cells, such as CD28+, CD4+, CDC, CD45RA+, and CD45RO+ T cells, can be further isolated by positive or negative selection techniques. For example, in one embodiment, T cells are isolated by incubation with anti-CD3/anti-CD28 (i.e., 3×28)-conjugated beads, such as DYNABEADS® M-450 CD3/CD28 T, or XCYTE DYNABEADS™ for a time period sufficient for positive selection of the desired T

cells. In one embodiment, the time period is about 30 minutes. In a further embodiment, the time period ranges from 30 minutes to 36 hours or longer and all integer values there between. In a further embodiment, the time period is at least 1, 2, 3, 4, 5, or 6 hours. In yet another embodiment, the time period is 10 to 24 hours. In one embodiment, the incubation time period is 24 hours. For isolation of T cells from patients with leukemia, use of longer incubation times, such as 24 hours, can increase cell yield. Longer incubation times can be used to isolate T cells in any situation where there are few T cells as compared to other cell types, such in isolating tumor infiltrating lymphocytes (TIL) from tumor tissue or from immunocompromised individuals. Further, use of longer incubation times can increase the efficiency of capture of CD8+ T cells.

[0077] Enrichment of a T cell population by negative selection can be accomplished with a combination of antibodies directed to surface markers unique to the negatively selected cells. In one embodiment, the method is cell sorting and/or selection via negative magnetic immunoadherence or flow cytometry that uses a cocktail of monoclonal antibodies directed to cell surface markers present on the cells negatively selected. For example, to enrich for CD4+ cells by negative selection, a monoclonal antibody cocktail typically includes antibodies to CD14, CD20, CD11b, CD16, HLA-DR, and CD8.

[0078] Further, monocyte populations (i.e., CD14+ cells) can be depleted from blood preparations by a variety of methodologies, including anti-CD14 coated beads or columns, or utilization of the phagocytotic activity of these cells to facilitate removal. Accordingly, in one embodiment, the invention uses paramagnetic particles of a size sufficient to be engulfed by phagocytotic monocytes. In certain embodiments, the paramagnetic particles are commercially available beads, for example, those produced by Life Technologies under the trade name Dynabeads™. In one embodiment, other non-specific cells are removed by coating the paramagnetic particles with “irrelevant” proteins (e.g., serum proteins or antibodies). Irrelevant proteins and antibodies include those proteins and antibodies or fragments thereof that do not specifically target the T cells to be isolated. In certain embodiments the irrelevant beads include beads coated with sheep anti-mouse antibodies, goat anti-mouse antibodies, and human serum albumin.

[0079] In brief, such depletion of monocytes is performed by preincubating T cells isolated from whole blood, apheresed peripheral blood, or tumors with one or more varieties of irrelevant or non-antibody coupled paramagnetic particles at any amount that allows for removal of monocytes (approximately a 20:1 bead:cell ratio) for about 30 minutes to 2 hours at 22 to 37 degrees C., followed by magnetic removal of cells which have attached to or engulfed the paramagnetic particles. Such separation can be performed using standard methods available in the art. For example, any magnetic separation methodology can be used including a variety of which are commercially available, (e.g., DYNAL® Magnetic Particle Concentrator (DYNAL MPC®)). Assurance of requisite depletion can be monitored by a variety of methodologies known to those of ordinary skill in the art, including flow cytometric analysis of CD14 positive cells, before and after depletion.

[0080] For isolation of a desired population of cells by positive or negative selection, the concentration of cells and surface (e.g., particles such as beads) can be varied. In certain embodiments, it can be desirable to significantly decrease the volume in which beads and cells are mixed together (i.e., increase the concentration of cells), to ensure maximum contact of cells and beads. For example, in one embodiment, a concentration of 2 billion cells/ml is used. In one embodiment, a concentration of 1 billion cells/ml is used. In a further embodiment, greater than 100 million cells/ml is used. In a further embodiment, a concentration of cells of 10, 15, 20, 25, 30, 35, 40, 45, or 50 million cells/ml is used. In yet another embodiment, a concentration of cells from 75, 80, 85, 90, 95, or 100 million cells/ml is used. In further embodiments, concentrations of 125 or 150 million cells/ml can be used. Using high concentrations can result in increased cell yield, cell activation, and cell expansion. Further, use of high cell concentrations allows more efficient capture of cells that can weakly express target antigens of interest, such as CD28-negative T cells, or from samples where there are many tumor cells present (e.g., leukemic blood, tumor tissue). Such populations of cells can have therapeutic value and would be desirable to obtain. For example, using high concentration of cells allows more efficient selection of CD8+ T cells that normally have weaker CD28 expression.

[0081] In a related embodiment, it can be desirable to use lower concentrations of cells. By significantly diluting the mixture of T cells and surface (e.g., particles such as beads),

interactions between the particles and cells is minimized. This selects for cells that express high amounts of desired antigens to be bound to the particles. For example, CD4⁺ T cells express higher levels of CD28 and are more efficiently captured than CD8⁺ T cells in dilute concentrations. In one embodiment, the concentration of cells used is 5×10^6 /ml. In other embodiments, the concentration used can be from about 1×10^5 /ml to 1×10^6 /ml, and any integer value in between.

[0082] T cells can also be frozen. The freeze and subsequent thaw step can provide a more uniform product by removing granulocytes and to some extent monocytes in the cell population. After a washing step to remove plasma and platelets, the cells can be suspended in a freezing solution. While many freezing solutions and parameters are known in the art and will be useful in this context, one method involves using PBS containing 20% DMSO and 8% human serum albumin, or other suitable cell freezing media, the cells then are frozen to -80°C at a rate of 1° per minute and stored in the vapor phase of a liquid nitrogen storage tank. Other methods of controlled freezing can be used as well as uncontrolled freezing immediately at -20°C . or in liquid nitrogen.

[0083] T cells for use in the present invention can also be antigen-specific T cells. For example, tumor-specific T cells can be used. In certain embodiments, antigen-specific T cells can be isolated from a patient of interest, such as a patient afflicted with a cancer, such as patient with a tumor. In some embodiments, the patient has melanoma.

[0084] In one embodiment neoepitopes are determined for a subject and T cells specific to these antigens are isolated. Antigen-specific cells for use in expansion can also be generated *in vitro* using any number of methods known in the art, for example, as described in U.S. Patent Publication No. US 20040224402 entitled, Generation And Isolation of Antigen-Specific T Cells, or in U.S. Pat. Nos. 6,040,177. Antigen-specific cells for use in the present invention can also be generated using any number of methods known in the art, for example, as described in Current Protocols in Immunology, or Current Protocols in Cell Biology, both published by John Wiley & Sons, Inc., Boston, Mass.

[0085] In a related embodiment, it can be desirable to sort or otherwise positively select (e.g. via magnetic selection) the antigen specific cells prior to or following one or two rounds of

expansion. Sorting or positively selecting antigen-specific cells can be carried out using peptide-MHC tetramers (Altman, *et al.*, 1996 *Science*. Oct. 4; 274(5284):94-6). In another embodiment the adaptable tetramer technology approach is used (Andersen *et al.*, 2012 *Nat Protoc*. 7:891-902). Tetramers are limited by the need to utilize predicted binding peptides based on prior hypotheses, and the restriction to specific HLAs. Peptide-MHC tetramers can be generated using techniques known in the art and can be made with any MHC molecule of interest and any antigen of interest as described herein. Specific epitopes to be used in this context can be identified using numerous assays known in the art. For example, the ability of a polypeptide to bind to MHC class I can be evaluated indirectly by monitoring the ability to promote incorporation of ¹²⁵I labeled β 2-microglobulin (β 2m) into MHC class I/ β 2m/peptide heterotrimeric complexes (see Parker *et al.* 1994, *J. Immunol.* 152:163.).

[0086] In some embodiments, the T cells are recombinantly modified to express a modified or chimeric receptor (e.g., chimeric antigen receptor (CAR) modified T cells).

[0087] In one embodiment, cells are directly labeled with an epitope-specific reagent for isolation by flow cytometry followed by characterization of phenotype and TCRs. In one embodiment, T cells are isolated by contacting the T cell specific antibodies. Sorting of antigen-specific T cells, or generally any cells of the present invention, can be carried out using any of a variety of commercially available cell sorters, including, but not limited to, MoFlo sorter (DakoCytomation, Fort Collins, Colo.), FACS Aria™, FACSAria™, FACSVantage™, BD™ LSR II, and FACSCalibur™ (BD Biosciences, San Jose, Calif.).

[0088] In one embodiment, the method comprises selecting cells that also express CD3. The method can comprise specifically selecting the cells in any suitable manner. Preferably, the selecting is carried out using flow cytometry. The flow cytometry can be carried out using any suitable method known in the art. The flow cytometry can employ any suitable antibodies and stains. Preferably, the antibody is chosen such that it specifically recognizes and binds to the particular biomarker being selected. For example, the specific selection of CD3, CD8, TIM-3, LAG-3, 4-1BB, or PD-1 can be carried out using anti-CD3, anti-CD8, anti-TIM-3, anti-LAG-3, anti-4-1BB, or anti-PD-1 antibodies, respectively. The antibody or antibodies can be conjugated to a bead (e.g., a magnetic bead) or to a fluorochrome. Preferably, the flow cytometry is

fluorescence-activated cell sorting (FACS). TCRs expressed on T cells can be selected based on reactivity to autologous tumors. Additionally, T cells that are reactive to tumors can be selected for based on markers using the methods described in patent publication Nos. WO2014133567 and WO2014133568, herein incorporated by reference in their entirety. Additionally, activated T cells can be selected for based on surface expression of CD107a.

[0089] In one embodiment, the method further comprises expanding the numbers of T cells in the enriched cell population. Such methods are described in U.S. Patent No. 8,637,307 and is herein incorporated by reference in its entirety. The T cells can be expanded before or after treatment of the cells with the PTD-MYC polypeptide. The numbers of T cells can be increased at least about 3-fold (or 4-, 5-, 6-, 7-, 8-, or 9-fold), more preferably at least about 10-fold (or 20-, 30-, 40-, 50-, 60-, 70-, 80-, or 90-fold), more preferably at least about 100-fold, more preferably at least about 1,000 fold, or most preferably at least about 100,000-fold. The numbers of T cells can be expanded using any suitable method known in the art. Exemplary methods of expanding the numbers of cells are described in patent publication No. WO 2003057171, U.S. Patent No. 8,034,334, and U.S. Patent Application Publication No. 2012/0244133, each of which is incorporated herein by reference.

[0090] In one embodiment, *ex vivo* T cell expansion can be performed by isolation of T cells and subsequent stimulation or activation followed by further expansion. In one embodiment of the invention, the T cells can be stimulated or activated by a single agent. In another embodiment, T cells are stimulated or activated with two agents, one that induces a primary signal and a second that is a co-stimulatory signal. Ligands useful for stimulating a single signal or stimulating a primary signal and an accessory molecule that stimulates a second signal can be used in soluble form. Ligands can be attached to the surface of a cell, to an Engineered Multivalent Signaling Platform (EMSP), or immobilized on a surface. In a one embodiment both primary and secondary agents are co-immobilized on a surface, for example a bead or a cell. In one embodiment, the molecule providing the primary activation signal can be a CD3 ligand, and the co-stimulatory molecule can be a CD28 ligand or 4-1BB ligand. In some embodiments, the cells are expanded by stimulation with one or more antigens, such as a melanoma tumor antigen or antigens derived from the patient's tumor.

[0091] In some embodiments, the isolated immune cells are immediately treated with the PTD-MYC fusion polypeptide following isolation. In other embodiments, the isolated immune cells are stored in a suitable buffer and frozen prior to treatment with the PTD-MYC fusion polypeptide. In some embodiments, the isolated immune cells are immediately treated with the PTD-MYC fusion polypeptide following isolation and the treated cells are stored in a suitable buffer and frozen until needed for administration to the patient.

[0092] In certain embodiments, the isolated immune cells (e.g., a mixed population immune cells or isolated types, such as tumor infiltrating lymphocytes) are contacted with a composition containing a PTD-MYC fusion polypeptide for a period of time sufficient to be taken up by the cells. In some embodiments, the immune cells are contacted with a composition containing a PTD-MYC fusion polypeptide for less than about 24 hours, less than about 23 hours, less than about 22 hours, less than about 21 hours, less than about 20 hours, less than about 19 hours, less than about 18 hours, less than about 17 hours, less than about 16 hours, less than about 15 hours, less than about 14 hours, less than about 13 hours, less than about 12 hours, less than about 11 hours, less than about 10 hours, less than about 9 hours, less than about 8 hours, less than about 7 hours, less than about 6 hours, less than about 5 hours, less than about 4 hours, less than about 3 hours, less than about 2 hours, or less than about 1 hour.

