COMPOUNDS AND COMPOSITIONS FOR TREATING HER2 POSITIVE TUMORS

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims the benefit of, and priority to, Chinese Patent Application Serial Nos. 201410011324.5, 201410011262.8, and 201410011362.0, all filed January 10, 2014, the entire disclosures of which are hereby incorporated by reference in their entireties.

FIELD OF THE INVENTION

[0002] The present invention relates to compounds for targeted immunotherapy, as well as compositions comprising the same. Further, the present invention relates to the use of the compounds in the treatment of diseases such as cancer.

BACKGROUND OF THE INVENTION

[0003] Therapeutic antibodies have been used in clinical applications for over twenty years. Currently, there are fifteen anti-tumor antibody drugs in clinic, including Rituxan (1997), Herceptin (1998), Mylotarg (2000), Campath (2001), Zevalin (2002), Bexxer (2003), Avastin (2004), Erbitux (2004), Vectibix (2006); Arzerra (2009); Benlysta (2011); Yervoy (2011); Adcetris (2011); Perjeta (2012); and Kadcyla (2013). These antibodies target mainly four molecules: EGFR, Her2, CD20 and VEGF.

[0004] In general, therapeutic antibodies kill tumor cells via three mechanisms (Scott AM, Wolchok JD, Old LJ. Antibody therapy of cancer. Nat Rev Cancer. (2012), 12 :278-87): (1) Direct antibody action, that is, blockade or agonist activity of ligand / receptor signaling, induction of apoptosis, and delivery of drugs or cytotoxic agents. Antibody receptor activation activity can produce direct tumor cell killing effect. For example, some antibodies can bind to receptors on the surface of tumor cells, activate the receptor, leading to apoptosis (e.g., in mitochondria). Antibodies can also mediate tumor cell killing by receptor-antagonistic activity. For example, certain antibodies can bind to cell surface receptors and block dimerization, kinase activation and downstream signaling, thereby inhibiting proliferation and promote apoptosis. Binding of antibodies to an enzyme can lead to neutralization, signal abrogation, and cell death. (2) Through immune-mediated cell killing mechanisms include complement-dependent cytotoxicity (CDC), antibody-dependent cell-mediated cytotoxicity (ADCC), T cell function regulation, etc. Immune-mediated killing of tumor cells can be accomplished through the following ways: induction of phagocytosis, complement activation, antibody-dependent cell-mediated cytotoxicity, genetically modified T cells being targeted to the tumor by single-chain variable fragment (scFv), through antibody-mediated antigenic cross presentation to dendritic cell to activate T cells, inhibition of T cell inhibitory

receptors, such as cytotoxic T lymphocyte-associated antigen 4 (CTLA4). Of them, the Fc portion of the antibody feature is especially important for CDC and ADCC-mediated tumor cell killing effect. (3) Specific effect of antibody on tumor vasculature and matrix, through trapping of vascular receptor antagonist or ligand to induce vascular and stromal cells ablation, including: stromal cell inhibition, delivery of toxins to stromal cells, and delivery of toxins to the vasculature. (Scott AM, Wolchok JD, Old LJ. Antibody therapy of cancer. Nat Rev Cancer. 2012, 12 (4):278-87).

[0005] Therapeutic monoclonal antibody drugs have advanced anti-cancer drug research and development. However, some issues still need further study to be solved, such as antibody immunogenicity, tolerance of long-term use of tumor target, and long-term effects of simple single blockade of signal transduction pathway. In short, a simple majority of antibodies are difficult to achieve long-term efficient inhibition and killing of tumor cells.

[0006] In 1964, "Nature" magazine presented the new idea of antibody - drug conjugates (ADC) technology, which in recent years have seen breakthroughs. ADC covalently links antibody with a highly toxic drug (toxin) through a chemical linker (linker). Antibody recognizes cancer cell surface antigen molecule, the endocytosis ADC brings it into cytoplasm, and in particular intracellular environment toxins released after hydrolysis of the linker kills cells.

[0007] Seattle Genetics has developed such drug Brentuximab Vedotin (trade name Adcetris) that has been approved by the FDA to market. It is monomethyl auristatin E (MMAE), a synthetic toxic anti-cancer drug, coupled with antibody targeting lymphoma cells specific CD30 molecule, with improved efficacy of killing tumor cells.

[0008] Currently, there are more than dozens of such ADC drugs in clinical trials. Among them, Genentech and Immunogen jointly developed trastuzumab coupled with maytansines as a drug named ado-trastuzumab emtansine (Kadcyla), also known as T-DM1, to treat breast cancer. In February 2013, the FDA has approved T-DM1 for human epidermal growth factor receptor 2 (Her2)-positive metastatic breast cancer. Maytansines is a small molecule toxin that can bind tubulin and prevent formation of microtubules by forming non-reducing dual - maleimide - propanediol complex. Trastuzumab acts on breast cancer and gastric cancer by targeting human Her2. It was approved for Her2-positive cancer. However, trastuzumab cannot promote apoptosis of all of the Her2 positive cells. T-DM1 combines the selective targeting Her2 receptor trastuzumab with the potent cytotoxic agent maytansine to kill tumor cells. T-DM1 antibody binds Her2 receptors, causing cellular internalization of the maytansines released from conjugates, thereby killing the tumor cells. T-DM1 has better overall efficacy and pharmacokinetic properties and low toxicity.

[0009] Traditional small molecule chemotherapeutic drugs have strong toxicity and pharmacokinetic advantages, but in the process of treatment of tumors may affect other physiological targets with serious side effects. Antibody - drug conjugates combines targeting function and small molecule drug with particular pharmacokinetics. The structure of antibody-drug conjugates is the attachment of a monoclonal antibody with targeting function to a compound with specific pharmacological properties. This technique requires the therapeutic antibody have binding specificity to a target, to be coupled to a molecule with therapeutic effect or other functions such as cyto-toxins. Many factors affect the effect of this type of antibodies, such as endocytosis of the coupled antibody, stability of the coupling, and release and killing activity of the toxins.

[0010] Toxin molecules currently being used include tubulin inhibitors Auristatin analogues monomethyl auristatin E, monomethyl auristatin F and maytansine. Monomethyl auristatin E is a synthetic microtubule polymer inhibitor that can inhibit microtubule aggregation, interfere tumor cell mitosis and induce apoptosis (Naumovski L and Junutula JR. Glembatumumab vedotin, a conjugate of an anti-glycoprotein non-metastatic melanoma protein B mAb and monomethyl auristatin E for treatment of melanoma and breast cancer. Curr Opin Mol Ther 2003; 12 (2): 248-57.Francisco JA, Cerveny CG et al. cAC10-vcMMAE, an anti-CD30-monomethyl auristatin E conjugate with potent and selective antitumor activity. Blood 102 (4): 1458-65. Monomethyl auristatin F is an anti-mitotic Auristatin derivative with a charged phenylalanine residue at C terminus. In comparison to uncharged MMAE it minimizes damage to cell signaling pathway and minimizes cytotoxicity. A large number of test with CD30 cells found that mAb-maleimidocaproyl-valine - citrulline-p-aminobenzyloxycarbonyl-MMAF (mAb-L1-MMAF) has a toxicity that is 2,200 times stronger than MMAF only (Doronina SO et al., Enhanced activity of monomethylauristatin F through monoclonal antibody delivery: effects of linker technology on efficacy and toxicity. Bioconjug Chem, 2006; 17 (1): p114-24). Maytansine is an antimitotic agent acting as an inhibitor of tubulin polymerization, thus interfering with formation of microtubules in the cell nucleus. Maytansine also inhibits DNA, RNA, and protein synthesis, with the greatest effect being seen on DNA synthesis.

[0011] Antibodies - drug conjugates have direct and indirect anti-cancer effect. The antibody blocks or activates ligand / receptor signaling, induces apoptosis, and at the same time can present or deliver payload drug directly or indirectly (such as a drug, toxin, small interfering RNA or radioisotope) to the tumor cells. Therapeutic antibody drug conjugate utilizes dual characteristics of the antibody and the coupled drug, first is the binding function that it specifically binds to the target molecule, second is the tumor cell killing function of the antibody itself, and the third is the particular effect of the conjugated drug. Current antibody - drug conjugates drugs are limited in how to kill tumor cells directly. However,

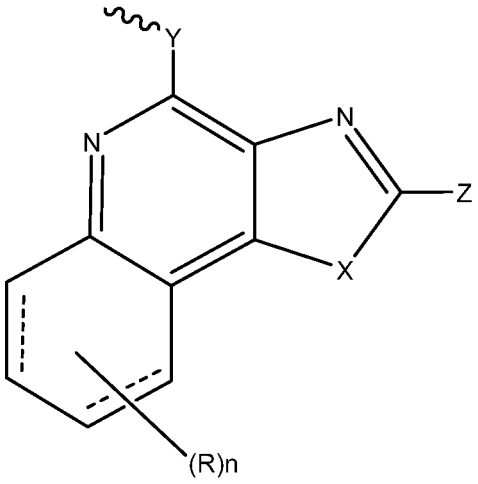
because of the tough requirement of technologies in antibody, linker molecule, toxin molecules, and conjugation, as well as the limitation of bringing toxins within the tumor microenvironment molecules, there are still some difficulties in actual clinical studies.

SUMMARY OF THE INVENTION

[0012] In one aspect, the present invention provides a compound having the structure of Formula (Ia):

TM-L-AM (Ia),

wherein TM is an antibody or a functional fragment thereof that specifically binds to Her2/Neu (e.g., anti-Her2 antibody), AM is an activating moiety that is represented by structure of formula (I):



(I)

wherein dashed line represents bond or absence of bond, ∿ is the point to be connected to the linker;

X is S or $-NR_1$, R_1 is $-W_0—W_1—W_2—W_3—W_4$,

W_0 is a bond, alkyl, alkenyl, alkynyl, alkoxy, or –alkyl-S-alkyl--,

W_1 is a bond, --O--, or $-NR_2$--, wherein R_2 is hydrogen, alkyl or alkenyl,

W_2 is a bond, --O--, --C(O)--, --C(S)--, or $-S(O)_2—$,

W_3 is a bond, $--NR_3$--, wherein R_3 is hydrogen, alkyl or alkenyl,

W_4 is hydrogen, alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, aryl, aryloxy, heteroaryl, or heterocyclyl, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, $--NH_2$, nitro, --alkyl-

hydroxyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --alkyl-O-R_4, --C(O)-R_4, --alkyl-C(O)-R_4, --alkyl-C(O)-O-R_4, --C(O)-O-R_4, --S-R_4, --$S(O)_2$-R_4, --NH-$S(O)_2$-R_4, --alkyl-S-R4, --alkyl-$S(O)_2$-R_4, --NHR_4, --NR_4R_4,--NH-alkyl-R_4, halogen, --CN, --NO_2, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

Z is hydrogen, alkyl, alkenyl, alkynyl, alkoxy, aryl, haloalkyl, heteroaryl, heterocyclyl, each of which can be optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, aryl, heteroaryl, heterocyclyl, halogen, cyano, nitro, --$N(R_5)_2$, --alkoxy-alkyl, --alkoxy-alkenyl, --C(O)-alkyl, --C(O)-O-alkyl, --O-C(O)-alkyl, --C(O)-$N(R_5)_2$, aryl, heteroaryl, --CO-aryl, and –CO-heteroaryl, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl;

R is hydrogen, alkyl, alkoxy, haloalkyl, halogen, aryl, heteroaryl, heterocyclyl, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, --NH_2, nitro, --alkyl-hydroxyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --alkyl-O-R_4, --C(O)-R_4, --C(O)-NH-R_4, --C(O)-NR_4R_4, --alkyl-C(O)-R_4, --alkyl-C(O)-O-R_4, --C(O)-O-R_4, --O-C(O)-R_4, --S-R_4, --C(O)-S-R_4, --S-C(O)-R_4, --$S(O)_2$-R_4, --NH-$S(O)_2$-R_4, --alkyl-S-R_4, --alkyl-$S(O)_2$-R_4, --NHR_4, --NR_4R_4,--NH-alkyl-R_4, halogen, --CN, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, alkoxy, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

n is 0, 1, 2, 3, or 4;

Y is –NR_6R_7, –$CR_6R_7R_8$, or –alkyl-NH_2, each of which can be optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, --NH_2, halogen, --$N(R_5)_2$, --alkoxy-alkyl, --alkoxy-alkenyl, --C(O)-alkyl, --C(O)-O-alkyl, --C(O)-$N(R_5)_2$, aryl, heteroaryl, --CO-aryl, and –CO-heteroaryl,

wherein R_6, R_7 and R_8 are independently hydrogen, alkyl, alkenyl, alkoxy, alkylamino, dialkylamino, alkylthio, arylthio, --alkyl-hydroxyl, --alkyl-C(O)-O-R_9, --alkyl-C(O)-R_9, or –alkyl-O-C(O)-R_9, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl, wherein R_9 is hydrogen, alkyl, alkenyl, halogen, or haloalkyl;

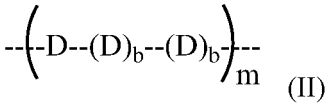
X and Z taken together may optionally form a (5-9)-membered ring;

or a pharmaceutically acceptable salt or solvate thereof.

[0013] In some embodiments, AM is a compound of formula (I) selected from 2-propylthiazolo[4,5-c]quinolin-4-amine, 1-(2-methylpropyl)-1H-imidazo[4,5-c]quinolin-4-amine, 4-amino-2-(ethoxymethyl)-

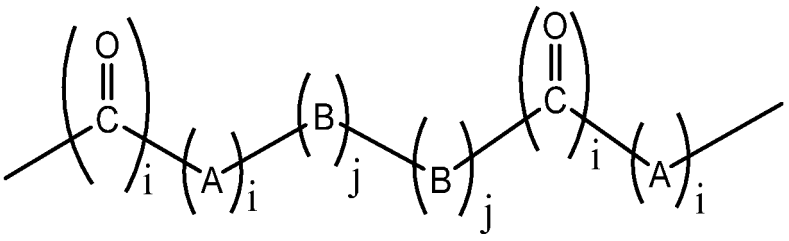
a,a-di-methyl-1H-imidazo[4,5-c]quinoline-1-ethanol, 1-(4-amino-2-ethylaminomethylimidazo-[4,5-c]quinolin-1-yl)-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl-]methanesulfonamide, 4-amino-2-ethoxymethyl-aa-dimethyl-6,7,8,9-tetrahydro-1h-imidazo[4,5-c]quinoline-1-ethanol, 4-amino-aa-dimethyl-2-methoxyethyl-1h-imidazo[4,5-c]quinoline-1-ethanol, 1-{2-[3-(benzyloxy)propoxy]ethyl}-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, N-[4-(4-amino-2-butyl-1H-imidazo[4,5- c][1,5]naphthyridin-1-yl)butyl]-n'-butylurea, N1-[2-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5] naphthyridin-1-yl)ethyl]-2-amino-4- methylpentanamide, N-(2-{2-[4-amino-2-(2-methoxyethyl)-1H- imidazo[4,5-c]quinolin-1-yl]ethoxy}ethyl)-n'- phenylurea, 1-(2-amino-2-methylpropyl)-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, 1-{4-[(3,5-dichlorophenyl)sulfonyl]butyl}-2-ethyl- 1H-imidazo[4,5-c]quinolin-4-amine, N-(2-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]ethoxy}ethyl)-n'-cyclohexylurea, N-{3-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]propyl}-n'-(3-cyanophenyl)thiourea, N-[3-(4-amino-2-butyl-1H-imidazo[4,5-c]quinolin-1- yl)-2,2-dimethylpropyl]benzamide, 2-butyl-1-[3-(methylsulfonyl)propyl]-1H- imidazo[4,5-c]quinolin-4-amine, N-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]-1,1-dimethylethyl}-2- ethoxyacetamide, 1-[4-amino-2-ethoxymethyl-7-(pyridin-4-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 1-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, N-{3-[4-amino-1-(2-hydroxy-2-methylpropyl)-2- (methoxyethyl)-1H-imidazo[4,5-c]quinolin-7-yl]phenyl}methanesulfonamide, 1-[4-amino-7-(5-hydroxymethylpyridin-3-yl)-2-(2- methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 3-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]propane-1,2-diol, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-propylurea, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-cyclopentylurea, 1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methyl]-2- (ethoxymethyl)-7-(4-hydroxymethylphenyl)-1H- imidazo[4,5-c]quinolin-4-amine, 4-[4-amino-2-ethoxymethyl-1-(2-hydroxy-2-methylpropyl)-1H-imidazo[4,5-c]quinolin-7-yl]-N- methoxy-N-methylbenzamide, 2-ethoxymethyl-N1-isopropyl-6,7,8,9-tetrahydro-1H-imidazo[4,5-c]quinoline-1,4-diamine, 1-[4-amino-2-ethyl-7-(pyridin-4-yl)-1H-imidazo[4,5- c]quinolin-1-yl]-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl]methanesulfonamide, or N-[4-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)butyl]-n'-cyclohexylurea.

[0014] In some embodiments, L is represented by structure of formula (II):



(II)

m is 1, 2, 3, 4, 5, or 6, each b independently is 0 or 1, and D is independently represented by structure of formula (III):



(III)

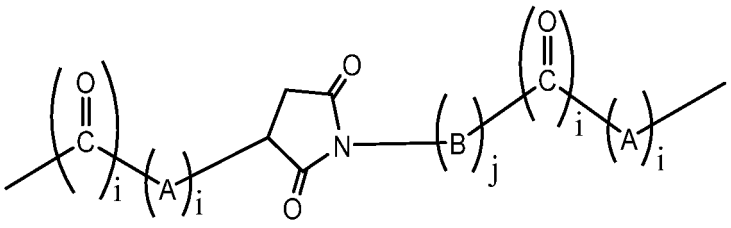
wherein each i independently is 0 or 1;

each j independently is 0, 1, 2, 3, 4, 5, or 6;

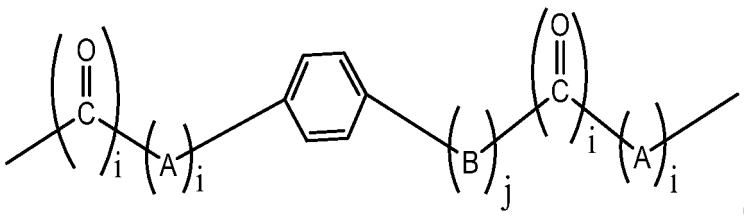
each A independently is S, O, or N-Ra, wherein Ra is hydrogen, alkyl, alkenyl, or alkoxy;

each B independently is alkyl, alkenyl, --O-alkyl--, --alkyl-O--, --S-alkyl--, --alkyl-S--, aryl, heteroaryl, heterocyclyl, or peptide, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --C(O)-R_4, --C(O)-O-R_4, --S-R_4, --$S(O)_2$-R_4, --NHR_4, --NH-alkyl-R_4, halogen, --CN, --NO_2, and –SH, wherein R_4 is alkyl, alkenyl, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl.

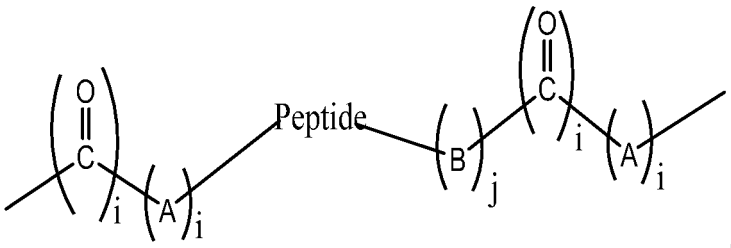
[0015] In some embodiments, the linker is presented by the following strucutres of formula (V)-(VII):



(V)



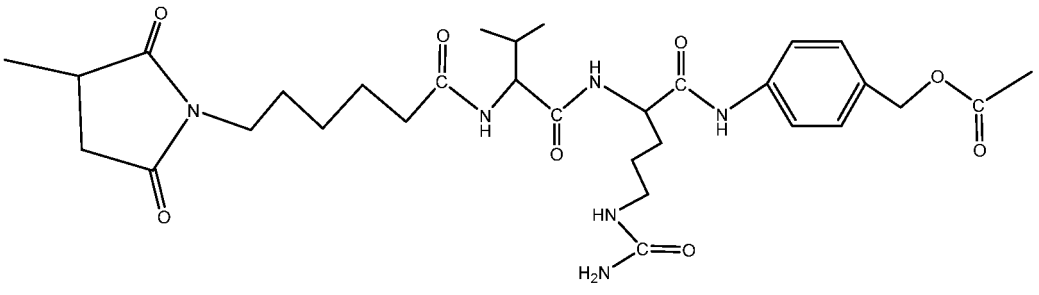
(VI)



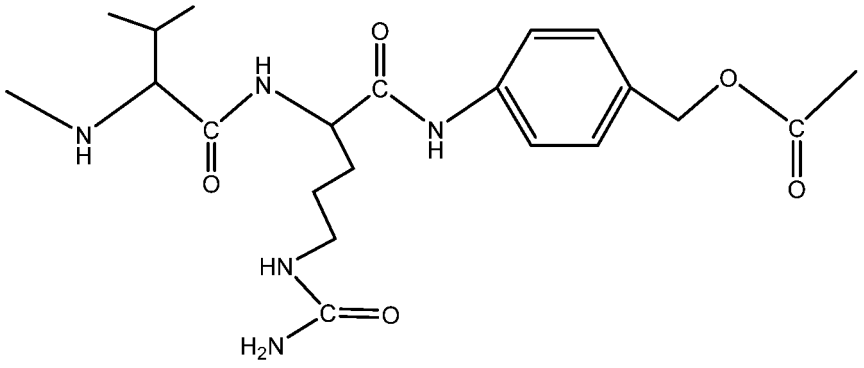
(VII)

A, B, i and j are defined above.

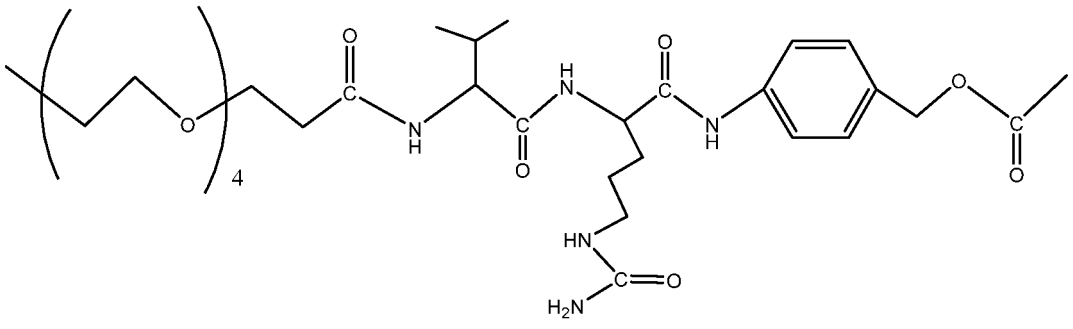
[0016] In some embodiments, the linker is selected from S1, S2, S3, S4, S5, S6, S7, –Gly-Phe-Leu-Gly-, -Ala-Leu-Ala-Leu-, -Phe-Arg-, -Phe-Lys-, -Val-Lys-, -Val-Ala-, or Val-Cit-, wherein S1-S7 are represented by the following structures:



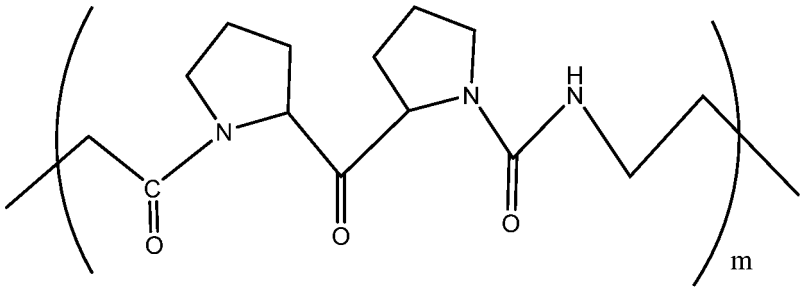
S1



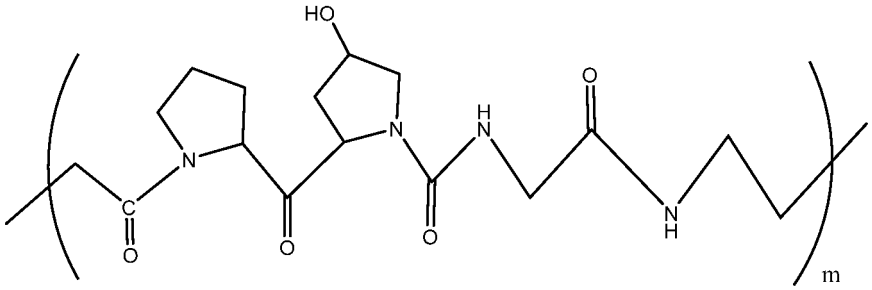
S2



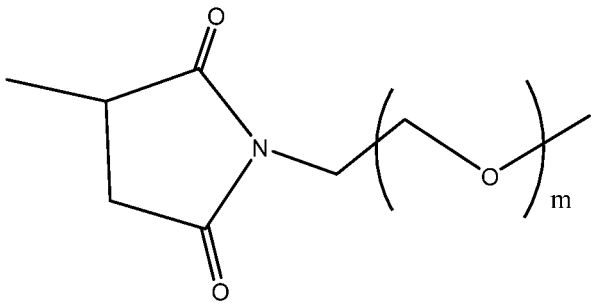
S3



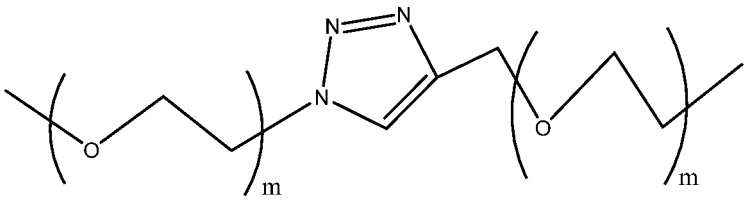
S4



S5



S6



S7,

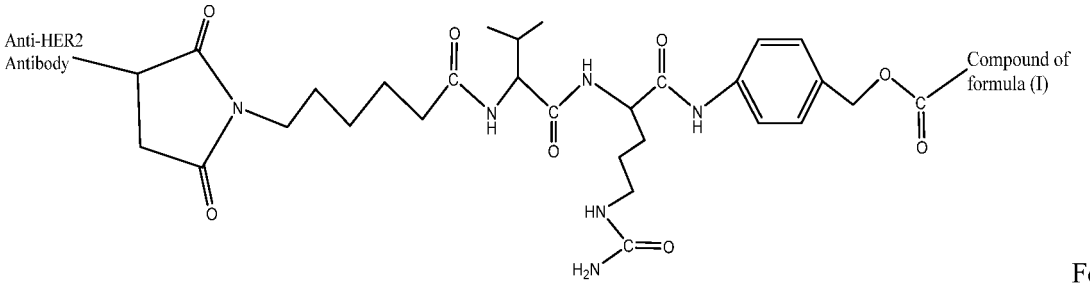
wherein each m is independently 1 to 20.

[0017] In some embodiments, the antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22).

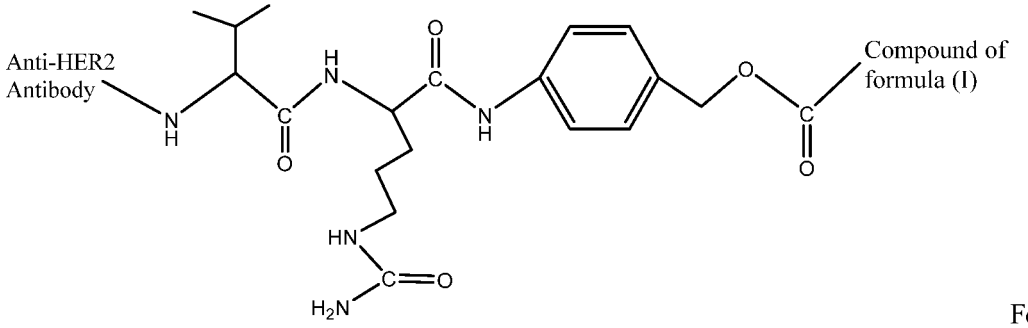
[0018] In some embodiments, TM is an anti-HER2 antibody; L is selected from S1, S2, S3, S4, S5, S6, S7, –Gly-Phe-Leu-Gly-, -Ala-Leu-Ala-Leu-, -Phe-Arg-, -Phe-Lys-, -Val-Lys-, -Val-Ala-, or Val-Cit-; AM is a compound of formula (I) selected from selected from 2-propylthiazolo[4,5-c]quinolin-4-amine, 1-(2-methylpropyl)-1H-imidazo[4,5-c]quinolin-4-amine, 4-amino-2-(ethoxymethyl)-a,a-di-methyl-1H-imidazo[4,5-c]quinoline-1-ethanol, 1-(4-amino-2-ethylaminomethylimidazo-[4,5-c]quinolin-1-yl)-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl-]methanesulfonamide, 4-amino-2-ethoxymethyl-aa-dimethyl-6,7,8,9-tetrahydro-1h-imidazo[4,5-c]quinoline-1-ethanol, 4-amino-aa-dimethyl-2-methoxyethyl-1h-imidazo[4,5-c]quinoline-1-ethanol, 1-{2-[3-(benzyloxy)propoxy]ethyl}-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, N-[4-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)butyl]-n'-butylurea, N1-[2-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5] naphthyridin-1-yl)ethyl]-2-amino-4- methylpentanamide, N-(2-{2-[4-amino-2-(2-methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]ethoxy}ethyl)-n'- phenylurea, 1-(2-amino-2-methylpropyl)-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, 1-{4-[(3,5-dichlorophenyl)sulfonyl]butyl}-2-ethyl-1H-imidazo[4,5-c]quinolin-4-amine, N-(2-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]ethoxy}ethyl)-n'-cyclohexylurea, N-{3-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]propyl}-n'-(3-cyanophenyl)thiourea, N-[3-(4-amino-2-butyl-1H-imidazo[4,5-c]quinolin-1- yl)-2,2-dimethylpropyl]benzamide, 2-butyl-1-[3-(methylsulfonyl)propyl]-1H- imidazo[4,5-c]quinolin-4-amine, N-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]-1,1-dimethylethyl}-2-ethoxyacetamide, 1-[4-amino-2-ethoxymethyl-7-(pyridin-4-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 1-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, N-{3-[4-amino-1-(2-hydroxy-2-methylpropyl)-2- (methoxyethyl)-1H-imidazo[4,5-c]quinolin-7-yl]phenyl}methanesulfonamide, 1-[4-amino-7-(5-hydroxymethylpyridin-3-yl)-2-(2-methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 3-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]propane-1,2-diol, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-propylurea, 1-[2-(4-amino-2-ethoxymethyl-1H-

imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-cyclopentylurea, 1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methyl]-2- (ethoxymethyl)-7-(4-hydroxymethylphenyl)-1H- imidazo[4,5-c]quinolin-4-amine, 4-[4-amino-2-ethoxymethyl-1-(2-hydroxy-2-methylpropyl)-1H-imidazo[4,5-c]quinolin-7-yl]-N- methoxy-N-methylbenzamide, 2-ethoxymethyl-N1-isopropyl-6,7,8,9-tetrahydro-1H-imidazo[4,5-c]quinoline-1,4-diamine, 1-[4-amino-2-ethyl-7-(pyridin-4-yl)-1H-imidazo[4,5- c]quinolin-1-yl]-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl]methanesulfonamide, or N-[4-(4-amino-2-butyl-1H-imidazo[4,5- c][1,5]naphthyridin-1-yl)butyl]-n'-cyclohexylurea, wherein the amine group on the quinoline ring is the point of connection with the linker. Preferably, the anti-HER2 antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); L is S1, S2 or S3. Also preferably the anti-HER2 antibody is trastuzumab (Herceptin); AM is Resiquimod or Imiquimod, wherein the amine group on the quinoline ring is the point of connection with the linker.