[0093] In certain embodiments, the immune cells are contacted with a composition containing a PTD-MYC fusion polypeptide for less than about 55 minutes, less than about 50 minutes, less than about 45 minutes, less than about 40 minutes, less than about 35 minutes, less than about 30 minutes, less than about 29 minutes, less than about 28 minutes, less than about 27 minutes, less than about 26 minutes, less than about 25 minutes, less than about 24 minutes, less than about 23 minutes, less than about 22 minutes, less than about 21 minutes, less than about 20 minutes, less than about 19 minutes, less than about 18 minutes, less than about 17 minutes, less than about 16 minutes, less than about 15 minutes, less than about 14 minutes, less than about 13 minutes, less than about 12 minutes, less than about 11 minutes, or less than about 10 minutes. In certain embodiments, the immune cells are contacted with a composition containing a PTD-MYC fusion polypeptide for about 1 hour.

[0094] In certain embodiments, the immune cells are contacted with a composition containing a PTD-MYC fusion polypeptide for 24 hours or longer. In certain embodiments, the immune cells are contacted with a composition containing a PTD-MYC fusion polypeptide for less than about 12 days, less than about 11 days, less than about 10 days, less than about 9 days, less than about 8 days, less than about 7 days, less than about 6 days, less than about 5 days, less than about 4 days, less than about 2 days, or less than about 1 day.

[0095] In certain embodiments that may be combined with any of the preceding embodiments, the cells are contacted with a MYC-fusion polypeptide at a concentration of 0.5 µg/ml to 500 µg/ml. 0.5 µg/ml, at least 0.6 µg/ml, at least 0.7 µg/ml, at least 0.8 µg/ml, at least 0.9 µg/ml, at least 1 µg/ml, at least 2 µg/ml, at least 3 µg/ml, at least 4 µg/ml, at least 5 µg/ml, at least 6 µg/ml, at least 7 µg/ml, at least 8 µg/ml, at least 9 µg/ml, at least 10 µg/ml, at least 15 µg/ml, at least 20 µg/ml, at least 25 µg/ml, at least 30 µg/ml, at least 35 µg/ml, at least 40 µg/ml, at least 45 µg/ml, at least 50 µg/ml, at least 55 µg/ml, at least 60 µg/ml, at least 65 µg/ml, at least 70 µg/ml, at least 75 µg/ml, at least 80 µg/ml, at least 85 µg/ml, at least 90 µg/ml, at least 95 µg/ml, or at least 100 µg/ml.

MYC fusion proteins

[0096] In some embodiments, the PTD-MYC fusion polypeptide comprises a protein transduction domain (PTD), a MYC polypeptide that promotes one or more of cell survival or proliferation, and optionally a protein tag domain, *e.g.*, one or more amino acid sequences that facilitate purification of the fusion protein. In some embodiments, a cell contacted with MYC polypeptide exhibits increased survival time (*e.g.*, as compared to an identical or similar cell of the same type that was not contacted with MYC), and/or increased proliferation (*e.g.*, as compared to an identical or similar cell of the same type that was not contacted with MYC).

[0097] In some embodiments, the fusion protein comprises (a) a protein transduction domain; and (b) a MYC polypeptide sequence. In some embodiments, the fusion peptide is a peptide of Formula (I):

protein transduction domain-MYC polypeptide sequence.

[0098] In some embodiments, a fusion peptide disclosed herein comprises (a) a protein transduction domain; (b) a MYC polypeptide sequence; and (c) one or more molecules that link

the protein transduction domain and the MYC polypeptide sequence. In some embodiments, the fusion peptide is a peptide of Formula (II):

protein transduction domain-X-MYC polypeptide sequence,

wherein -X- is molecule that links the protein transduction domain and the MYC polypeptide sequence. In some embodiments, -X- is at least one amino acid.

[0099] In some embodiments, a fusion peptide disclosed herein comprises (a) a protein transduction domain; (b) a MYC polypeptide sequence; (c) at least two protein tags; and (d) optionally linker(s). In some embodiments, the fusion peptide is a peptide of Formula (III-VI):

protein transduction domain-X-MYC polypeptide sequence-X-protein tag 1-X-protein tag 2
(Formula (III)), or

protein transduction domain-MYC polypeptide sequence-X-protein tag 1-X-protein tag 2
(Formula (IV)), or

protein transduction domain-MYC polypeptide sequence-protein tag 1-X-protein tag 2
(Formula (V)), or

protein transduction domain-MYC polypeptide sequence-protein tag 1-protein tag 2
(Formula (VI)),

wherein -X- is a linker. In some embodiments, -X- is one or more amino acids.

[0100] In some embodiments, a fusion peptide disclosed herein comprises (a) a protein transduction domain; (b) a MYC polypeptide sequence; (c) a 6-histidine tag; (d) a V5 epitope tag; and (e) optionally linker(s). In some embodiments, the fusion peptide is a peptide of Formula (VII-XIV):

protein transduction domain-X-MYC polypeptide sequence-X-6-histidine tag-X-V5 epitope tag
(Formula (VII)), or

protein transduction domain-MYC polypeptide sequence-X-6-histidine tag-X-V5 epitope tag
(Formula (VIII)), or

protein transduction domain-MYC polypeptide sequence-6-histidine tag-X-V5 epitope tag
(Formula (IX)), or

protein transduction domain-MYC polypeptide sequence-6-histidine tag-V5 epitope tag
(Formula (X)),

protein transduction domain-X-MYC polypeptide sequence-X-V5 epitope tag-X-6-histidine tag

(Formula (XI)), or

protein transduction domain-MYC polypeptide sequence-X-V5 epitope tag-X-6-histidine tag

(Formula (XII)), or

protein transduction domain-MYC polypeptide sequence-V5 epitope tag-X-6-histidine tag

(Formula (XIII)), or

protein transduction domain-MYC polypeptide sequence-V5 epitope tag-6-histidine tag

(Formula (XIV)),

wherein -X- is a linker. In some embodiments, -X- is one or more amino acids.

[0101] As noted above, in some embodiments, the MYC fusion protein comprises one or more linker sequences. The linker sequences can be employed to link the protein transduction domain, MYC polypeptide sequence, V5 epitope tag and/or 6-histidine tag of the fusion protein. In some embodiments, the linker comprises one or more amino acids. In some embodiments, the amino acid sequence of the linker comprises KGELNSKLE. In some embodiments, the linker comprises the amino acid sequence of RTG.

[0102] Protein Transduction Domain (PTD)

[0103] In some embodiments, the MYC fusion protein includes a protein transduction domain. Peptide transport provides an alternative for delivery of small molecules, proteins, or nucleic acids across the cell membrane to an intracellular compartment of a cell. One non-limiting example and well-characterized protein transduction domain (PTD) is a TAT-derived peptide. Frankel *et al.*, (see, e.g., U.S. Pat. No. 5,804,604, U.S. Pat. No. 5,747,641, U.S. Pat. No. 5,674,980, U.S. Pat. No. 5,670,617, and U.S. Pat. No. 5,652,122) demonstrated transport of a cargo protein (β -galactosidase or horseradish peroxidase) into a cell by conjugating a peptide containing amino acids 48–57 of TAT to the cargo protein. In some embodiments, TAT comprises an amino acid sequence of MRKKRRQRRR (SEQ ID NO: 7).

[0104] Another non-limiting example of a PTD is penetratin. Penetratin can transport hydrophilic macromolecules across the cell membrane (Derossi *et al.*, *Trends Cell Biol.*, 8:84–87 (1998) incorporated herein by reference in its entirety). Penetratin is a 16 amino acid peptide

that corresponds to amino acids 43–58 of the homeodomain of Antennapedia, a *Drosophila* transcription factor which is internalized by cells in culture.

[0105] Yet another non-limiting example of a PTD is VP22. VP22, a tegument protein from Herpes simplex virus type 1 (HSV-1), has the ability to transport proteins and nucleic acids across a cell membrane (Elliot *et al.*, *Cell* 88:223–233, 1997, incorporated herein by reference in its entirety). Residues 267–300 of VP22 are necessary but cannot be sufficient for transport. Because the region responsible for transport function has not been identified, the entire VP22 protein is commonly used to transport cargo proteins and nucleic acids across the cell membrane (Schwarze *et al.*, *Trends Pharmacol Sci*, 21:45–48, 2000).

[0106] In some embodiments, the PTD-MYC fusion polypeptide includes a protein transduction domain. By way of example, but not by way of limitation, in some embodiments, the protein transduction domain comprises the protein transduction domain of one or more of TAT, penetratin, VP22, vpr, EPTD, R9, R15, VP16, and Antennapedia. In some embodiments, the protein transduction domain comprises the protein transduction domain of one or more of TAT, penetratin, VP22, vpr, and EPTD. In some embodiments, the protein transduction domain comprises the protein transduction domain of at least one of TAT, penetratin, VP22, vpr, EPTD, R9, R15, VP16, and Antennapedia. In some embodiments, the protein transduction domain comprises a synthetic protein transduction domain (*e.g.*, polyarginine or PTD-5). In particular embodiments, the protein transduction domain comprises a TAT protein transduction domain. In some embodiments, the protein transduction domain is covalently linked to the MYC polypeptide. In some embodiments, the protein transduction domain is linked to the MYC polypeptide via a peptide bond. In some embodiments, the protein transduction domain is linked to the MYC polypeptide via a linker sequence. In some embodiments, the linker comprises a short amino acid sequence. By way of example, but not by way of limitation, in some embodiments, the linker sequences is 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10 amino acids in length.

[0107] The MYC fusion protein of the present technology can be arranged in any desired order. For example, in some embodiments, the MYC fusion protein can be arranged in order of a) the protein transduction domain linked in frame to the MYC polypeptide, b) the MYC polypeptide linked in frame to the V5 domain, and c) the V5 domain linked in frame to the 6-

histidine epitope tag. In some embodiments, the MYC fusion protein has an order of components of a) the MYC polypeptide linked in frame to the protein transduction domain, b) the protein transduction domain linked in frame to the V5 domain, and c) the V5 domain linked in frame to the 6-histidine epitope tag. In some embodiments, additional amino acid sequences can be included between each of the sequences. In some embodiments, additional amino acids can be included at the start and/or end of the polypeptide sequences.

[0108] In some embodiments, the protein transduction domain is a TAT protein transduction domain. In some embodiments, the protein transduction domain is TAT_[48-57]. In some embodiments, the protein transduction domain is TAT_[57-48].

[0109] Protein Tag Domains

[0110] In some embodiments, the MYC fusion protein comprises a protein tag domain that comprises one or more amino acid sequences that facilitate purification of the fusion protein. In some embodiments, the protein tag domain comprises one or more of a polyhistidine tag, and an epitope tag. By way of example, but not by way of limitation, exemplary tags include one or more of a V5, a histidine-tag (*e.g.*, a 6-histidine tag), HA (hemagglutinin) tags, FLAG tag, CBP (calmodulin binding peptide), CYD (covalent yet dissociable NorpD peptide), StrepII, or HPC (heavy chain of protein C). In some embodiments, the protein tag domain comprise about 10 to 20 amino acids in length. In some embodiments, the protein tag domain comprises 2 to 40 amino acids in length, for example 6-20 amino acids in length. In some embodiments, two of the above listed tags (for example, V5 and the HIS-tag) are used together to form the protein tag domain.

[0111] In some embodiments, the histidine tag is a 6-histidine tag. In some embodiments, the histidine tag comprises the sequence HHHHHH (SEQ ID NO:8). In some embodiments, the fusion peptide disclosed herein comprises a V5 epitope tag. In some embodiments, the V5 tag comprises the amino acid sequence of: GKPIPNPLLGLDST (SEQ ID NO:9). In some embodiments, the V5 tag comprises the amino acid sequence of IPNPLLGLD (SEQ ID NO:10).

[0112] The protein tags can be added to the fusion protein disclosed herein by any suitable method. By way of example, but not by way of limitation, in some embodiments, a TAT-MYC polypeptide sequence is cloned into an expression vector encoding one or more protein tags, *e.g.*,

a polyHis-tag and/or a V5 tag. In some embodiments, a polyhistidine tag and/or a V5 tag is added by PCR (*i.e.*, the PCR primers comprise a polyhistidine sequence and/ or V5 sequence).

[0113] Construction of PTD-MYC fusion polypeptides

[0114] PTD-MYC fusion polypeptides (*e.g.*, TAT-MYC fusion polypeptide) disclosed herein can be constructed by methods well known in the art. By way of example, but not by way of limitation, a nucleotide sequence encoding a TAT-MYC fusion polypeptide can be generated by PCR. In some embodiments, a forward primer for a human MYC sequence comprises an in frame N-terminal 9-amino-acid sequence of the TAT protein transduction domain (*e.g.*, RKKRRQRRR). In some embodiments, a reverse primer for a human MYC sequence is designed to remove the stop codon. In some embodiments, the PCR product is cloned into any suitable expression vector. In some embodiments, the expression vector comprises a polyhistidine tag and a V5 tag.

[0115] In some embodiments, a fusion peptide disclosed herein comprises (a) TAT, and (b) c-MYC . In some embodiments, a fusion peptide disclosed herein comprises (a) TAT_[48-57], and (b) c-MYC . In some embodiments, a fusion peptide disclosed herein comprises (a) TAT_[57-48], and (b) c-MYC .

[0116] In some embodiments, a fusion peptide disclosed herein comprises (a) TAT, (b) c-MYC , (c) linker(s), (d) V5 tag, and (e) 6-histidine tag. In some embodiments, a fusion peptide disclosed herein comprises (a) TAT_[48-57], (b) c-MYC , (c) linker(s), (d) V5 tag, and (e) 6-histidine tag. In some embodiments, a fusion peptide disclosed herein comprises (a) TAT_[57-48], (b) c-MYC , (c) linker(s), (d) V5 tag, and (e) 6-histidine tag.

[0117] In some embodiments, the PTD-MYC fusion polypeptide comprises SEQ ID NO: 1; in some embodiments, the PTD-MYC fusion polypeptide is SEQ ID NO: 1.

MRKKRRQRRRPLNVSFTNRNYDL DYDSVQPYFYCDEENFYQQQQQSELPAPSEDI
WKKFELLPTPPLSPSRRLCSPSYVAVTPFSLRGDNDGGGGSFSTADQLEMVTELLGG
DMVNQSFICDPDDETFIKNIIHQDCMWSGFSAAKLVSEKLASYQAARKDSGSPNPARG
HSVCSTSSLYLQDL SAAASECIDPSVFPYPLNDSSSPKSCASQDSSAFSPSSD SLLSSTES
SPQGSPEPLVLHEETPPTTSSDSEEEQEDEEEIDVVSVEKRQAPGKRSESGSPSAGGH SKP
PHSPLVLKRCHVSTHQHNYAAPSTRKDYPAAKRVKLDSVRVLRQISNNRKCTSPRSSD

TEENVKRRTHNVLERQRRNELKRSFFALRDQIPELENNEKAPKVVLKKATAYILSVQAE
EQKLISEEDLLRKRRRELKHKLEQLRKGELNSKLEGKPIPNNLLGLDSTRTGHHHHHHH
(SEQ ID NO: 1).

[0118] The fusion protein can be modified during or after synthesis to include one or more functional groups. By way of example but not by way of limitation, the protein can be modified to include one or more of an acetyl, phosphate, acetate, amide, alkyl, and/or methyl group. This list is not intended to be exhaustive, and is exemplary only. In some embodiments, the protein includes at least one acetyl group.

[0119] A PTD-MYC fusion polypeptide can be generated by any suitable method known the art, e.g. by recombinant protein expression in a cell, such as a bacterial cell, an insect cell, or mammalian cell. In some embodiments, a PTD-MYC fusion polypeptide is recombinantly produced by microbial fermentation. In some embodiments microbial fermentation is performed in a fermentation volume of from about 1 to about 10,000 liters, for example, a fermentation volume of about 10 to about 1000 liters. The fermentation can utilize any suitable microbial host cell and culture medium. In exemplary embodiments, *E. coli* is utilized as the microbial host cell. In alternative embodiments, other microorganisms can be used, e.g., *S. cerevisiae*, *P. pastoris*, *Lactobacilli*, *Bacilli* and *Aspergilli*. In an exemplary embodiment the microbial host cell is BL-21 StarTM *E. coli* strain (Invitrogen). In an exemplary embodiment the microbial host cell is BLR DE3 *E.coli* strain.

[0120] In some embodiments the host cells are modified to provide tRNAs for rare codons, which are employed to overcome host microbial cell codon bias to improve translation of the expressed proteins. In exemplary embodiments, the host cells (e.g., *E. coli*) transformed with a plasmid, such as pRARE (CamR), which express tRNAs for AGG, AGA, AUA, CUA, CCC, GGA codons. Additional, suitable plasmids or constructs for providing tRNAs for particular codons are known in the art and can be employed in the methods provided.

[0121] Integrative or self-replicative vectors can be used for the purpose of introducing the PTD-MYC fusion polypeptide expression cassette into a host cell of choice. In an expression cassette, the coding sequence for the PTD-MYC fusion polypeptide is operably linked to promoter, such as an inducible promoter. Inducible promoters are promoters that initiate

increased levels of transcription from DNA under their control in response to some change in culture conditions, *e.g.*, the presence or absence of a nutrient or a change in temperature. In some embodiments, the nucleic acid encoding the PTD-MYC fusion polypeptide is codon optimized for bacterial expression.

[0122] Exemplary promoters that are recognized by a variety of potential host cells are well known. These promoters can be operably linked to PTD-MYC fusion polypeptide-encoding DNA by removing the promoter from the source DNA, if present, by restriction enzyme digestion and inserting the isolated promoter sequence into the vector. Promoters suitable for use with microbial hosts include, but are not limited to, the β -lactamase and lactose promoter systems (Chang *et al.*, (1978) *Nature*, 275:617-624; Goeddel *et al.*, (1979) *Nature*, 281: 544), alkaline phosphatase, a tryptophan (trp) promoter system (Goeddel (1980) *Nucleic Acids Res.* 8: 4057; EP 36,776), and hybrid promoters such as the tac promoter (deBoer *et al.*, (1983) *Proc. Natl. Acad. Sci. USA* 80: 21-25). Any promoter for suitable for expression by the selected host cell can be used. Nucleotide sequences for suitable are published, thereby enabling a skilled worker operably to ligate them to DNA encoding PTD-MYC fusion polypeptide (see, *e.g.*, Siebenlist *et al.*, (1980) *Cell* 20: 269) using linkers or adaptors to supply any required restriction sites. In exemplary embodiments, promoters for use in bacterial systems can contain a Shine-Dalgarno (S.D.) sequence operably linked to the coding sequence. In some embodiments, the inducible promoter is the lacZ promoter, which is induced with Isopropyl β -D-1-thiogalactopyranoside (IPTG), as is well-known in the art. Promoters and expression cassettes can also be synthesized *de novo* using well known techniques for synthesizing DNA sequences of interest. In an exemplary embodiment, the expression vector for expression of the PTD-MYC fusion polypeptides herein is pET101/D-Topo (Invitrogen).

[0123] For expression of the PTD-MYC fusion polypeptides, the microbial host containing the expression vector encoding the PTD-MYC fusion polypeptide is typically grown to high density in a fermentation reactor. In some embodiments, the reactor has controlled feeds for glucose. In some embodiments, a fermenter inoculum is first cultured in medium supplemented with antibiotics (*e.g.*, overnight culture). The fermenter inoculum is then used to inoculate the fermenter culture for expression of the protein. At an OD600 of at least about 15, usually at least

about 20, at least 25, at least about 30 or higher, of the fermenter culture, expression of the recombinant protein is induced. In exemplary embodiments, where the inducible promoter is the lacZ promoter, IPTG is added to the fermentation medium to induce expression of the PTD-MYC fusion polypeptide. Generally, the IPTG is added to the fermenter culture at an OD₆₀₀ which represents logarithmic growth phase.

[0124] In certain embodiments of the methods provided, induced protein expression is maintained for around about 2 to around about 5 hours post induction, and can be from around about 2 to around about 3 hours post-induction. Longer periods of induction may be undesirable due to degradation of the recombinant protein. The temperature of the reaction mixture during induction is preferably from about 28°C to about 37°C, usually from about 30°C to about 37°C. In particular embodiments, induction is at about 37°C.

[0125] The PTD-MYC fusion polypeptide is typically expressed as cytosolic inclusion bodies in microbial cells. To harvest inclusion bodies, a cell pellet is collected by centrifugation of the fermentation culture following induction, frozen at -70°C or below, thawed and resuspended in disruption buffer. The cells are lysed by conventional methods, *e.g.*, sonication, homogenization, etc. The lysate is then resuspended in solubilization buffer, usually in the presence of urea at a concentration effective to solubilize proteins, *e.g.*, from around about 5M, 6M, 7M, 8M, 9M or greater. Resuspension may require mechanically breaking apart the pellet and stirring to achieve homogeneity. In some embodiments, the cell pellet is directly resuspended in urea buffer and mixed until homogenous. In some embodiments, the resuspension/solubilization buffer is 8M Urea, 50 mM Phosphate pH 7.5 and the suspension is passed through a homogenizer.

[0126] In some embodiments, the homogenized suspension is sulfonylated. For example, in some embodiments, the homogenized suspension is adjusted to include 200 mM Sodium Sulfite and 10 mM Sodium Tetrathionate. The solution is then mixed at room temperature until homogeneous. The mixed lysate is then mixed for an additional period of time to complete the sulfonylation (*e.g.*, at 2-8°C for ≥ 12 hours). The sulfonylated lysate was then centrifuged for an hour. The supernatant containing the sulfonylated PTD-MYC fusion polypeptides is then

collected by centrifugation and the cell pellet discarded. The supernatant is then passed through a filter, *e.g.*, 0.22 μ m membrane filter to clarify the lysate.

[0127] The solubilized protein is then purified. Purification methods may include affinity chromatography, reverse phase chromatography, gel exclusion chromatography, and the like. In some embodiments, affinity chromatography is used. For example, the protein is provided with an epitope tag or histidine 6 tag for convenient purification. In the present methods, exemplary PTD-MYC fusion polypeptide comprise histidine 6 tag for purification using Ni affinity chromatography using Ni- resin.

[0128] In exemplary embodiments, the Ni- resin column is equilibrated in a buffer containing urea. In some embodiments, the equilibration buffer is 6M Urea, 50 mM Phosphate, 500 mM NaCl, and 10% Glycerol solution. The sulfonlated and clarified supernatant comprising the PTD-MYC fusion polypeptide is then loaded onto the Ni- resin column. The column is then washed with a wash buffer, *e.g.*, 6M Urea, 50mM Phosphate, 10% Glycerol, 500 mM NaCl, pH 7.5. The column was then washed with sequential wash buffers with decreasing salt concentration. For example, exemplary subsequent washed can include 6M Urea, 50mM Phosphate, 10% Glycerol, and 2M NaCl, pH 7.5, followed another wash of 6M Urea, 50mM Phosphate, 10% Glycerol, 50mM NaCl, and 30mM Imidazole, pH 7.5.

[0129] Following sequential application of the wash buffers the PTD-MYC fusion polypeptide is eluted from the column by addition of elution buffer, *e.g.*, 6M Urea, 50mM Phosphate, 10% Glycerol, and 50mM NaCl, pH 7.5 with a gradient from 100 to 300 mM Imidazole, and collecting fractions. The protein containing fractions to be pooled are then filtered through a 0.22 μ m membrane. Assessment of protein yield can be measured using any suitable method, *e.g.*, spectrophotometry at UV wavelength 280.

[0130] In some embodiments, one or more additional purification methods can be employed to further purify the isolated PTD-MYC fusion polypeptides. In exemplary embodiments, the pooled fractions from the Ni-Sepharose chromatography step are further purified by anion exchange chromatography using a Q-Sepharose resin. In some embodiments, the pool is prepared for loading onto the Q-Sepharose column by diluting the samples to the conductivity of the Q sepharose buffer (17.52 +/-1 mS/cm) with the second wash buffer (*e.g.*, 6M Urea, 50mM

Phosphate, 10% Glycerol, 2M NaCl, pH 7.5) from the Ni Sepharose chromatography step. The diluted pool is then loaded onto the Q-Sepharose column, followed by two chase steps using a chase buffer (*e.g.*, 6M Urea, 50mM Phosphate, 300mM NaCl, and 10% Glycerol), with further sequential applications of the chase buffer until the UV trace reaches baseline, indicating that the protein has eluted from the column.

Methods of Treatment

[0131] The PTD-MYC fusion polypeptide-modified immune cells are administered for the treatment of a cancer in a patient. In some embodiments, the patient has a solid tumor. In some embodiments, the patient has a carcinoma, adenoma, adenocarcinoma, blastoma, sarcoma, or lymphoma. In some embodiments, the patient has a metastatic tumor. In some embodiments, the patient has received one or more agents for the treatment of the cancer prior to administration of the PTD-MYC fusion polypeptide-modified immune cells. In some embodiments, the cancer is a relapsed or refractory cancer. In some embodiments, the cancer is resistant to one or more agents for the treatment of the cancer.