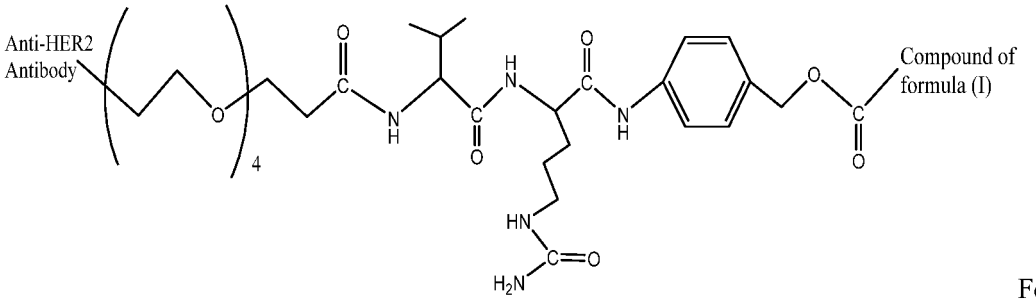
[0019] In another aspect, the present invention provides a compopund having the structures of Formula A-C:



Formula A,



Formula B,



Formula C;

or a pharmaceutically acceptable salt or solvate thereof.

[0020] In one embodiment of Formula A-C, the anti-HER2 antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); the compound of formula (I) is selected from 2-propylthiazolo[4,5-c]quinolin-4-amine, 1-(2-methylpropyl)-1H-imidazo[4,5-c]quinolin-4-amine, 4-amino-2-(ethoxymethyl)-a,a-di-methyl-1H-imidazo[4,5-c]quinoline-1-ethanol, 1-(4-amino-2-ethylaminomethylimidazo-[4,5-c]quinolin-1-yl)-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl-]methanesulfonamide, 4-amino-2-ethoxymethyl-aa-dimethyl-6,7,8,9-tetrahydro-1h-imidazo[4,5-c]quinoline-1-ethanol, 4-amino-aa-dimethyl-2-methoxyethyl-1h-imidazo[4,5-c]quinoline-1-ethanol, 1-{2-[3-(benzyloxy)propoxy]ethyl}-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, N-[4-(4-amino-2-butyl-1H-imidazo[4,5- c][1,5]naphthyridin-1-yl)butyl]-n'-butylurea, N1-[2-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5] naphthyridin-1-yl)ethyl]-2-amino-4- methylpentanamide, N-(2-{2-[4-amino-2-(2-methoxyethyl)-1H- imidazo[4,5-c]quinolin-1-yl]ethoxy}ethyl)-n'- phenylurea, 1-(2-amino-2-methylpropyl)-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, 1-{4-[(3,5-dichlorophenyl)sulfonyl]butyl}-2-ethyl- 1H-imidazo[4,5-c]quinolin-4-amine, N-(2-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]ethoxy}ethyl)-n'-cyclohexylurea, N-{3-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]propyl}-n'-(3-cyanophenyl)thiourea, N-[3-(4-amino-2-butyl-1H-imidazo[4,5-c]quinolin-1- yl)-2,2-dimethylpropyl]benzamide, 2-butyl-1-[3-(methylsulfonyl)propyl]-1H- imidazo[4,5-c]quinolin-4-amine, N-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]-1,1-dimethylethyl}-2- ethoxyacetamide, 1-[4-amino-2-ethoxymethyl-7-(pyridin-4-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 1-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, N-{3-[4-amino-1-(2-hydroxy-2-methylpropyl)-2- (methoxyethyl)-1H-imidazo[4,5-c]quinolin-7-yl]phenyl}methanesulfonamide, 1-[4-amino-7-(5-hydroxymethylpyridin-3-yl)-2-(2- methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 3-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]propane-1,2-diol, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-propylurea, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-cyclopentylurea, 1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methyl]-2- (ethoxymethyl)-7-(4-hydroxymethylphenyl)-1H- imidazo[4,5-c]quinolin-4-amine, 4-[4-amino-2-ethoxymethyl-1-(2-hydroxy-2-methylpropyl)-1H-imidazo[4,5-c]quinolin-7-yl]-N- methoxy-N-methylbenzamide, 2-ethoxymethyl-N1-isopropyl-6,7,8,9-tetrahydro-1H-imidazo[4,5-c]quinoline-1,4-diamine, 1-[4-amino-2-ethyl-7-(pyridin-4-yl)-1H-imidazo[4,5- c]quinolin-1-yl]-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl]methanesulfonamide, or N-[4-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)butyl]-n'-cyclohexylurea, wherein the amine group on the quinoline ring is the point of connection with the linker. Also in one embodiment of Formula A-C, the anti-HER2 antibody is

trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); the compound of formula (I) is Resiquimod or Imiquimod, wherein the amine group on the quinoline ring is the point of connection with the linker. Also in one embodiment of Formula A-C, the anti-HER2 antibody is trastuzumab (Herceptin); AM is is Resiquimod or Imiquimod, wherein the amine group on the quinoline ring is the point of connection with the linker.

[0021] In some embodiments, the present invention provides a pharmaceutical composition comprising a compound as provided herein, or a pharmaceutically acceptable salt thereof, and/or one or more pharmaceutically acceptable carriers.

[0022] In some embodiments, the pharmaceutical composition further comprises an additional therapeutic agent. In some embodiments, the additional therapeutic agent is an anticancer agent. In some embodiments, the additional therapeutic agent is an antimetabolite, an inhibitor of topoisomerase I and II, an alkylating agent, a microtubule inhibitor, an antiandrogen agent, a GNRh modulator or mixtures thereof. In some embodiments, the additional therapeutic agent is selected from the group consisting of tamoxifen, raloxifene, anastrozole, exemestane, letrozole, imatanib, paclitaxel, cyclophosphamide, lovastatin, minosine, gemcitabine, cytarabine, 5- fluorouracil, methotrexate, docetaxel, goserelin, vincristine, vinblastine, nocodazole, teniposide etoposide, gemcitabine, epothilone, vinorelbine, camptothecin, daunorubicin, actinomycin D, mitoxantrone, acridine, doxorubicin, epirubicin, or idarubicin.

[0023] In yet another aspect, the present invention provides a method of inhibiting proliferation of a HER2 positive tumor cell comprising administering to said tumor cell a compound of the present invention.

[0024] In some embodiments, the present invention provides a method of treating a HER2 positive tumor/cancer in a subject comprising administering to the subject the compounds of the present invention. In some embodiments, the the tumor/cell is selected from esophageal, stomach, colon, rectal, , pancreatic, lung, breast, cervix uteri, corpus uteri, ovary, bladder, head and neck,endometrial, osteosarcoma, prostate, neuroblastoma,

INCORPORATION BY REFERENCE

[0025] All publications, patents, and patent applications mentioned in this specification are herein incorporated by reference to the same extent as if each individual publication, patent, or patent application was specifically and individually indicated to be incorporated by reference.

BRIEF DESCRIPTION OF THE DRAWINGS

[0026] The novel features of the invention are set forth with particularity in the appended claims. A better understanding of the features and advantages of the present invention will be obtained by reference to the following detailed description that sets forth illustrative embodiments, in which the principles of the invention are utilized, and the accompanying drawings of which:

[0027] **Figure 1** depicts conjugated compound characteristics. Binding of Trastuzumab MC-vc-resiquimod and Trastuzu -resiquimod conjugates to Her2 expressing L cells is compared with unconjugated Trastuzumab (as a reference), control human IgG, and irrelevant human IgG1 antibody, as determined by FACS analysis.

[0028] **Figure 2** depicts in vitro antibody-dependent cellular cytotoxicity (ADCC) analysis of Trastuzumab, Trastuzumab MC-vc-TLRL and Trastuzumab MC-TLRL conjugates. PBMCs (effector cells) were added with SKBR3 cells (target cells) into 96-well plates (ratio targets/effectors 1/60) for 17h containing control human IgG1, trastuzumab, or Trastuzumab MC-vc-TLRL and Trastuzumab MC-TLRL conjugates at different concentrations. All experiments were performed in duplicate. Analysis was based on the negative SKBR3 population determination. The dead cells were stained by LDH.

[0029] **Figure 3A** depicts the percentages of DCs before and post enrichment. The numbers in upper two plots represent the percentages of DCs (HLA-DR+Lin-) of total cells before and after lineage depletion. The numbers in lower plots represent percentages of mDC (CD11C +CD123-) and pDC (CD123+CD11C-) of total DCs before and after lineage depletion. **Figures 3B and 3C** depicts analysis of cytokine production by purified human DCs. Purified human DCs were plated in a 96-well plate and cultured with allogeneic untreated (medium) or treated different concentration of TLRL, or Trastuzumab MC-vc-TLRL and Trastuzumab MC-TLRL directly for 20-22h in 37°C incubator. The supernatant were collected and human IFN-α, IL-6, IL-12(p70) and TNF-α were analyzed by ELISA. Data are given as mean ± SD of triplicate cultures and are representative of independent experiments from two of three healthy donors.

[0030] **Figures 4A-4C** depict therapeutic efficacy of Trastuzumab MC-vc-TLRL and Trastuzumab MC-TLRL in PDX gastric tumor model. BALB/c nu/nu nude mice 6-8 week-old female mice bearing subcutaneous patient derived gastric tumor were treated intravenously with 10mg/kg of Trastuzumab MC-vc-TLRL, Trastuzumab MC-TLRL, or unconjugated Trastuzumab or saline (12 mice per group). Treatment was performed weekly for a period of 45 days. Therapy was initiated when tumors reached a size on average of 170 mm^3. **Figure 4A:** Data represents mean tumor volumes (mean ± SD). Tumor growth curves were stopped when tumors reached a size of 2000 mm^3. **Figure 4B:** Body weight

variations of the mice during and after therapy. No detectable weight loss was observed. **Figure 4C:** Survival curves of treatment and control groups; substantial prolongation for mice which received Trastuzumab MC-vc-TLRL, or unconjugated Trastuzumab.

DETAILED DESCRIPTION OF THE INVENTION

[0031] Several aspects of the invention are described below with reference to example applications for illustration. It should be understood that numerous specific details, relationships, and methods are set forth to provide a full understanding of the invention. One having ordinary skill in the relevant art, however, will readily recognize that the invention can be practiced without one or more of the specific details or with other methods. The present invention is not limited by the illustrated ordering of acts or events, as some acts may occur in different orders and/or concurrently with other acts or events.

[0032] Furthermore, not all illustrated acts or events are required to implement a methodology in accordance with the present invention.

[0033] The terminology used herein is for the purpose of describing particular embodiments only and is not intended to be limiting of the invention. 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. Furthermore, to the extent that the terms "including", "includes", "having", "has", "with", or variants thereof are used in either the detailed description and/or the claims, such terms are intended to be inclusive in a manner similar to the term "comprising".

[0034] The term "about" or "approximately" means within an acceptable error range for the particular value as determined by one of ordinary skill in the art, which will depend in part on how the value is measured or determined, i.e., the limitations of the measurement system. For example, "about" can mean within 1 or more than 1 standard deviation, per the practice in the art. Alternatively, "about" can mean a range of up to 20%, preferably up to 10%, more preferably up to 5%, and more preferably still up to 1% of a given value. Alternatively, particularly with respect to biological systems or processes, the term can mean within an order of magnitude, preferably within 5-fold, and more preferably within 2-fold, of a value. Where particular values are described in the application and claims, unless otherwise stated the term "about" meaning within an acceptable error range for the particular value should be assumed.

Definitions and Abbreviations

[0035] Unless defined otherwise, all technical and scientific terms used herein generally have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Generally, the nomenclature used herein and the laboratory procedures in cell culture, molecular genetics, organic chemistry and nucleic acid chemistry and hybridization are those well-known and commonly

employed in the art. Standard techniques are used for nucleic acid and peptide synthesis. The techniques and procedures are generally performed according to conventional methods in the art and various general references, which are provided throughout this document. The nomenclature used herein and the laboratory procedures in analytical chemistry, and organic synthetic described below are those well-known and commonly employed in the art. Standard techniques, or modifications thereof, are used for chemical syntheses and chemical analyses.

[0036] The term "alkyl," by itself or as part of another substituent, means, unless otherwise stated, a straight or branched chain, or cyclic hydrocarbon radical, or combination thereof, which may be fully saturated, mono- or polyunsaturated and can include di- and multivalent radicals, having the number of carbon atoms designated (*i.e.* C_1-C_{10} means one to ten carbons). Examples of saturated hydrocarbon radicals include, but are not limited to, groups such as methyl, ethyl, n-propyl, isopropyl, n-butyl, t-butyl, isobutyl, sec-butyl, cyclohexyl, (cyclohexyl)methyl, cyclopropylmethyl, homologs and isomers of, for example, n-pentyl, n-hexyl, n-heptyl, n-octyl, and the like. An unsaturated alkyl group is one having one or more double bonds or triple bonds. Examples of unsaturated alkyl groups include, but are not limited to, vinyl, 2-propenyl, crotyl, 2-isopentenyl, 2-(butadienyl), 2,4-pentadienyl, 3-(1,4-pentadienyl), ethynyl, 1- and 3-propynyl, 3-butynyl, and the higher homologs and isomers. The term "alkyl," unless otherwise noted, is also meant to include those derivatives of alkyl defined in more detail below, such as "heteroalkyl." Alkyl groups, which are limited to hydrocarbon groups, are termed "homoalkyl".

[0037] The term "alkylene" by itself or as part of another substituent means a divalent radical derived from an alkane, as exemplified, but not limited, by $-CH_2CH_2CH_2CH_2-$, and further includes those groups described below as "heteroalkylene." Typically, an alkyl (or alkylene) group will have from 1 to 24 carbon atoms, with those groups having 10 or fewer carbon atoms being preferred in the present invention. A "lower alkyl" or "lower alkylene" is a shorter chain alkyl or alkylene group, generally having eight or fewer carbon atoms.

[0038] The terms "alkoxy," "alkylamino" and "alkylthio" (or thioalkoxy) are used in their conventional sense, and refer to those alkyl groups attached to the remainder of the molecule via an oxygen atom, an amino group, or a sulfur atom, respectively.

[0039] The term "heteroalkyl," by itself or in combination with another term, means, unless otherwise stated, a stable straight or branched chain, or cyclic hydrocarbon radical, or combinations thereof, consisting of the stated number of carbon atoms and at least one heteroatom selected from the group consisting of O, N, Si and S, and wherein the nitrogen and sulfur atoms may optionally be oxidized and the nitrogen heteroatom may optionally be quaternized. The heteroatom(s) O, N and S and Si may be placed at any interior position of the heteroalkyl group or at the position at which the alkyl group is

attached to the remainder of the molecule. Examples include, but are not limited to, -CH_2-CH_2-O-CH_3, -CH_2-CH_2-NH-CH_3, -CH_2-CH_2-N(CH_3)-CH_3, -CH_2-S-CH_2-CH_3, -CH_2-CH_2,-S(O)-CH_3, -CH_2-CH_2-S(O)$_2$-CH_3, -CH=CH-O-CH_3, -Si(CH_3)$_3$, -CH_2-CH=N-OCH_3, and –CH=CH-N(CH_3)-CH_3. Up to two heteroatoms may be consecutive, such as, for example, -CH_2-NH-OCH_3 and –CH_2-O-Si(CH_3)$_3$. Similarly, the term "heteroalkylene" by itself or as part of another substituent means a divalent radical derived from heteroalkyl, as exemplified, but not limited by, -CH_2-CH_2-S-CH_2-CH_2- and –CH_2-S-CH_2-CH_2-NH-CH_2-. For heteroalkylene groups, heteroatoms can also occupy either or both of the chain termini (*e.g.*, alkyleneoxy, alkylenedioxy, alkyleneamino, alkylenediamino, and the like). Still further, for alkylene and heteroalkylene linking groups, no orientation of the linking group is implied by the direction in which the formula of the linking group is written. For example, the formula –C(O)$_2$R'- represents both –C(O)$_2$R'- and –R'C(O)$_2$-.

[0040] In general, an "acyl substituent" is also selected from the group set forth above. As used herein, the term "acyl substituent" refers to groups attached to, and fulfilling the valence of a carbonyl carbon that is either directly or indirectly attached to the polycyclic nucleus of the compounds of the present invention.

[0041] The terms "cycloalkyl" and "heterocycloalkyl", by themselves or in combination with other terms, represent, unless otherwise stated, cyclic versions of "alkyl" and "heteroalkyl", respectively. Additionally, for heterocycloalkyl, a heteroatom can occupy the position at which the heterocycle is attached to the remainder of the molecule. Examples of cycloalkyl include, but are not limited to, cyclopentyl, cyclohexyl, 1-cyclohexenyl, 3-cyclohexenyl, cycloheptyl, and the like. Examples of heterocycloalkyl include, but are not limited to, 1 –(1,2,5,6-tetrahydropyridyl), 1-piperidinyl, 2-piperidinyl, 3-piperidinyl, 4-morpholinyl, 3-morpholinyl, tetrahydrofuran-2-yl, tetrahydrofuran-3-yl, tetrahydrothien-2-yl, tetrahydrothien-3-yl, 1 –piperazinyl, 2-piperazinyl, and the like.

[0042] The terms "halo" or "halogen," by themselves or as part of another substituent, mean, unless otherwise stated, a fluorine, chlorine, bromine, or iodine atom. Additionally, terms such as "haloalkyl," are meant to include monohaloalkyl and polyhaloalkyl. For example, the term "halo(C_1-C_4)alkyl" is mean to include, but not be limited to, trifluoromethyl, 2,2,2-trifluoroethyl, 4-chlorobutyl, 3-bromopropyl, and the like.

[0043] As used herein, the term "haloalkyl" refers to an alkyl as defined herein, that is substituted by one or more halo groups as defined herein. Preferably the haloalkyl can be monohaloalkyl, dihaloalkyl or polyhaloalkyl including perhaloalkyl. A monohaloalkyl can have one iodo, bromo, chloro or fluoro within the alkyl group. Dihaloalkyl and polyhaloalkyl groups can have two or more of the same halo atoms or a combination of different halo groups within the alkyl. Preferably, the polyhaloalkyl contains up to 12, 10,

or 8, or 6, or 4, or 3, or 2 halo groups. Non-limiting examples of haloalkyl include fluoromethyl, difluoromethyl, trifluoromethyl, chloromethyl, dichioromethyl, trichioromethyl, pentafluoroethyl, heptafluoropropyl, difluorochioromethyl, dichiorofluoromethyl, difluoroethyl, difluoropropyl, dichloroethyl and dichioropropyl. A perhaloalkyl refers to an alkyl having all hydrogen atoms replaced with halo atoms.

[0044] As used herein, the term "heteroaryl" refers to a 5-14 membered monocyclic- or bicyclic- or fused polycyclic-ring system, having 1 to 8 heteroatoms selected from N, O, S or Se. Preferably, the heteroaryl is a 5-10 membered ring system. Typical heteroaryl groups include 2- or 3-thienyl, 2- or 3-furyl, 2- or 3-pyrrolyl, 2-, 4-, or 5-imidazolyl, 3-, 4-, or 5- pyrazolyl, 2-, 4-, or 5-thiazolyl, 3-, 4-, or 5-isothiazolyl, 2-, 4-, or 5-oxazolyl, 3-, 4-, or 5-isoxazolyl, 3- or 5-1,2,4-triazolyl, 4- or 5-1,2, 3-triazolyl, tetrazolyl, 2-, 3-, or 4-pyridyl, 3- or 4-pyridazinyl, 3-, 4-, or 5-pyrazinyl, 2-pyrazinyl, 2-, 4-, or 5-pyrimidinyl.

[0045] The term “heteroaryl” also refers to a group in which a heteroaromatic ring is fused to one or more aryl, cycloaliphatic, or heterocycloalkyl rings, where the radical or point of attachment is on the heteroaromatic ring. Nonlimiting examples include but are not limited to 1-, 2-, 3-, 5-, 6-, 7-, or 8-indolizinyl, 1-, 3-, 4-, 5-, 6-, or 7-isoindolyl, 2-, 3-, 4-, 5-, 6-, or 7-indolyl, 2-, 3-, 4-, 5-, 6-, or 7-indazolyl, 2-, 4-, 5-, 6-, 7-, or 8- purinyl, 1-, 2-, 3-, 4-, 6-, 7-, 8-, or 9-quinolizinyl, 2-, 3-, 4-, 5-, 6-, 7-, or 8-quinoliyl, 1-, 3-, 4-, 5-, 6-, 7-, or 8-isoquinoliyl, 1-, 4-, 5-, 6-, 7-, or 8-phthalazinyl, 2-, 3-, 4-, 5-, or 6-naphthyridinyl, 2-, 3-, 5-, 6-, 7-, or 8-quinazolinyl, 3-, 4-, 5-, 6-, 7-, or 8-cinnolinyl, 2-, 4-, 6-, or 7-pteridinyl, 1-, 2-, 3-, 4-, 5-, 6-, 7-, or 8-4aH carbazolyl, 1-, 2-, 3-, 4-, 5-, 6-, 7-, or 8-carbzaolyl, 1-, 3-, 4-, 5-, 6-, 7-, 8-, or 9-carbolinyl, 1-, 2-, 3-, 4-, 6-, 7-, 8-, 9-, or 10-phenanthridinyl, 1- , 2-, 3-, 4-, 5-, 6-, 7-, 8-, or 9-acridinyl, 1-, 2-, 4-, 5-, 6-, 7-, 8-, or 9-perimidinyl, 2-, 3-, 4-, 5-, 6-, 8-, 9-, or 10-phenathrolinyl, 1-, 2- , 3-, 4-, 6-, 7-, 8-, or 9-phenazinyl, 1-, 2-, 3-, 4-, 6-, 7-, 8-, 9-, or 10-phenothiazinyl, 1-, 2-, 3-, 4-, 6-, 7-, 8-, 9-, or 10-phenoxazinyl, or l-, 3-, 4-, 5-, 6-, 7-, 8-, 9-, or 10- benzisoqinolinyl, 2-, 3-, 4-, or 5-thieno[2,3-b]furanyl, 2-, 3-, 5-, 6-, 7-, 8-, 9-, 10 -, or 11-7H-pyrazino[2,3-c]carbazolyl,2-, 3-, 5-, 6-, or 7-2H- furo[3,2-b]-pyranyl, 2-, 3-, 4-, 5-, 7-, or 8-5H-pyrido[2,3-d]-o-oxazinyl, 1-, 3-, or 5-1H-pyrazolo[4,3-d]-oxazolyl, 2-, 4-, or 54H-imidazo[4,5-d] thiazolyl, 3-, 5-, or 8-pyrazino[2,3-d]pyridazinyl, 2-, 3-, 5-, or 6- imidazo[2,1-b] thiazolyl, 1-, 3-, 6-, 7-, 8-, or 9-furo[3,4-c]cinnolinyl, 1-, 2-, 3-, 4-, 5-, 6-, 8-, 9-, 10, or 11-4H-pyrido[2,3-c]carbazolyl, 2-, 3-, 6-, or 7-imidazo[1,2-b][1,2,4]triazinyl, 7-benzo[b]thienyl, 2-, 4-, 5- , 6-, or 7-benzoxazolyl, 2-, 4-, 5-, 6-, or 7-benzimidazolyl, 2-, 4-, 4-, 5-, 6-, or 7-benzothiazolyl, 1-, 2-, 4-, 5-, 6-, 7-, 8-, or 9- benzoxapinyl, 2-, 4-, 5-, 6-, 7-, or 8-benzoxazinyl, 1-, 2-, 3-, 5-, 6-, 7-, 8-, 9-, 10-, or 11-1H-pyrrolo[1,2-b][2]benzazapinyl. Typical fused heteroaryl groups include, but are not limited to 2-, 3-, 4-, 5-, 6-, 7-, or 8-quinolinyl, 1-, 3-, 4-, 5-, 6-, 7-, or 8-isoquinolinyl, 2-, 3-, 4-, 5-, 6-, or 7-indolyl, 2-, 3-, 4-, 5-,

6-, or 7-benzo[b]thienyl, 2-, 4-, 5- , 6-, or 7-benzoxazolyl, 2-, 4-, 5-, 6-, or 7-benzimidazolyl, 2-, 4-, 5-, 6-, or 7-benzothiazolyl.

[0046] As used herein, the term "heterocyclyl" or "heterocyclo" refers to an optionally substituted, fully saturated or unsaturated, aromatic or nonaromatic cyclic group, e.g., which is a 4- to 7-membered monocyclic, 7- to 12-membered bicyclic or 10- to 15-membered tricyclic ring system, which has at least one heteroatom in at least one carbon atom-containing ring. Each ring of the heterocyclic group containing a heteroatom may have 1, 2 or 3 heteroatoms selected from nitrogen atoms, oxygen atoms and sulfur atoms, where the nitrogen and sulfur heteroatoms may also optionally be oxidized. The heterocyclic group may be attached at a heteroatom or a carbon atom.

[0047] Exemplary monocyclic heterocyclic groups include pyrrolidinyl, pyrrolyl, pyrazolyl, oxetanyl, pyrazolinyl, imidazolyl, imidazolinyl, imidazolidinyl, triazolyl, oxazolyl, oxazolidinyl, isoxazolinyl, isoxazolyl, thiazolyl, thiadiazolyl, thiazolidinyl, isothiazolyl, isothiazolidinyl, furyl, tetrahydrofuryl, thienyl, oxadiazolyl, piperidinyl, piperazinyl, 2-oxopiperazinyl, 2-oxopiperidinyl, 2-oxopyrrolodinyl, 2-oxoazepinyl, azepinyl, 4-piperidonyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, tetrahydropyranyl, morpholinyl, thiamorpholinyl, thiamorpholinyl sulfoxide, thiamorpholinyl sulfone, 1,3-dioxolane and tetrahydro-1,1-dioxothienyl, 1,1,4-trioxo-1,2,5-thiadiazolidin-2-yl and the like.

[0048] Exemplary bicyclic heterocyclic groups include indolyl, dihydroidolyl, benzothiazolyl, benzoxazinyl, benzoxazolyl, benzothienyl, benzothiazinyl, quinuclidinyl, quinolinyl, tetrahydroquinolinyl, decahydroquinolinyl, isoquinolinyl, tetrahydroisoquinolinyl, decahydroisoquinolinyl, benzimidazolyl, benzopyranyl, indolizinyl, benzofuryl, chromonyl, coumarinyl, benzopyranyl, cinnolinyl, quinoxalinyl, indazolyl, pyrrolopyridyl, furopyridinyl (such as furo[2,3-c]pyridinyl, furo[3,2-b]-pyridinyl] or furo[2,3-b]pyridinyl), dihydroisoindolyl, 1,3-dioxo-1,3-dihydroisoindol-2-yl, dihydroquinazolinyl (such as 3,4-dihydro-4-oxo-quinazolinyl), phthalazinyl and the like.

[0049] Exemplary tricyclic heterocyclic groups include carbazolyl, dibenzoazepinyl, dithienoazepinyl, benzindolyl, phenanthrolinyl, acridinyl, phenanthridinyl, phenoxazinyl, phenothiazinyl, xanthenyl, carbolinyl and the like.

[0050] The term "heterocyclyl" further refers to heterocyclic groups as defined herein substituted with 1, 2 or 3 substituents selected from the groups consisting of the following:

(a) alkyl;

(b) hydroxy (or protected hydroxy);

(c) halo;

(d) oxo, i.e., =O;

(e) amino, alkylamino or dialkylamino;

(f) alkoxy;

(g) cycloalkyl;

(h) carboxy;

(i) heterocyclooxy, wherein heterocyclooxy denotes a heterocyclic group bonded through an oxygen bridge;

(j) alkyl-O-C(O)--;

(k) mercapto;

(l) nitro;

(m) cyano;

(n) sulfamoyl or sulfonamido;

(o) aryl;

(p) alkyl-C(O)-O--;

(q) aryl-C(O)-O--;

(r) aryl-S--;

(s) aryloxy;

(t) alkyl-S--;

(u) formyl, i.e., HC(O)--;

(v) carbamoyl;

(w) aryl-alkyl--; and

(x) aryl substituted with alkyl, cycloalkyl, alkoxy, hydroxy, amino, alkyl-C(O)-NH--, alkylamino, dialkylamino or halogen.

[0051] As used herein, the term "alkenyl" refers to a straight or branched hydrocarbon group having 2 to 20 carbon atoms and that contains at least one double bonds. The alkenyl groups preferably have about 2 to 8 carbon atoms.

[0052] The term "aryl" means, unless otherwise stated, a polyunsaturated, aromatic, hydrocarbon substituent, which can be a single ring or multiple rings (preferably from 1 to 3 rings), which are fused together or linked covalently. The term "heteroaryl" refers to aryl groups (or rings) that contain from one to four heteroatoms selected from N, O, and S, wherein the nitrogen and sulfur atoms are optionally oxidized, and the nitrogen atom(s) are optionally quaternized. A heteroaryl group can be attached to the remainder of the molecule through a heteroatom. Non-limiting examples of aryl and heteroaryl groups include phenyl, 1-naphthyl, 2-naphthyl, 4-biphenyl, 1-pyrrolyl, 2-pyrrolyl, 3-pyrrolyl, 3-pyrazolyl, 2-imidazolyl, 4-imidazolyl, pyrazinyl, 2-oxazolyl, 4-oxazolyl, 2-phenyl-4-oxazolyl, 5-oxazolyl, 3-isoxazolyl, 4-isoxazolyl, 5-isoxazolyl, 2-thiazolyl, 4-thiazolyl, 5-thiazolyl, 2-furyl, 3-furyl, 2-thienyl, 3-thienyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 2-pyrimidyl, 4-pyrimidyl, 5-benzothiazolyl, purinyl, 2-benzimidazolyl, 5-indolyl, 1-isoquinolyl, 5-isoquinolyl, 2-quinoxalinyl, 5-quinoxalinyl, 3-quinolyl, and 6-quinolyl. Substituents for each of the above noted aryl and heteroaryl ring systems are selected from the group of acceptable substituents described below.

[0053] For brevity, the term "aryl" when used in combination with other terms (*e.g.*, aryloxy, arylthioxy, arylalkyl) includes both aryl and heteroaryl rings as defined above. Thus, the term "arylalkyl" is meant to include those radicals in which an aryl group is attached to an alkyl group (*e.g.*, benzyl, phenethyl, pyridylmethyl and the like) including those alkyl groups in which a carbon atom (*e.g.*, a methylene group) has been replaced by, for example, an oxygen atom (*e.g.*, phenoxymethyl, 2-pyridyloxymethyl, 3-(1-naphthyloxy)propyl, and the like).

[0054] Each of the above terms (*e.g.*, "alkyl," "heteroalkyl," "aryl" and "heteroaryl") include both substituted and unsubstituted forms of the indicated radical. Preferred substituents for each type of radical are provided below.

[0055] Substituents for the alkyl, and heteroalkyl radicals (including those groups often referred to as alkylene, alkenyl, heteroalkylene, heteroalkenyl, alkynyl, cycloalkyl, heterocycloalkyl, cycloalkenyl, and heterocycloalkenyl) are generally referred to as "alkyl substituents" and "heteroakyl substituents," respectively, and they can be one or more of a variety of groups selected from, but not limited to: -OR', =O, =NR', =N-OR', -NR'R", -SR', -halogen, -SiR'R"R"', -OC(O)R', -C(O)R', -CO_2R', -CONR'R", -OC(O)NR'R", -NR"C(O)R', -NR'-C(O)NR"R"', -NR"$C(O)_2$R', -NR-C(NR'R"R"')=NR"", -NR-C(NR'R")=NR"', -S(O)R', -$S(O)_2$R', -$S(O)_2$NR'R", -$NRSO_2$R', -CN and -NO_2 in a number ranging from zero to (2m'+1), where m' is the total number of carbon atoms in such radical. R', R", R"' and R"" each preferably independently refer to hydrogen, substituted or unsubstituted heteroalkyl, substituted or unsubstituted aryl, e.g., aryl substituted with 1-3 halogens, substituted or unsubstituted alkyl, alkoxy or thioalkoxy groups, or arylalkyl groups. When a compound

of the invention includes more than one R group, for example, each of the R groups is independently selected as are each R', R", R"' and R"" groups when more than one of these groups is present. When R' and R" are attached to the same nitrogen atom, they can be combined with the nitrogen atom to form a 5-, 6-, or 7-membered ring. For example, -NR'R" is meant to include, but not be limited to, 1-pyrrolidinyl and 4-morpholinyl. From the above discussion of substituents, one of skill in the art will understand that the term "alkyl" is meant to include groups including carbon atoms bound to groups other than hydrogen groups, such as haloalkyl (*e.g.*, -CF_3 and –CH_2CF_3) and acyl (*e.g.*, -C(O)CH_3, -C(O)CF_3, -C(O)CH_2OCH_3, and the like).