[0132] Exemplary tumors in humans for methods of treatment provided herein include, but are not limited to, melanoma, bladder tumor, breast tumor, prostate tumor, carcinoma, basal cell carcinoma, biliary tract cancer, bladder cancer, bone cancer, brain cancer, CNS cancer, glioma tumor, cervical cancer, choriocarcinoma, colon and rectum cancer, connective tissue cancer, cancer of the digestive system, endometrial cancer, esophageal cancer, eye cancer, cancer of the head and neck, gastric cancer, intra-epithelial neoplasm, kidney cancer, larynx cancer, leukemia, liver cancer, lung cancer, lymphoma, Hodgkin's lymphoma, Non-Hodgkin's lymphoma, myeloma, neuroblastoma, oral cavity cancer, ovarian cancer, pancreatic cancer, retinoblastoma, rhabdomyosarcoma, rectal cancer, renal cancer, cancer of the respiratory system, sarcoma, skin cancer, stomach cancer, testicular cancer, thyroid cancer, uterine cancer, and cancer of the urinary system. In some embodiments, the cancer is a metastatic cancer. In some embodiments, the cancer is a relapsed or refractory cancer.

[0133] In some embodiments, administration of the PTD-MYC fusion polypeptide-modified immune cells inhibits growth of a tumor or reduces the volume of a tumor. In some embodiments, administration of the PTD-MYC fusion polypeptide-modified immune cells to a

subject having a cancer alleviates one or more symptoms of the cancer. In some embodiments, administration of the PTD-MYC fusion polypeptide-modified immune cells to a subject having cancer increases the overall survival of the subject. In some embodiments, administration of the PTD-MYC fusion polypeptide-modified immune cells to a subject having cancer increases the regression of the cancer.

[0134] The administration of the PTD-MYC fusion polypeptide-modified immune cells (e.g. PTD-MYC fusion polypeptide treated tumor infiltrating lymphocytes) according to the methods provided herein can be carried out in any suitable manner for administering cells to a subject, including but not limited to injection, transfusion, implantation or transplantation. In some embodiments, the PTD-MYC fusion polypeptide-modified immune cells are administered to a patient subcutaneously, intradermally, intratumorally, intranodally, intramedullary, intramuscularly, intrathecally, by intravenous or intralymphatic injection, or intraperitoneally. In some embodiments, the PTD-MYC fusion polypeptide-immune cells are administered into a cavity formed by the resection of tumor tissue (*i.e.*, intracavity delivery) or directly into a tumor prior to resection (*i.e.* intratumoral delivery). In one embodiment, the MYC-fusion polypeptide-immune cells are administered by intravenous injection.

[0135] In addition to the PTD-MYC fusion polypeptide-modified immune cells, compositions for administration can comprise any other agents such as pharmaceutically acceptable carriers, buffers, excipients, adjuvants, additives, antiseptics, filling, stabilizing and/or thickening agents, and/or any components normally found in corresponding products. Selection of suitable ingredients and appropriate manufacturing methods for formulating the compositions for particular routes of administration generally known in the art.

[0136] The adoptive cell therapeutic composition comprising PTD-MYC fusion polypeptide-modified immune cells can be in any form, such as solid, semisolid or liquid form, suitable for administration. A formulation can be selected from a group consisting of, but not limited to, solutions, emulsions, suspensions, tablets, pellets and capsules. The compositions are not limited to a certain formulation, instead the composition can be formulated into any known pharmaceutically acceptable formulation. The pharmaceutical compositions may be produced by any conventional processes known in the art.

[0137] In some embodiments, the administration of the MYC-fusion polypeptide-modified immune cells comprises administering of 10^4 - 10^{10} of the cells per kg body weight, including 10^5 to 10^6 cells/kg body weight, including all integer values of cell numbers within those ranges. In some embodiments, the cells are administered with or without a course of lymphodepletion, for example with cyclophosphamide.

[0138] The MYC-fusion polypeptide-modified immune cells can be administered in one or more doses. In one embodiment, the therapeutically effective amount of PTD-MYC fusion polypeptide-modified immune cells are administered as a single dose. In some embodiments, administering a single dose of the PTD-MYC fusion polypeptide-modified immune cells has a therapeutic effect. In another embodiment, the effective amount of MYC-fusion polypeptide-modified immune cells are administered as more than one dose over a period time. Timing of administration is within the judgment of managing physician and depends on various factors, including, but not limited to the age, gender, or clinical condition of the patient and characteristics of the cancer, including type, degree or location of cancer. While individual needs vary, determination of optimal ranges of effective amounts of a MYC-fusion polypeptide-modified immune cell for treatment of a particular disease or conditions are within the skill of one in the art.

[0139] PTD-MYC fusion polypeptide-modified immune cells can be administered for example from 1 to 10 times in the first 2 weeks, 3 weeks, 4 weeks, monthly or during the treatment period. In some embodiments, PTD-MYC fusion polypeptide-modified immune cells are administered 2, 3, 4, 5, 6, 7, 8, 9, or 10 times. In some embodiments, PTD-MYC fusion polypeptide-modified immune cells are administered weekly, every 2 weeks, every 3 weeks or monthly.

[0140] A therapeutically effective amount means an amount which provides a therapeutic or prophylactic benefit. The dosage administered will be dependent upon the age, health and weight of the recipient, kind of concurrent treatment, if any, frequency of treatment and the nature of the effect desired.

[0141] In some embodiments, a patient receiving PTD-MYC modified immune cells are first pretreated with one or more cytokines and/or other immunomodulatory agents. In some

embodiments, a patient receiving PTD-MYC modified immune cells is lymphodepleted prior to administration of the PTD-MYC modified immune cells. The purpose of lymphodepletion is to make room for the infused lymphocytes, in particular by eliminating regulatory T cells and other non-specific T cells which compete for homeostatic cytokines.

[0142] In some embodiments, the PTD-MYC modified immune cells are administered with an additional therapeutic agent. In some embodiments, additional therapeutic agent is administered prior to, simultaneously with, intermittently with, or following treatment with the PTD-MYC modified immune cells. In some embodiments, the additional therapeutic agent is an immunomodulator, such as an interleukin (e.g. IL-2, IL-7, IL-12), a cytokine, a chemokine, or and immunomodulatory drug. In some embodiments, the cytokine is selected from among cytokine is selected from a group consisting of interferon alpha, interferon beta, interferon gamma, complement C5a, IL-2, TNF α , CD40L, IL12, IL-23, IL15, IL17, CCL1, CCL11, CCL12, CCL13, CCL14-1, CCL14-2, CCL14-3, CCL15-1, CCL15-2, CCL16, CCL17, CCL18, CCL19, CCL19, CCL2, CCL20, CCL21, CCL22, CCL23-1, CCL23-2, CCL24, CCL25-1, CCL25-2, CCL26, CCL27, CCL28, CCL3, CCL3L1, CCL4, CCL4L1, CCL5 (=RANTES), CCL6, CCL7, CCL8, CCL9, CCR10, CCR2, CCR5, CCR6, CCR7, CCR8, CCRL1, CCRL2, CX3CL1, CX3CR, CXCL1, CXCL10, CXCL11, CXCL12, CXCL13, CXCL14, CXCL15, CXCL16, CXCL2, CXCL3, CXCL4, CXCL5, CXCL6, CXCL7, CXCL8, CXCL9, CXCL9, CXCR1, CXCR2, CXCR4, CXCR5, CXCR6, CXCR7 and XCL2. In some embodiments, the additional therapeutic agent is an anticancer agent, such as chemotherapy or radiation therapy.

[0143] In some embodiments, the modified immune cells administered for the treatment of cancer are T cells with genetically modified antigen receptors, including chimeric antigen receptor (CAR)-T cells. Various strategies can, for example, be employed to genetically modify T cells by altering the specificity of the T cell receptor (TCR), for example, by introducing new TCR α and β chains with selected peptide specificity (see, e.g., U.S. Patent No. 8,697,854; PCT Patent Publications: WO2003020763, WO2004033685, WO2004044004, WO2005114215, WO2006000830, WO2008038002, WO2008039818, WO2004074322, WO2005113595, WO2006125962, WO2013166321, WO2013039889, WO2014018863, WO2014083173; U.S. Patent No. 8,088,379). Chimeric antigen receptors (CARs) can be used in order to generate

immunoresponsive cells, such as T cells, specific for selected targets, such as malignant cells, with a wide variety of receptor chimera constructs having been described (*see*, e.g. U.S. Patent Nos. 5,843,728; 5,851,828; 5,912,170; 6,004,811; 6,284,240; 6,392,013; 6,410,014; 6,753,162; 8,211,422; and, PCT Publication WO9215322). Methods for the preparation of CAR T cells are known in the art and can be used in combination with the methods provided herein to generate modified CAR T cells comprising a MYC fusion polypeptide (e.g. PTD) as described herein.

[0144] In general, CARs are comprised of an extracellular domain, a transmembrane domain, and an intracellular domain, wherein the extracellular domain comprises an antigen-binding domain that is specific for a predetermined target. While the antigen-binding domain of a CAR is often an antibody or antibody fragment (e.g., a single chain variable fragment, scFv), the binding domain is not particularly limited so long as it results in specific recognition of a target. For example, in some embodiments, the antigen-binding domain may comprise a receptor, such that the CAR is capable of binding to the ligand of the receptor. Alternatively, the antigen-binding domain may comprise a ligand, such that the CAR is capable of binding the endogenous receptor of that ligand.

[0145] In some embodiments, the T cells expressing a desired CAR are selected through co-culture with γ -irradiated activating and propagating cells (AaPC), which co-express the cancer antigen and co-stimulatory molecules. In some embodiments, the engineered CAR T-cells are expanded, for example by co-culture on AaPC in presence of soluble factors, such as IL-2 and IL-21. This expansion can for example be carried out so as to provide memory CAR⁺ T cells. In this way, CAR T cells can be provided that have specific cytotoxic activity against antigen-bearing tumors (optionally in conjunction with production of desired chemokines such as interferon- γ).

[0146] In some embodiments, the CAR T-cells are contacted with a PTD-MYC fusion polypeptide provided herein *in vitro* to generation a modified CAR T cells for the treatment of a cancer. The modified CAR T cells can be administered according any suitable method, including the methods for administration of the PTD-MYC fusion polypeptide-modified immune cells as described above.

Kits

[0147] Pharmaceutical compositions comprising MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells provided herein can be assembled into kits or pharmaceutical systems for use in treating a cancer. Kits according to this embodiment can comprise a carrier means, such as a box, carton, tube, having in close confinement therein one or more containers, such as vials, tubes, ampoules, bottles, syringes, or bags. The kits can also comprise associated instructions for using the MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells.

[0148] In some embodiments, the kit comprises an effective amount of an adoptive cell therapy, such as MYC-fusion polypeptide-modified immune cells. In some embodiments, the kit comprises one or more reagents for the detection of the administered MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells. In some embodiments, the kit comprises cells for treatment with a MYC-fusion polypeptide provided herein, for example, hematopoietic stem cells, donor leukocytes, T cells, or NK cells. In some embodiments, the kit further comprises an effective amount of a therapeutic agent to be administered in combination with MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells provided herein. In some embodiments, therapeutic agent is an anti-cancer agent.

[0149] Kits provided herein also can include a device for administering MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells provided herein to a subject. Any of a variety of devices known in the art for administering polypeptides and cells to a subject can be included in the kits provided herein. Exemplary devices include a hypodermic needle, an intravenous needle, a catheter, a needle-less injection, but are not limited to, a hypodermic needle, an intravenous needle, a catheter, a needle-less injection device, an inhaler and a liquid dispenser such as an eyedropper. Typically the device for administering the MYC-fusion polypeptides and/or MYC-fusion polypeptide-modified immune cells of the kit will be compatible with the desired method of administration of the composition. For example, a composition to be delivered intravenously can be included in a kit with a hypodermic needle and a syringe.

EXAMPLES

Example 1. Immune cells treated with TAT-MYC to generate TAT-MYC-treated lymphocytes for immunotherapy of melanoma tumors

[0150] In this example, the ability of a PTD-MYC fusion polypeptide comprising the protein transduction domain of HIV-1 transactivation protein (TAT) and MYC to modulate an immune response against melanoma cells *in vivo* was examined. Specifically, the ability of lymphoid cells, derived from melanoma-bearing mice and treated with TAT-MYC, to treat mice harboring melanoma tumors was studied. The object of these studies was to determine whether immune cells derived from melanoma bearing mice and treated with TAT-MYC to generate TAT-MYC lymphocytes would be an effective treatment for melanoma tumors upon transplantation into melanoma bearing mice.

[0151] *Materials and Methods*

[0152] C57BL/6J is the most widely used inbred strain and the first to have its genome sequenced. Although this strain is refractory to many tumors, it is a permissive background for maximal expression of most mutations. C57BL/6J mice are resistant to audiogenic seizures, have a relatively low bone density, and develop age-related hearing loss. They are also susceptible to diet-induced obesity, type 2 diabetes, and atherosclerosis. Macrophages from this strain are resistant to the effects of anthrax lethal toxin.