[0056] Similar to the substituents described for the alkyl radical, the aryl substituents and heteroaryl substituents are generally referred to as "aryl substituents" and "heteroaryl substituents," respectively and are varied and selected from, for example: halogen, -OR', =O, =NR', =N-OR', -NR'R", -SR', -halogen, -SiR'R"R"', -OC(O)R', -C(O)R', -CO_2R', -CONR'R", -OC(O)NR'R", -NR"C(O)R', -NR'-C(O)NR"R"', -NR"C(O)$_2$R', -NR-C(NR'R")=NR"', -S(O)R', -S(O)$_2$R', -S(O)$_2$NR'R", -$NRSO_2$R', -CN and –NO_2, -R', -N_3, -$CH(Ph)_2$, fluoro(C_1-C_4)alkoxy, and fluoro(C_1-C_4)alkyl, in a number ranging from zero to the total number of open valences on the aromatic ring system; and where R', R", R"' and R"" are preferably independently selected from hydrogen, (C_1-C_8)alkyl and heteroalkyl, unsubstituted aryl and heteroaryl, (unsubstituted aryl)-(C_1-C_4)alkyl, and (unsubstituted aryl)oxy-(C_1-C_4)alkyl. When a compound of the invention includes more than one R group, for example, each of the R groups is independently selected as are each R', R", R"' and R"" groups when more than one of these groups is present.

[0057] Two of the aryl substituents on adjacent atoms of the aryl or heteroaryl ring may optionally be replaced with a substituent of the formula –T-C(O)-(CRR')$_q$-U-, wherein T and U are independently –NR-, -O-, -CRR'- or a single bond, and q is an integer of from 0 to 3. Alternatively, two of the substituents on adjacent atoms of the aryl or heteroaryl ring may optionally be replaced with a substituent of the formula –A-(CH_2)$_r$-B-, wherein A and B are independently –CRR'-, -O-, -NR-, -S-, -S(O)-, -S(O)$_2$-, -S(O)$_2$NR'- or a single bond, and r is an integer of from 1 to 4. One of the single bonds of the new ring so formed may optionally be replaced with a double bond. Alternatively, two of the substituents on adjacent atoms of the aryl or heteroaryl ring may optionally be replaced with a substituent of the formula –(CRR')$_s$-X-(CR"R"')$_d$-, where s and d are independently integers of from 0 to 3, and X is –O-, -NR'-, -S-, -S(O)-, -S(O)$_2$-, or –S(O)$_2$NR'-. The substituents R, R', R" and R"' are preferably independently selected from hydrogen or substituted or unsubstituted (C_1-C_6) akyl.

[0058] As used herein, the term "heteroatom" includes oxygen (O), nitrogen (N), sulfur (S), phosphorus (P) and silicon (Si).

[0059] As used herein, the term "aryloxy" refers to both an -O-aryl and an -O-heteroaryl group, wherein aryl and heteroaryl are defined herein.

[0060] As used herein, the term "pharmaceutically acceptable salts" refers to salts that retain the biological effectiveness and properties of the compounds of this invention and, which are not biologically or otherwise undesirable. In many cases, the compounds of the present invention are capable of forming acid and/or base salts by virtue of the presence of amino and/or carboxyl groups or groups similar thereto (e.g., phenol or hydroxyamic acid). Pharmaceutically acceptable acid addition salts can be formed with inorganic acids and organic acids. Inorganic acids from which salts can be derived include, for example, hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and the like. Organic acids from which salts can be derived include, for example, acetic acid, propionic acid, glycolic acid, pyruvic acid, oxalic acid, maleic acid, malonic acid, succinic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, *p*- toluenesulfonic acid, salicylic acid, and the like. Pharmaceutically acceptable base addition salts can be formed with inorganic and organic bases. Inorganic bases from which salts can be derived include, for example, sodium, potassium, lithium, ammonium, calcium, magnesium, iron, zinc, copper, manganese, aluminum, and the like; particularly preferred are the ammonium, potassium, sodium, calcium and magnesium salts. Organic bases from which salts can be derived include, for example, primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines, basic ion exchange resins, and the like, specifically such as isopropylamine, trimethylamine, diethylamine, triethylamine, tripropylamine, and ethanolamine. The pharmaceutically acceptable salts of the present invention can be synthesized from a parent compound, a basic or acidic moiety, by conventional chemical methods. Generally, such salts can be prepared by reacting free acid forms of these compounds with a stoichiometric amount of the appropriate base (such as Na, Ca, Mg, or K hydroxide, carbonate, bicarbonate, or the like), or by reacting free base forms of these compounds with a stoichiometric amount of the appropriate acid. Such reactions are typically carried out in water or in an organic solvent, or in a mixture of the two. Generally, non-aqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred, where practicable. Lists of additional suitable salts can be found, e.g., in *Remington's Pharmaceutical Sciences,* 20th ed., Mack Publishing Company, Easton, Pa., (1985), which is herein incorporated by reference.

[0061] As used herein, the term "pharmaceutically acceptable carrier/excipient" includes any and all solvents, dispersion media, coatings, surfactants, antioxidants, preservatives (e.g., antibacterial agents, antifungal agents), isotonic agents, absorption delaying agents, salts, drugs, drug stabilizers, binders, excipients, disintegration agents, lubricants, sweetening agents, flavoring agents, dyes, such like materials

and combinations thereof, as would be known to one of ordinary skill in the art (see, for example, Remington's Pharmaceutical Sciences, 18th Ed. Mack Printing Company, 1990, pp. 1289- 1329, incorporated herein by reference). Except in so far as any conventional carrier is incompatible with the active ingredient, its use in the therapeutic or pharmaceutical compositions is contemplated.

[0062] As used herein, the term "subject" refers to an animal. Preferably, the animal is a mammal. A subject also refers to for example, primates (e.g., humans), cows, sheep, goats, horses, dogs, cats, rabbits, rats, mice, fish, birds and the like. In a preferred embodiment, the subject is a human.

Compounds and Compositions

[0063] In one aspect, the present invention provides a compound having the structure of Formula (Ia):

TM-L-AM (Ia),

wherein TM is a tartgeting moiety, such as an antibody or a functional fragment thereof, that specifically binds to Her2/Neu (e.g., anti-Her2 antibody), AM is an activating moiety that activates a dendritic cell, natural killer cell, or a tumor cell, or a combination thereof, L is a linker.

[0064] By "activating moiety" herein is meant a molecule or agent that is capable of stimulating or enhancing the body's immune system or tumor cells. In general, the activating moiety acts, directly or indirectly, on toll like receptors, nucleotide-oligomerization domain-like receptors, RIG-I-Like receptors, c-type lectin receptors, or cytosolic DNA Sensors, or a combination thereof.

[0065] In some embodiments, the activating moiety activates human immune cells, including but not limited to dendritic cells, macrophages, monocytes, myeloid-derived suppressor cells, NK cells, B cells, T cells, or tumor cells, or a combination thereof.

[0066] Dendritic cells are the most powerful antigen-presenting cells. Dendritic cells play an essential role for the initiation of both innate and adaptive immune responses. Dendritic cells also play a key role in the induction and maintenance of immune tolerance.

[0067] By "dendritic cells" (DC) herein is meant a heterogeneous cell population including two main subtypes: namely, myeloid DC (mDC) and plasmacytoid DC (pDC) (Steinman et al., 1979, J. Exp. Med., 149, 1-16). These two blood DC subsets were originally differentiated by their expression of CD11c (integrin complement receptor) and CD123 (IL-3Rα). Each of the pDC and mDC populations constitutes between about 0.2 to about 0.6% of the PBMC population in humans.

[0068] By "pDC" herein is meant plasmacytoid dendritic cells and they represent a subtype of dendritic cells found in the blood and peripheral lymphoid organs. These cells express the surface markers CD123, BDCA-2(CD303) and BDCA-4(CD304) and HLA-DR, but do not express CD11c, CD14, CD3, CD20 or

CD56, which distinguishes them from conventional dendritic cells, monocytes, T-cells, B cells and NK cells. As components of the innate immune system, these cells express intracellular Toll-like receptors 7 and 9, which enable the detection of viral and bacterial nucleic acids, such as ssRNA or CpG DNA motifs. Upon stimulation and subsequent activation, these cells produce large amounts of Type I interferon (mainly IFN-α and IFN-β) and Type III interferon (e.g., IFN-λ), which are critical pleiotropic anti-viral compounds mediating a wide range of effects. By generating a large number of type I interferon, cytokines and chemokines, plasmacytoid dendritic cells are widely involved in the body's innate and adaptive immune responses. They can regulate NK cells, T cells, B cells and other cells involved in immune response intensity, duration, and response mode, thus play a very important function in tumor, infection and autoimmune disease. (Liu YJ. IPC: professional type 1 interferon-producing cells and plasmacytoid dendritic cell precursors. Annu Rev Immunol. 2005; 23:275-306. Gilliet M, Cao W, Liu YJ. Plasmacytoid dendritic cells: sensing nucleic acids in viral infection and autoimmune diseases. Nat Rev Immunol. 2008 Aug; 8 (8) :594-606).

[0069] By "mDC" herein is meant myeloid dendritic cells and they represent a subtype of circulating dendritic cellsfound in blood and peripheral lymphoid organs. These cells express the surface markers CD11c, CD1a, HLA-DR and either BDCA-1 (CD1c) or BDCA-3 (CD141). They do not express BDCA-2 or CD123, which distinguishes them from pDC. mDC also do not express CD3, CD20 or CD56. As components of the innate immune system, mDC express Toll-like receptors (TLR), including TLR2, 3, 4, 5, 6 and 8, which enable the detection of bacterial and viral components. Upon stimulation and subsequent activation, these cells are the most potent antigen presenting cells to activate antigen-specific CD4 as well as CD8 T cells. In addition, mDCs has the ability to produce large amounts of IL-12 and IL23, which is critical for the induction of Th1-mediated or Th17 cell-mediated immunity.

[0070] Study found that many solid tumors such as breast cancer and head and neck cancer, ovarian cancer has pDC's invasion (Treilleux I, Blay JY, Bendriss-Vermare N et al. Dendritic cell infiltration and prognosis of early stage breast cancer. Clin Cancer Res 2004; 10:7466-7474. Hartmann E, Wollenberg B, Rothenfusser S et al. Identification and functional analysis of tumor-infiltrating plasmacytoid dendritic cells in head and neck cancer. Cancer Res 2003; 63:6478-6487. Zou WP, Machelon V, Coulomb-L'Hermin A, et al. Stromal-derived factor-1 in human tumors recruits and alters the function of plasmacytoid precursor dendritic cells. Nat Med 2001; 7:1339-1346) and factors secreted by tumor cells inhibit DC maturation. (Gabrilovich DI, Corak J, Ciernik IF et al. Decreased antigen presentation by dendritic cells in patients with breast cancer. Clin Cancer Res 1997; 3:483-490. Bell D, Chomarat P, Broyles D et al. In breast carcinoma tissue, immature dendritic cells reside within the tumor, whereas mature dendritic cells are located in peritumoral areas. J Exp Med 1999; 190:1417-1425. Menetrier-Caux

C, Montmain G, Dieu MC et al. Inhibition of the differentiation of dendritic cells from CD34 (+) progenitors by tumor cells: role of interleukin-6 and macrophage colony-stimulating factor. Blood 1998; 92:4778-4791). These immature DC cells did not play a role in promoting anti-tumor immunity. By contrast, DCs within the tumor microenvironment promote tumor growth by inhibiting antitumor immunity and by promoting angiogenesis. There is evidence that Toll-like receptor 7 agonist Imiquimod, and Toll-like receptor 9 agonist CpG drugs can stimulate pDC within the tumor microenvironment to inhibit tumor development. (Dummer R, Urosevic M, Kempf W et al. Imiquimod in basal cell carcinoma: how does it work? Br J Dermatol 2003; 149:57-58. Miller RL, Gerster JF, Owens ML et al Imiquimod applied topically: a novel immune response modifier and new class of drug. Int J Immunopharmacol 1999; 21:1-14. Hofmann MA, Kors C, Audring H et al Phase 1 evaluation of intralesionally injected TLR9-agonist PF-3512676 in patients with basal cell carcinoma or metastatic melanoma. J Immunother 2008; 31:520-527).

[0071] Natural killer (NK) cells are a type of cytotoxic lymphocyte that constitutes a major component of the immune system. NK cells are a subset of peripheral blood lymphocytes defined by the expression of CD56 or CD 16 and the absence of the T cell receptor (CD3). They recognize and kill transformed cell lines without priming in an MHC-unrestricted fashion. NK cells play a major role in the rejection of tumors and cells infected by viruses. The process by which an NK cell recognizes a target cell and delivers a sufficient signal to trigger target lysis is determined by an array of inhibitory and activating receptors on the cell surface. NK discrimination of self from altered self involves inhibitory receptor recognition of MHC-I molecules and non-MHC ligands like CD48 and Clr-1b. NK recognition of infected or damaged cells (altered self) is coordinated through stress induced ligands (e.g., MICA, MICB, Rae1, H60, Mult1) or virally encoded ligands (e.g., m157, hemagluttinin) recognized by various activating receptors, including NKG2D, Ly49H and NKp46/Ncr1.

[0072] NK cells represent the predominant lymphoid cell in the peripheral blood for many months after allogeneic or autologous stem cell transplant and they have a primary role in immunity to pathogens during this period (Reittie et al (1989) Blood 73: 1351-1358; Lowdell et al (1998) Bone Marrow Transplant 21: 679-686). The role of NK cells in engraftment, graft- versus-host disease, anti-leukemia activity and post-transplant infection is reviewed in Lowdell (2003) Transfusion Medicine 13:399-404.

[0073] Human NK cells mediate the lysis of tumor cells and virus-infected cells via natural cytotoxicity and antibody-dependent cellular cytotoxicity (ADCC).

[0074] Human NK cells are controlled by positive and negative cytolytic signals. Negative (inhibitory) signals are transduced by C-lectin domain containing receptors CD94/NKG2A and by some Killer Immunoglobulin- like Receptors (KIRs). The regulation of NK lysis by inhibitory signals is known as

the "missing self" hypothesis in which specific HLA-class I alleles expressed on the target cell surface ligate inhibitory receptors on NK cells. The down- regulation of HLA molecules on tumor cells and some virally infected cells (e.g. CMV) lowers this inhibition below a target threshold and the target cells may become susceptible to NK cell- mediated lysis if the target cells also carry NK-priming and activating molecules. TLR7, TLR8 or TLR9 agonists can activate both mDC and pDCs to produce type I IFNs and express costimulatory molecules such as GITR-ligand, which subsequently activate NK cells to produce IFN-g and potently promote NK cell killing function.

[0075] Inhibitory receptors fall into two groups, those of the Ig-superfamily called Killer Immunoglobulin- like Receptors (KIRs) and those of the lectin family, the NKG2, which form dimers with CD94 at the cell surface. KIRs have a 2- or 3-domain extracellular structure and bind to HLA-A, -B or -C. The NKG2/CD94 complexes ligate HLA-E.

[0076] Inhibitory KIRs have up to 4 intracellular domains which contain ITIMs and the best characterized are KIR2DL1, KIR2DL2 and KIR2DL3 which are known to bind HLA-C molecules. KIR2DL2 and KIR2DL3 bind the group 1 HLA-C alleles while KIR2DL1 binds to group 2 alleles. Certain leukemia/lymphoma cells express both group 1 and 2 HLA-C alleles and are known to be resistant to NK-mediated cell lysis.

[0077] With regards to positive activating signals, ADCC is thought to be mediated via CD 16, and a number of triggering receptors responsible for natural cytotoxicity have been identified, including CD2, CD38, CD69, NKRP-I, CD40, B7-2, NK-TR, NKp46, NKp30 and NKp44. In addition, several KIR molecules with short intracytoplasmic tails are also stimulatory. These KIRs (KIR2DS1, KIR2DS2 and KIR2DS4) are known to bind to HLA-C; their extracellular domains being identical to their related inhibitory KIRs. The activatory KIRs lack the ITIMs and instead associate with DAP 12 leading to NK cell activation. The mechanism of control of expression of inhibitory versus activatory KIRs remains unknown.

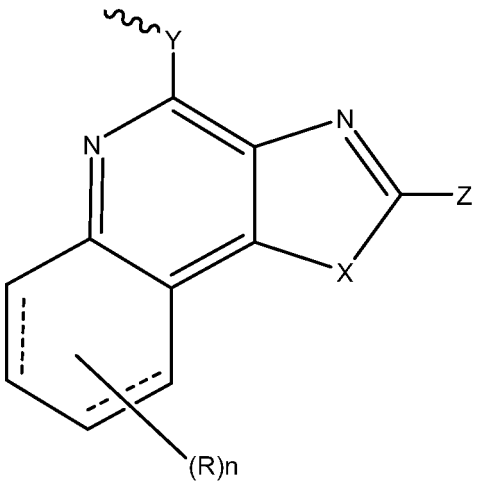
[0078] Several reports have described the expression of TLRs in mouse or human cancer or cancer cell lines. For example, TLR1 to TLR6 are expressed by colon, lung, prostate, and melanoma mouse tumor cell lines (Huang B, et al. Toll-like receptors on tumor cells facilitate evasion of immune surveillance. Cancer Res. 2005;65(12):5009–5014.), TLR3 is expressed in human breast cancer cells (Salaun B, Coste I, Rissoan MC, Lebecque SJ, Renno T. TLR3 can directly trigger apoptosis in human cancer cells. J Immunol. 2006;176(8):4894–4901.), hepatocarcinoma and gastric carcinoma cells express TLR2 and TLR4 (Huang B, et al. Listeria monocytogenes promotes tumor growth via tumor cell toll-like receptor 2 signaling. Cancer Res. 2007;67(9):4346–4352), and TLR9 (Droemann D, et al. Human lung cancer cells express functionally active Toll-like receptor 9. Respir Res. 2005;6:1.) and TLR4 (He W, Liu Q, Wang L,

Chen W, Li N, Cao X. TLR4 signaling promotes immune escape of human lung cancer cells by inducing immunosuppressive cytokines and apoptosis resistance. Mol Immunol. 2007;44(11):2850–2859.) are expressed by human lung cancer cells. TLR7 and TLR8 are found in tumor cells of human lung cancer (Cherfils-Vicini J, Platonova S, Gillard M, Laurans L, Validire P, Caliandro R, Magdeleinat P, Mami-Chouaib F, Dieu-Nosjean MC, Fridman WH, Damotte D, Sautès-Fridman C, Cremer I. J. Clin Invest. 2010;120(4):1285–1297).

[0079] TLR are a family of proteins that sense a microbial product and/or initiates an adaptive immune response. TLRs activate a dendritic cell (DC). TLRs are conserved membrane spanning molecules containing an ectodomain of leucine-rich repeats, a transmembrane domain and an intracellular TIR (Toll/interleukin receptor) domain. TLRs recognize distinct structures in microbes, often referred to as "PAMPs" (pathogen associated molecular patterns). Ligand binding to TLRs invokes a cascade of intra-cellular signaling pathways that induce the production of factors involved in inflammation and immunity.

[0080] In some embodiments, the activating moiety is a TLR7 and/or TLR8 agonist. TLR7 and TLR8 are phylogenetically and structurally related. TLR7 is selectively expressed by human pDCs and B cells. TLR8 is predominantly expressed mDCs, monocytes, macrophages and myeloid suppressor cells. TLR7-specific agonists activate plasmacytoid DCs (pDCs) to produce large amounts of type 1 IFNs and expressing high levels of costimulatory molecules that promote activation of T cells, NK cells, B cells and mDCs. TLR8-specific agonists activate myeloid DCs, monocytes, macrophages or myeloid-derived suppressor cells to produce large amounts of type 1 IFN, IL-12 and IL-23, and express high levels of MHC class I, MHC class II and costimulatory molecules that promote the activation of antigen specific CD4 and CD8+ T cells.

[0081] In some embodiments, the activating moiety is a TLR7 and/or TLR8 agonist that is represented by structure of formula (I):



(I)

wherein dashed line represents bond or absence of bond, ∿∿ is the point to be connected to the linker;

X is S or $-NR_1$, R_1 is $-W_0—W_1—W_2—W_3—W_4$,

W_0 is a bond, alkyl alkenyl, alkynyl, alkoxy, or –alkyl-S-alkyl--,

W_1 is a bond, --O--, or $-NR_2$--, wherein R_2 is hydrogen, alkyl or alkenyl,

W_2 is a bond, --O--, --C(O)--, --C(S)--, or $-S(O)_2—$,

W_3 is a bond, $--NR_3$--, wherein R_3 is hydrogen, alkyl or alkenyl,

W_4 is hydrogen, alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, aryl, aryloxy, heteroaryl, or heterocyclyl, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, $--NH_2$, nitro, --alkyl-hydroxyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, $--O-R_4$, --O-alkyl-R_4, --alkyl-O-R_4, --C(O)-R_4, --alkyl-C(O)-R_4, --alkyl-C(O)-O-R_4, --C(O)-O-R_4, --S-R_4, $--S(O)_2-R_4$, $--NH-S(O)_2-R_4$, --alkyl-S-R4, --alkyl-$S(O)_2-R_4$, $--NHR_4$, $--NR_4R_4$,--NH-alkyl-R_4, halogen, --CN, $--NO_2$, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

Z is hydrogen, alkyl, alkenyl, alkynyl, alkoxy, aryl, haloalkyl, heteroaryl, heterocyclyl, each of which can be optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, aryl, heteroaryl, heterocyclyl, halogen, cyano, nitro, $--N(R_5)_2$, --alkoxy-alkyl, --alkoxy-alkenyl, --C(O)-alkyl, --C(O)-O-alkyl, --O-C(O)-alkyl, $--C(O)-N(R_5)_2$, aryl, heteroaryl, --CO-aryl, and –CO-heteroaryl, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl;

R is hydrogen, alkyl, alkoxy, haloalkyl, halogen, aryl, heteroaryl, heterocyclyl, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, --NH_2, nitro, --alkyl-hydroxyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --alkyl-O-R_4, --C(O)-R_4, --C(O)-NH-R_4, --C(O)-NR_4R_4, --alkyl-C(O)-R_4, --alkyl-C(O)-O-R_4, --C(O)-O-R_4, --O-C(O)-R_4, --S-R_4, --C(O)-S-R_4, --S-C(O)-R_4, --$S(O)_2$-R_4, --NH-$S(O)_2$-R_4, --alkyl-S-R_4, --alkyl-$S(O)_2$-R_4, --NHR_4, --NR_4R_4,--NH-alkyl-R_4, halogen, --CN, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, alkoxy, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

n is 0, 1, 2, 3, or 4;

Y is –NR_6R_7, –$CR_6R_7R_8$, or –alkyl-NH_2, each of which can be optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, --NH_2, halogen, --$N(R_5)_2$, --alkoxy-alkyl, --alkoxy-alkenyl, --C(O)-alkyl, --C(O)-O-alkyl, --C(O)-$N(R_5)_2$, aryl, heteroaryl, --CO-aryl, and –CO-heteroaryl,

wherein R_6, R_7 and R_8 are independently hydrogen, alkyl, alkenyl, alkoxy, alkylamino, dialkylamino, alkylthio, arylthio, --alkyl-hydroxyl, --alkyl-C(O)-O-R_9, --alkyl-C(O)-R_9, or –alkyl-O-C(O)-R_9, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl, wherein R_9 is hydrogen, alkyl, alkenyl, halogen, or haloalkyl;

X and Z taken together may optionally form a (5-9)-membered ring;

or a pharmaceutically acceptable salt or solvate thereof.

[0082] In some embodiments, X of formula (I) is S.

[0083] In some embodiments, X of formula (I) is –NR_1, R_1 is alkyl, --alkyl-W_4, --alkyl-O-W_4, --alkyl-NH-C(O)-W_4, --alkoxy-NH-C(O)-W_4, --alkyl-NH-C(O)-NH-W_4, --alkoxy-NH-C(O)-NH-W_4, --alkyl-$S(O)_2$-W_4, or --alkyl-NH-C(S)-W_4, wherein W_4 is defined above.

[0084] In some embodiments, Z of formula (I) is hydrogen, alkyl, alkoxy, aryl, heteroaryl, haloalkyl, each of which is optionally substituted by one to three substituents selected from the group consisting of hydroxyl, alkyl, aryl, heteroaryl, heterocyclyl, cyano, --alkoxy-alkyl, nitro, and –$N(R_5)_2$, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl.

[0085] In some embodiments, Y of formula (I) is –NH_2, --alkyl-NH_2, each of which is optionally substituted by one to three substituents selected from the group consisting of alkyl, alkoxy, alkenyl, and alkynyl.

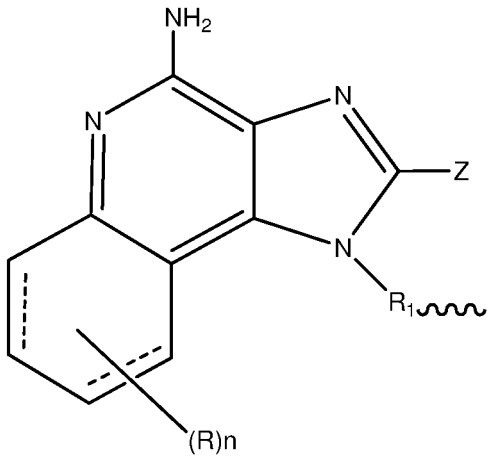
[0086] In some embodiments, n of formula (I) is 1 or 2.

[0087] In some embodiments, R of formula (I) is aryl or heteroaryl each of which is optionally substituted by one to three substituents selected from the group consisting of hydroxyl, alkoxy, --alkyl-hydroxyl, --O-R_4, --O-alkyl-R_4, --alkyl-O-R_4, --C(O)-R_4, --C(O)-NH-R_4, --C(O)-NR_4R_4, --alkyl-C(O)-R_4, --alkyl-C(O)-O-R_4, --C(O)-O-R_4, --O-C(O)-R_4, --S-R_4, --C(O)-S-R_4, --S-C(O)-R_4, --$S(O)_2$-R_4, --NH-$S(O)_2$-R_4, --alkyl-S-R_4, --alkyl-$S(O)_2$-R_4, --NHR_4, --NR_4R_4,--NH-alkyl-R_4, halogen, --CN, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, alkoxy, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl.

[0088] In another aspect, the present invention provides a compound having the structure of Formula (Ib):

TM-L-AM (Ib),

wherein TM is a targeting moiety, L is a linker, AM is an activating moiety that is represented by structure of formula (I):



(Ic)

wherein dashed line represents bond or absence of bond, ∿∿ is the point to be connected to the linker;

R_1 is $–W_0—W_1—W_2—W_3—W_4$,

W_0 is a bond, alkyl alkenyl, alkynyl, alkoxy, or –alkyl-S-alkyl--,

W_1 is a bond, --O--, or $–NR_2$--, wherein R_2 is hydrogen, alkyl or alkenyl,

W_2 is a bond, --O--, --C(O)--, --C(S)--, or $–S(O)_2—$,

W_3 is a bond, $--NR_3$--, wherein R_3 is hydrogen, alkyl or alkenyl,

W_4 is hydrogen, alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, aryl, aryloxy, heteroaryl, or heterocyclyl, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, $--NH_2$, nitro, --alkyl-hydroxyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --alkyl-O-R_4, --C(O)-

R_4, --alkyl-C(O)-R_4, --alkyl-C(O)-O-R_4, --C(O)-O-R_4, --S-R_4, --$S(O)_2$-R_4, --NH-$S(O)_2$-R_4, --alkyl-S-R4, --alkyl-$S(O)_2$-R_4, --NHR_4, --NR_4R_4,--NH-alkyl-R_4, halogen, --CN, --NO_2, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

Z is hydrogen, alkyl, alkenyl, alkynyl, alkoxy, aryl, haloalkyl, heteroaryl, heterocyclyl, each of which can be optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, aryl, heteroaryl, heterocyclyl, halogen, cyano, nitro, --$N(R_5)_2$, --alkoxy-alkyl, --alkoxy-alkenyl, --C(O)-alkyl, --C(O)-O-alkyl, --O-C(O)-alkyl, --C(O)-$N(R_5)_2$, aryl, heteroaryl, --CO-aryl, and –CO-heteroaryl, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl;

R is hydrogen, alkyl, alkoxy, haloalkyl, halogen, aryl, heteroaryl, heterocyclyl, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, --NH_2, nitro, --alkyl-hydroxyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --alkyl-O-R_4, --C(O)-R_4, --C(O)-NH-R_4, --C(O)-NR_4R_4, --alkyl-C(O)-R_4, --alkyl-C(O)-O-R_4, --C(O)-O-R_4, --O-C(O)-R_4, --S-R_4, --C(O)-S-R_4, --S-C(O)-R_4, --$S(O)_2$-R_4, --NH-$S(O)_2$-R_4, --alkyl-S-R_4, --alkyl-$S(O)_2$-R_4, --NHR_4, --NR_4R_4,--NH-alkyl-R_4, halogen, --CN, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, alkoxy, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

n is 0, 1, 2, 3, or 4;

Y is –NR_6R_7, –$CR_6R_7R_8$, or –alkyl-NH_2, each of which can be optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, --NH_2, halogen, --$N(R_5)_2$, --alkoxy-alkyl, --alkoxy-alkenyl, --C(O)-alkyl, --C(O)-O-alkyl, --C(O)-$N(R_5)_2$, aryl, heteroaryl, --CO-aryl, and –CO-heteroaryl,

wherein R_6, R_7 and R_8 are independently hydrogen, alkyl, alkenyl, alkoxy, alkylamino, dialkylamino, alkylthio, arylthio, --alkyl-hydroxyl, --alkyl-C(O)-O-R_9, --alkyl-C(O)-R_9, or –alkyl-O-C(O)-R_9, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl, wherein R_9 is hydrogen, alkyl, alkenyl, halogen, or haloalkyl;

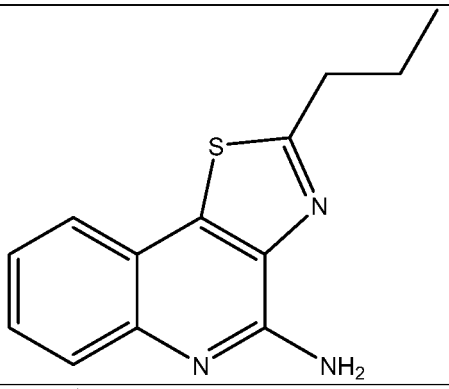
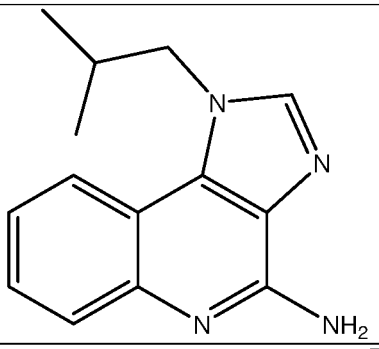
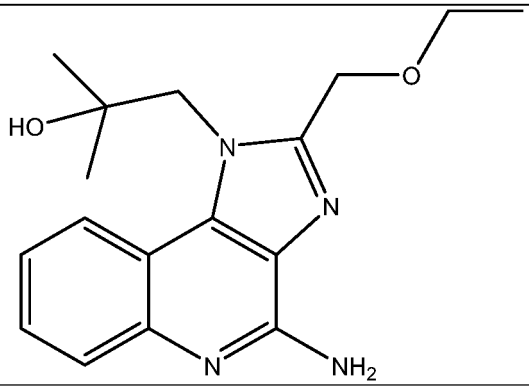
X and Z taken together may optionally form a (5-9)-membered ring;

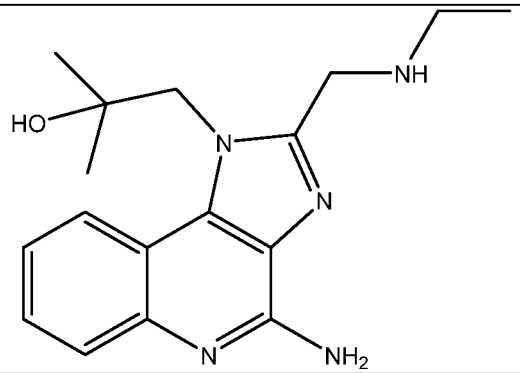
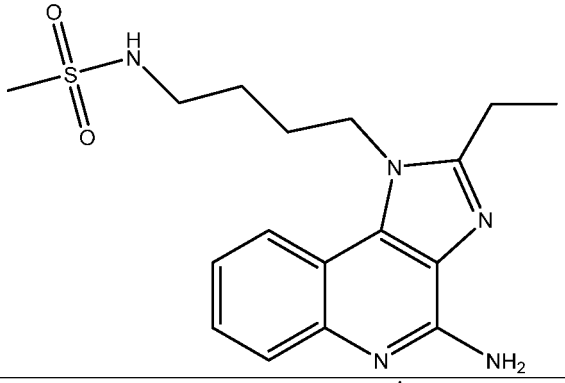
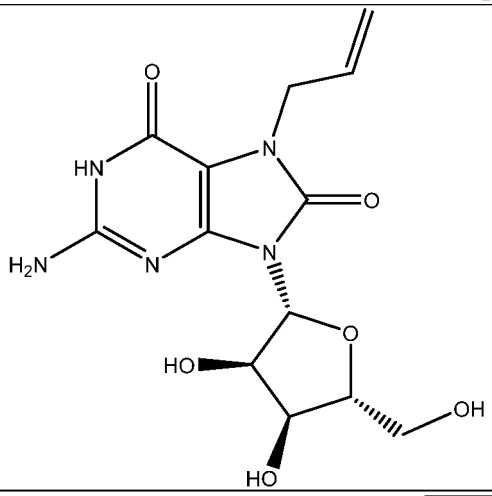
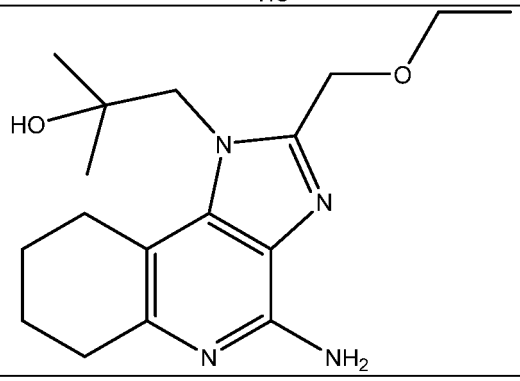
or a pharmaceutically acceptable salt or solvate thereof.

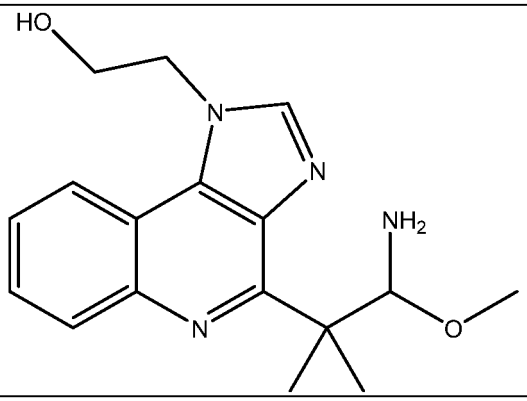
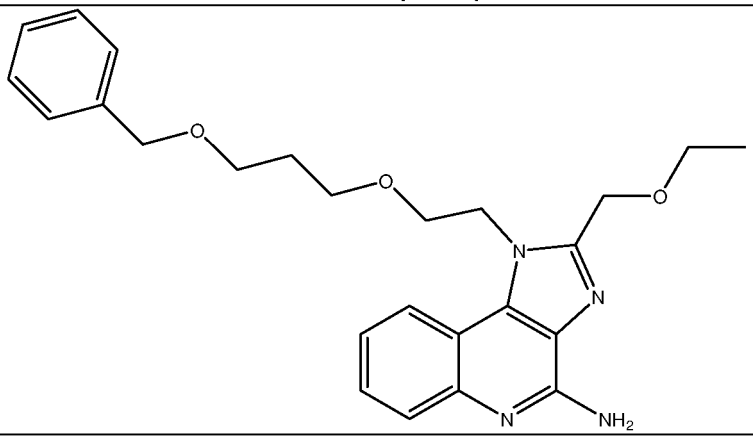
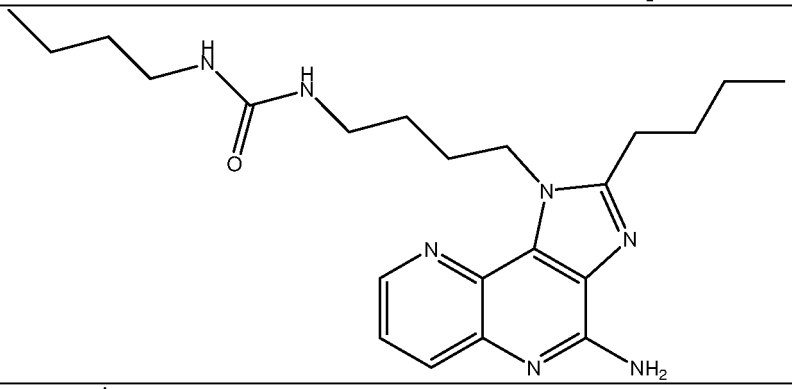
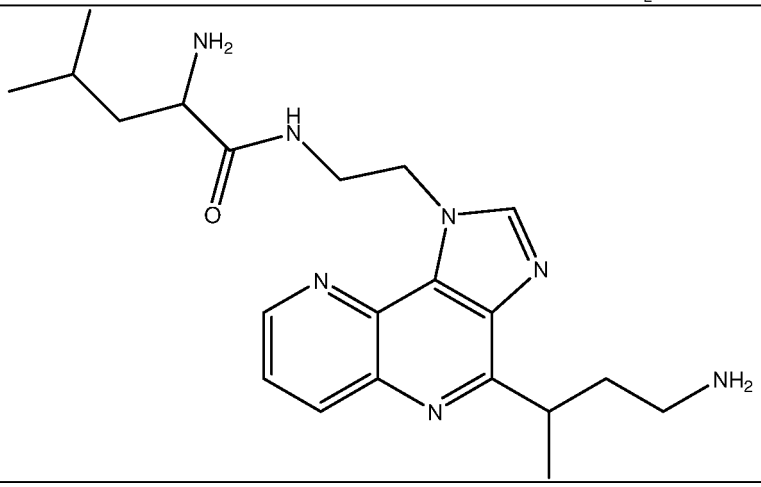
[0089] In some embodiments, the activating moiety is a TLR7 and/or TLR8 agonist that is selected from Table 1. The compounds in Table 1 are described and characterized in more details in US4,689,338,

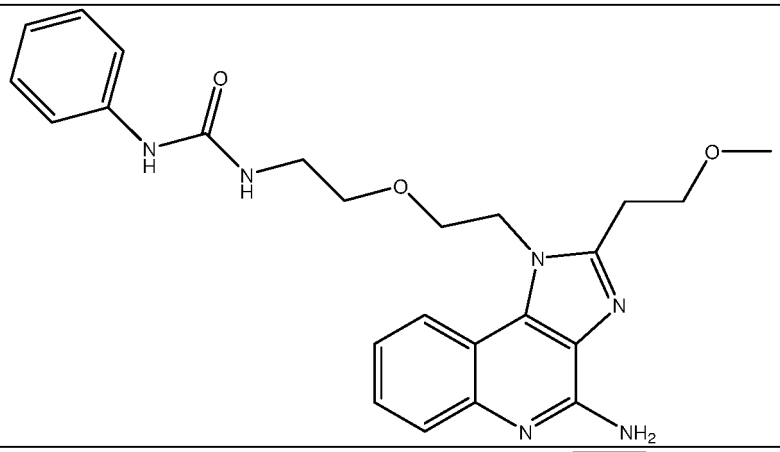
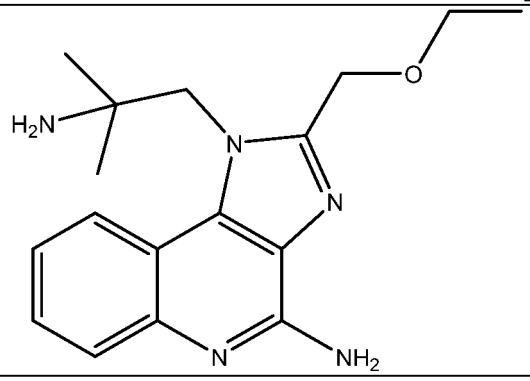
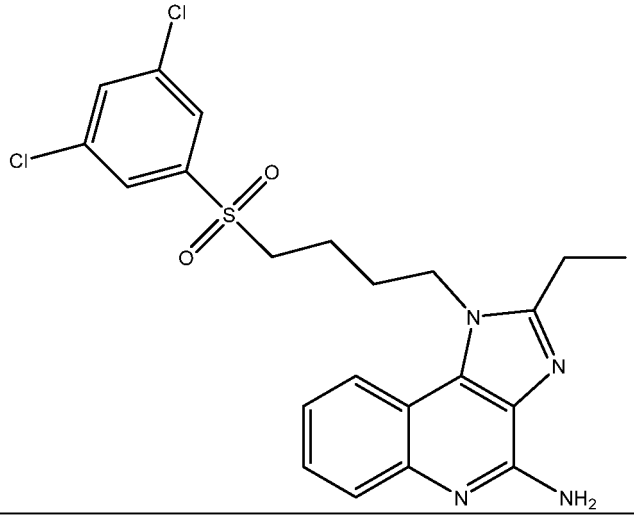
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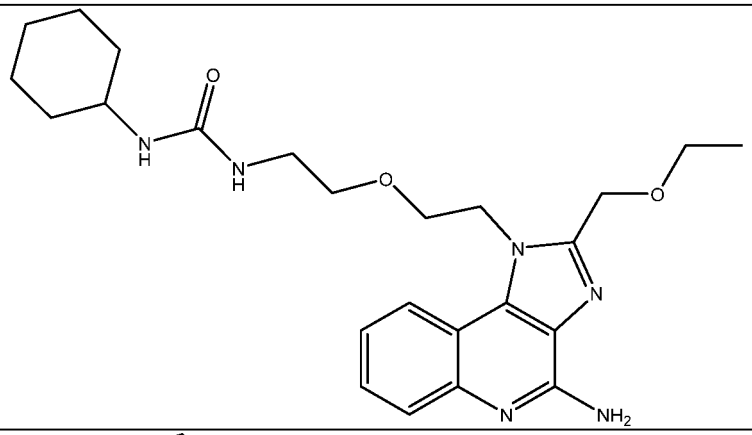
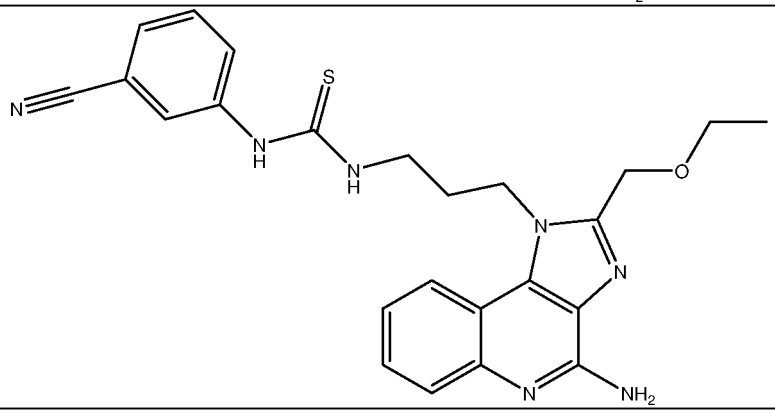
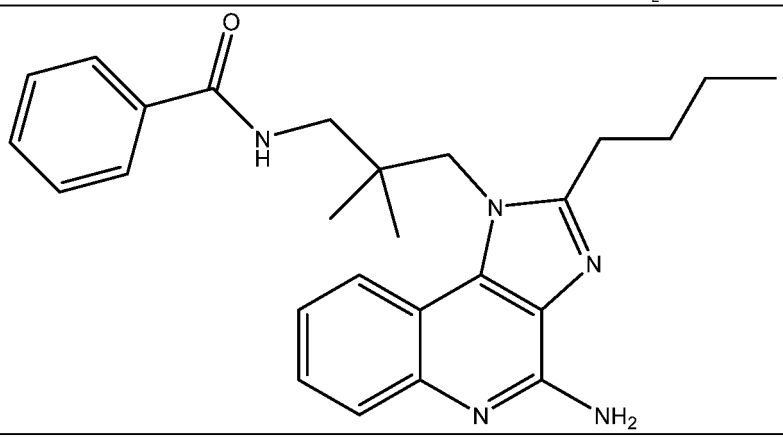
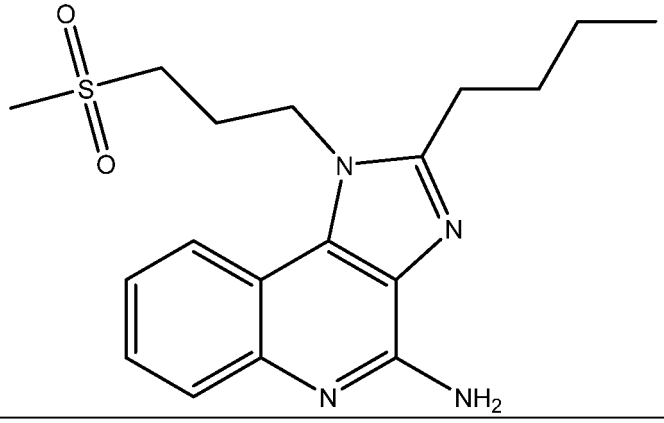
Table 1 Representative TLR7 and/or TLR8 Agonists

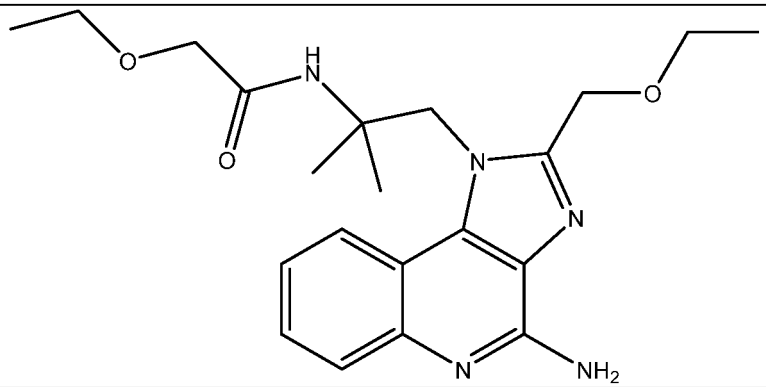
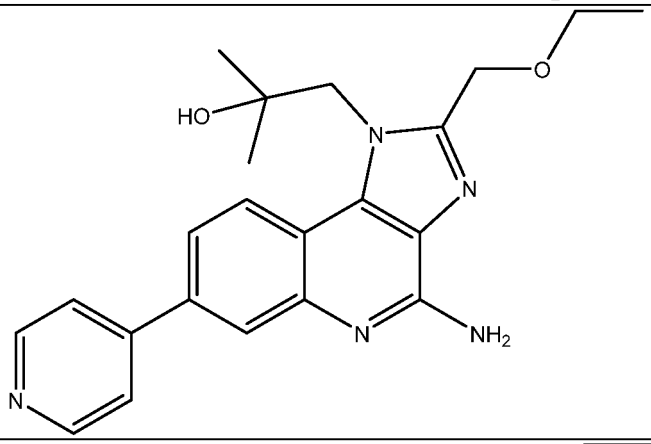
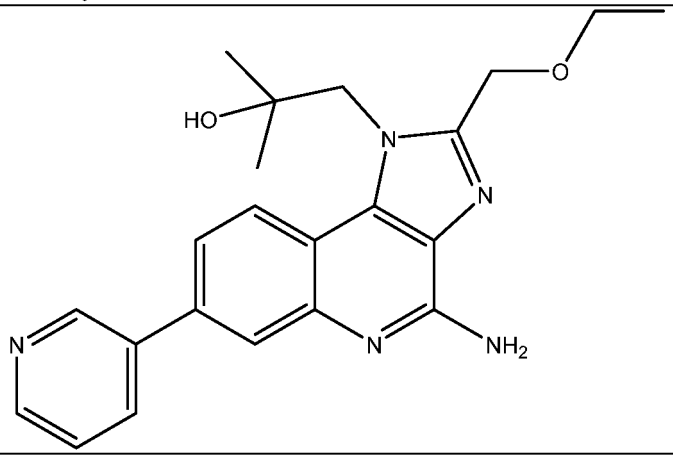
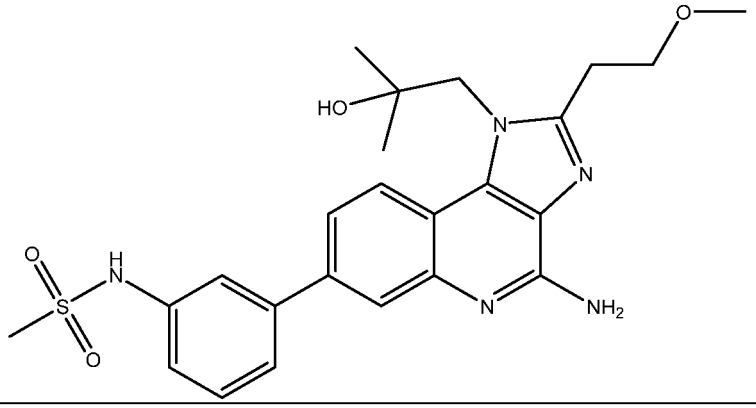
Name	Structure
2-propylthiazolo[4,5-c]quinolin-4-amine (CL075)	
1-(2-methylpropyl)-1H-imidazo[4,5-c]quinolin-4-amine (Imiquimod)	
4-amino-2-(ethoxymethyl)-a,a-dimethyl-1H-imidazo[4,5-c]quinoline-1-ethanol (Resiquimod)	

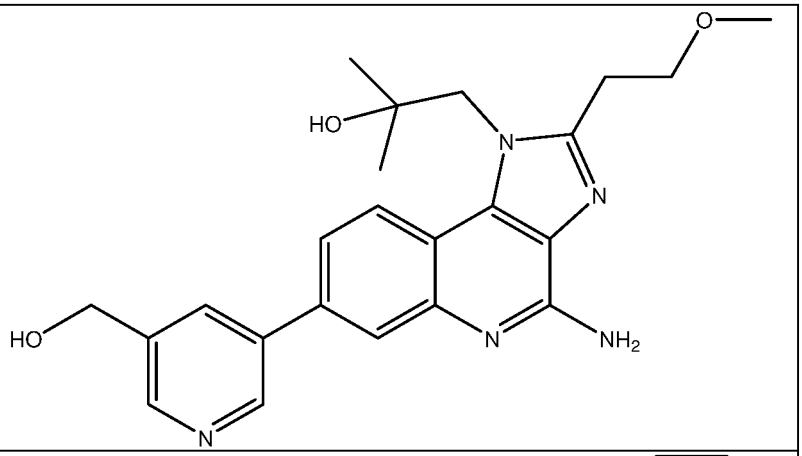
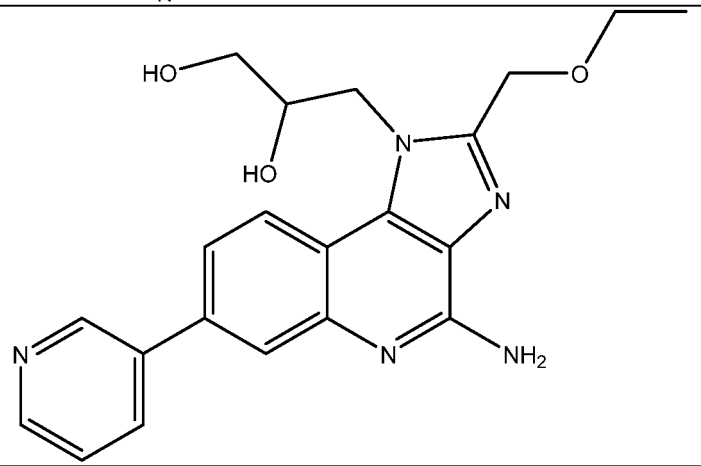
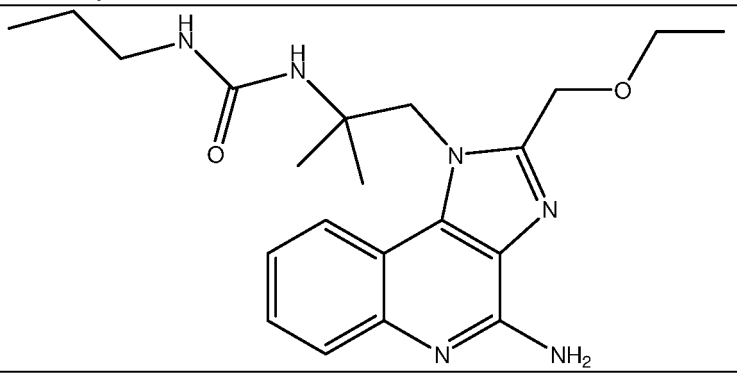
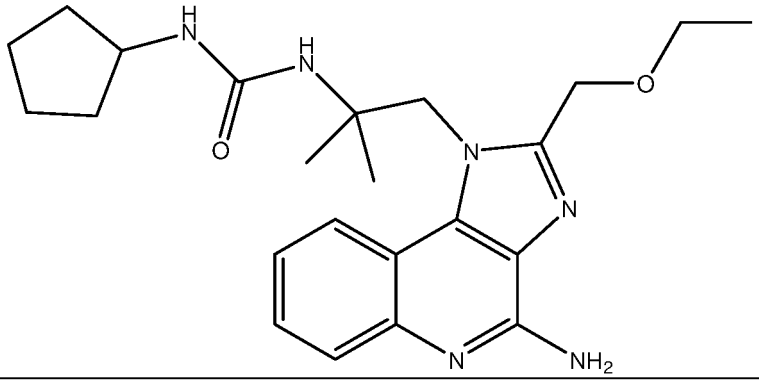
1-(4-amino-2-ethylaminomethylimidazo-[4,5-c]quinolin-1-yl)-2-methylpropan-2-ol (Gardiquimod)	
N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl-]methanesulfonamide (CM001)	
7-allyl-7,8-dihydro-8-oxo-guanosine (Loxoribine)	
4-amino-2-ethoxymethyl-aa-dimethyl-6,7,8,9-tetrahydro-1h-imidazo[4,5-c]quinoline-1-ethanol ol	

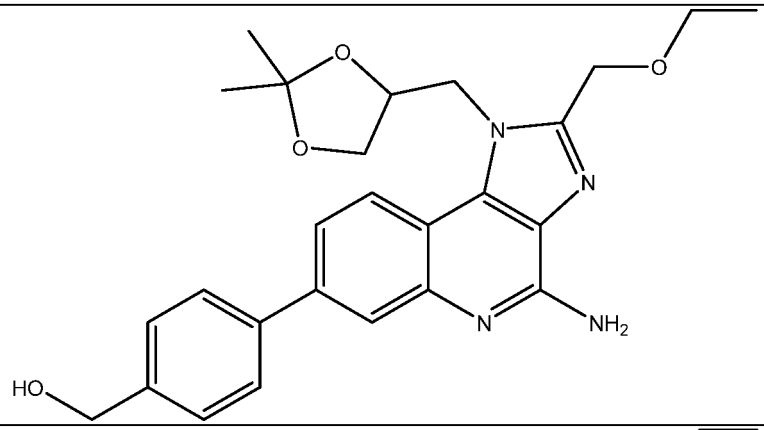
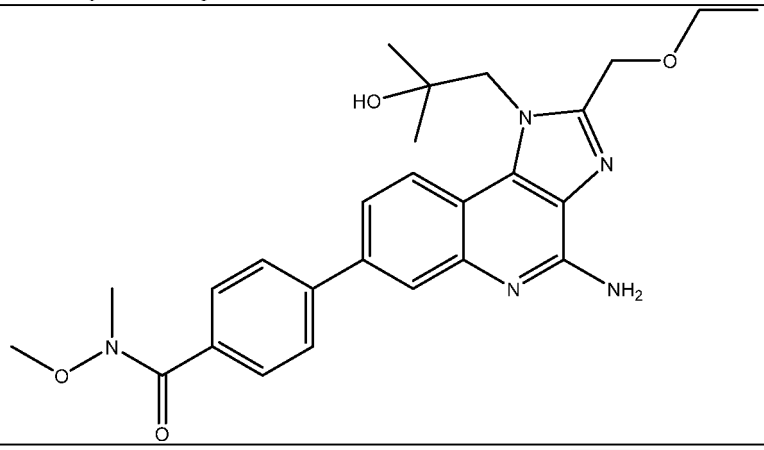
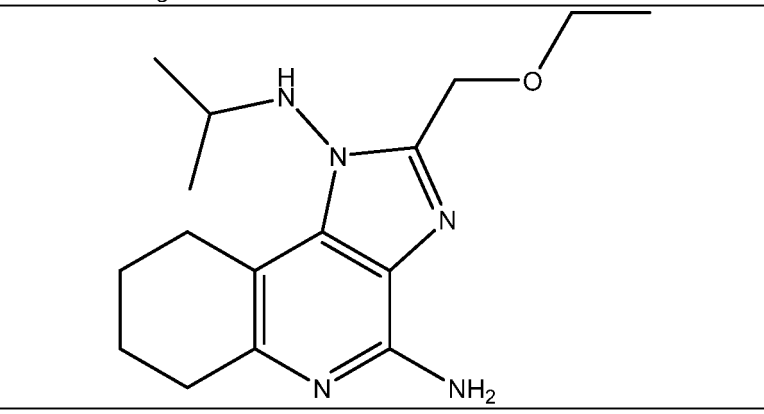
4-amino-aa-dimethyl-2-methoxyethyl-1h-imidazo[4,5-c]quinoline-1-ethanol	
1-(2-(3-(benzyloxy)propoxy)ethyl)-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine	
N-[4-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)butyl]-n'-butylurea	
N1-[2-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)ethyl]-2-amino-4-methylpentanamide	

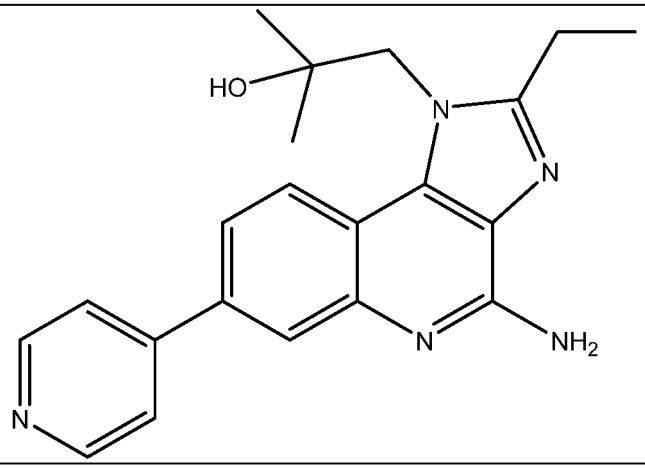
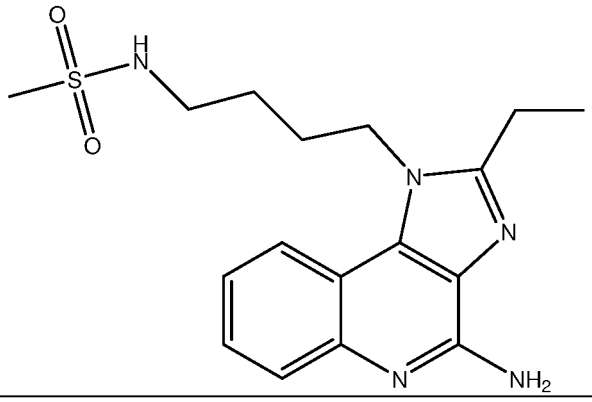
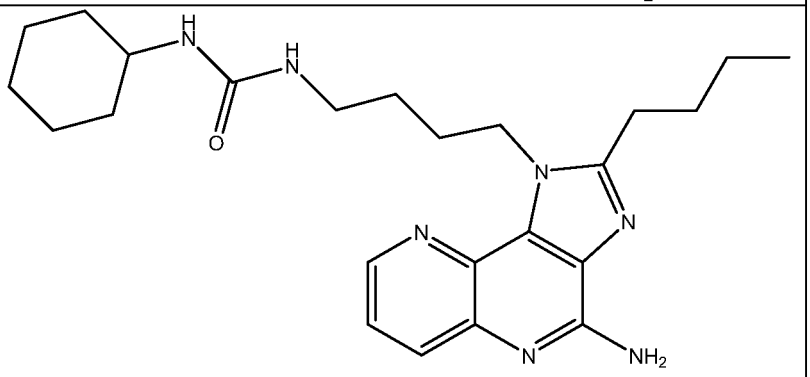
N-(2-{2-[4-amino-2-(2-methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]ethoxy}ethyl)-n'-phenylurea	
1-(2-amino-2-methylpropyl)-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine	
1-{4-[(3,5-dichlorophenyl)sulfonyl]butyl}-2-ethyl-1H-imidazo[4,5-c]quinolin-4-amine	

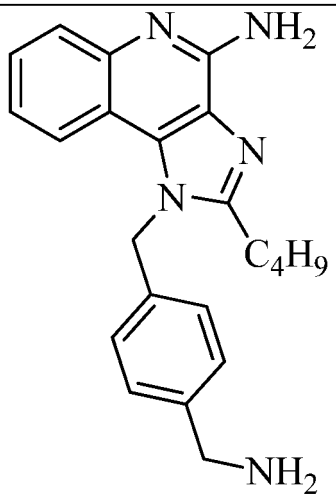
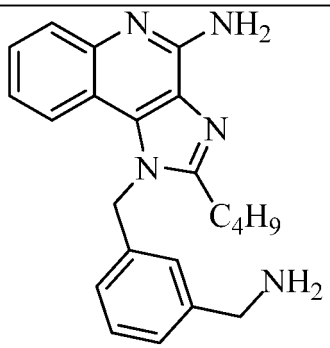
N-(2-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-1-yl]ethoxy}ethyl)-n'-cyclohexylurea	
N-{3-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-1-yl]propyl}-n'-(3-cyanophenyl)thiourea	
N-[3-(4-amino-2-butyl-1H-imidazo[4,5-c]quinolin-1-yl)-2,2-dimethylpropyl]benzamide	
2-butyl-1-[3-(methylsulfonyl)propyl]-1H-imidazo[4,5-c]quinolin-4-amine	

N-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-1-yl]-1,1-dimethylethyl}-2-ethoxyacetamide	
1-[4-amino-2-ethoxymethyl-7-(pyridin-4-yl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol	
1-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol	
N-{3-[4-amino-1-(2-hydroxy-2-methylpropyl)-2-(methoxyethyl)-1H-imidazo[4,5-c]quinolin-7-yl]phenyl}methanesulfonamide	

1-[4-amino-7-(5-hydroxymethylpyridin-3-yl)-2-(2-methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol	
3-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H-imidazo[4,5-c]quinolin-1-yl]propane-1,2-diol	
1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5-c]quinolin-1-yl)-1,1-dimethylethyl]-3-propylurea	
1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5-c]quinolin-1-yl)-1,1-dimethylethyl]-3-cyclopentylurea	

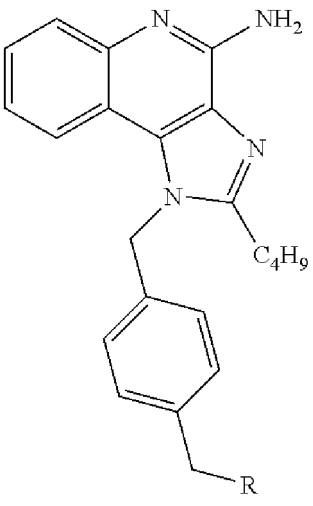
1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methyl]-2-(ethoxymethyl)-7-(4-hydroxymethylphenyl)-1H-imidazo[4,5-c]quinolin-4-amine	
4-[4-amino-2-ethoxymethyl-1-(2-hydroxy-2-methylpropyl)-1H-imidazo[4,5-c]quinolin-7-yl]-N-methoxy-N-methylbenzamide	
2-ethoxymethyl-N1-isopropyl-6,7,8,9-tetrahydro-1H-imidazo[4,5-c]quinoline-1,4-diamine	

1-[4-amino-2-ethyl-7-(pyridin-4-yl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol	
N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl]methanesulfonamide	
N-[4-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)butyl]-n'-cyclohexylurea	

	 **p-IMDQ**
	 **m-IMDQ**

[0090] Preferably, AM is is Resiquimod or Imiquimod.