[0153] *Treatment Groups*

[0154] Fifteen C57BL/6 mice (Jackson Laboratory Stock# 000664) weighing approximately 25g and harboring melanoma tumors were generated and divided into 3 cohorts of 5 animals, one cohort of one mouse as a no treatment control, one cohort treated with Lymphoid cells derived from tumor-bearing mice and treated with control TAT-fusion protein, and one cohort treated with TAT-MYC lymphocytes.

[0155] *Generation of tumor-bearing donor mice and preparation of donor cells*

[0156] B16-F10 melanoma cells (ATCC CRL 6475, mouse skin melanoma) for implantation were cultured in D10 media (DMEM, 10% FBS, Pen/Strep (10,000 units per/ml) (Gibco Cat#

15140); L-glutamine (200mM) (Gibco Cat# 25030); MEM Non-essential Amino Acids (Gibco Cat# 11140)).

[0157] The C57BL/6j mice (Jackson Laboratory #003548) were implanted with 1×10^4 B16-F10 melanoma cells in 250 μ L PBS via tail vein injection. Prior to injection, each test mouse was placed under a 250W heat lamp for 1-2 minutes and then injected intravenously with the melanoma cells. At 14 days post-transplant, lymph nodes from the injected mice were harvested and ground with the plunger of a 10 mL syringe.

[0158] For the first study, lymph nodes were harvested from 5 mice. For the second lymph nodes were harvested from 10 mice. The cells were washed with C10, collected and spun at $260 \times g$ for 5 min. After discarding the supernatant, the cells were resuspended in 10mL sterile TAC, spun at $260 \times g$ for 5 minutes. After discarding the supernatant, the cells were resuspended in 2mL of sterile filtered PBS with 5% BSA.

[0159] The lymph node cells were treated with TAT-MYC to generate TAT-MYC lymphocytes or treated with a control TAT-Fusion protein. The cells were split into 2, 15mL conical tubes (1mL each), treated with 1mL of 25ug/ml of a control protein (TAT-CRE for experiment 1, TAT-GFP for experiment 2) or 1mL of 25ug/ml of TAT-MYC lot C18. After one hour of room temp incubation, each tube was washed with sterile PBS three times, transferred to 5mL sterile tubes and placed on ice.

[0160] The test mice were prepared by injecting 1×10^4 B16-F10 melanoma cells in 250uL PBS into the tail vein for each cohort of 5 C57BL/6j mice. After injection, the mice were observed once daily. Changes in body weight, food consumption, activity, and mortality were monitored. At 7 days post-transplant, TAT-MYC lymphocytes or control lymphoid cells were then transplanted into melanoma cell injected mice.

[0161] Symptoms were monitored daily. The mice were euthanized when severe symptoms presented and deaths were recorded. Mice were either found dead or euthanized if found with severe symptoms such as heavy breathing, hunched back and immobility. Day of death was recorded with day of treatment as Day 0.

[0162] The results from Experiments 1 and 2 are shown in Figures 1 and 2, respectively. As shown in the figures, treating melanoma-bearing mice with TAT-MYC lymphocytes (TBX-

3400) generated by contacting mouse lymphoid cells derived from melanoma bearing mice with TAT-MYC, significantly improved the overall survival of the mice compared to transplanting lymphoid cells treated with control TAT-Fusion protein. These results suggest that TAT-MYC treatment of immune cells are useful in the treatment of melanoma using adoptive cell transfer.

Example 2. Dose Response Effect of TAT-MYC-treated lymphocytes for immunotherapy of melanoma tumors

[0163] In this example, the therapeutic effects of different amounts administered TAT-MYC-treated lymphocytes for immunotherapy of melanoma tumors was examined. This experiment was performed as described above in Example 1, except that several different doses of the TAT-MYC-treated lymphocytes were injected and compared. Two experiments were performed. In the first experiment, Experiment 3, TAT-MYC lymphocytes were administered to the melanoma-bearing mice via tail vein injection according to the following dosing groups: 3.0×10^6 cells/kg, 6.0×10^6 cells/kg, 14.0×10^6 cells/kg, and 70.0×10^6 cells/kg. For the control groups, the mice were administered 70.0×10^6 TAT-Cre treated or no cells (NT). In the second experiment, Experiment 4, TAT-MYC lymphocytes were administered to the melanoma-bearing mice via tail vein injection according to the following dosing groups: 4.0×10^3 cells/kg, 4.0×10^4 cells/kg, 4.0×10^5 cells/kg, 4.0×10^6 cells/kg and 4.0×10^7 cells/kg. For the control groups, the mice were administered 4.0×10^6 TAT-Cre treated or no cells (NT). The results from Experiments 3 and 4 are shown in Figures 3 and 4, respectively. As shown in the figures, treating melanoma-bearing mice with increasing amounts of TAT-MYC lymphocytes (TBX-3400) led to a significantly improved overall survival rate in both experiments. These experiments demonstrate the both the reproducibility and efficacy of TAT-MYC lymphocytes for treating melanoma-bearing subjects.

Example 3. Immune cells treated with TAT-MYC to generate TAT-MYC-treated lymphocytes for immunotherapy of colon cancer

[0164] In this example, the ability of a PTD-MYC fusion polypeptide comprising the protein transduction domain of HIV-1 transactivation protein (TAT) and MYC to modulate an immune response against colon cancer cells *in vivo* was examined. Specifically, the ability of lymphoid

cells, derived from colon tumor-bearing mice and treated with TAT-MYC, to treat mice harboring colon cancer cell-derived tumors was studied. The object of these studies was to determine whether immune cells derived from colon tumor-bearing mice and treated with TAT-MYC to generate TAT-MYC lymphocytes would be an effective treatment for colon cancer upon transplantation into colon tumor-bearing mice.

[0165] *Materials and Methods*

[0166] MC-38 murine colon adenocarcinoma cells (Kerafast #ENH204) for implantation were cultured in D10 media (DMEM, 10% FBS, Pen/Strep (10,000 units per/ml) (Gibco Cat# 15140); L-glutamine (200mM) (Gibco Cat# 25030); MEM Non-essential Amino Acids (Gibco Cat# 11140)).

[0167] Nine donor C57BL/6J mice (Jackson Laboratory Stock# 000664) weighing approximately 25g were implanted with 1×10^6 MC-38 murine colon adenocarcinoma cells in 250 μ L PBS via tail vein injection. One week later, 18 recipient C57BL/6J mice were implanted with 1×10^6 MC-38 murine colon adenocarcinoma cells.

[0168] At 14 days post-transplant of the donor mice, lymph nodes from the nine injected donor mice were harvested and ground with the plunger of a 10 mL syringe. The cells were washed with C10, collected and spun at $260 \times g$ for 5 min. After discarding the supernatant, the cells were resuspended in 10mL sterile TAC, spun at $260 \times g$ for 5 minutes. After discarding the supernatant, the cells were resuspended in 2mL of sterile filtered PBS with 5% BSA. The lymph node cells were treated with TAT-MYC to generate TAT-MYC lymphocytes or treated with a control TAT-Fusion protein. The cells were split into 2, 15mL conical tubes (1mL each), treated with 1mL of 25ug/ml of a control TAT-fusion protein (TAT-CRE) or 1mL of 25ug/ml of TAT-MYC lot C18. After one hour of room temp incubation, each tube was washed with sterile PBS three times, transferred to 5mL sterile tubes and placed on ice. 125,000 lymph cells were then injected in 250uL PBS into the tail vein of each recipient mouse (equivalent to 5×10^6 cells/kg) to create 3 cohorts of 6 mice each: one cohort of no treatment control, one cohort treated with lymphoid cells derived from tumor-bearing mice and treated with control TAT-fusion protein, and one cohort treated with lymphoid cells derived from tumor-bearing mice and treated with TAT-MYC fusion protein (TAT-MYC lymphocytes).

[0169] After injection, the mice were observed once daily. Changes in body weight, food consumption, activity, and mortality were monitored. Symptoms were monitored daily. The mice were euthanized when severe symptoms presented and deaths were recorded. Mice were either found dead or euthanized if found with severe symptoms such as heavy breathing, hunched back and immobility. Day of death was recorded with day of treatment as Day 0.

[0170] *Results*

[0171] The results from this experiment are shown in Figure 5. At day 27 post treatment, the first no treatment and TAT-CRE control mice were found dead. By day 34 post treatment, all no treatment and TAT-CRE control mice were either found dead or needed to be euthanized. By contrast, the first TAT-MYC lymphocyte treated mouse to die occurred at day 32 post treatment. By day 52, only 3 of the TAT-MYC lymphocyte treated mice had died.

[0172] Accordingly, as shown in the figure, treating colon tumor-bearing mice with TAT-MYC lymphocytes generated by contacting mouse lymphoid cells derived from colon tumor-bearing mice with TAT-MYC, significantly ($p < 0.0019$) improved the overall survival of the mice compared to transplanting lymphoid cells treated with control TAT-MYC fusion protein. These results suggest that TAT-MYC treatment of immune cells are useful in the treatment of colon cancer using adoptive cell transfer.

Example 4. Dose Response Effect of TAT-MYC-treated lymphocytes for immunotherapy of melanoma tumors

[0173] In this example, the therapeutic effects of different amounts administered TAT-MYC-treated lymphocytes for immunotherapy of melanoma tumors was examined. This experiment was performed as described above in Example 3, except that several different doses of the TAT-MYC-treated lymphocytes were injected and compared. In this experiment, TAT-MYC lymphocytes were administered to the melanoma-bearing mice via tail vein injection according to the following dosing groups (5 tumor bearing mice per group): 5.0×10^3 cells/kg, 5.0×10^4 cells/kg, 5.0×10^5 cells/kg, 5.0×10^6 cells/kg, 5.0×10^7 cells/kg, and 5.0×10^8 cells/kg, cells/kg. For the control groups, the mice were administered no cells (NT) (8 mice), or 5.0×10^7 cells/kg was administered to non-tumor bearing mice (5 mice). The results from this experiment are shown in

Figure 6. As shown in the figure, treating melanoma-bearing mice with increasing amounts of TAT-MYC lymphocytes (TBX-3400) led to a significantly improved overall survival rate. In the untreated cohort, 75% of the animals died by Day 19 and the remaining untreated mice died by day 41 post tumor transplant. All of the mice receiving 5×10^3 cells/kg died by Day 19. In contrast, 60% of animals receiving 5×10^5 cells/kg and 100% of animals receiving 5×10^6 or 5×10^7 cells/kg survived throughout the study. Additionally, 100% of the non-tumor bearing mice receiving 5×10^7 cells/kg survived throughout the study. All the mice that received 5×10^8 cells/kg presented with thinning of their fur. 40% of the mice receiving 5×10^8 cells/kg died by day 41. The remaining 60% of mice that received 5×10^8 cells/kg survived throughout the study.

[0174] These experiments demonstrate the both the reproducibility and efficacy of TAT-MYC lymphocytes for treating colon tumor-bearing subjects in an otherwise fatal MC-38 colorectal cancer tumor model.

Example 5. Immune cells treated with TAT-MYC to generate TAT-MYC-treated lymphocytes for immunotherapy of solid tumors

[0175] In this example, the ability of a PTD-MYC fusion polypeptide comprising the protein transduction domain of HIV-1 transactivation protein (TAT) and MYC to modulate an immune response against additional tumor cell types *in vivo* is examined. Specifically, the ability of lymphoid cells, derived from tumor-bearing mice and treated with TAT-MYC, to treat mice harboring solid tumors is studied. The object of these studies is to determine whether immune cells derived from tumor bearing mice and treated with TAT-MYC to generate TAT-MYC lymphocytes is an effective treatment for solid tumors upon transplantation into tumor bearing mice.

[0176] Numerous mouse xenograft models that use transplantation of cancer cell lines are available in the art and can be employed in the present example to assess treatment of a solid tumor. Non-limiting examples of cancers and available cell lines for the generation of xenograft tumors are listed in the table below.

Histotype	Cell Line
Adrenal	H295R

Bladder	SW780
Brain	SF-295, SK-N-AS, U87 MG, U251
Breast	BT474, HCC-1806, HCC-1954, JIMT-1, MCF-7, MDA-MB-231, HCC70
Colon	CL-34, COLO 205, DLD-1, HCT 116, HCT-15, HT-29, LoVo, LS-174T, LS411N, RKO, SW48, SW480, SW620
Epithelial	A-431
Ewing's Sarcoma	RD-ES
Gastric	N87, SNU-5, MKN-45
Head and Neck	FaDu
Leukemia	HL-60, MOLM-13, MOLT-4, MV4-11, RS4;11, SET2, THP-1, HEL92.1.7, K-562, Kasumi-1
Liver	Hep3B, HuH-7, SNU-398
Lung (Non-small cell)	A-427, A549, H1299, H1975, H226, H23, H292, H460, H520, H522, H647, H727, H810, HCC-44, NCI-H2122, SK-MES-1, COR-L23, NCI-H1963, H2009, H1581, H1993, H441
Lung (Small cell)	H69, H82, H211, H526, SHP-77, DMS 114
Lymphoma	Daudi, DoHH-2, Granta 519, JEKO-1, KARPAS-299, MOLT-4, Raji B, Ramos, REC-1, RL, SU-DHL-4, WSU-DLCL2, Mino, Namalwa
Melanoma	A2058, A375, IGR-37, SK-MEL-5, UACC-62, CHL-1, COLO 800, IGR-1, IGR-37
Mesothelioma	MSTO-211H
Multiple Myeloma	H929, OPM-2, RPMI 8226
Neuroblastoma	SKNAS
Ovarian	A2780, IGR0V1, OVCAR-3, OVCAR-5, SK-OV-3, TOV-21G, OV-90
Pancreas	BxPc-3, HPAC, HPAF II, KP4, MIA PaCa-2, PANC-1
Prostate	22Rv.1, PC3, DU145, LNCaP, VCaP
Renal	786-0, G-401, G-402
Sarcoma	HT-1080, SJSA-1
Thyroid	8505C, FTC-238, K1
Uterine	ECC-1, MFE-280, HEC-1-A, HEC-1-B

[0177] *Materials and Methods*

[0178] Fifteen C57BL/6 mice (Jackson Laboratory Stock# 000664) weighing approximately 25g and harboring solid tumors are generated and divided into 3 cohorts of 5 animals, one cohort of one mouse as a no treatment control, one cohort treated with Lymphoid cells derived from tumor-bearing mice and treated with control TAT-fusion protein, and one cohort treated with TAT-MYC lymphocytes.

[0179] *Generation of tumor-bearing donor mice and preparation of donor cells*

[0180] Solid tumor cells for implantation are cultured in D10 media (DMEM, 10% FBS, Pen/Strep (10,000 units per/ml) (Gibco Cat# 15140); L-glutamine (200mM) (Gibco Cat# 25030); MEM Non-essential Amino Acids (Gibco Cat# 11140)).

[0181] The C57BL/6j mice (Jackson Laboratory #003548) are implanted with 1×10^4 tumor cells in 250 μ L PBS via tail vein injection. Prior to injection, each test mouse was placed under a 250W heat lamp for 1-2 minutes and then injected intravenously with the tumor cells. At 14 days post-transplant, lymph nodes from the injected mice are harvested and ground with the plunger of a 10 mL syringe.

[0182] For a first study, lymph nodes are harvested from 5 mice. For a second study, lymph nodes were harvested from 10 mice. The cells are washed with C10, collected and spun at $260 \times g$ for 5 min. After discarding the supernatant, the cells are resuspended in 10mL sterile TAC, spun at $260 \times g$ for 5 minutes. After discarding the supernatant, the cells are resuspended in 2mL of sterile filtered PBS with 5% BSA.

[0183] The lymph node cells are treated with TAT-MYC to generate TAT-MYC lymphocytes or treated with a control TAT-Fusion protein. The cells are split into 2, 15mL conical tubes (1mL each), treated with 1mL of 25ug/ml of a control protein (TAT-CRE for experiment 1, TAT-GFP for experiment 2) or 1mL of 25ug/ml of TAT-MYC lot C18. After one hour of room temp incubation, each tube is washed with sterile PBS three times, transferred to 5mL sterile tubes and placed on ice.

[0184] The test mice are prepared by injecting 1×10^4 tumor cells in 250uL PBS into the tail vein for each cohort of 5 C57BL/6j mice. After injection, the mice are observed once daily. Changes in body weight, food consumption, activity, and mortality are monitored. At 7 days post-transplant, TAT-MYC lymphocytes or control lymphoid cells are then transplanted into tumor cell injected mice.

[0185] Symptoms are monitored daily. The mice are euthanized when severe symptoms presented and deaths are recorded. Mice are either found dead or euthanized if found with severe symptoms such as heavy breathing, hunched back and immobility. Day of death is recorded with day of treatment as Day 0.

[0186] It is expected that treating tumor-bearing mice with TAT-MYC lymphocytes (TBX-3400) generated by contacting mouse lymphoid cells derived from tumor bearing mice with TAT-MYC, will significantly improve the overall survival of the mice compared to transplanting lymphoid cells treated with control TAT-Fusion protein. These results will show that that TAT-MYC treatment of immune cells are useful in the treatment of solid tumors using adoptive cell transfer.

Example 6. Dose Response Effect of TAT-MYC-treated lymphocytes for immunotherapy of solid tumors

[0187] In this example, the therapeutic effects of different amounts administered TAT-MYC-treated lymphocytes for immunotherapy of solid tumors are examined. This experiment is performed as described above, except that several different doses of the TAT-MYC-treated lymphocytes are injected and compared. Two experiments are performed. In the first experiment, TAT-MYC lymphocytes are administered to the tumor-bearing mice via tail vein injection according to the following dosing groups: 3.0×10^6 cells/kg, 6.0×10^6 cells/kg, 14.0×10^6 cells/kg, and 70.0×10^6 cells/kg. For the control groups, the mice are administered 70.0×10^6 TAT-Cre treated or no cells (NT). In the second experiment, Experiment 4, TAT-MYC lymphocytes are administered to the tumor-bearing mice via tail vein injection according to the following dosing groups: 4.0×10^3 cells/kg, 4.0×10^4 cells/kg, 4.0×10^5 cells/kg, 4.0×10^6 cells/kg and 4.0×10^7 cells/kg. For the control groups, the mice are administered 4.0×10^6 TAT-Cre treated or no cells (NT). It is expected that treating tumor-bearing mice with increasing amounts of TAT-MYC lymphocytes (TBX-3400) will led to a significantly improved overall survival rate in both experiments. These experiments will also demonstrate the reproducibility and efficacy of TAT-MYC lymphocytes for treating tumor-bearing subjects.

[0188] While preferred embodiments of the present disclosure have been shown and described herein, it will be obvious to those skilled in the art that such embodiments are provided by way of example only. Numerous variations, changes, and substitutions will now occur to

those skilled in the art without departing from the disclosure. It should be understood that various alternatives to the embodiments of the disclosure described herein *may* be employed in practicing the disclosure. It is intended that the following claims define the scope of the disclosure and that methods and structures within the scope of these claims and their equivalents be covered thereby.

[0189] All patents, patent applications, provisional applications, and publications referred to or cited herein are incorporated by reference in their entirety, including all figures and tables, to the extent they are not inconsistent with the explicit teachings of this specification.

[0190] Other embodiments are set forth within the following claims.

WHAT IS CLAIMED IS:

1. A composition comprising:
 - (a) a MYC fusion peptide, comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence; and
 - (b) one or more primary immune cells isolated from a donor subject that has a solid tumor, wherein the one or more primary immune cells are reactive against a tumor-specific antigen.
2. The composition of claim 1, wherein the solid tumor is a carcinoma, adenoma, adenocarcinoma, blastoma, sarcoma, or lymphoma.
3. The composition of claim 1, wherein the solid tumor is a metastatic tumor.
4. The composition of claim 1, wherein the solid tumor is a basal cell carcinoma, biliary tract cancer, bladder cancer, breast cancer, cervical cancer, choriocarcinoma, CNS cancer, colon cancer, colorectal cancer, connective tissue cancer, cancer of the digestive system, endometrial cancer, esophageal cancer, eye cancer, gastric cancer, glial cell tumor, head and neck cancer, hepatoma, hepatic carcinoma, Hodgkin's lymphoma, Non-Hodgkin's lymphoma, intra-epithelial neoplasm, kidney cancer, larynx cancer, liver cancer, small-cell lung cancer, non-small cell lung cancer, melanoma, myeloma, neuroblastoma, oral cavity cancer, ovarian cancer, pancreatic cancer, prostate cancer, rectal cancer, renal cancer, cancer of the respiratory system, retinoblastoma, rhabdomyosarcoma, salivary gland carcinoma, squamous cell cancer, stomach cancer, testicular cancer, thyroid cancer, uterine cancer, cancer of the urinary system, or a vulvar cancer.
5. The composition of any one of claims 1-4, wherein the MYC fusion peptide comprises SEQ ID NO: 1.

6. The composition of any one of claims 1-5, wherein the one or more immune cells have antitumor activity against the solid tumor cells.

7. The composition of any one of claims 1-6, wherein the one or more immune cells comprises one or more lymphocytes.

8. The composition of claim 7, wherein the one or more lymphocytes comprises a T cell, a B cell, an NK cell, or any combination thereof.

9. The composition of any one of claims 7-8, wherein the one or more lymphocytes is a tumor-infiltrating lymphocyte, T-cell receptor modified lymphocyte, or a chimeric antigen receptor modified lymphocyte.

10. The composition of claim 9, wherein the tumor-infiltrating lymphocyte has a CD8+CD25+ signature or a CD4+CD25+ signature.

11. The composition of any one of claims 1-10, wherein the one or more immune cells comprises a detectable moiety.

12. A method for treating a cancer in a subject, comprising administering one or more modified immune cells to the subject in need thereof, wherein the one or more modified immune cells comprise a MYC fusion peptide comprising (i) a protein transduction domain; (ii) a MYC polypeptide sequence and are reactive to a tumor-specific antigen.

13. The method of claim 12, wherein the one or more modified immune cells are derived from primary immune cells isolated from the subject.

14. The method of claim 12, wherein the one or more modified immune cells are derived from primary immune cells isolated from a separate donor subject having the same type of cancer.

15. The method of any one of claims 12-14, wherein the cancer is carcinoma or a sarcoma.

16. The method of any one of claims 12-14, wherein the cancer is metastatic cancer.
17. The composition any one of claims 12-14, wherein the cancer is a basal cell carcinoma, biliary tract cancer, bladder cancer, breast cancer, cervical cancer, choriocarcinoma, CNS cancer, colon cancer, colorectal cancer, connective tissue cancer, cancer of the digestive system, endometrial cancer, esophageal cancer, eye cancer, gastric cancer, glial cell tumor, head and neck cancer, hepatoma, hepatic carcinoma, Hodgkin's lymphoma, Non-Hodgkin's lymphoma, intra-epithelial neoplasm, kidney cancer, larynx cancer, liver cancer, small-cell lung cancer, non-small cell lung cancer, melanoma, myeloma, neuroblastoma, oral cavity cancer, ovarian cancer, pancreatic cancer, prostate cancer, rectal cancer, renal cancer, cancer of the respiratory system, retinoblastoma, rhabdomyosarcoma, salivary gland carcinoma, squamous cell cancer, stomach cancer, testicular cancer, thyroid cancer, uterine cancer, cancer of the urinary system, or a vulvar cancer.
18. The method of any one of claims 13-17, wherein the one or more modified immune cells are prepared by contacting the primary immune cells *in vitro* with the MYC fusion peptide following isolation.
19. The method of any one of claims 13-17, further comprising expanding the primary immune cells *in vitro* prior to prior to or following contacting with the MYC fusion peptide.
20. The method of any one of claims 12-19, wherein the MYC fusion peptide comprises SEQ ID NO: 1.
21. The method of any one of claims 12-20, wherein the one or more modified immune cells have antitumor activity against cancer cells in the subject.
22. The method of any one of claims 12-21, wherein the one or more modified immune cells comprise one or more anergic immune cells.

23. The method of any one of claims 12-22, wherein the one or more immune cells comprises one or more lymphocytes.

24. The method of claim 23, wherein the one or more lymphocytes comprises a T cell, a B cell, an NK, or any combination thereof.

25. The method of claim 23, wherein the one or more lymphocytes is a tumor-infiltrating lymphocyte, T-cell receptor modified lymphocyte, or a chimeric antigen receptor modified lymphocyte.

26. The method of claim 25, wherein the lymphocyte has a $CD8^{+}CD28^{-}CD152^{-}$ signature, a $CD8^{+}CD25^{+}$ signature, or a $CD4^{+}CD25^{+}$ signature.

27. The method of any one of claims 13-26, wherein method further comprises isolating the primary immune cells from the donor subject.

28. The method of any one of claims 12-27, wherein the one or more modified immune cells are administered intravenously, intraperitoneally, subcutaneously, intramuscularly, or intratumorally.

29. The method of any one of claims 12-28, further comprising lymphodepleting the subject prior to administration of the one or more modified immune cells.

30. The method of any one of claims 12-29, further comprising administering a cytokine to the subject.

31. The method of any one of claims 12-30, wherein the subject is a human or an animal.

32. The method of any one of claims 12-31, further comprising administering an additional cancer therapy.

33. A method for preparing modified immune cells for cancer therapy, comprising contacting one or more immune cells *in vitro* with a MYC fusion polypeptide, wherein the immune cells are from a donor that has been exposed to one or more tumor antigens and wherein the MYC fusion peptide comprises (i) a protein transduction domain; (ii) a MYC polypeptide sequence and are reactive to a tumor-specific antigen.