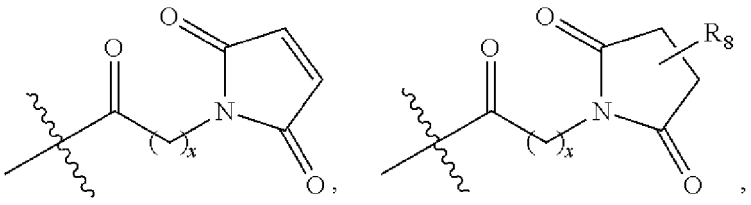
[0091] In some embodiments, the AM comprises an imidazoquinoline derivatives having the structure of fomular (Id):



(Id)

wherein R is selected from the group consisting of: —NH(R_5) and isothiocyanate (—NCS);

R_5 is selected from the group consisting of hydrogen (—H), acetyl , —CO-tert-Bu (-Boc), —CO—$(CH_2)_x$—R_6, C_1-C_{16} alkyl, —CO-4-(phenylboronic acid), —C(S)—NH—$(CH_2)_x$—NH—$(CH_2)_x$—NH—$(CH_2)_x$—NH_2,



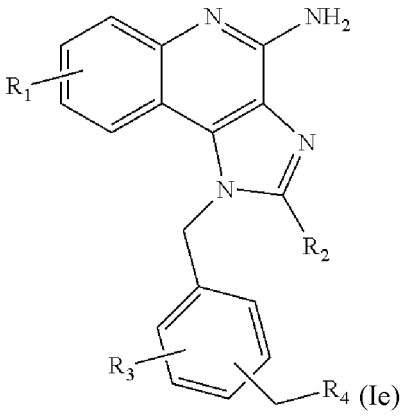
R_6 is selected from the group consisting of hydrogen, alkyne, azido, carboxylic acid, and —CONH—$(CH_2)_x$—O—$(CH_2)_x$—O—$(CH_2)_x$—O—$(CH_2)_x$—R_7;

R_7 is selected from the group consisting of amino, isothiocyanate, and —NH—CO—$(CH_2)_x$—CO_2H;

R_8 is selected from a peptide antigen moiety or a protein antigen moiety; and

x is any integer from 1 to 10.

[0092] In some embodiments, the AM comprises an imidazoquinoline derivatives having the structure of fomular (Ie):

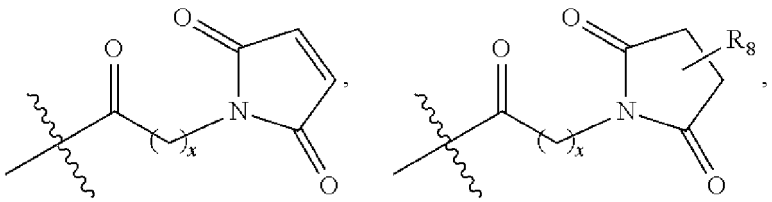


wherein, R_1 and R_3 are each independently selected from the group consisting of hydrogen, halogen, nitro, —NH_2, azido, hydroxyl, —CF_3, carboylic acid, and —CO_2R_2;

R_2 is a C_2-C_5 alkyl, and

R_4 selected from the group consisting of: —NH(R_5) and isothiocyanate;

R_5 is selected from the group consisting of hydrogen, acetyl, —CO-tert-Bu (-Boc), —CO—$(CH_2)_x$—R_6, C_1-C_{16} alkyl, —CO-4-(phenylboronic acid),, —C(S)—NH—$(CH_2)_x$—NH—$(CH_2)_x$—NH—$(CH_2)_x$—NH_2,



R_6 is selected from the group consisting of hydrogen, alkyne, azido, carboxylic acid, and —CONH—$(CH_2)_x$—O—$(CH_2)_x$—O—$(CH_2)_x$—O—$(CH_2)_x$—R_7;

R_7 is selected from the group consisting of amino, isothiocyanate, and —NH—CO—$(CH_2)_x$—CO_2H;

R_8 is selected from a peptide antigen moiety or a protein antigen moiety; and

x is any integer from 1 to 10. US Pat. Appl. Pub. No. 20140256922 A1, the discloodure of which is incorporated by reference in its entirety.

[0093] In general, when the AM comprises an imidazoquinoline derivatives having the structure of fomular (Id) or formular (Ie), the AM is attached the the linker at positions, such as at the NH_2 or R of formular (Id), or the NH_2 or R4 of formular (Ie).

Targeting Moiety

[0094] In general, the compounds of the present invention comprise a target moiety.

[0095] By "targeting moiety (TM)" or "targeting agent" here in is meant a molecule, complex, or aggregate, that binds specifically or selectively to a target molecule, cell, particle, tissue or aggregate, which generally is referred to as a "target" or a "marker," and these are discussed in further detail herein.

[0096] In some embodiments, the targeting moiety comprises an immunoglobulin, a protein, a peptide, a small molecule, a nanoparticle, or a nucleic acid.

[0097] Exemplary targeting agents such as antibodies (e.g., chimeric, humanized and human), ligands for receptors, lecitins, and saccharides, and substrate for certain enzymes are recognized in the art and are useful without limitation in practicing the present invention. Other targeting agents include a class of compounds that do not include specific molecular recognition motifs include nanoparticles, macromolecules such as poly(ethylene glycol), polysaccharide, and polyamino acids which add molecular mass to the activating moiety. The additional molecular mass affects the pharmacokinetics of the activating moiety, e.g., serum half-life.

[0098] In some embodiments, a targeting moiety is an antibody, antibody fragment, bispecific antibody or other antibody-based molecule or compound. However, other examples of targeting moieties are known in the art and may be used, such as aptamers, avimers, receptor-binding ligands, nucleic acids,

biotin-avidin binding pairs, binding peptides or proteins, etc. The terms "targeting moiety" and "binding moiety" are used synonymously herein.

[0099] By "target" or "marker" herein is meant any entity that is capable of specifically binding to a particular targeting moiety, such as Her2/Neu.

[0100] In some embodiments, the targeting moiety is capable of binding to Her2/Neu specifically or preferably in comparison to a non-target.

[0101] By "specifically binds" or "preferably binds" herein is meant that the binding between two binding partners (e.g., between a targeting moiety and its binding partner) is selective for the two binding partners and can be discriminated from unwanted or non-specific interactions. For example, the ability of an antigen-binding moiety to bind to a specific antigenic determinant can be measured either through an enzyme- linked immunosorbent assay (ELISA) or other techniques familiar to one of skill in the art, e.g. surface plasmon resonance technique (analyzed on a BIAcore instrument) (Liljeblad et al., Glyco J 17, 323-329 (2000)), and traditional binding assays (Heeley, Endocr Res 28, 217-229 (2002)). The terms "anti- [antigen] antibody" and "an antibody that binds to [antigen]" refer to an antibody that is capable of binding the respective antigen with sufficient affinity such that the antibody is useful as a diagnostic and/or therapeutic agent in targeting the antigen. In some embodiments, the extent of binding of an anti-[antigen] antibody to an unrelated protein is less than about 10% of the binding of the antibody to the antigen as measured, e.g., by a radioimmunoassay (RIA). In some embodiments, an antibody that binds to [antigen] has a dissociation constant (KD) of $< I\mu M$, < 100 nM, < 10 nM, < 1 nM, < 0.1 nM, < 0.01 nM, or < 0.001 nM (e.g. 10^{-8} M or less, e.g. from 10^{-8} M to 10^{-13} M, e.g., from 10^{-9} M to 10^{-13} M). It is understood that the above definition is also applicable to antigen-binding moieties that bind to an antigen.

[0102] In some embodiments, the targeting moiety comprises an antibody, or a functional fragment thereof.

[0103] By immunoglobulin" or "antibody" herein is meant a full-length (i.e., naturally occurring or formed by normal immunoglobulin gene fragment recombinatorial processes) immunoglobulin molecule (e.g., an IgG antibody) or an immunologically active (i.e., specifically binding) portion of an immunoglobulin molecule, like an antibody fragment. An antibody or antibody fragment may be conjugated or otherwise derivatized within the scope of the claimed subject matter. Such antibodies include IgGl , lgG2a, IgG3, IgG4 (and IgG4 subforms), as well as IgA isotypes.

[0104] The term "antibody" herein is used in the broadest sense and encompasses various antibody structures, including but not limited to monoclonal antibodies, polyclonal antibodies, multispecific antibodies (e.g. bispecific antibodies), and antibody fragments so long as they exhibit the desired antigen-

binding activity and comprise an Fc region or a region equivalent to the Fc region of an immunoglobulin The terms "full-length antibody", "intact antibody", "and "whole antibody" are used herein interchangeably to refer to an antibody having a structure substantially similar to a native antibody structure or having heavy chains that contain an Fc region as defined herein.

[0105] By "native antibodies" herein is meant naturally occurring immunoglobulin molecules with varying structures. For example, native IgG antibodies are heterotetrameric glycoproteins of about 150,000 daltons, composed of two identical light chains and two identical heavy chains that are disulfide-bonded. From N- to C-terminus, each heavy chain has a variable region (VH), also called a variable heavy domain or a heavy chain variable domain, followed by three constant domains (CHI, CH2, and CH3), also called a heavy chain constant region. Similarly, from N- to C-terminus, each light chain has a variable region (VL), also called a variable light domain or a light chain variable domain, followed by a constant light (CL) domain, also called a light chain constant region. The light chain of an antibody may be assigned to one of two types, called kappa (κ) and lambda (λ), based on the amino acid sequence of its constant domain.

[0106] By "antibody fragment" herein is meant a molecule other than an intact antibody that comprises a portion of an intact antibody that binds the antigen to which the intact antibody binds. Examples of antibody fragments include but are not limited to Fv, Fab, Fab', Fab'-SH, F(ab')2, diabodies, linear antibodies, single-chain antibody molecules (e.g. scFv), single-domain antibodies, and multispecific antibodies formed from antibody fragments. For a review of certain antibody fragments, see Hudson et al., Nat Med 9, 129-134 (2003). For a review of scFv fragments, see e.g. Pliickthun, in The Pharmacology of Monoclonal Antibodies, vol. 113, Rosenburg and Moore eds., Springer- Verlag, New York, pp. 269-315 (1994); see also WO 93/16185; and U.S. Patent Nos. 5,571,894 and 5,587,458. For discussion of Fab and F(ab')2 fragments comprising salvage receptor binding epitope residues and having increased in vivo half-life, see U.S. Patent No. 5,869,046. Diabodies are antibody fragments with two antigen-binding sites that may be bivalent or bispecific. See, for example, EP 404,097; WO 1993/01161; Hudson et al., Nat Med 9, 129- 134 (2003); and Hollinger et al., Proc Natl Acad Sci USA 90, 6444-6448 (1993). Triabodies and tetrabodies are also described in Hudson et al., Nat Med 9, 129-134 (2003). Single-domain antibodies are antibody fragments comprising all or a portion of the heavy chain variable domain or all or a portion of the light chain variable domain of an antibody. In certain embodiments, a single-domain antibody is a human single-domain antibody (Domantis, Inc., Waltham, MA; see e.g. U.S. Patent No. 6,248,516 B l). Antibody fragments can be made by various techniques, including but not limited to proteolytic digestion of an intact antibody as well as production by recombinant host cells (e.g. E. coli or phage), as described herein.

[0107] By "antigen binding domain" herein is meant the part of an antibody that comprises the area which specifically binds to and is complementary to part or all of an antigen. An antigen binding domain may be provided by, for example, one or more antibody variable domains (also called antibody variable regions). Particularly, an antigen binding domain comprises an antibody light chain variable region (VL) and an antibody heavy chain variable region (VH).

[0108] By "variable region" or "variable domain" herein is meant the domain of an antibody heavy or light chain that is involved in binding the antibody to antigen. The variable domains of the heavy chain and light chain (VH and VL, respectively) of a native antibody generally have similar structures, with each domain comprising four conserved framework regions (FRs) and three hypervariable regions (HVRs). See, e.g., Kindt et al., Kuby Immunology, 6th ed., W.H. Freeman and Co., page 91 (2007). A single VH or VL domain may be sufficient to confer antigen-binding specificity.

[0109] By "hypervariable region" or "HVR" herein is meant each of the regions of an antibody variable domain which are hypervariable in sequence and/or form structurally defined loops ""hypervariable loops"). Generally, native four-chain antibodies comprise six HVRs; three in the VH (HI, H2, H3), and three in the VL (LI, L2, L3). HVRs generally comprise amino acid residues from the hypervariable loops and/or from the complementarity determining regions (CDRs), the latter being of highest sequence variability and/or involved in antigen recognition. With the exception of CDRl in VH, CDRs generally comprise the amino acid residues that form the hypervariable loops. Hypervariable regions (HVRs) are also referred to as "complementarity determining regions" (CDRs), and these terms are used herein interchangeably in reference to portions of the variable region that form the antigen binding regions. This particular region has been described by Kabat et al., U.S. Dept. of Health and Human Services, Sequences of Proteins of Immunological Interest (1983) and by Chothia et al., J Mol Biol 196:901-917 (1987), where the definitions include overlapping or subsets of amino acid residues when compared against each other. Nevertheless, application of either definition to refer to a CDR of an antibody or variants thereof is intended to be within the scope of the term as defined and used herein. The exact residue numbers which encompass a particular CDR will vary depending on the sequence and size of the CDR. Those skilled in the art can routinely determine which residues comprise a particular CDR given the variable region amino acid sequence of the antibody.

[0110] The antibody of the present invention can be chimeric antibodies, humanized antibodies, human antibodies, or antibody fusion proteins.

[0111] By "chimeric antibody" herein is meant a recombinant protein that contains the variable domains of both the heavy and light antibody chains, including the complementarity determining regions (CDRs) of an antibody derived from one species, preferably a rodent antibody, more preferably a murine antibody,

while the constant domains of the antibody molecule are derived from those of a human antibody. For veterinary applications, the constant domains of the chimeric antibody may be derived from that of other species, such as a subhuman primate, cat or dog.

[0112] By "humanized antibody" herein is meant a recombinant protein in which the CDRs from an antibody from one species; e.g., a rodent antibody, are transferred from the heavy and light variable chains of the rodent antibody into human heavy and light variable domains. The constant domains of the antibody molecule are derived from those of a human antibody. In some embodiments, specific residues of the framework region of the humanized antibody, particularly those that are touching or close to the CDR sequences, may be modified, for example replaced with the corresponding residues from the original rodent, subhuman primate, or other antibody.

[0113] By "human antibody" herein is meant an antibody obtained, for example, from transgenic mice that have been "engineered" to produce specific human antibodies in response to antigenic challenge. In this technique, elements of the human heavy and light chain locus are introduced into strains of mice derived from embryonic stem cell lines that contain targeted disruptions of the endogenous heavy chain and light chain loci. The transgenic mice can synthesize human antibodies specific for human antigens, and the mice can be used to produce human antibody-secreting hybridomas. Methods for obtaining human antibodies from transgenic mice are described by Green et al, Nature Genet. 7: 13 (1994), Lonberg et al, Nature 368:856 (1994), and Taylor et al, Int. Immun. 6:579 (1994). A fully human antibody also can be constructed by genetic or chromosomal transfection methods, as well as phage display technology, all of which are known in the art. See for example, McCafferty et al, Nature 348:552-553 (1990) for the production of human antibodies and fragments thereof in vitro, from immunoglobulin variable domain gene repertoires from unimmunized donors. In this technique, antibody variable domain genes are cloned in-frame into either a major or minor coat protein gene of a filamentous bacteriophage, and displayed as functional antibody fragments on the surface of the phage particle. Because the filamentous particle contains a single-stranded DNA copy of the phage genome, selections based on the functional properties of the antibody also result in selection of the gene encoding the antibody exhibiting those properties. In this way, the phage mimics some of the properties of the B cell. Phage display can be performed in a variety of formats, for their review, see e.g. Johnson and Chiswell, Current Opinion in Structural Biology 3:5564-571 (1993). Human antibodies may also be generated by in vitro activated B cells. See U.S. Patent Nos. 5,567,610 and 5,229,275, which are incorporated herein by reference in their entirety.

[0114] By "antibody fusion protein" herein is meant a recombinantly-produced antigen- binding molecule in which two or more of the same or different natural antibody, single-chain antibody or antibody fragment segments with the same or different specificities are linked. A fusion protein

comprises at least one specific binding site. Valency of the fusion protein indicates the total number of binding arms or sites the fusion protein has to antigen(s) or epitope(s); i.e., monovalent, bivalent, trivalent or mutlivalent. The multivalency of the antibody fusion protein means that it can take advantage of multiple interactions in binding to an antigen, thus increasing the avidity of binding to the antigen, or to different antigens. Specificity indicates how many different types of antigen or epitope an antibody fusion protein is able to bind; i.e., monospecific, bispecific, trispecific, multispecific. Using these definitions, a natural antibody, e.g., an IgG, is bivalent because it has two binding arms but is monospecific because it binds to one type of antigen or epitope. A monospecific, multivalent fusion protein has more than one binding site for the same antigen or epitope. For example, a monospecific diabody is a fusion protein with two binding sites reactive with the same antigen. The fusion protein may comprise a multivalent or multispecific combination of different antibody components or multiple copies of the same antibody component. The fusion protein may additionally comprise a therapeutic agent.

[0115] In some embodiments, the targeting moiety comprises a probody, such as those disclosed in US Patent Nos,: 8,518,404;8,513,390; and US Pat. Appl. Pub. Nos.; 20120237977A1, 20120149061A1, 20130150558A1, the disclosures of which are incorporated by reference in their entireties.

[0116] Probodies are monoclonal antibodies that are selectively activated within the cancer microenvironment, focusing the activity of therapeutic antibodies to tumors and sparing healthy tissue.

[0117] In general, the porbody comprises at least an antibody or antibody fragment thereof (collectively referred to as "AB"), capable of specifically binding a target, wherein the AB is modified by a masking moiety (MM). When the AB is modified with a MM and is in the presence of the target, specific binding of the AB to its target is reduced or inhibited, as compared to the specific binding of the AB not modified with an MM or the specific binding of the parental AB to the target. The dissociation constant (Kd) of the MM towards the AB is generally greater than the Kd of the AB towards the target. When the AB is modified with a MM and is in the presence of the target, specific binding of the AB to its target can be reduced or inhibited, as compared to the specific binding of the AB not modified with an MM or the specific binding of the parental AB to the target. When an AB is coupled to or modified by a MM, the MM can 'mask' or reduce, or inhibit the specific binding of the AB to its target. When an AB is coupled to or modified by a MM, such coupling or modification can effect a structural change which reduces or inhibits the ability of the AB to specifically bind its target.

[0118] In some embodiments, the probody is an activatable antibodies (AAs) where the AB modified by an MM can further include one or more cleavable moieties (CM). Such AAs exhibit activatable/switchable binding, to the AB's target. AAs generally include an antibody or antibody fragment (AB), modified by or coupled to a masking moiety (MM) and a modifiable or cleavable moiety

(CM). In some embodiments, the CM contains an amino acid sequence that serves as a substrate for a protease of interest. In other embodiments, the CM provides a cysteine-cysteine disulfide bond that is cleavable by reduction. In yet other embodiments the CM provides a photolytic substrate that is activatable by photolysis.

[0119] The CM and AB of the AA may be selected so that the AB represents a binding moiety for a target of interest, and the CM represents a substrate for a protease that is co-localized with the target at a treatment site in a subject. Alternatively or in addition, the CM is a cysteine-cysteine disulfide bond that is cleavable as a result of reduction of this disulfide bond. AAs contain at least one of a protease-cleavable CM or a cysteine-cysteine disulfide bond, and in some embodiments include both kinds of CMs. The AAs can alternatively or further include a photolabile substrate, activatable by a light source. The AAs disclosed herein find particular use where, for example, a protease capable of cleaving a site in the CM is present at relatively higher levels in target-containing tissue of a treatment site (for example diseased tissue; for example for therapeutic treatment or diagnostic treatment) than in tissue of non-treatment sites (for example in healthy tissue). The AAs disclosed herein also find particular use where, for example, a reducing agent capable of reducing a site in the CM is present at relatively higher levels in target-containing tissue of a treatment or diagnostic site than in tissue of non-treatment non-diagnostic sites. The AAs disclosed herein also find particular use where, for example, a light source, for example, by way of laser, capable of photolysing a site in the CM is introduced to a target-containing tissue of a treatment or diagnostic site.

[0120] In some embodiments, AAs can provide for reduced toxicity and/or adverse side effects that could otherwise result from binding of the AB at non-treatment sites if the AB were not masked or otherwise inhibited from binding its target. Where the AA contains a CM that is cleavable by a reducing agent that facilitates reduction of a disulfide bond, the ABs of such AAs may selected to exploit activation of an AB where a target of interest is present at a desired treatment site characterized by elevated levels of a reducing agent, such that the environment is of a higher reduction potential than, for example, an environment of a non-treatment site.

[0121] In general, an AA can be designed by selecting an AB of interest and constructing the remainder of the AA so that, when conformationally constrained, the MM provides for masking of the AB or reduction of binding of the AB to its target. Structural design criteria to be taken into account to provide for this functional feature.

[0122] In some embodiments, the TM is a monoclonal anti-HER2 antibody. HER2 is a transmembrane tyrosine kinase receptor encoded by the neu proto-oncogene. HER2 heterodimerizes with HER1, HER3 or HER4, and transmits signals that lead to cell proliferation and survival. Overexpression of HER2 on

the cell surface is correlated with tumor or cancer. Therefore, therapeutics directed at HER2 can be helpful for patients with HER2 positive cancers or tumors. These therapeutic anti-HER2 antibodies mediate their anti-tumor/cancer activities against HER2 positive tumors/cancers by a combination of mechanisms, including promoting modulation of the receptor (internalization and/or "shedding"), induction of apoptosis, antibody-mediated cellular cytotoxicity (ADCC), and presentation of the antigenic determinants of opsonized cells to antigen-presenting cells.

[0123] In some embodiments, the anti-HER2 antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22). Both trastuzumab and pertuzumab are monoclonal antibodies that have anti-tumor activities against HER2 positive tumor or cancer cells. Margetuximab (or MGAH22) is a next-generation, Fc-optimized monoclonal antibody (mAb) that targets HER2. Clinical trials have shown that margetuximab is effective for the tumor or cancer cells that express moderate levels of HER2.

[0124] Herceptin (Trastuzumab) is a humanized monoclonal antibody that acts on human epidermal growth factor receptor extracellular domain of Her2, which is expressed in 25% -30% of breast cancer. It is believed that Trastuzumab has anti-tumor effect through (1) down-regulation Her2 receptor, inhibition of Her2 intracellular signaling transduction pathways and induction of apoptosis; (2) immune mechanisms related antibody dependent ADCC and CDC to kill tumor cells; (3) enhance the effects of chemotherapy.

[0125] In some embodiments, the targeting moiety comprises a Fab, Fab', F(ab')2, single domain antibody, T and Abs dimer, Fv, scFv, dsFv, ds-scFv, Fd, linear antibody, minibody, diabody, bispecific antibody fragment, bibody, tribody, sc-diabody, kappa (lamda) body, BiTE, DVD-Ig, SIP, SMIP, DART, or an antibody analogue comprising one or more CDRs.

[0126] In some embodiments, the targeting moiety comprises extracellular domains (ECD) or soluble form of PD-1, CTLA4, CD47, BTLA, KIR, TIM3, 4-1BB, and LAG3, full length of partial of a surface ligand Amphiregulin, Betacellulin, EGF, Ephrin, Epigen, Epiregulin, IGF, Neuregulin, TGF, TRAIL, or VEGF.

[0127] In some embodiments, the targeting moiety comprises a particle (target particle), preferably a nanoparticle, optionally a targeted nanoparticle that attached to a targeting molecule that can binds specifically or preferably to a target. In some embodiments, the targeting particle by itself guides the compound of the present invention (such as by enrichment in tumor cells or tissue) and there is no additional targeting molecules attached therein.

[0128] By "nanoparticle" herein is meant any particle having a diameter of less than 1000 nm. In some embodiments, a therapeutic agent and/or targeting molecule can be associated with the polymeric matrix.

In some embodiments, the targeting molecule can be covalently associated with the surface of a polymeric matrix. In some embodiments, covalent association is mediated by a linker. In some embodiments, the therapeutic agent can be associated with the surface of, encapsulated within, surrounded by, and/or dispersed throughout the polymeric matrix. US Pat. No. 8,246,968, which is incorporated in its entirety.

[0129] In general, nanoparticles of the present invention comprise any type of particle. Any particle can be used in accordance with the present invention. In some embodiments, particles are biodegradable and biocompatible. In general, a biocompatible substance is not toxic to cells. In some embodiments, a substance is considered to be biocompatible if its addition to cells results in less than a certain threshold of cell death. In some embodiments, a substance is considered to be biocompatible if its addition to cells does not induce adverse effects. In general, a biodegradable substance is one that undergoes breakdown under physiological conditions over the course of a therapeutically relevant time period (e.g., weeks, months, or years). In some embodiments, a biodegradable substance is a substance that can be broken down by cellular machinery. In some embodiments, a biodegradable substance is a substance that can be broken down by chemical processes. In some embodiments, a particle is a substance that is both biocompatible and biodegradable. In some embodiments, a particle is a substance that is biocompatible, but not biodegradable. In some embodiments, a particle is a substance that is biodegradable, but not biocompatible.

[0130] In some embodiments, particles are greater in size than the renal excretion limit (e.g. particles having diameters of greater than 6 nm). In some embodiments, particles are small enough to avoid clearance of particles from the bloodstream by the liver (e.g. particles having diameters of less than 1000 nm). In general, physiochemical features of particles should allow a targeted particle to circulate longer in plasma by decreasing renal excretion and liver clearance.

[0131] It is often desirable to use a population of particles that is relatively uniform in terms of size, shape, and/or composition so that each particle has similar properties. For example, at least 80%, at least 90%, or at least 95% of the particles may have a diameter or greatest dimension that falls within 5%, 10%, or 20% of the average diameter or greatest dimension. In some embodiments, a population of particles may be heterogeneous with respect to size, shape, and/or composition.

[0132] A variety of different particles can be used in accordance with the present invention. In some embodiments, particles are spheres or spheroids. In some embodiments, particles are spheres or spheroids. In some embodiments, particles are flat or plate-shaped. In some embodiments, particles are cubes or cuboids. In some embodiments, particles are ovals or ellipses. In some embodiments, particles are cylinders, cones, or pyramids.

[0133] In some embodiments, particles are microparticles (e.g. microspheres). In general, a "microparticle" refers to any particle having a diameter of less than 1000 μm. In some embodiments, particles are picoparticles (e.g. picospheres). In general, a "picoparticle" refers to any particle having a diameter of less than 1 nm. In some embodiments, particles are liposomes. In some embodiments, particles are micelles.

[0134] Particles can be solid or hollow and can comprise one or more layers (e.g., nanoshells, nanorings). In some embodiments, each layer has a unique composition and unique properties relative to the other layer(s). For example, particles may have a core/shell structure, wherein the core is one layer and the shell is a second layer. Particles may comprise a plurality of different layers. In some embodiments, one layer may be substantially cross-linked, a second layer is not substantially cross-linked, and so forth. In some embodiments, one, a few, or all of the different layers may comprise one or more therapeutic or diagnostic agents to be delivered. In some embodiments, one layer comprises an agent to be delivered, a second layer does not comprise an agent to be delivered, and so forth. In some embodiments, each individual layer comprises a different agent or set of agents to be delivered.

[0135] In some embodiments, a particle is porous, by which is meant that the particle contains holes or channels, which are typically small compared with the size of a particle. For example a particle may be a porous silica particle, e.g., a mesoporous silica nanoparticle or may have a coating of mesoporous silica (Lin et al., 2005, J. Am. Chem. Soc., 17:4570). Particles may have pores ranging from about 1 nm to about 50 nm in diameter, e.g., between about 1 and 20 nm in diameter. Between about 10% and 95% of the volume of a particle may consist of voids within the pores or channels.

[0136] Particles may have a coating layer. Use of a biocompatible coating layer can be advantageous, e.g., if the particles contain materials that are toxic to cells. Suitable coating materials include, but are not limited to, natural proteins such as bovine serum albumin (BSA), biocompatible hydrophilic polymers such as polyethylene glycol (PEG) or a PEG derivative, phospholipid-(PEG), silica, lipids, polymers, carbohydrates such as dextran, other nanoparticles that can be associated with inventive nanoparticles etc. Coatings may be applied or assembled in a variety of ways such as by dipping, using a layer-by-layer technique, by self-assembly, conjugation, etc. Self-assembly refers to a process of spontaneous assembly of a higher order structure that relies on the natural attraction of the components of the higher order structure (e.g., molecules) for each other. It typically occurs through random movements of the molecules and formation of bonds based on size, shape, composition, or chemical properties.

[0137] Examples of polymers include polyalkylenes (e.g. polyethylenes), polycarbonates (e.g. poly(1,3-dioxan-2one)), polyanhydrides (e.g. poly(sebacic anhydride)), polyhydroxyacids (e.g. poly(β-hydroxyalkanoate)), polyfumarates, polycaprolactones, polyamides (e.g. polycaprolactam), polyacetals,

polyethers, polyesters (e.g. polylactide, polyglycolide), poly(orthoesters), polyvinyl alcohols, polyurethanes, polyphosphazenes, polyacrylates, polymethacrylates, polycyanoacrylates, polyureas, polystyrenes, and polyamines. In some embodiments, polymers in accordance with the present invention include polymers which have been approved for use in humans by the U.S. Food and Drug Administration (FDA) under 21 C.F.R. §177.2600, including but not limited to polyesters (e.g. polylactic acid, polyglycolic acid, poly(lactic-co-glycolic acid), polycaprolactone, polyvalerolactone, poly(1,3-dioxan-2one)); polyanhydrides (e.g. poly(sebacic anhydride)); polyethers (e.g., polyethylene glycol); polyurethanes; polymethacrylates; polyacrylates; and polycyanoacrylates.

[0138] In some embodiments, particles can be non-polymeric particles (e.g. metal particles, quantum dots, ceramic particles, polymers comprising inorganic materials, bone-derived materials, bone substitutes, viral particles, etc.). In some embodiments, a therapeutic or diagnostic agent to be delivered can be associated with the surface of such a non-polymeric particle. In some embodiments, a non-polymeric particle is an aggregate of non-polymeric components, such as an aggregate of metal atoms (e.g. gold atoms). In some embodiments, a therapeutic or diagnostic agent to be delivered can be associated with the surface of and/or encapsulated within, surrounded by, and/or dispersed throughout an aggregate of non-polymeric components.

[0139] Particles (e.g. nanoparticles, microparticles) may be prepared using any method known in the art. For example, particulate formulations can be formed by methods as nanoprecipitation, flow focusing fluidic channels, spray drying, single and double emulsion solvent evaporation, solvent extraction, phase separation, milling, microemulsion procedures, microfabrication, nanofabrication, sacrificial layers, simple and complex coacervation, and other methods well known to those of ordinary skill in the art. Alternatively or additionally, aqueous and organic solvent syntheses for monodisperse semiconductor, conductive, magnetic, organic, and other nanoparticles have been described (Pellegrino et al., 2005, Small, 1:48; Murray et al., 2000, Ann. Rev. Mat. Sci., 30:545; and Trindade et al., 2001, Chem. Mat., 13:3843).

[0140] Methods for making microparticles for delivery of encapsulated agents are described in the literature (see, e.g., Doubrow, Ed., "Microcapsules and Nanoparticles in Medicine and Pharmacy," CRC Press, Boca Raton, 1992; Mathiowitz et al., 1987, J. Control. Release, 5:13; Mathiowitz et al., 1987, Reactive Polymers, δ: 275; and Mathiowitz et al., 1988, J. Appl. Polymer Sci., 35:755).

[0141] In some embodiments, the targeting moiety comprises an nucleic acid targeting moiety.

[0142] In general, a nucleic acid targeting moiety is any polynucleotide that binds to a component associated with an organ, tissue, cell, extracellular matrix component, and/or intracellular compartment (the target).