34. The method of claim 33, wherein the one or more modified immune cells are derived from primary immune cells isolated from a subject having cancer.

35. The method of any one of claims 33-34, further comprising expanding the primary immune cells *in vitro* prior to or following contacting with the MYC fusion peptide.

36. The method of any one of claims 33-35, wherein the MYC fusion peptide comprises SEQ ID NO: 1.

37. The method of any one of claims 33-34, wherein the one or more modified immune cells have antitumor activity.

38. The method of any one of claims 33-37, wherein the one or more immune cells comprises a T cell, a B cell, an NK, or any combination thereof.

39. The method of any one of claims 33-37, wherein the one or more immune cells is a tumor-infiltrating lymphocyte, T-cell receptor modified lymphocyte, or a chimeric antigen receptor modified lymphocyte.

40. A method for increasing the efficacy of adoptive cell therapy or T-cell therapy in a subject comprising administering the composition of any of claims 1-11.

41. The composition of any of claims 1-11 for use in treating a cancer.

42. Use of a composition of any one of claims 1-11 in the manufacture of a medicament for the treatment of a cancer.

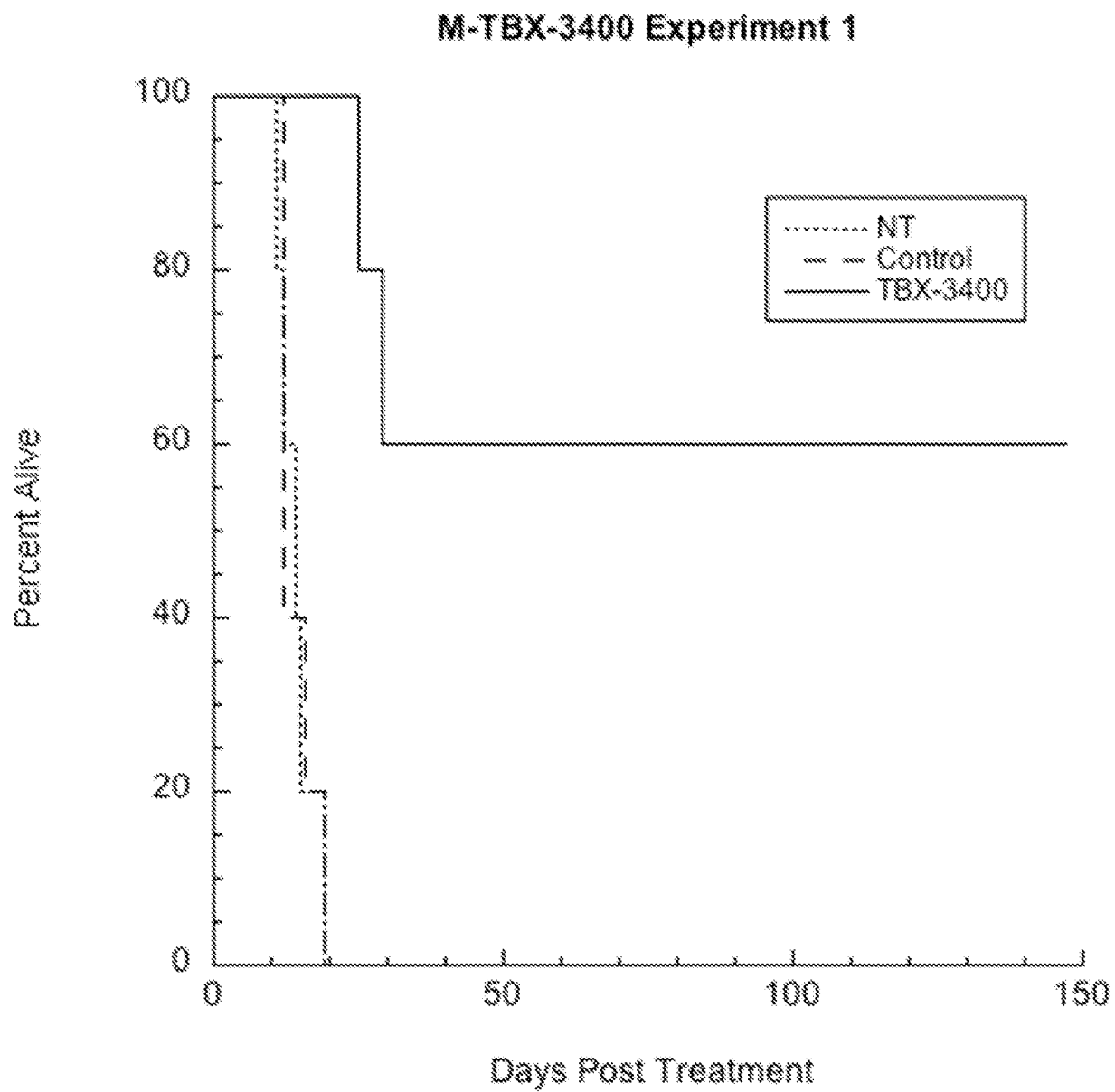
FIG. 1

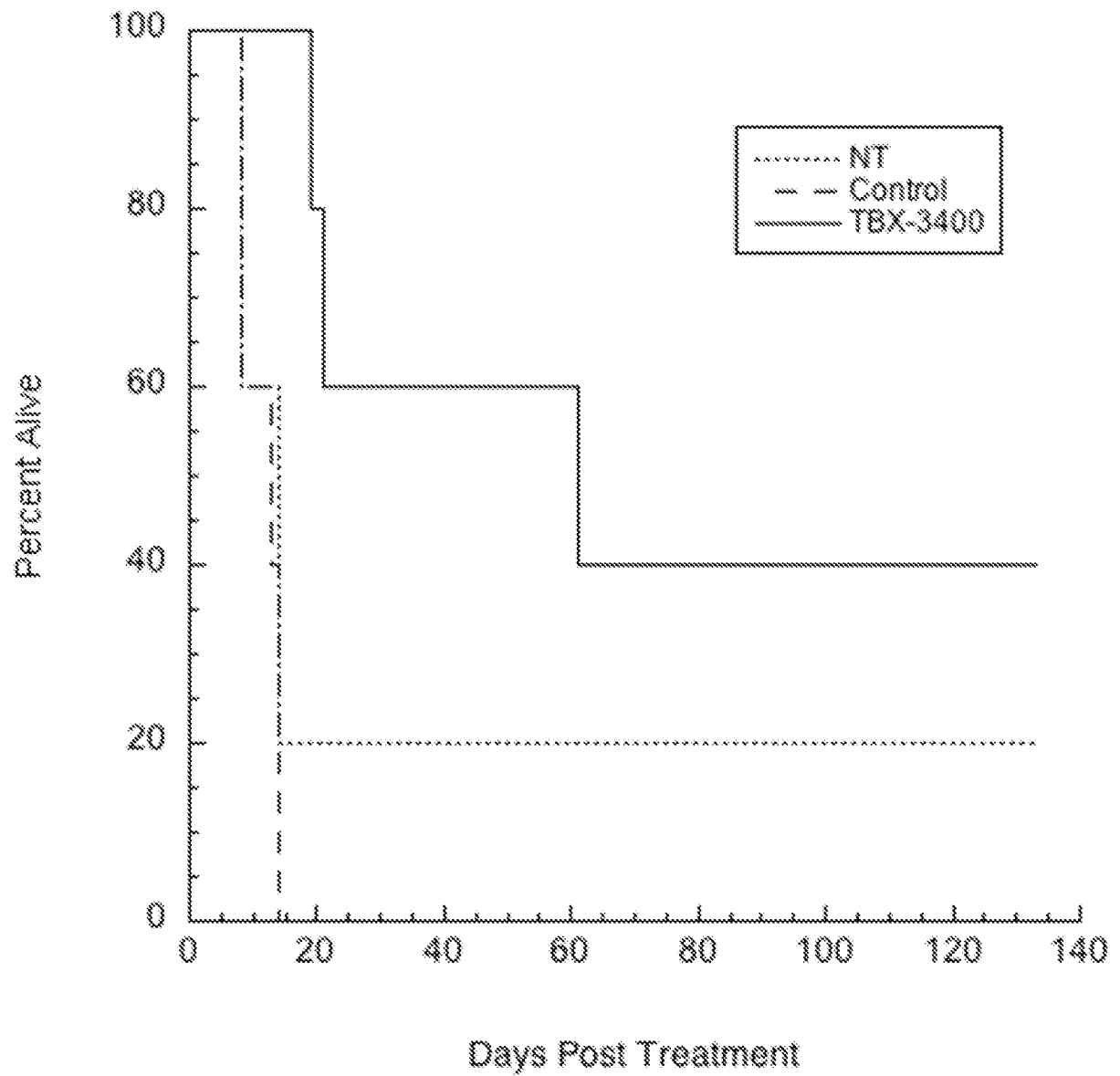
FIG. 2**M-TBX-3400 Experiment 2**

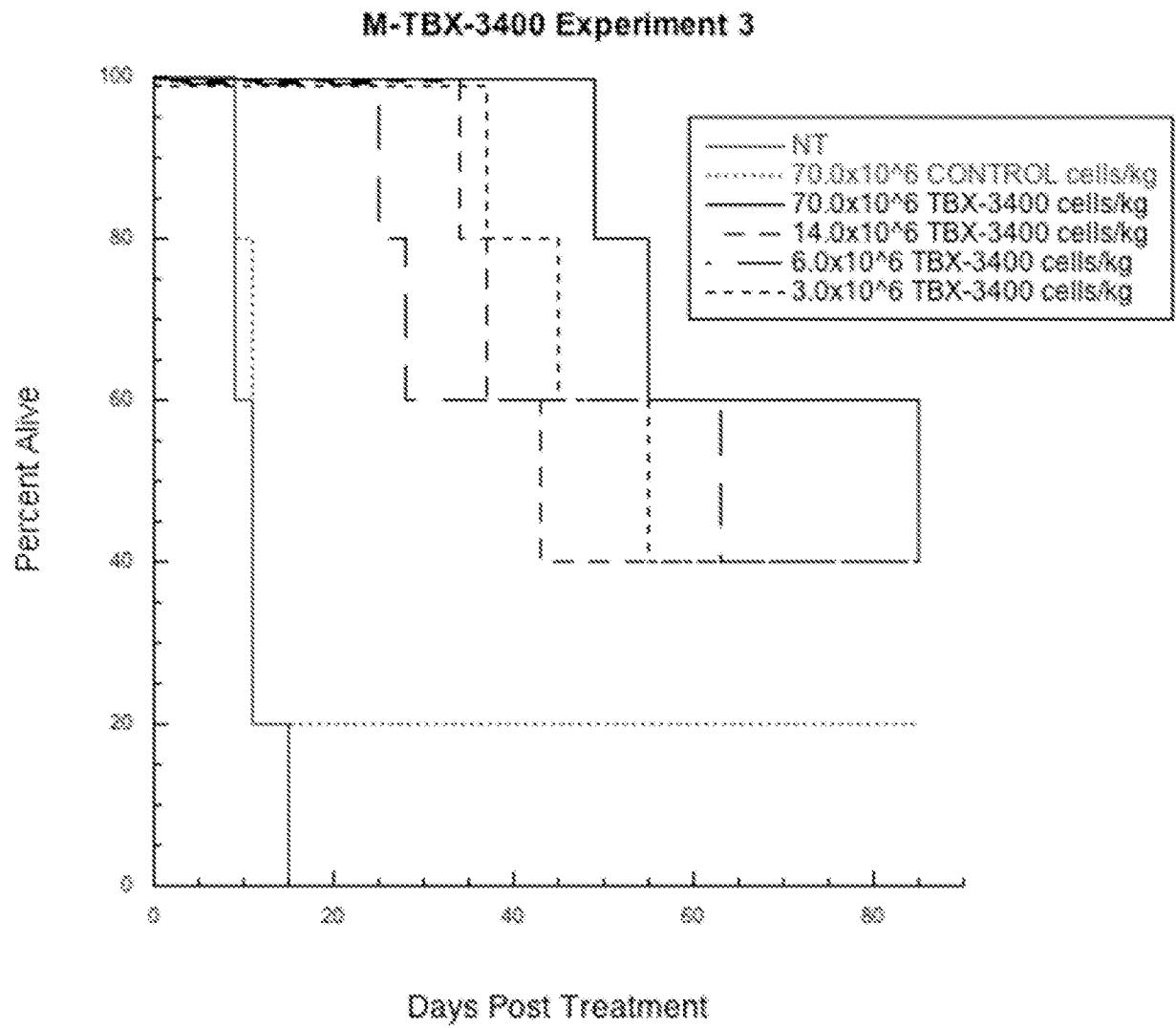
FIG. 3

FIG. 4

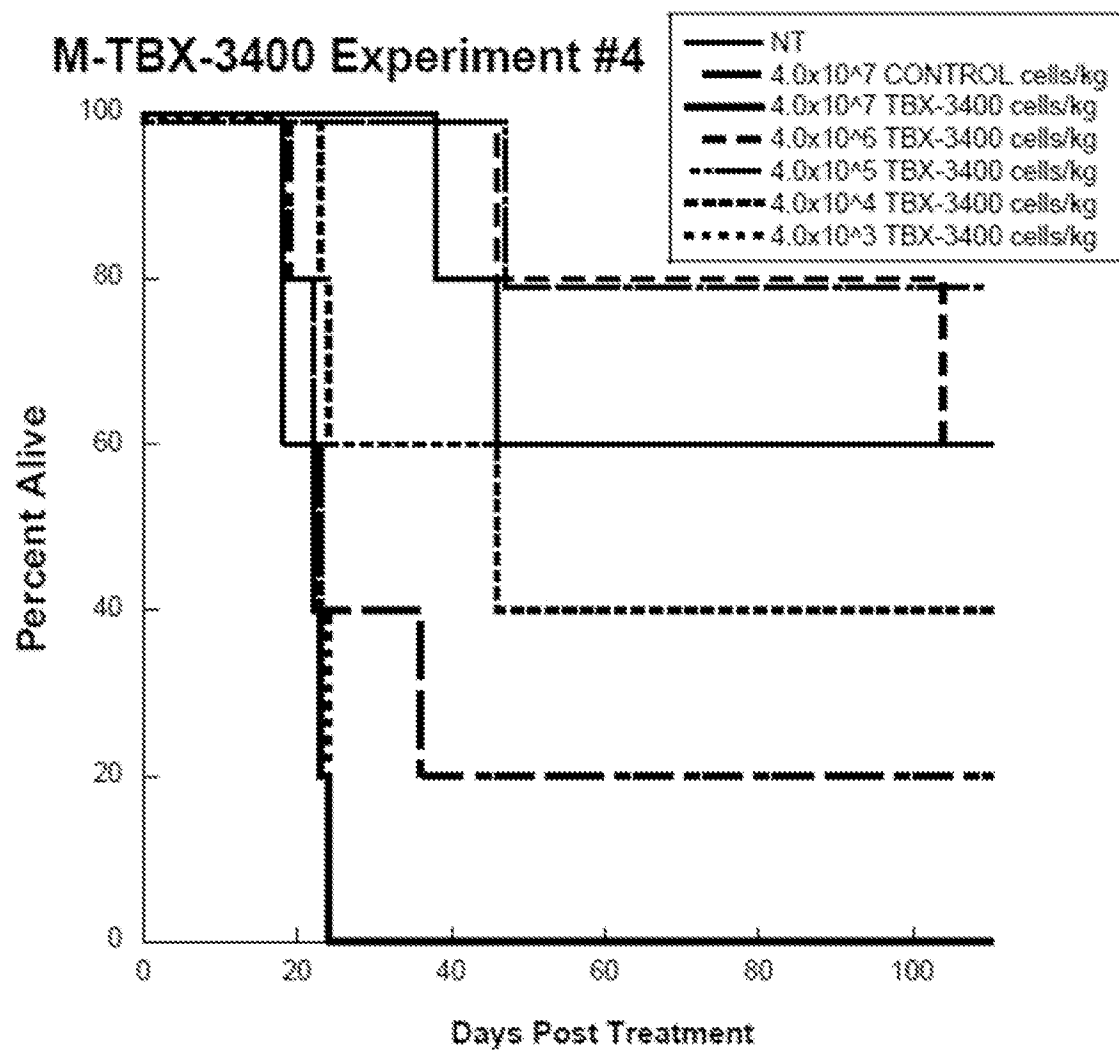


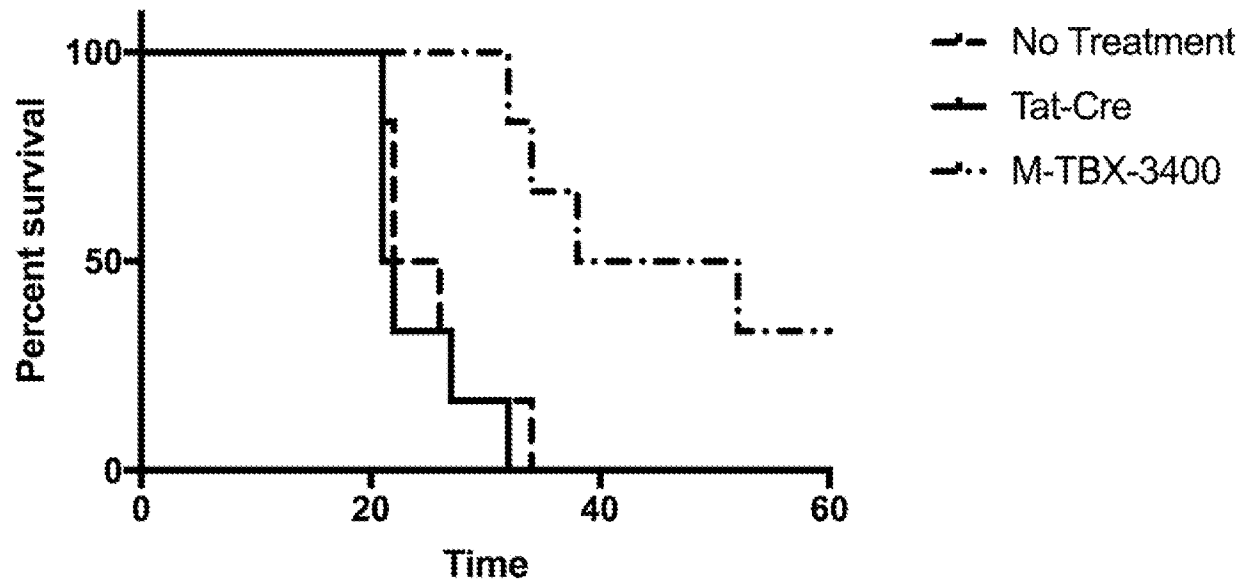
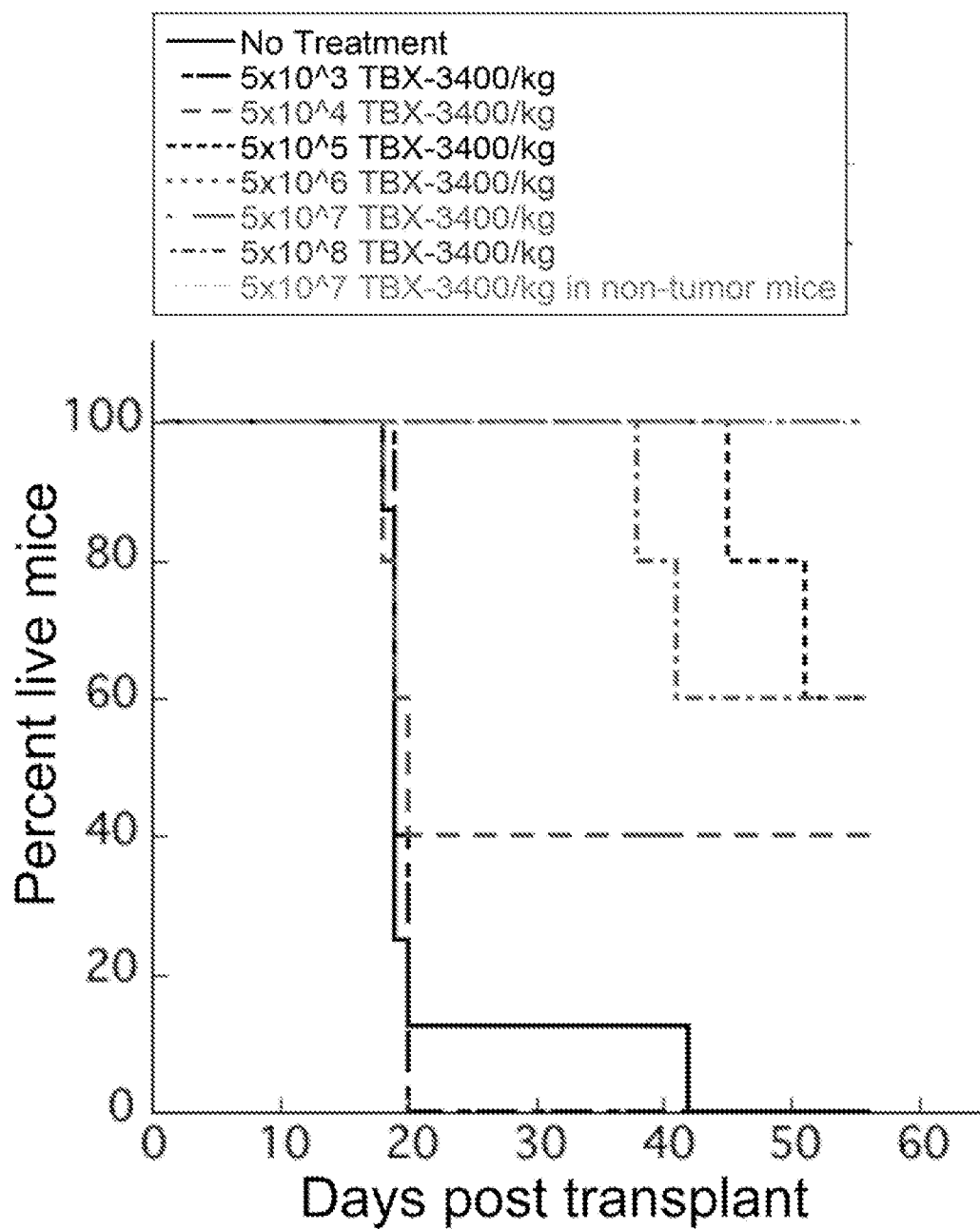
FIG. 5**MC-38 Exp #1**

FIG. 6



INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2018/044740

Box No. I Nucleotide and/or amino acid sequence(s) (Continuation of item 1.c of the first sheet)

1. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international search was carried out on the basis of a sequence listing:

a. ☒ forming part of the international application as filed:

☒ in the form of an Annex C/ST.25 text file.

☐ on paper or in the form of an image file.

b. ☐ furnished together with the international application under PCT Rule 13*ter*.1(a) for the purposes of international search only in the form of an Annex C/ST.25 text file.

c. ☐ furnished subsequent to the international filing date for the purposes of international search only:

☐ in the form of an Annex C/ST.25 text file (Rule 13*ter*.1(a)).

☐ on paper or in the form of an image file (Rule 13*ter*.1(b) and Administrative Instructions, Section 713).

2. ☐ In addition, in the case that more than one version or copy of a sequence listing has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that forming part of the application as filed or does not go beyond the application as filed, as appropriate, were furnished.

3. Additional comments:

SEQ ID NOs: 1-13 were searched.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2018/044740

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☒ Claims Nos.: 6-11, 18-32, 36, 38-42
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US2018/044740

A. CLASSIFICATION OF SUBJECT MATTER

IPC(8) - C07K 14/47; C12N 5/06 (2018.01)

CPC - C07K 14/4702; C07K 2319/10; C12N 5/0634; C12N 2501/606 (2018.08)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

See Search History document

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

USPC - 424/573; 424/93.21; 424/192.1 (keyword delimited)

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

See Search History document

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2014/0356392 A1 (TAIGA BIOTECHNOLOGIES, INC.) 04 December 2014 (04.12.2014) entire document	33, 35, 37
Y		1-4, 12-17, 34
Y	US 2015/0164950 A1 (TAIGA BIOTECHNOLOGIES, INC.) 18 June 2015 (18.06.2015) entire document	1-4, 12-17, 34
Y	US 2012/0251563 A1 (NICCHITTA et al) 04 October 2012 (04.10.2012) entire document	3, 16
P, X	US 2018/0036396 A1 (TAIGA BIOTECHNOLOGIES, INC.) 08 February 2018 (08.02.2018) entire document	1-5, 12-17, 33-35, 37
A	US 6,358,739 B1 (BAETGE et al) 19 March 2002 (19.03.2002) entire document	1-5, 12-17, 33-35, 37
A	US 2006/0222657 A1 (DOWDY et al) 05 October 2006 (05.10.2006) entire document	1-5, 12-17, 33-35, 37
A	US 2012/0027792 A1 (PAVLAKIS et al) 02 February 2012 (02.02.2012) entire document	1-5, 12-17, 33-35, 37

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

26 September 2018

Date of mailing of the international search report

16 OCT 2018

Name and mailing address of the ISA/US

Mail Stop PCT, Attn: ISA/US, Commissioner for Patents

P.O. Box 1450, Alexandria, VA 22313-1450

Facsimile No. 571-273-8300

Authorized officer

Blaine R. Copenheaver

PCT Helpdesk: 571-272-4300

PCT OSP: 571-272-7774