[0143] In some embodiments, nucleic acid targeting moieties are aptamers.

[0144] An aptamer is typically a polynucleotide that binds to a specific target structure that is associated with a particular organ, tissue, cell, extracellular matrix component, and/or intracellular compartment. In general, the targeting function of the aptamer is based on the three-dimensional structure of the aptamer. In some embodiments, binding of an aptamer to a target is typically mediated by the interaction between the two- and/or three-dimensional structures of both the aptamer and the target. In some embodiments, binding of an aptamer to a target is not solely based on the primary sequence of the aptamer, but depends on the three-dimensional structure(s) of the aptamer and/or target. In some embodiments, aptamers bind to their targets via complementary Watson-Crick base pairing which is interrupted by structures (e.g. hairpin loops) that disrupt base pairing.

[0145] In some embodiments, nucleic acid targeting moieties are spiegelmers (PCT Publications WO 98/08856, WO 02/100442, and WO 06/117217). In general, spiegelmers are synthetic, mirror-image nucleic acids that can specifically bind to a target (i.e. mirror image aptamers). Spiegelmers are characterized by structural features which make them not susceptible to exo- and endo-nucleases.

[0146] One of ordinary skill in the art will recognize that any nucleic acid targeting moiety (e.g. aptamer or spiegelmer) that is capable of specifically binding to a target can be used in accordance with the present invention. In some embodiments, nucleic acid targeting moieties to be used in accordance with the present invention may target a marker associated with a disease, disorder, and/or condition. In some embodiments, nucleic acid targeting moieties to be used in accordance with the present invention may target cancer-associated targets. In some embodiments, nucleic acid targeting moieties to be used in accordance with the present invention may target tumor markers. Any type of cancer and/or any tumor marker may be targeted using nucleic acid targeting moieties in accordance with the present invention. To give but a few examples, nucleic acid targeting moieties may target markers associated with prostate cancer, lung cancer, breast cancer, colorectal cancer, bladder cancer, pancreatic cancer, endometrial cancer, ovarian cancer, bone cancer, esophageal cancer, liver cancer, stomach cancer, brain tumors, cutaneous melanoma, and/or leukemia.

[0147] Nucleic acids of the present invention (including nucleic acid nucleic acid targeting moieties and/or functional RNAs to be delivered, e.g., RNAi-inducing entities, ribozymes, tRNAs, etc., described in further detail below) may be prepared according to any available technique including, but not limited to chemical synthesis, enzymatic synthesis, enzymatic or chemical cleavage of a longer precursor, etc. Methods of synthesizing RNAs are known in the art (see, e.g., Gait, M. J. (ed.) Oligonucleotide synthesis: a practical approach, Oxford [Oxfordshire], Washington, D.C.: IRL Press, 1984; and Herdewijn, P. (ed.)

Oligonucleotide synthesis: methods and applications, Methods in molecular biology, v. 288 (Clifton, N.J.) Totowa, N.J.: Humana Press, 2005).

[0148] The nucleic acid that forms the nucleic acid nucleic acid targeting moiety may comprise naturally occurring nucleosides, modified nucleosides, naturally occurring nucleosides with hydrocarbon linkers (e.g., an alkylene) or a polyether linker (e.g., a PEG linker) inserted between one or more nucleosides, modified nucleosides with hydrocarbon or PEG linkers inserted between one or more nucleosides, or a combination of thereof. In some embodiments, nucleotides or modified nucleotides of the nucleic acid nucleic acid targeting moiety can be replaced with a hydrocarbon linker or a polyether linker provided that the binding affinity and selectivity of the nucleic acid nucleic acid targeting moiety is not substantially reduced by the substitution (e.g., the dissociation constant of the nucleic acid nucleic acid targeting moiety for the target should not be greater than about 1×10^{-3} M).

[0149] It will be appreciated by those of ordinary skill in the art that nucleic acids in accordance with the present invention may comprise nucleotides entirely of the types found in naturally occurring nucleic acids, or may instead include one or more nucleotide analogs or have a structure that otherwise differs from that of a naturally occurring nucleic acid. U.S. Pat. Nos. 6,403,779; 6,399,754; 6,225,460; 6,127,533; 6,031,086; 6,005,087; 5,977,089; and references therein disclose a wide variety of specific nucleotide analogs and modifications that may be used. See Crooke, S. (ed.) Antisense Drug Technology: Principles, Strategies, and Applications (1st ed), Marcel Dekker; ISBN: 0824705661; 1st edition (2001) and references therein. For example, 2′-modifications include halo, alkoxy and allyloxy groups. In some embodiments, the 2′-OH group is replaced by a group selected from H, OR, R, halo, SH, SR, NH2, NHR, NR2 or CN, wherein R is C1-C6 alkyl, alkenyl, or alkynyl, and halo is F, Cl, Br, or I. Examples of modified linkages include phosphorothioate and 5′-N-phosphoramidite linkages.

[0150] Nucleic acids comprising a variety of different nucleotide analogs, modified backbones, or non-naturally occurring internucleoside linkages can be utilized in accordance with the present invention. Nucleic acids of the present invention may include natural nucleosides (i.e., adenosine, thymidine, guanosine, cytidine, uridine, deoxyadenosine, deoxythymidine, deoxyguanosine, and deoxycytidine) or modified nucleosides. Examples of modified nucleotides include base modified nucleoside (e.g., aracytidine, inosine, isoguanosine, nebularine, pseudouridine, 2,6-diaminopurine, 2-aminopurine, 2-thiothymidine, 3-deaza-5-azacytidine, 2′-deoxyuridine, 3-nitorpyrrole, 4-methylindole, 4-thiouridine, 4-thiothymidine, 2-aminoadenosine, 2-thiothymidine, 2-thiouridine, 5-bromocytidine, 5-iodouridine, inosine, 6-azauridine, 6-chloropurine, 7-deazaadenosine, 7-deazaguanosine, 8-azaadenosine, 8-azidoadenosine, benzimidazole, M1-methyladenosine, pyrrolo-pyrimidine, 2-amino-6-chloropurine, 3-methyl adenosine, 5-propynylcytidine, 5-propynyluridine, 5-bromouridine, 5-fluorouridine, 5-

methylcytidine, 7-deazaadenosine, 7-deazaguanosine, 8-oxoadenosine, 8-oxoguanosine, O(6)-methylguanine, and 2-thiocytidine), chemically or biologically modified bases (e.g., methylated bases), modified sugars (e.g., 2′-fluororibose, 2′-aminoribose, 2′-azidoribose, 2′-O-methylribose, L-enantiomeric nucleosides arabinose, and hexose), modified phosphate groups (e.g., phosphorothioates and 5′-N-phosphoramidite linkages), and combinations thereof. Natural and modified nucleotide monomers for the chemical synthesis of nucleic acids are readily available. In some cases, nucleic acids comprising such modifications display improved properties relative to nucleic acids consisting only of naturally occurring nucleotides. In some embodiments, nucleic acid modifications described herein are utilized to reduce and/or prevent digestion by nucleases (e.g. exonucleases, endonucleases, etc.). For example, the structure of a nucleic acid may be stabilized by including nucleotide analogs at the 3′ end of one or both strands order to reduce digestion.

[0151] Modified nucleic acids need not be uniformly modified along the entire length of the molecule. Different nucleotide modifications and/or backbone structures may exist at various positions in the nucleic acid. One of ordinary skill in the art will appreciate that the nucleotide analogs or other modification(s) may be located at any position(s) of a nucleic acid such that the function of the nucleic acid is not substantially affected. To give but one example, modifications may be located at any position of a nucleic acid targeting moiety such that the ability of the nucleic acid targeting moiety to specifically bind to the target is not substantially affected. The modified region may be at the 5′-end and/or the 3′-end of one or both strands. For example, modified nucleic acid targeting moieties in which approximately 1-5 residues at the 5′ and/or 3′ end of either of both strands are nucleotide analogs and/or have a backbone modification have been employed. The modification may be a 5′ or 3′ terminal modification. One or both nucleic acid strands may comprise at least 50% unmodified nucleotides, at least 80% unmodified nucleotides, at least 90% unmodified nucleotides, or 100% unmodified nucleotides.

[0152] Nucleic acids in accordance with the present invention may, for example, comprise a modification to a sugar, nucleoside, or internucleoside linkage such as those described in U.S. Patent Application Publications 2003/0175950, 2004/0192626, 2004/0092470, 2005/0020525, and 2005/0032733. The present invention encompasses the use of any nucleic acid having any one or more of the modification described therein. For example, a number of terminal conjugates, e.g., lipids such as cholesterol, lithocholic acid, aluric acid, or long alkyl branched chains have been reported to improve cellular uptake. Analogs and modifications may be tested using, e.g., using any appropriate assay known in the art, for example, to select those that result in improved delivery of a therapeutic or diagnostic agent, improved specific binding of an nucleic acid targeting moiety to a target, etc. In some embodiments, nucleic acids in accordance with the present invention may comprise one or more non-natural nucleoside

linkages. In some embodiments, one or more internal nucleotides at the 3′-end, 5′-end, or both 3′- and 5′-ends of the nucleic acid targeting moiety are inverted to yield a linkage such as a 3′-3′ linkage or a 5′-5′ linkage.

[0153] In some embodiments, nucleic acids in accordance with the present invention are not synthetic, but are naturally-occurring entities that have been isolated from their natural environments.

[0154] Any method can be used to design novel nucleic acid targeting moieties (see, e.g., U.S. Pat. Nos. 6,716,583; 6,465,189; 6,482,594; 6,458,543; 6,458,539; 6,376,190; 6,344,318; 6,242,246; 6,184,364; 6,001,577; 5,958,691; 5,874,218; 5,853,984; 5,843,732; 5,843,653; 5,817,785; 5,789,163; 5,763,177; 5,696,249; 5,660,985; 5,595,877; 5,567,588; and 5,270,163; and U.S. Patent Application Publications 2005/0069910, 2004/0072234, 2004/0043923, 2003/0087301, 2003/0054360, and 2002/0064780). The present invention provides methods for designing novel nucleic acid targeting moieties. The present invention further provides methods for isolating or identifying novel nucleic acid targeting moieties from a mixture of candidate nucleic acid targeting moieties.

[0155] Nucleic acid targeting moieties that bind to a protein, a carbohydrate, a lipid, and/or a nucleic acid can be designed and/or identified. In some embodiments, nucleic acid targeting moieties can be designed and/or identified for use in the complexes of the invention that bind to proteins and/or characteristic portions thereof, such as tumor-markers, integrins, cell surface receptors, transmembrane proteins, intercellular proteins, ion channels, membrane transporter proteins, enzymes, antibodies, chimeric proteins etc. In some embodiments, nucleic acid targeting moieties can be designed and/or identified for use in the complexes of the invention that bind to carbohydrates and/or characteristic portions thereof, such as glycoproteins, sugars (e.g., monosaccharides, disaccharides and polysaccharides), glycocalyx (i.e., the carbohydrate-rich peripheral zone on the outside surface of most eukaryotic cells) etc. In some embodiments, nucleic acid targeting moieties can be designed and/or identified for use in the complexes of the invention that bind to lipids and/or characteristic portions thereof, such as oils, saturated fatty acids, unsaturated fatty acids, glycerides, hormones, steroids (e.g., cholesterol, bile acids), vitamins (e.g. vitamin E), phospholipids, sphingolipids, lipoproteins etc. In some embodiments, nucleic acid targeting moieties can be designed and/or identified for use in the complexes of the invention that bind to nucleic acids and/or characteristic portions thereof, such as DNA nucleic acids; RNA nucleic acids; modified DNA nucleic acids; modified RNA nucleic acids; and nucleic acids that include any combination of DNA, RNA, modified DNA, and modified RNA; etc.

[0156] Nucleic acid targeting moieties (e.g. aptamers or spiegelmers) may be designed and/or identified using any available method. In some embodiments, nucleic acid targeting moieties are designed and/or identified by identifying nucleic acid targeting moieties from a candidate mixture of nucleic acids.

Systemic Evolution of Ligands by Exponential Enrichment (SELEX), or a variation thereof, is a commonly used method of identifying nucleic acid targeting moieties that bind to a target from a candidate mixture of nucleic acids.

[0157] Nucleic acid targeting moieties that bind selectively to any target can be isolated by the SELEX process, or a variation thereof, provided that the target can be used as a target in the SELEX process.

Linker

[0158] In general, the compound of the invention comprises a linker that links the targeting moiety and the activating moiety. Though, in some compound there is no linker and the activating moiety and the targeting moiety is linked directly.

[0159] By "linker" herein is meant a moiety that connects a first molecule to a second molecule through chemical bonds. In linkers of the invention, the connection can be severed so as to release a biologically active form of the first and/or second molecule. A preferred example of a linker is a moiety that comprises a bond that is stable at neutral pH but is readily cleaved under conditions of low pH. Particularly preferred examples of linkers are moieties that comprise a bond that is stable at pH values between 7 and 8 but is readily cleaved at pH values between 4 and 6. Another example of a linker is a moiety that comprises a bond that is readily cleaved in the presence of an enzyme. Preferred examples of such enzyme-sensitive linkers are peptides comprising a recognition sequence for an endosomal peptidase. Another example of a linker is a redox potential-sensitive linker that is stable under conditions of low reduction potential (e.g., low thiol or glutathione concentration) but cleaved under conditions of high reduction potential (e.g., high thiol or glutathione concentration). Preferred examples of such redox potential-sensitive linkers include disulfides and sulfenamides. Particularly preferred examples include substituted aryl-alkyl disulfides in which the aryl group is substituted with sterically-demanding and electron-withdrawing or electron-donating substitutents, so as to control the sensitivity of the disulfide linkage towards reaction with thiol. Another example of a linker is a moiety that comprises a bond that is readily cleaved upon exposure to radiation. Preferred examples of such radiation-sensitive linkers are 2-nitrobenzyl ethers that are cleaved upon exposure to light. Particularly preferred examples of linkers are moieties that mask the biological activity of one of the two linked molecules until the linkage is severed.

[0160] In some embodiments, the compound of the invention comprises a linker that is selected from the group consisting of a hydrazine group, a polypeptide, a disulfide group, and a thioether group.

[0161] By "hydrazine group" or "hydrazine linker" or "self-cyclizing hydrazine linker" herein is meant a linker moiety that, upon a change in condition, such as a shift in pH, will undergo a cyclization reaction and form one or more rings. The hydrazine moiety is converted to a hydrazone when attached. This

attachment can occur, for example, through a reaction with a ketone group on the L4 moiety. Therefore, the term hydrazine linker can also be used to describe the linker of the current invention because of this conversion to a hydrazone upon attachment.

[0162] By "five-membered hydrazine linker" or "5-membered hydrazine linker" herein is meant hydrazine-containing molecular moieties that, upon a change in condition, such as a shift in pH, will undergo a cyclization reaction and form one or more 5-membered rings. Alternatively, this five-membered linker may similarly be described as a five-membered hydrazine linker or a 5-membered hydrazine linker.

[0163] By "six-membered hydrazine linker" or "6-membered hydrazine linker" herein is meant hydrazine-containing molecular moieties that, upon a change in condition such as a shift in pH, will undergo a cyclization reaction and form one or more 6-membered rings. This six membered linker may similarly be described as a six-membered hydrazine linker or a 6-membered hydrazine linker.

[0164] By "cyclization reaction" herein is meant the cyclization of a peptide, hydrazine, or disulfide linker, indicates the cyclization of that linker into a ring and initiates the separation of the drug-ligand complex. This rate can be measured ex situ, and is completed when at least 90%, 95%, or 100% of the product is formed.

[0165] In some embodiments, the compound of the present invention comprises a linker region between the targeting moiety and the activating moiety, and the linker is cleavable by a cleaving agent that is present in the intracellular environment (e.g., within a lysosome or endosome or caveolea). The linker can be, e.g., a peptidyl linker that is cleaved by an intracellular peptidase or protease enzyme, including, but not limited to, a lysosomal or endosomal protease. Typically, the peptidyl linker is at least two amino acids long or at least three amino acids long. Cleaving agents can include cathepsins B and D and plasmin, all of which are known to hydrolyze dipeptide drug derivatives resulting in the release of active drug inside target cells (see, e.g., Dubowchik and Walker, 1999, Pharm. Therapeutics 83:67-123). Most typical are peptidyl linkers that are cleavable by enzymes that are present in targeted cells or tissues. For example, a peptidyl linker that is cleavable by the thiol-dependent protease cathepsin-B, which is highly expressed in cancerous tissue, can be used (e.g., a Phe-Leu or a Gly-Phe-Leu-Gly) linker). Other such linkers are described, e.g., in U.S. Pat. No. 6,214,345. In some embodiments, the peptidyl linker cleavable by an intracellular protease is a Val-Cit linker or a Phe-Lys linker (see, e.g., U.S. Pat. No. 6,214,345, which describes the synthesis of doxorubicin with the val-cit linker). One advantage of using intracellular proteolytic release of the therapeutic agent is that the agent is typically attenuated when conjugated and the serum stabilities of the conjugates are typically high.

[0166] In some embodiments, the cleavable linker is pH-sensitive, i.e., sensitive to hydrolysis at certain pH values. Typically, the pH-sensitive linker is hydrolyzable under acidic conditions. For example, an acid-labile linker that is hydrolyzable in the lysosome (e.g., a hydrazone, semicarbazone, thiosemicarbazone, cis-aconitic amide, orthoester, acetal, ketal, or the like) can be used. (See, e.g., U.S. Pat. Nos. 5,122,368; 5,824,805; 5,622,929; Dubowchik and Walker, 1999, Pharm. Therapeutics 83:67-123; Neville et al., 1989, Biol. Chem. 264:14653-14661.) Such linkers are relatively stable under neutral pH conditions, such as those in the blood, but are unstable at below pH 5.5 or 5.0, the approximate pH of the lysosome. In certain embodiments, the hydrolyzable linker is a thioether linker (such as, e.g., a thioether attached to the therapeutic agent via an acylhydrazone bond (see, e.g., U.S. Pat. No. 5,622,929)).

[0167] In yet other embodiments, the linker is cleavable under reducing conditions (e.g., a disulfide linker). A variety of disulfide linkers are known in the art, including, for example, those that can be formed using SATA (N-succinimidyl-5-acetylthioacetate), SPDP (N-succinimidyl-3-(2-pyridyldithio)propionate), SPDB (N-succinimidyl-3-(2-pyridyldithio)butyrate) and SMPT (N-succinimidyl-oxycarbonyl-alpha-methyl-alpha-(2-pyridyl-dithio)toluene), SPDB and SMPT (See, e.g., Thorpe et al., 1987, Cancer Res. 47:5924-5931; Wawrzynczak et al., In Immunoconjugates: Antibody Conjugates in Radioimagery and Therapy of Cancer (C. W. Vogel ed., Oxford U. Press, 1987. See also U.S. Pat. No. 4,880,935.)

[0168] In yet other specific embodiments, the linker is a malonate linker (Johnson et al., 1995, Anticancer Res. 15:1387-93), a maleimidobenzoyl linker (Lau et al., 1995, Bioorg-Med-Chem. 3(10):1299-1304), or a 3′-N-amide analog (Lau et al., 1995,Bioorg-Med-Chem. 3(10):1305-12).

[0169] Typically, the linker is not substantially sensitive to the extracellular environment. As used herein, "not substantially sensitive to the extracellular environment," in the context of a linker, means that no more than about 20%, typically no more than about 15%, more typically no more than about 10%, and even more typically no more than about 5%, no more than about 3%, or no more than about 1% of the linkers, in a sample of compounds of the present invention, are cleaved when the compounds of the present invention present in an extracellular environment (e.g., in plasma). Whether a linker is not substantially sensitive to the extracellular environment can be determined, for example, by incubating independently with plasma both (a) the compound of the invention (the "Compound sample") and (b) an equal molar amount of unconjugated antibody or therapeutic agent (the "control sample") for a predetermined time period (e.g., 2, 4, 8, 16, or 24 hours) and then comparing the amount of unconjugated antibody or therapeutic agent present in the Compound sample with that present in control sample, as measured, for example, by high performance liquid chromatography.

[0170] In other, non-mutually exclusive embodiments, the linker promotes cellular internalization. In certain embodiments, the linker promotes cellular internalization when conjugated to the activating moiety. In yet other embodiments, the linker promotes cellular internalization when conjugated to both the targeting moiety and the activating moiety.

[0171] A variety of linkers that can be used with the present compositions and methods are described in WO 2004010957 entitled "Drug Conjugates and Their Use for Treating Cancer, An Autoimmune Disease or an Infectious Disease", US20120141509A1, and US20120288512A1 (the disclosure of which is incorporated by reference herein).

[0172] In certain embodiments, the linker unit has the following general formula:

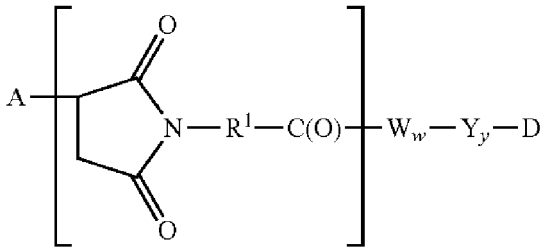
-Ta-Ww-Yy-

wherein -T- is a stretcher unit; a is 0 or 1; each -W- is independently an amino acid unit; w is independently an integer ranging from 2 to 12; -Y- is a spacer unit; and y is 0, 1 or 2.

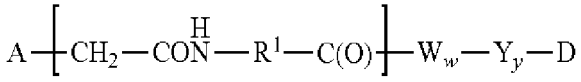
The Stretcher Unit

[0173] The stretcher unit (-T-), when present, links the targeting moiety to an amino acid unit (-W-). Useful functional groups that can be present on a targeting moiety, such as an antibody, either naturally or via chemical manipulation include, but are not limited to, sulfhydryl, amino, hydroxyl, the anomeric hydroxyl group of a carbohydrate, and carboxyl. Suitable functional groups are sulfhydryl and amino. Sulfhydryl groups can be generated by reduction of the intramolecular disulfide bonds of an antibody. Alternatively, sulfhydryl groups can be generated by reaction of an amino group of a lysine moiety of an antibody with 2-iminothiolane (Traut's reagent) or other sulfhydryl generating reagents. In some embodiments, the antibody is a recombinant antibody and is engineered to carry one or more lysines. In other embodiments, the recombinant antibody is engineered to carry additional sulfhydryl groups, e.g., additional cysteines.

[0174] In some embodiments, the stretcher unit forms a bond with a sulfur atom of the antibody. The sulfur atom can be derived from a sulfhydryl (-SH) group of a reduced antibody (A). Representative stretcher units of these embodiments are depicted within the square brackets of Formulas (IIa) and (IIb), wherein A-, -W-, -Y-, -D, w and y are as defined above and R^1 is selected from —C_1-C_{10} alkylene-, —C_3-C_8 carbocyclo-, —O—(C_1-C_8 alkyl)-, -arylene-, —C_1-C_{10} alkylene-arylene-, -arylene-C_1-C_{10} alkylene-, —C_1-C_{10} alkylene-(C_3-C_8 carbocyclo)-, —(C_3-C_8 carbocyclo)-C_1-C_{10}alkylene-, —C_3-C_8 heterocyclo-, —C_1-C_{10} alkylene-(C_3-C_8 heterocyclo)-, —(C_3-C_8 heterocyclo)-C_1-C_{10} alkylene-, —(CH_2CH_2O)r—, and —(CH_2CH_2O)r—CH_2—; and r is an integer ranging from 1-10.

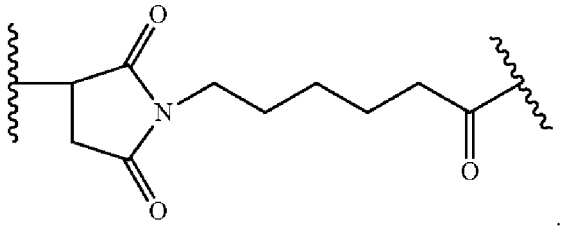


(IIa)

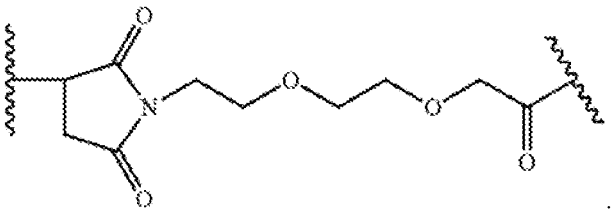


(IIb)

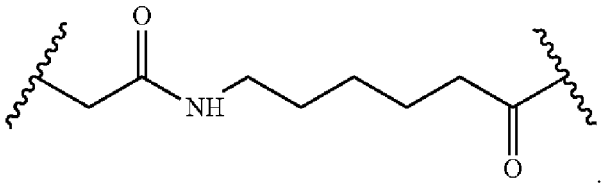
[0175] An illustrative stretcher unit is that of formula (IIa) where R^1 is $—(CH_2)_5—$:



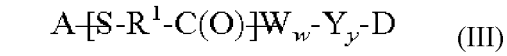
[0176] Another illustrative stretcher unit is that of formula (IIa) where R^1 is $—(CH_2CH_2O)r—CH_2—$ and r is 2:



[0177] Still another illustrative stretcher unit is that of formula (IIb) where R^1 is $—(CH_2)_5—$:

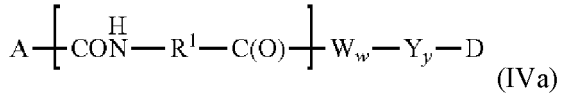


[0178] In certain other specific embodiments, the stretcher unit is linked to the antibody unit (A) via a disulfide bond between a sulfur atom of the antibody unit and a sulfur atom of the stretcher unit. A representative stretcher unit of this embodiment is depicted within the square brackets of Formula (III), wherein R^1, A-, -W-, -Y-, -D, w and y are as defined above.

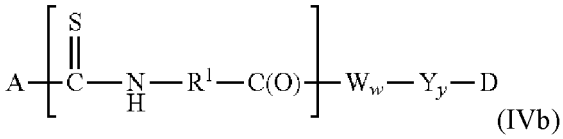


(III)

[0179] In other specific embodiments, the reactive group of the stretcher contains a reactive site that can be reactive to an amino group of an antibody. The amino group can be that of an arginine or a lysine. Suitable amine reactive sites include, but are not limited to, activated esters such as succinimide esters, 4-nitrophenyl esters, pentafluorophenyl esters, anhydrides, acid chlorides, sulfonyl chlorides, isocyanates and isothiocyanates. Representative stretcher units of these embodiments are depicted within the square brackets of Formulas (IVa) and (IVb), wherein R^1, A-, -W-, -Y-, -D, w and y are as defined above;

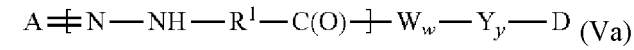


(IVa)

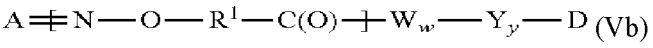


(IVb)

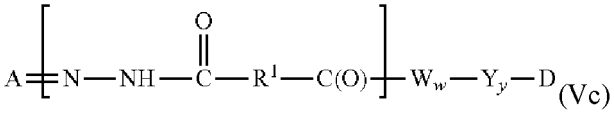
[0180] In yet another aspect, the reactive function of the stretcher contains a reactive site that is reactive to a modified carbohydrate group that can be present on an antibody. In some embodiments, the antibody is glycosylated enzymatically to provide a carbohydrate moiety. The carbohydrate may be mildly oxidized with a reagent such as sodium periodate and the resulting carbonyl unit of the oxidized carbohydrate can be condensed with a stretcher that contains a functionality such as a hydrazide, an oxime, a reactive amine, a hydrazine, a thiosemicarbazide, a hydrazine carboxylate, and an arylhydrazide such as those described by Kaneko et al., 1991, Bioconjugate Chem 2:133-41. Representative stretcher units of this embodiment are depicted within the square brackets of Formulas (Va)-(Vc), wherein R^1, A-, -W-, -Y-, -D, w and y are as defined above.



(Va)



(Vb)

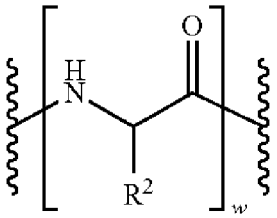


(Vc)

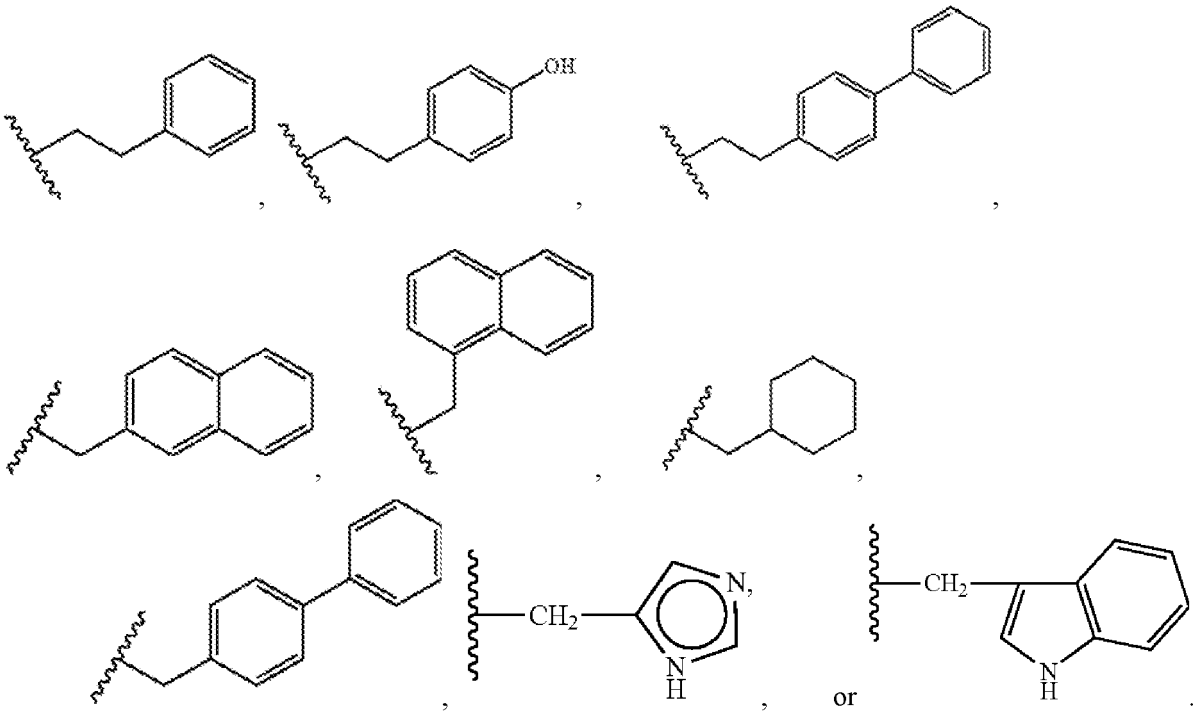
The Amino Acid Unit

[0181] The amino acid unit (-W-) links the stretcher unit (-T-) to the Spacer unit (-Y-) if the Spacer unit is present, and links the stretcher unit to the cytotoxic or cytostatic agent (Activating Moiety; D) if the spacer unit is absent. -Ww- is a dipeptide, tripeptide, tetrapeptide, pentapeptide, hexapeptide, heptapeptide, octapeptide, nonapeptide, decapeptide, undecapeptide or dodecapeptide unit. Each -W- unit

independently has the formula denoted below in the square brackets, and w is an integer ranging from 2 to 12:

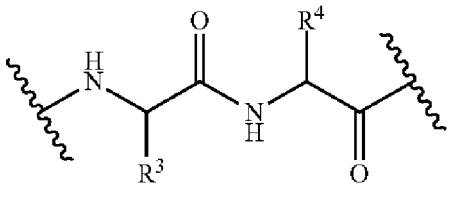


wherein R^2 is hydrogen, methyl, isopropyl, isobutyl, sec-butyl, benzyl, p-hydroxybenzyl, —CH_2OH, —$CH(OH)CH_3$, —$CH_2CH_2SCH_3$, —CH_2CONH_2, —CH_2COOH, —$CH_2CH_2CONH_2$, —CH_2CH_2COOH, —$(CH_2)_3NHC(=NH)NH_2$, —$(CH_2)_3NH_2$, —$(CH_2)_3NHCOCH_3$, —$(CH_2)_3NHCHO$, —$(CH_2)_4NHC(=NH)NH_2$, —$(CH_2)_4NH_2$, —$(CH_2)_4NHCOCH_3$, —$(CH_2)_4NHCHO$, —$(CH_2)_3NHCONH_2$, —$(CH_2)_4NHCONH_2$, —$CH_2CH_2CH(OH)CH_2NH_2$, 2-pyridylmethyl-, 3-pyridylmethyl-, 4-pyridylmethyl-, phenyl, cyclohexyl,



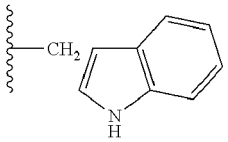
[0182] The amino acid unit of the linker unit can be enzymatically cleaved by an enzyme including, but not limited to, a tumor-associated protease to liberate the activating moiety (-D) which is protonated in vivo upon release to provide an activating molecule (D).

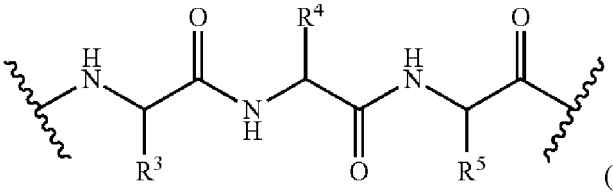
[0183] Illustrative W_w units are represented by formulas (VI)-(VIII):



(VI)

wherein R^3 and R^4 are as follows:

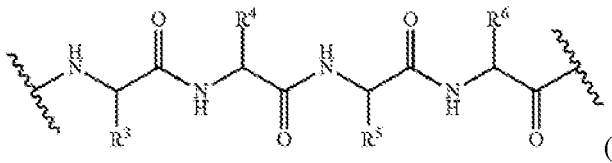
R^3	R^4
Benzyl	$(CH_2)_4NH_2$;
Methyl	$(CH_2)_4NH_2$;
Isopropyl	$(CH_2)_4NH_2$;
Isopropyl	$(CH_2)_3NHCONH_2$;
Benzyl	$(CH_2)_3NHCONH_2$;
Isobutyl	$(CH_2)_3NHCONH_2$;
sec-butyl	$(CH_2)_3NHCONH_2$;
	$(CH_2)_3NHCONH_2$;
Benzyl	methyl; and
Benzyl	$(CH_2)_3NHC(=NH)NH_2$;



(VII)

wherein R^3, R^4 and R^5 are as follows:

R^3	R^4	R^5
Benzyl	benzyl	$(CH_2)_4NH_2$;
Isopropyl	benzyl	$(CH_2)_4NH_2$; and
H	benzyl	$(CH_2)_4NH_2$;



(VIII)

wherein R^3, R^4, R^5 and R^6 are as follows:

R^3	R^4	R^5	R^6
H	Benzyl	isobutyl	H; and
methyl	Isobutyl	methyl	isobutyl.

[0184] Suitable amino acid units include, but are not limited to, units of formula (VI) where: R^3 is benzyl and R^4 is —$(CH_2)_4NH_2$; R_3 is isopropyl and R^4 is —$(CH_2)_4NH_2$; or R^3 is isopropyl and R^4 is —$(CH_2)_3NHCONH_2$. Another suitable amino acid unit is a unit of formula (VII), where: R^3 is benzyl, R^4 is benzyl, and R^5 is —$(CH_2)_4NH_2$. -Ww-units can be designed and optimized in their selectivity for enzymatic cleavage by a particular tumor-associated protease. The suitable -Ww- units are those whose cleavage is catalyzed by the proteases, cathepsin B, C and D, and plasmin.

[0185] In some embodiments, -Ww- is a dipeptide, tripeptide or tetrapeptide unit.

[0186] Where R^2, R^3, R^4, R^5 or R^6 is other than hydrogen, the carbon atom to which R^2, R^3, R^4, R^5 or R^6 is attached is chiral. Each carbon atom to which R^2, R^3, R^4, R^5 or R^6 is attached is independently in the (S) or (R) configuration.

[0187] In some embodiments, the amino acid unit is a phenylalanine-lysine dipeptide (Phe-Lys or FK linker). In some embodiments, the amino acid unit is a valine-citrulline dipeptide (Val-Cit or VC linker). In some embodiments, the amino acid unit is 5-aminovaleric acid, homo phenylalanine lysine, tetraisoquinolinecarboxylate lysine, cyclohexylalanine lysine, isonepecotic acid lysine, beta-alanine lysine, glycine serine valine glutamine, or isonepecotic acid.

[0188] The amino acid unit can comprise natural amino acids. In other embodiments, the Amino Acid unit can comprise non-natural amino acids.

The Spacer Unit

[0189] The spacer unit (-Y-), when present, links an amino acid unit to the drug unit. Spacer units are of two general types: self-immolative and non self-immolative. A non self-immolative spacer unit is one in which part or all of the spacer unit remains bound to the activating moiety unit after enzymatic cleavage of an amino acid unit from the TM-linker-AM conjugate or the drug-linker compound. Examples of a

non self-immolative spacer unit include, but are not limited to a (glycine-glycine) spacer unit and a glycine spacer unit. When a TM-linker-AM conjugate containing a glycine-glycine spacer unit or a glycine spacer unit undergoes enzymatic cleavage via a tumor-cell associated-protease, a cancer-cell-associated protease or a lymphocyte-associated protease, a glycine-glycine-drug moiety or a glycine-drug moiety is cleaved from A-T-Ww-. To liberate the AM, an independent hydrolysis reaction should take place within the target cell to cleave the glycine-drug unit bond.

[0190] In a typical embodiment, -Yy- is a p-aminobenzyl ether which can be substituted with Qm where Q is —C_1-C_8 alkyl, —C1-C_8alkoxy, -halogen, -nitro or -cyano; and m is an integer ranging from 0-4.

[0191] In some embodiments, a non self-immolative spacer unit (-Y-) is -Gly-Gly-.

[0192] In some embodiments, a non self-immolative the spacer unit (-Y-) is -Gly-.

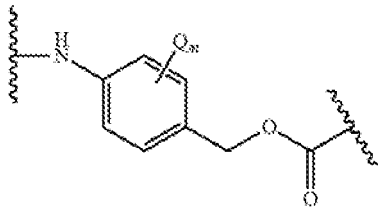
[0193] In one embodiment, the AM-linker compound or an TM-linker-AM conjugate lacks a spacer unit (y=0).

[0194] Alternatively, an TM-linker-AM conjugate containing a self-immolative spacer unit can release the AM (D) without the need for a separate hydrolysis step. In these embodiments, -Y- is a p-aminobenzyl alcohol (PAB) unit that is linked to -Ww- via the nitrogen atom of the PAB group, and connected directly to -D via a carbonate, carbamate or ether group.

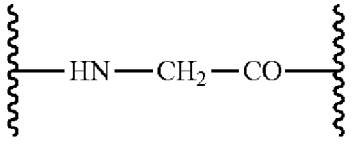
[0195] Other examples of self-immolative spacers include, but are not limited to, aromatic compounds that are electronically equivalent to the PAB group such as 2-aminoimidazol-5-methanol derivatives (see Hay et al., 1999, Bioorg. Med. Chem. Lett.9:2237 for examples) and ortho or para-aminobenzylacetals. Spacers can be used that undergo facile cyclization upon amide bond hydrolysis, such as substituted and unsubstituted 4-aminobutyric acid amides (Rodrigues et al., 1995, Chemistry Biology 2:223), appropriately substituted bicyclo[2.2.1] and bicyclo[2.2.2] ring systems (Storm et al., 1972, J. Amer. Chem. Soc. 94:5815) and 2-aminophenylpropionic acid amides (Amsberry et al., 1990, J. Org. Chem. 55:5867) Elimination of amine-containing drugs that are substituted at the α-position of glycine (Kingsbury, et al., 1984, J. Med. Chem. 27:1447) are also examples of self-immolative spacer strategies that can be applied to the TM-linker-AM conjugates.

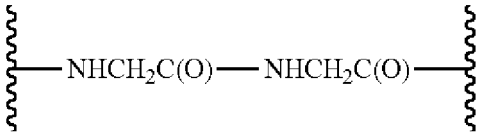
[0196] In an alternate embodiment, the spacer unit is a branched bis(hydroxymethyl)styrene (BHMS) unit, which can be used to incorporate moieties.

[0197] Typical spacer units (-Yy-) are represented by Formulas (IX)-(XI):


(IX)

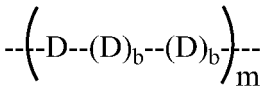
where Q is C_1-C_8 alkyl, C_1-C_8 alkoxy, halogen, nitro or cyano; and m is an integer ranging from 0-4;


(X), and

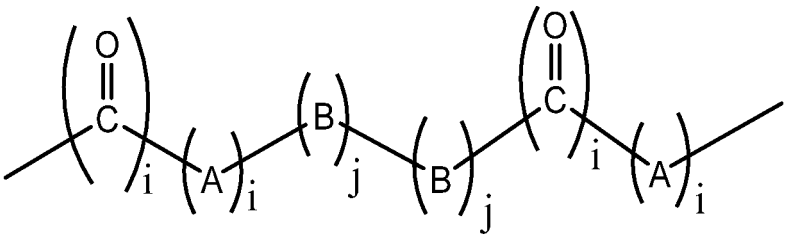

(XI).

[0198] In some embodiments, the linker is enzymatic cleavable. In some embodiments, the linker is not enzymatic cleavable.

[0199] In some embodiments, the linker is presented by the following structure of formula ((II):


(II)

m is 1, 2, 3, 4, 5, or 6, each b independently is 0 or 1, and D is independently represented by structure of formula (III):


(III)

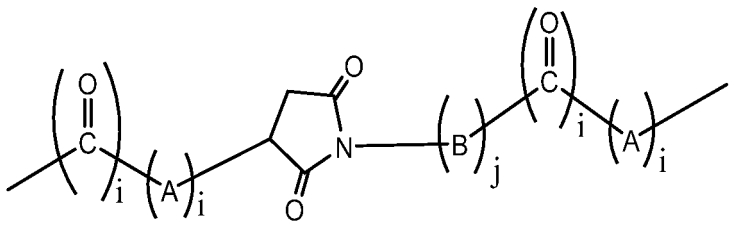
wherein each i independently is 0 or 1;

each j independently is 0, 1, 2, 3, 4, 5, or 6;

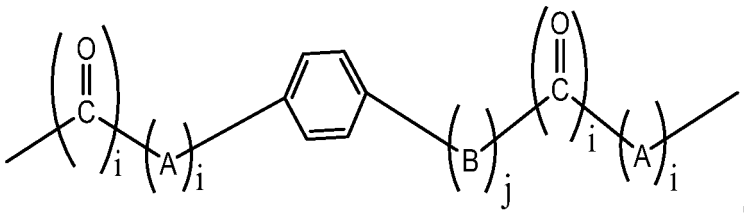
each A independently is S, O, or N-Ra, wherein Ra is hydrogen, alkyl, alkenyl, or alkoxy;

each B independently is alkyl, alkenyl, --O-alkyl--, --alkyl-O--, --S-alkyl--, --alkyl-S--, aryl, heteroaryl, heterocyclyl, or peptide, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --C(O)-R_4, --C(O)-O-R_4, --S-R_4, --$S(O)_2$-R_4, --NHR_4, --NH-alkyl-R_4, halogen, --CN, --NO_2, and –SH, wherein R_4 is alkyl, alkenyl, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

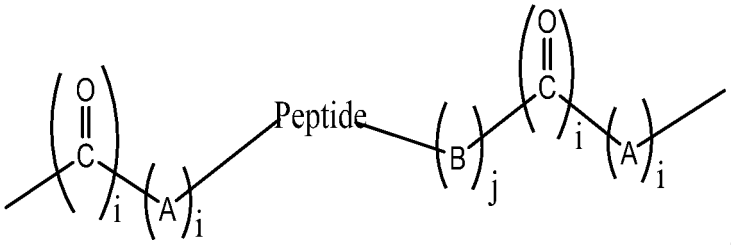
[0200] In some embodiments, the linker is presented by the following strucutres of formula (V)-(VII):



(V)



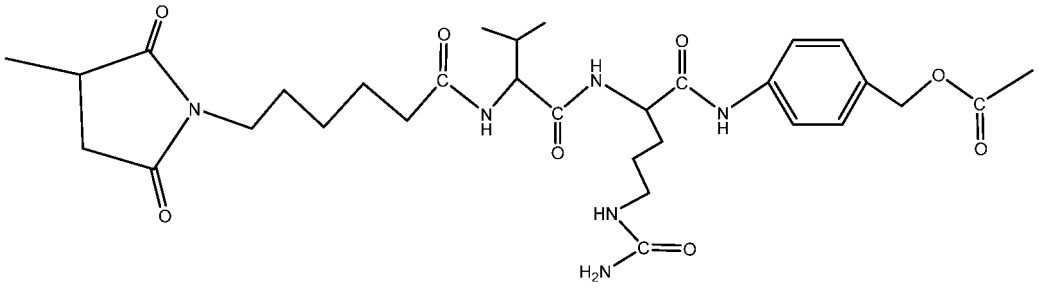
(VI)



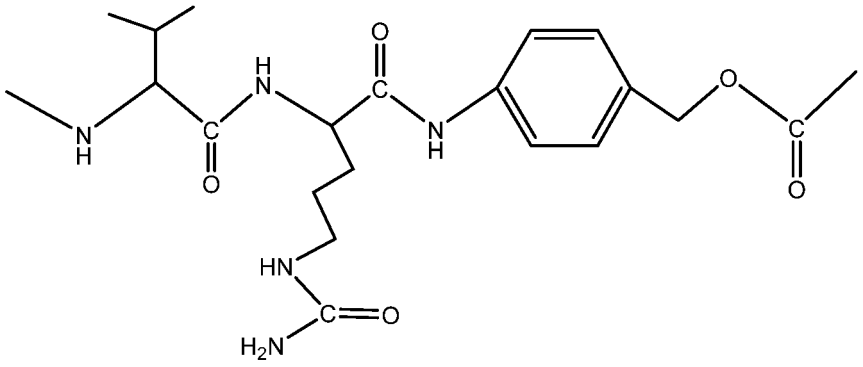
(VII)

A, B, i and j are defined above.

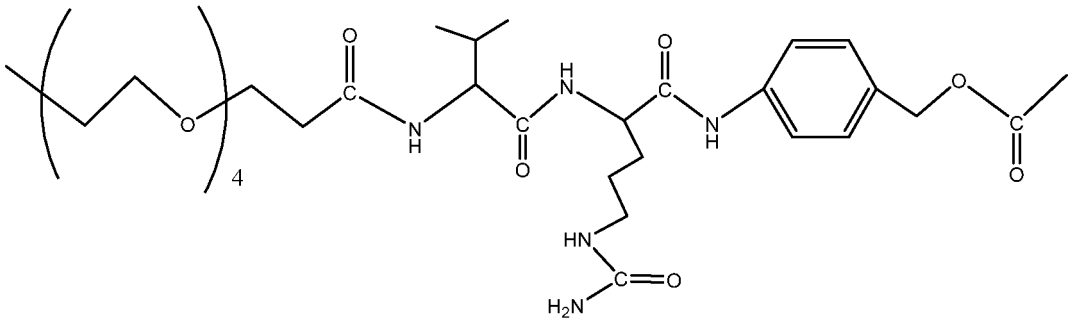
[0201] In some embodiments, the linker is selected from S1, S2, S3, S4, S5, S6, S7, –Gly-Phe-Leu-Gly-, -Ala-Leu-Ala-Leu-, -Phe-Arg-, -Phe-Lys-, -Val-Lys-, -Val-Ala-, or Val-Cit-, wherein S1-S7 are represented by the following structures:



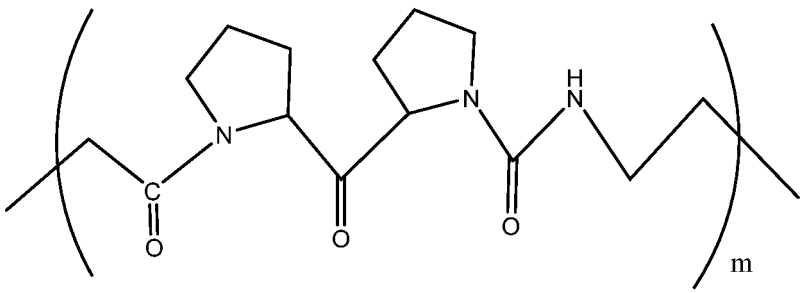
S1



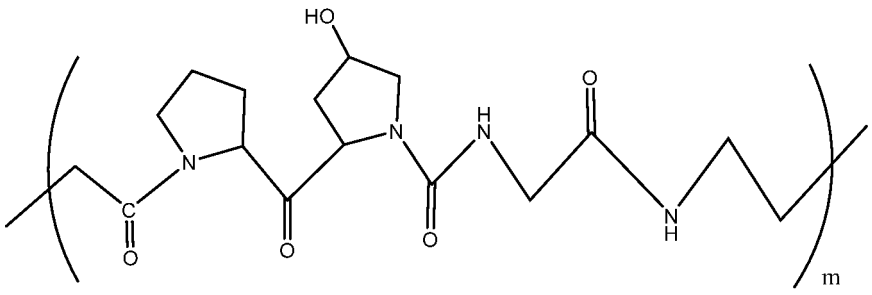
S2



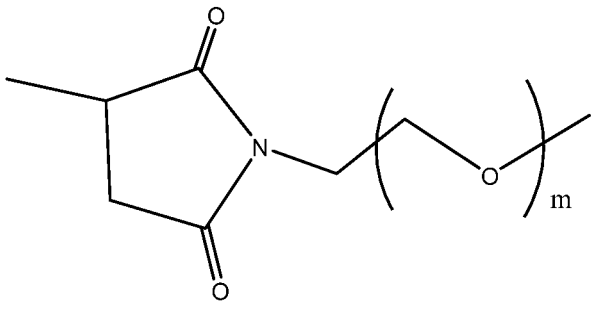
S3



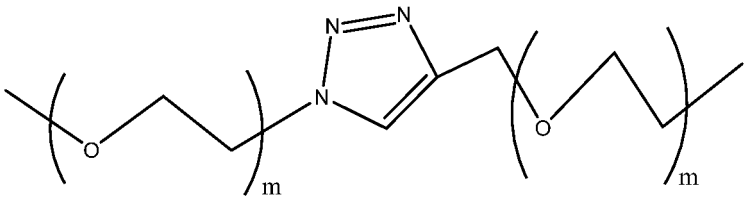
S4



S5



S6

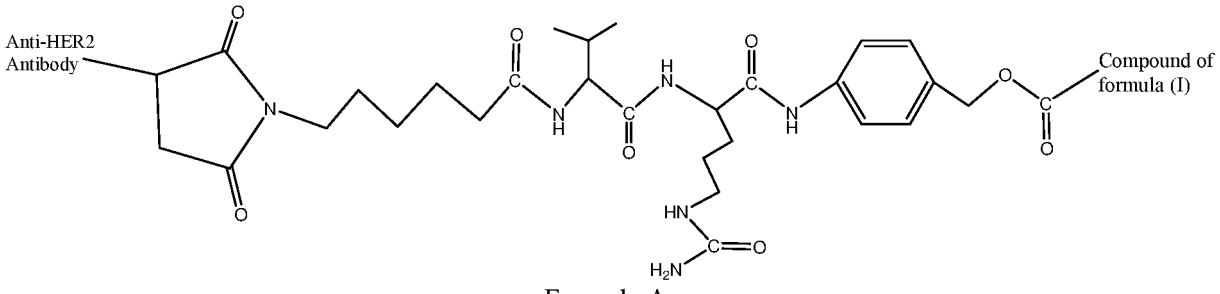


S7,

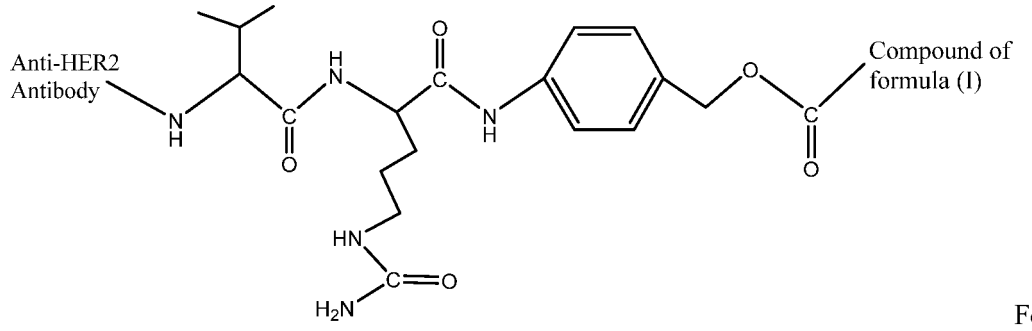
wherein each m is independently 1 to 20. Preferably m is 1 to 3, 1 to 5, 1 to 10, or 2 to 5.

[0202] Accordingly, the present invention provides a compound of formula (Ia), wherein TM is an anti-HER2 antibody; L is selected from S1, S2, S3, S4, S5, S6, S7, –Gly-Phe-Leu-Gly-, -Ala-Leu-Ala-Leu-, -Phe-Arg-, -Phe-Lys-, -Val-Lys-, -Val-Ala-, or Val-Cit-; AM is a compound of formula (I). In one embosiment, TM is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); AM is a compound selected from Table 1, wherein the amine group on the quinoline ring is the point of connection with the linker. Also in one embodiment, TM is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); L is S1, S2 or S3; AM is a compound selected from Table 1, wherein the amine group on the quinoline ring is the point of connection with the linker. Also in one embodiment, TM is trastuzumab (Herceptin); L is S1, S2 or S3; AM is is Resiquimod or Imiquimod, wherein the amine group on the quinoline ring is the point of connection with the linker.

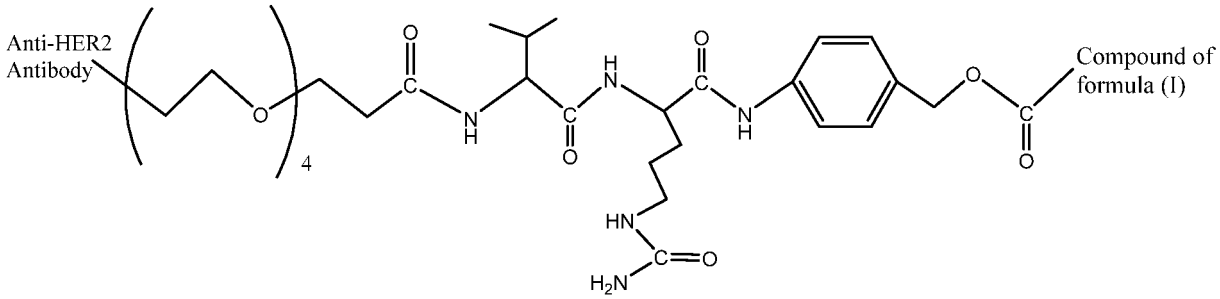
[0203] Yet in another embodiment, the compound of the present invention is represented by the following structures Formula A-C:



Formula A,



Formula B, and

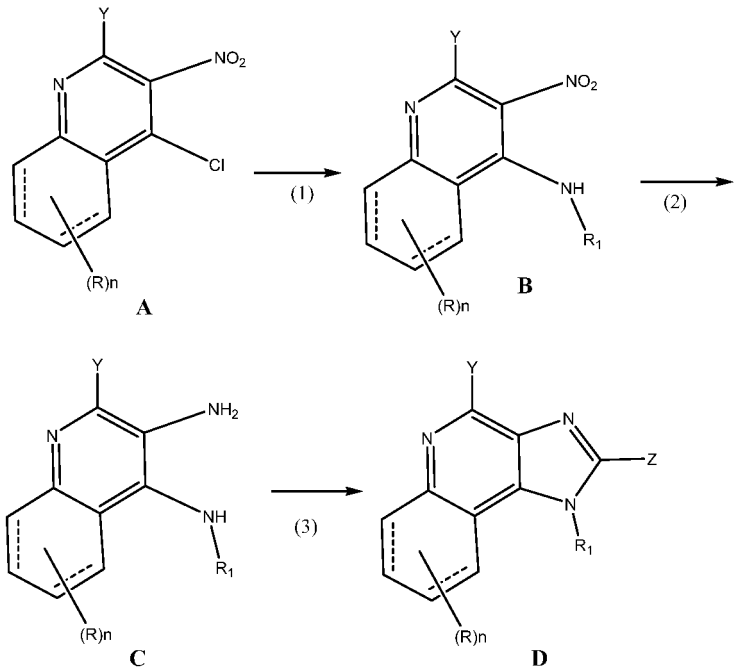


Formula C.

[0204] In one embodiment of Formula A-C, the anti-HER2 antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); the compound of formula (I) is selected from Table 1, wherein the amine group on the quinoline ring is the point of connection with the linker. Also in one embodiment of Formula A-C, the anti-HER2 antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); the compound of formula (I) is Resiquimod or Imiquimod, wherein the amine group on the quinoline ring is the point of connection with the linker. Also in one embodiment of Formula A-C, the anti-HER2 antibody is trastuzumab (Herceptin); AM is is Resiquimod or Imiquimod, wherein the amine group on the quinoline ring is the point of connection with the linker.

Preparation of the Compounds

[0205] In general, the activating moiety represented by the structures of formula (I) can be made using the synthetic procedures outlined below. In step (1), a 4-chloro-3-nitroquinoline of formula A is reacted with an amine of formula R_1NH_2 to provide a 3-nitroquinoline-4-amine of formula B. In step 2, the 3-nitroquinoline-4-amine of formula B is reduced to provide a quinoine-3-4-diamine of formula C. In step 3, the quinoine-3-4-diamine of formula C is reacted with a carboxylic acid or an equivalent thereof to providea 1H-imidazo[4,5c]quinoline of formula D.



[0206] Alternatively, the compounds of formula (I) can be prepared according to synthetic methods described in US6,331,539B1, US6,451,810B1, US7,157,452 and US7,301027B2.

[0207] In another aspect, the compounds of the formula (Ia) can be prepared by using a linker to connect with both a targeting moiety and an activating moiety. The linker uses its reactive sites to bind to the targeting and activating moieties. In some embodiments, the binding is through forming covalent bonds between the linker and the targeting and activating moieties. In some embodiments, the reactive sites are nucleophilic groups. In some embodiments, the reactive sites are electrophilic groups. Useful nucleophilic groups n a linker include but are not limited to, hydrazide, oxime, amino, hydrazine, thiosemicarbazone, hydrazine carboxylate and arylhydrazide groups. Useful electrophilic groups include but are not limited to, maleimide, carbonate and haloacetamide groups.

Pharmaceutical Formulations and Administration

[0208] The present invention further relates to a pharmaceutical formulation comprising a compound of the invention or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable carriers.

[0209] The compounds described herein including pharmaceutically acceptable carriers such as addition salts or hydrates thereof, can be delivered to a patient using a wide variety of routes or modes of administration. Suitable routes of administration include, but inhalation, transdermal, oral, rectal, transmucosal, intestinal and parenteral administration, including intramuscular, subcutaneous and

intravenous injections. Preferably, the compouds of the invention comprising an antibody or antibody fragment as the targeting moiety are administered parenterally, more preferably intravenously.

[0210] As used herein, the terms "administering" or "administration" are intended to encompass all means for directly and indirectly delivering a compound to its intended site of action.

[0211] The compounds described herein, or pharmaceutically acceptable salts and/or hydrates thereof, may be administered singly, in combination with other compounds of the invention, and/or in cocktails combined with other therapeutic agents. Of course, the choice of therapeutic agents that can be co-administered with the compounds of the invention will depend, in part, on the condition being treated.

[0212] For example, when administered to patients suffering from a disease state caused by an organism that relies on an autoinducer, the compounds of the invention can be administered in cocktails containing agents used to treat the pain, infection and other symptoms and side effects commonly associated with the disease. Such agents include, e.g., analgesics, antibiotics, etc.

[0213] When administered to a patient undergoing cancer treatment, the compounds may be administered in cocktails containing anti-cancer agents and/or supplementary potentiating agents. The compounds may also be administered in cocktails containing agents that treat the side-effects of radiation therapy, such as anti-emetics, radiation protectants, etc.

[0214] Supplementary potentiating agents that can be co-administered with the compounds of the invention include,e.g., tricyclic anti-depressant drugs (e.g., imipramine, desipramine, amitriptyline, clomipramine, trimipramine, doxepin, nortriptyline, protriptyline, amoxapine and maprotiline); non-tricyclic and anti-depressant drugs (e.g., sertraline, trazodone and citalopram); Ca+2 antagonists (e.g., verapamil, nifedipine, nitrendipine and caroverine); amphotericin; triparanol analogues (e.g., tamoxifen); antiarrhythmic drugs (e.g., quinidine); antihypertensive drugs (e.g., reserpine); thiol depleters (e.g., buthionine and sulfoximine); and calcium leucovorin.

[0215] The active compound(s) of the invention are administered per se or in the form of a pharmaceutical composition wherein the active compound(s) is in admixture with one or more pharmaceutically acceptable carriers, excipients or diluents. Pharmaceutical compositions for use in accordance with the present invention are typically formulated in a conventional manner using one or more physiologically acceptable carriers comprising excipients and auxiliaries, which facilitate processing of the active compounds into preparations which, can be used pharmaceutically. Proper formulation is dependent upon the route of administration chosen.

[0216] For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

[0217] For oral administration, the compounds can be formulated readily by combining the active compound(s) with pharmaceutically acceptable carriers well known in the art. Such carriers enable the compounds of the invention to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, and suspensions for oral ingestion by a patient to be treated. Pharmaceutical preparations for oral use can be obtained solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxyniethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

[0218] Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures. Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

[0219] Pharmaceutical preparations, which can be used orally, include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added. All formulations for oral administration should be in dosages suitable for such administration.

[0220] For buccal administration, the compositions may take the form of tablets or lozenges formulated in conventional manner.

[0221] For administration by inhalation, the compounds for use according to the present invention are conveniently delivered in the form of an aerosol spray presentation from pressurized packs or a nebulizer, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges

of e.g., gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch.

[0222] The compounds may be formulated for parenteral administration by injection, e.g., by bolus injection or continuous infusion. Injection is a preferred method of administration for the compositions of the current invention. Formulations for injection may be presented in unit dosage form, e.g., in ampoules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

[0223] Pharmaceutical formulations for parenteral administration include aqueous solutions of the active compounds in water-soluble form. Additionally, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances, which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents, which increase the solubility of the compounds to allow for the preparation of highly, concentrated solutions. For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks's solution, Ringer's solution, or physiological saline buffer.

[0224] Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g., sterile pyrogen-free water, before use.

[0225] The compounds may also be formulated in rectal compositions such as suppositories or retention enemas,e.g., containing conventional suppository bases such as cocoa butter or other glycerides.

[0226] In addition to the formulations described previously, the compounds may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation or transcutaneous delivery (e.g., subcutaneously or intramuscularly), intramuscular injection or a transdermal patch. Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials (e.g., as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

[0227] The pharmaceutical compositions also may comprise suitable solid or gel phase carriers or excipients. Examples of such carriers or excipients include calcium carbonate, calcium phosate, various sugars, starches, cellulose derivatives, gelatin, and polymers such as polyethylene glycols.

[0228] A preferred pharmaceutical composition is a composition formulated for injection such as intravenous injection and includes about 0.01 % to about 100% by weight of the compound of the present invention, based upon 100% weight of total pharmaceutical composition. The drug-ligand conjugate may be an antibody-cytotoxin conjugatewhere the antibody has been selected to target a particular cancer.

[0229] In some embodiments, the pharmaceutical composition of the present invention further comprises an additional therapeutic agent.

[0230] In some embodiments, the additional therapeutic agent is an anticancer agent.

[0231] In some embodiments, the additional anticancer agent is selected from an antimetabolite, an inhibitor of topoisomerase I and II, an alkylating agent, a microtubule inhibitor, an antiandrogen agent, a GNRh modulator or mixtures thereof.

[0232] In some embodiments, the additional therapeutic agent is a chemotherapeutic agent.

[0233] By "chemotherapeutic agent" herein is meant a chemical compound useful in the treatment of cancer. Examples are but not limited to: Gemcitabine, Irinotecan, Doxorubicin, 5-Fluorouracil, Cytosine arabinoside ("Ara-C"), Cyclophosphamide, Thiotepa, Busulfan, Cytoxin, TAXOL, Methotrexate, Cisplatin, Melphalan, Vinblastine and Carboplatin.

[0234] In some embodiments, the second chemotherapeutic agent is selected from the group consisting of tamoxifen, raloxifene, anastrozole, exemestane, letrozole, imatanib, paclitaxel, cyclophosphamide, lovastatin, minosine, gemcitabine, cytarabine, 5- fluorouracil, methotrexate, docetaxel, goserelin, vincristine, vinblastine,nocodazole, teniposide etoposide, gemcitabine, epothilone, vinorelbine, camptothecin, daunorubicin, actinomycin D, mitoxantrone, acridine, doxorubicin, epirubicin, or idarubicin.

Kits

[0235] In another aspect, the present invention provides kits containing one or more of the compounds or compositions of the invention and directions for using the compound or composition. In an exemplary embodiment, the invention provides a kit for conjugating a linker arm of the invention to another molecule. The kit includes the linker, and directions for attaching the linker to a particular functional group. The kit may also include one or more of a cytotoxic drug, a targeting agent, a detectable label, pharmaceutical salts or buffers. The kit may also include a container and optionally one or more vial, test tube, flask, bottle, or syringe. Other formats for kits will be apparent to those of skill in the art and are within the scope of the present invention.

Medical Use

[0236] The compounds of the present invention comprise an anti-HER2 antibody, a linker and an activating moeity represent by formula (I), such asbeing TLR7/8 agonists. The present invention provides a new class of chemical cytotoxic agent in the context of antibody drug conjugates—TLR7/8 agonsits that modulate immune response, subsequently activate dendritic and natural killer cells and promote anti-tumor activities. It has been surprisingly discovered that the compounds of the present invention show superior anti-tumor activities. In a typical treatment with the compounds of the present invention, the antibody part acting like a missle, delivers the activating moiety part to the HER2 expressing tumor/cancer cells where the activating moeity part acts against the tumor/cancer cells directly or indirectly. Such treatment benefits from a reduced toxic side-effect profile and an improved pharmacokinetic profile. Furthermore, the slow release of activating moeity from the carrier antibody results in sustained high intratumoral levels and lower plasma concentrations of the activating moeity .

[0237] Accordingly, in another aspect, the present invention provides a method of inhibiting proliferation of a HER2 positive tumor/cancer call comprising admnisgtering to said tumor cell the compounds of the present invention. In some embodiments, the tumor can be metastasis or non-metastasis.

[0238] By "cancer" or "tumor" herein is meant the pathological condition in humans that is characterized by unregulated cell proliferation. Examples include but are not limited to: carcinoma, lymphoma, blastoma, and leukemia. More particular examples of cancers include but are not limited to: lung (small cell and non-small cell), breast, prostate, carcinoid, bladder, gastric, pancreatic, liver (hepatocellular), hepatoblastoma, colorectal, head and neck squamous cell carcinoma, esophageal, ovarian, cervical, endometrial, mesothelioma, melanoma, sarcoma, osteosarcoma, liposarcoma, thyroid, desmoids, chronic myelocytic leukemia (AML), and chronic myelocytic leukemia (CML).

[0239] By "inhibiting" or "treating" or "treatment" herein is meant to reduction, therapeutic treatment and prophylactic or preventative treatment, wherein the objective is to reduce or prevent the aimed pathologic disorder or condition. In one example, following administering of a compound of the present invention, a cancer patient may experience a reduction in tumor size. "Treatment" or "treating" includes (1) inhibiting a disease in a subject experiencing or displaying the pathology or symptoms of the disease, (2) ameliorating a disease in a subject that is experiencing or displaying the pathology or symptoms of the disease, and/or (3) affecting any measurable decrease in a disease in a subject or patient that is experiencing or displaying the pathology or symptoms of the disease. To the extent a compound of the present invention may prevent growth and/or kill cancer cells, it may be cytostatic and/or cytotoxic.

[0240] By "therapeutically effective amount" herein is meant an amount of a compound provided herein effective to "treat" a disorder in a subject or mammal. In the case of cancer, the therapeutically effective amount of the drug may reduce the number of cancer cells, reduce the tumor size, inhibit cancer cell

infiltration into peripheral organs, inhibit tumor metastasis, inhibit tumor growth to certain extent, and/or relieve one or more of the symptoms associated with the cancer to some extent.

[0241] In another aspect, the present invention provides a method of treating a HER2 positive tumor/cancer in a subject comprising administering to the subject a therapeutically effective amount of the compounds of the present invention. In some embodiments, the tumor or cancer can be at any stage, e.g., early or advanced, such as a stage I, II, III, IV or V tumor or cancer. In some embodiments, the tumor or cancer can be metastatic or non-metastatic. In the context of metastasis, the methods of the present invention can reduce or inhibit metastasis of a primary tumor or cancer to other sites, or the formation or establishment of metastatic tumors or cancers at other sites distal from the primary tumor or cancer therapy. Thus, the methods of the present invention include, among other things, 1) reducing or inhibiting growth, proliferation, mobility or invasiveness of tumor or cancer cells that potentially or do develop metastases (e.g., disseminated tumor cells, DTC); 2) reducing or inhibiting formation or establishment of metastases arising from a primary tumor or cancer to one or more other sites, locations or regions distinct from the primary tumor or cancer; 3) reducing or inhibiting growth or proliferation of a metastasis at one or more other sites, locations or regions distinct from the primary tumor or cancer after a metastasis has formed or has been established; and 4) reducing or inhibiting formation or establishment of additional metastasis after the metastasis has been formed or established.

[0242] In some embodiments, the tumor or cancer is solid or liquid cel mass. A "solid" tumor refers to cancer, neoplasia or metastasis that typically aggregates together and forms a mass. Specific non-limiting examples include breast, ovarian, uterine, cervical, stomach, lung, gastric, colon, bladder, glial, and endometrial tumors/cancers, etc.. A "liquid tumor," which refers to neoplasia that is dispersed or is diffuse in nature, as they do not typically form a solid mass. Particular examples include neoplasia of the reticuloendothelial or hematopoietic system, such as lymphomas, myelomas and leukemias. Non-limiting examples of leukemias include acute and chronic lymphoblastic, myeolblastic and multiple myeloma. Typically, such diseases arise from poorly differentiated acute leukemias, e.g., erythroblastic leukemia and acute megakaryoblastic leukemia. Specific myeloid disorders include, but are not limited to, acute promyeloid leukemia (APML), acute myelogenous leukemia (AML) and chronic myelogenous leukemia (CML). Lymphoid malignancies include, but are not limited to, acute lymphoblastic leukemia (ALL), which includes B-lineage ALL (B-ALL) and T-lineage ALL (T-ALL), chronic lymphocytic leukemia (CLL), prolymphocyte leukemia (PLL), hairy cell leukemia (HLL) and Waldenstroem's macroglobulinemia (WM). Specific malignant lymphomas include, non-Hodgkin lymphoma and variants, peripheral T cell lymphomas, adult T cell leukemia/lymphoma (ATL), cutaneous T-cell lymphoma (CTCL), large granular lymphocytic leukemia (LGF), Hodgkin's disease and Reed-Sternberg disease.

[0243] In some embodiments, the methods of the present invention can be practiced with other treatments or therapies (e.g., surgical resection, radiotherapy, ionizing or chemical radiation therapy, chemotherapy, immunotherapy, local or regional thermal (hyperthermia) therapy, or vaccination). Such other treatments or therapies can be administered prior to, substantially contemporaneously with (separately or in a mixture), or following administration of the compounds of the present invention.

[0244] In some embodiments, the methods of the present invention comprise administering a therapeutically effective amount of a compound of the present invention in combination with an additional therapeutic agent. In some embodiments, the additional therapeutic agent is an anticancer/antitumor agent. In some embodiments, the additional therapeutic agent is an antimetabolite, an inhibitor of topoisomerase I and II, an alkylating agent, a microtubule inhibitor, an antiandrogen agent, a GNRh modulator or mixtures thereof. In some embodiemnts, the additional therapeutic agent is selected from the group consisting of tamoxifen, raloxifene, anastrozole, exemestane, letrozole, imatanib, paclitaxel, cyclophosphamide, lovastatin, minosine, gemcitabine, cytarabine, 5- fluorouracil, methotrexate, docetaxel, goserelin, vincristine, vinblastine,nocodazole, teniposide etoposide, gemcitabine, epothilone, vinorelbine, camptothecin, daunorubicin, actinomycin D, mitoxantrone, acridine, doxorubicin, epirubicin, or idarubicin.

[0245] Administration "in combination with" one or more additional therapeutic agents includes simultaneous (concurrent) and consecutive administration in any order. As used herein, the term "pharmaceutical combination" refers to a product obtained from mixing or combining active ingredients, and includes both fixed and non-fixed combinations of the active ingredients. The term "fixed combination" means that the active ingredients, e.g. a compound of Formula (1) and a co-agent, are both administered to a patient simultaneously in the form of a single entity or dosage. The term "non-fixed combination" means that the active ingredients, e.g. a compound of Formula (1) and a co-agent, are both administered to a patient as separate entities either simultaneously, concurrently or sequentially with no specific time limits, wherein such administration provides therapeutically effective levels of the active ingredients in the body of the patient. The latter also applies to cocktail therapy, e.g. the administration of three or more active ingredients.

[0246] In some embodiments, the diseases condition is tumor or cancer. In some embodiments, the cancer or tumor is selected from stomach, colon, rectal, liver, pancreatic, lung, breast, cervix uteri, corpus uteri, ovary, testis, bladder, renal, brain/CNS, head and neck, throat, Hodgkin's disease, non-Hodgkin's lymphoma, multiple myeloma, leukemia, melanoma, non-melanoma skin cancer, acute lymphocytic leukemia, acute myelogenous leukemia, Ewing's sarcoma, small cell lung cancer, choriocarcinoma,

rhabdomyosarcoma, Wilms' tumor, neuroblastoma, hairy cell leukemia, mouth/pharynx, oesophagus, larynx, kidney cancer or lymphoma.

[0247] In some embodiments, the disease condition comprises abnormal cell proliferation, such as a pre-cancerous lesion.

[0248] The current invention is particularly useful for the treatment of cancer and for the inhibition of the multiplication of a tumor cell or cancer cell in an animal. Cancer, or a precancerous condition, includes a tumor, metastasis, or any disease or disorder characterized by uncontrolled cell growth, can be treated or prevented by administration the drug-ligand complex of the current invention. The compound delivers the activating moiety to a tumor cell or cancer cell. In some embodiments, the targeting moiety specifically binds to or associates with a cancer-cell or a tumor-cell-associated antigen. Because of its close proximity to the ligand, after being internalized, the activating moiety can be taken up inside a tumor cell or cancer cell through, for example, receptor-mediated endocytosis. The antigen can be attached to a tumor cell or cancer cell or can be an extracellular matrix protein associated with the tumor cell or cancer cell. Once inside the cell, the linker is hydrolytically or enzymatically cleaved by a tumor-cell or cancer-cell-associated proteases, thereby releasing the activating moiety. The released activating moiety is then free to diffuse and induce or enhance immune activity of immune cells or tumor cells. In an alternative embodiment, the activating moiety is cleaved from the compound tumor microenvironment, and the drug subsequently penetrates the cell.

[0249] Representative examples of precancerous conditions that may be targeted by the compounds of the present invention, include: metaplasia, hyperplysia, dysplasia, colorectal polyps, actinic ketatosis, actinic cheilitis, human papillomaviruses, leukoplakia, lychen planus and Bowen's disease.

[0250] Representative examples of cancers or tumors that may be targeted by compounds of the present invention include: lung cancer, colon cancer, prostate cancer, lymphoma, melanoma, breast cancer, ovarian cancer, testicular cancer, CNS cancer, renal cancer, kidney cancer, pancreatic cancer, stomach cancer, oral cancer, nasal cancer, cervical cancer and leukemia. It will be readily apparent to the ordinarily skilled artisan that the particular targeting moiety used in the compound can be chosen such that it targets the activating moiety to the tumor tissue to be treated with the drug (i.e., a targeting agent specific for a tumor-specific antigen is chosen). Examples of such targeting moiety are well known in the art, examples of which include anti-Her2 for treatment of breast cancer, anti-CD20 for treatment of lymphoma, anti-PSMA for treatment of prostate cancer and anti-CD30 for treatment of lymphomas, including non-Hodgkin's lymphoma.

[0251] In some embodiments, the abnormal proliferation is of cancer cells.

[0252] In some embodiments, the cancer is selected from the group consisting of: breast cancer, colorectal cancer, diffuse large B-cell lymphoma, endometrial cancer, follicular lymphoma, gastric cancer, glioblastoma, head and neck cancer, hepatocellular cancer, lung cancer, melanoma, multiple myeloma, ovarian cancer, pancreatic cancer, prostate cancer, and renal cell carcinoma.

[0253] In some embodiments, the present invention provides a compound for use in killing a cell. The compound is administered to the cell in an amount sufficient to kill said cell. In an exemplary embodiment, the compound is administered to a subject bearing the cell. In a further exemplary embodiment, the administration serves to retard or stop the growth of a tumor that includes the cell (e.g., the cell can be a tumor cell). For the administration to retard the growth, the rate of growth of the cell should be at least 10% less than the rate of growth before administration. Preferably, the rate of growth will be retarded at least 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, or completely stopped.

[0254] Additionally, the present invention provides a compound or a pharmaceutical composition of the present invention for use as a medicament. The present invention also provides a compound or a pharmaceutical composition for killing, inhibiting or delaying proliferation of a tumor or cancer cell, or for treating a disease wherein TLR7 and/or TLR8 are implicated.

Effective Dosages

[0255] Pharmaceutical compositions suitable for use with the present invention include compositions wherein the active ingredient is contained in a therapeutically effective amount, i.e., in an amount effective to achieve its intended purpose. The actual amount effective for a particular application will depend, inter alia, on the condition being treated. Determination of an effective amount is well within the capabilities of those skilled in the art, especially in light of the detailed disclosure herein.

[0256] For any compound described herein, the therapeutically effective amount can be initially determined from cell culture assays. Target plasma concentrations will be those concentrations of active compound(s) that are capable of inhibition cell growth or division. In preferred embodiments, the cellular activity is at least 25% inhibited. Target plasma concentrations of active compound(s) that are capable of inducing at least about 30%, 50%, 75%, or even 90% or higher inhibition of cellular activity are presently preferred. The percentage of inhibition of cellular activity in the patient can be monitored to assess the appropriateness of the plasma drug concentration achieved, and the dosage can be adjusted upwards or downwards to achieve the desired percentage of inhibition.

[0257] As is well known in the art, therapeutically effective amounts for use in humans can also be determined from animal models. For example, a dose for humans can be formulated to achieve a circulating concentration that has been found to be effective in animals. The dosage in humans can be

adjusted by monitoring cellular inhibition and adjusting the dosage upwards or downwards, as described above.

[0258] A therapeutically effective dose can also be determined from human data for compounds which are known to exhibit similar pharmacological activities. The applied dose can be adjusted based on the relative bioavailability and potency of the administered compound as compared with the known compound.

[0259] Adjusting the dose to achieve maximal efficacy in humans based on the methods described above and other methods as are well-known in the art is well within the capabilities of the ordinarily skilled artisan.

[0260] In the case of local administration, the systemic circulating concentration of administered compound will not be of particular importance. In such instances, the compound is administered so as to achieve a concentration at the local area effective to achieve the intended result.

[0261] For use in the prophylaxis and/or treatment of diseases related to abnormal cellular proliferation, a circulating concentration of administered compound of about 0.001 μM to 20 μM is preferred, with about 0.01 μM to 5 μM being preferred.

[0262] Patient doses for oral administration of the compounds described herein, typically range from about 1 mg/day to about 10,000 mg/day, more typically from about 10 mg/day to about 1,000 mg/day, and most typically from about 50 mg/day to about 500 mg/day. Stated in terms of patient body weight, typical dosages range from about 0.01 to about 150 mg/kg/day, more typically from about 0.1 to about 15 mg/kg/day, and most typically from about 1 to about 10 mg/kg/day, , for example 5 mg/kg/day or 3 mg/kg/day.

[0263] In at least some embodiments, patient doses that retard or inhibit tumor growth can be 1 μmol/kg/day or less. For example, the patient doses can be 0.9, 0.6, 0.5, 0.45, 0.3, 0.2, 0.15, or 0.1 μmol/kg/day or less (referring to moles of the drug). Preferably, the antibody with drug conjugates retards growth of the tumor when administered in the daily dosage amount over a period of at least five days.

[0264] For other modes of administration, dosage amount and interval can be adjusted individually to provide plasma levels of the administered compound effective for the particular clinical indication being treated. For example, in one embodiment, a compound according to the invention can be administered in relatively high concentrations multiple times per day. Alternatively, it may be more desirable to administer a compound of the invention at minimal effective concentrations and to use a less frequent administration regimen. This will provide a therapeutic regimen that is commensurate with the severity of the individual's disease.

[0265] Utilizing the teachings provided herein, an effective therapeutic treatment regimen can be planned which does not cause substantial toxicity and yet is entirely effective to treat the clinical symptoms demonstrated by the particular patient. This planning should involve the careful choice of active compound by considering factors such as compound potency, relative bioavailability, patient body weight, presence and severity of adverse side effects, preferred mode of administration and the toxicity profile of the selected agent.

[0266] While preferred embodiments of the present invention 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 invention. It should be understood that various alternatives to the embodiments of the invention described herein may be employed in practicing the invention. It is intended that the following claims define the scope of the invention and that methods and structures within the scope of these claims and their equivalents be covered thereby.

EXAMPLES

[0267] The present invention is further exemplified, but not limited, by the following and Examples that illustrate the preparation of the compounds of the invention.

Example 1

Generation of her2 transfected L cell lines

[0268] **Reagents:** L cell was from ATCC (Manassas, VA; Cat No CRL2648), Her2/pEZ-Lv105 cDNA was purchased from GeneCopoeia, (Cat No Z2866), Glucose DMEM, L-glutamine, Lipofectamine 2000 (Invitrogen; Carlsbad, CA).

[0269] To make cell lines for screening the conjugated Trastuzumab, L cell expressing the tagged Her2 were generated. The Her2/pEZ-Lv105 construct was transfected into L cells (that grown in high glucose DMEM+10% FBS+2 mM L-glutamine cells) by the standard Lipofectamine 2000 protocol.

Binding Analysis (FACS Analysis)

[0270] To determine binding capability of conjugated Trastuzumab, FACS analysis of L cells expressing human her2 were performance. Briefly, approximately 10^6 -cells of L cells with transient transfected her2 in 100 µl were incubated with varying amounts of Trastuzumab conjugated antibody, PBS or secondary antibody alone or irrelevant huIgG were used as a negative control. After washing, cells were re-suspended in FACS buffer and incubated 30 min at room temperature with 20 µL mouse anti-human IgG conjugated to phycoerythrin (Mu anti-Human-PE) secondary antibody in a 100 µL reaction volume. After

washing, cells were fixed in 200 μL of 2% paraformaldehyde/PBS and flow cytometery performed. The same procedure was used for irrelevant human IgG antibody as an isotype control to set baseline PMT per cell line. Flow cytomery was performed on a BD FACSCalibur® and geometric mean fluorescence intensity was recorded for each sample. Recorded data was analyzed using FlowJo software. Results are shown in Figure 1.

Generation of antibody with toll like receptor ligand conjugates

[0271] **Reagents:** Trastuzumab (Roche/Genentech Corporation, South San Francisco, CA), ; MC-val-cit-PAB linked with resiquimod (MC-vc-PAB-TLRL) or MC- linked with resiquimod (MC-TLRL) (Synthesized in Contract Research Organization, China), Sodium borate, Sodium chloride, Dithiothreitol (DTT), Sephadex G25, DTPA, DTNB, and Maleimidocaproyl-monomethyl (Sigma-Aldrich, Milwaukee, Wisconsin).

[0272] The drugs used for generation of antibody TLR ligand conjugates included Trastuzumab and resiquimod (TLRL). The linkers used for generation of the TLAC were cleavable linker MC-vc-PAB or non cleavable linker MC.

Preparation of Trastuzumab MC-TLRL

[0273] Trastuzumab was purified from HERCEPTIN® by buffer-exchange at 20 mg/mL, and antibody dissolved in 500 mM sodium borate and 500 mM sodium chloride at pH 8.0 is treated with an excess of 100 mM dithiothreitol (DTT). After incubation at 37°C for about 30 minutes, the buffer is exchanged by elution over Sephadex G25 resin and eluted with PBS with 1 mM DTPA. The thiol/Ab value is checked by determining the reduced antibody concentration from the absorbance at 280 nm of the solution and the thiol concentration by reaction with DTNB (Sigma-Aldrich, Milwaukee, Wisconsin) and determination of the absorbance at 412 nm. The reduced antibody dissolved in PBS is chilled on ice. The drug linker reagent, Mc- linked with resiquimod, dissolved in DMSO, is diluted in acetonitrile and water at known concentration, and added to the chilled reduced antibody in PBS. After about one hour, an excess of maleimide is added to quench the reaction and cap any unreacted antibody thiol groups. In some case, coupling to lysines of immunoglobulin with standard procedure was performed. The reaction mixture is concentrated by centrifugal ultrafiltration and conjugated antibody is purified and desalted by elution through G25 resin in PBS, filtered through 0.2 μm filters under sterile conditions, and frozen for storage.

Preparation of Trastuzumab MC-vc-TLRL

[0274] Antibodies were linked to TLRL through the cysteine by Maleimidocaproyl-valine-citruline (vc)-p-aminobenzyloxycarbonyl (MC-vc-PAB). The MC-vc-PAB linker is cleavable by intercellular proteases such as cathepsin B and when cleaved, releases free drug (Doronina et al., Nat. Biotechnol., 21: 778-784

(2003)) while the MC linker is resistant to cleavage by intracellular proteases. Purified Trastuzumab was dissolved in 500 mM sodium borate and 500 mM sodium chloride at pH 8.0 and further treated with an excess of 100 MM dithiothreitol (DTT). After incubation at 37°C for about 30 minutes, the buffer is exchanged by elution over Sephadex G25 resin and eluted with PBS with 1 mM DTPA. The thiol/Ab value was checked by determining the reduced antibody concentration from the absorbance at 280 nm of the solution and the thiol concentration by reaction with DTNB (Sigma-Aldrich, Milwaukee, Wisconsin) and determination of the absorbance at 412 nm (extinction coefficient =13600 cm^{-1} M^{-1}). The reduced antibody dissolved in PBS was chilled on ice. The MC-val-cit-PAB-PNP linked with resiquimod in DMSO, was dissolved in acetonitrile and water, and added to the chilled reduced antibody in PBS. After one hour incubation, an excess of maleimide was added to quench the reaction and cap any unreacted antibody thiol groups. In some case, coupling to lysines of immunoglobulin with standard procedure was performed. The reaction mixture was concentrated by centrifugal ultrafiltration and the antibody drug conjugate, was purified and desalted by elution through G25 resin in PBS, filtered through 0.2 μm filters under sterile conditions, and frozen for storage.

[0275] Typically a conjugation reaction of antibody with MC-TLRL or MC-vc-TLRL results in a heterogeneous mixture comprising antibodies different numbers of attached, conjugated TLRL drugs, i.e. drug loading where drug is a distribution from 1 to about 8. Thus, antibody MC-TLRL, or antibody MC-vc-TLRL, includes isolated, purified species molecules as well as mixtures of average drug loading from 1 to 8. By control process, drug loads were in the range of 3-5. The average number of TLRL drug moieties per antibody antibody in compound preparations from conjugation reactions may be characterized by conventional means such as mass spectroscopy, ELISA assay, electrophoresis, and HPLC. The quantitative distribution of antibody MC-TLRL or antibody MC-vc-TLRL in terms of drug may also be determined. By ELISA, the averaged value of drug payload number in a particular preparation of antibody with TLRL conjugation may be determined (Hamblett et al (2004) Clinical Cancer Res. 10:7063-7070; Sanderson et al (2005) Clinical Cancer Res. 11:843-852). However, the distribution of drug values is not discernible by the antibody-antigen binding and detection limitation of ELISA. Also, ELISA assay for detection of antibody-drug conjugates does not determine where the drug moieties are attached to the antibody, such as the heavy chain or light chain fragments, or the particular amino acid residues. In some instances, separation, purification, and characterization of homogeneous Trastuzumab MC-TLRL or Trastuzumab MC-vc-TLRL where drug is a certain value from Trastuzumab MC-TLRL or Trastuzumab MC-vc-TLRL with other drug loadings may be achieved by means such as reverse phase HPLC or electrophoresis.

Antibody-dependent cell-mediated cytotoxicity assay (ADCC)

[0276] **Reagents:** SKBR3 cells (ATCC, catalog #HTB-30), McCoy's 5A (Invitrogen, Cat No.22400, Lot No.747809); RPMI-1640 (Invitrogen, Cat No.11835, Lot No.764956); FCS (Hyclone, Cat No.SH30084.03, Lot No.GRH0054); Ficoll-Hypaque (Amersham Biosciences, Piscataway, NJ; Cat No 17-1440-02), 96 well plate (Costar, Cat No.3599, Cat No.3916); Trypan blue (Invitrogen Cat No 15250-061); LDH kit (Promega, Cat No.G7891), ELISA reader MD5 (Molecule device), Humanized antibody Herceptin® (Genentech Corporation, South San Francisco, CA; trade name Trastuzumab) was reconstituted with water to make a 10 mg/ml stock solution before use.

[0277] To determine if Trastuzumab conjugated with a high payload of TLRL would still have effector function, antibody-dependent cell-mediated cytotoxicity (ADCC) activities of the several different form of conjugation were tested and compared with the ADCC activity of Trastuzumab reference material which has been shown to have significantly enhanced ADCC activity. ADCC assays were carried out using peripheral blood mononuclear cells (PBMCs) from healthy donors as effector cells, and a human SKBR3 cell line as target cells. On first day, 1 x 10^4/100uL of SKBR3 cells was seeded into the 96 well plates followed by incubation at 37°C with 5% CO2 for 48 hours.

[0278] On third days, human PBMC was prepared fresh from human blood obtained from healthy volunteer donors. Human venous blood samples were collected in ammonium citrate (ACD-A) tubes. The tubes were inverted several times, and whole blood was transferred into one 50 ml conical tube. Blood was diluted 1:3 with PBS in 2% FBS. Ficoll-Hypaque was dispensed slowly underneath blood/PBS mixture. Samples were centrifuged at room temperature at 2400 rpm for 30 minutes before removing the upper layer (plasma/PBS) by aspiration. Buffy coats were collected with sterile pipettes and pooled into a 50 ml conical tube. If visible clumps of WBC remained below the buffy coat, all of the WBC material was collected, with care not to remove excess Ficoll-Hypaque. Sterile PBS-2% FBS was added and mixed by inversion. The diluted PBMC suspensions was centrifuged at 250 x g at room temp for 20 minutes, and the cell pellet was collected. The PBMC pellet was suspended in RPMI-1640 medium with 2% heat-inactivated FBS, washed and the number of viable cells was checked by Trypan blue exclusion. 1.2 x 10^7 cells/mL in density were prepared and ready to be used. Meanwhile the final concentrations of antibodies the ranged from 12, 4, 1.2, 0.4, 0.12, 0.04, 0.012, 0.004, to 0μg/mL were well prepared in RPMI-1640.

[0279] Serial dilutions of test and control antibodies then were added to wells containing the target cells, PBMC effector cells in 50 μL of RPMI-1640 medium were added to each well with effector:target cell ratio of 60:1 and incubated for an additional 17 hours. The plates were centrifuged at the end of incubation and the supernatants were assayed for lactate dehydrogenase (LDH) activity using a LDH measuring kit. Cell lysis was quantified through absorbance at 490 nm using a microplate reader. The absorbance of wells containing only the target cells served as the control for background, whereas

wells containing target cells lysed with Triton-X100 provided maximum signal available. Antibody-independent cellular cytotoxicity was measured in wells containing target and effector cells without the addition of antibody. Cytotoxicity was calculated according to the following equation:

% Cytotoxicity = 100% x [A490nm (Sample) – A490 (target cell) – A490 (effect cells]/[A490nm (Target cells lysed) –A490nm (target cells)]

[0280] The mean ADCC values from duplicates of sample dilutions were plotted against the antibody concentration, and the EC50 values and the maximum extent of ADCC (%) were generated by fitting into Prism 5 (GraphPad).

Enrichment of human dendritic cells (DCs) from PBMC

[0281] Human PBMC was prepared from Buffy coats obtained from healthy volunteer donors by Ficoll centrifugation. Dendritic cells were enriched by using negative depletion with magnetic beads (Miltenyi Biotec) with mixture of anti-CD3, CD19, CD20, CD14, and CD16 antibodies from human PBMC. The enrichment of DCs was stained with goat anti-mouse FITC (lineages), HLA-DR- APCCy7, CD123-BV421 and CD11C-APC. The stained cells were analyzed on BD LSR Fortessa. The anti- CD3, CD4, CD11C, CD19, CD14, CD16, CD123 monoclonal antibody were purchased from BD Biosciences or Biolgend.

[0282] Figure 3(a) shows the percentages of DCs before and post enrichment. The numbers in upper two plots represent the percentages of DCs (HLA-DR+Lin-) of total cells before and after lineage depletion. The numbers in lower plots represent percentages of mDC (CD11C +CD123-) and pDC (CD123+CD11C-) of total DCs before and after lineage depletion.

Stimulation of enriched human DCs and cytokines expression

[0283] 1-2 x 10^5 enriched DCs were plated in a 96-well plate in 100 μl media, 100 μl diluted stimulators were add to the plate and cultured for 20-22h in 37°C incubator. The supernatant were collected and human IFN-α, IL-6, IL-12(p70) and TNF-α were analyzed by ELISA (Mabtech AB).

Statistical analysis

[0284] The significance of all comparisons was calculated using a Student's two-tailed t test assuming unequal variance between mock and sample groups, and results considered significant when $p<0.05$. Correlations between parameters were assessed using Spearman's rank correlation test, P values <0.05 were consider to be statistically significant.

In vivo tumor cell killing assay using Trastuzumab TLRL conjugates

[0285] For development of patient-derived gastric carcinoma xenograft (PDX) mouse models, female Balb/c nude mice (from SLAC, Shanghai, China) of 6-8 weeks old were used for tumor fragment implantation. Animals were fed with normal nude mice diet and housed in SPF animal facility in accordance with the Guide for Care and Use of Laboratory Animals and regulations of the Institutional Animal Care and Use Committee. Patent STO#69 gastric tumor fragments of about 15-30 mm^3 in size were implanted (s.c.) into right flanks of Balb/c nude mice.

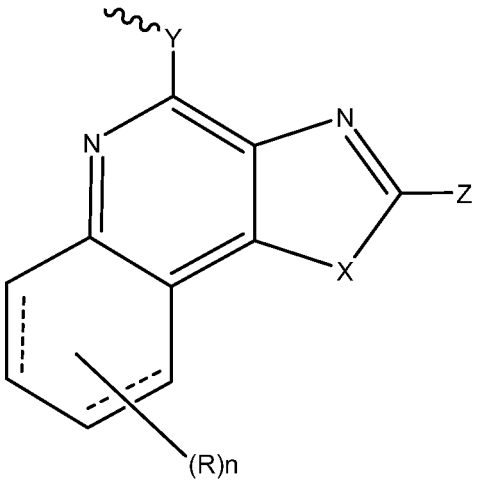
[0286] Drugs were administered by i.v. route with 5~20 mg/kg of antibody or reference drug, QWKx3. Tumors were measured once a week by caliper to determine its subcutaneous growth. Tumors were measured twice a week in two dimensions with calipers. Tumor volume was calculated using following formula: tumor volume = (length x width2) x 0.5. Average tumor volumes or body weights were plotted using graph program Prism 5 (GraphPad). An endpoint for efficacy study was set at 30-45 days post-first treatment or when a tumor size reached above 2000 mm3, whichever came first. If a mouse lose more than 20% of body weight or is very sick and cannot get to adequate food or water, it will be removed from the study and euthanized. Tumors were collected from mice at the end point, half frozen in LN_2 and half fixed in formalin for preparing FFPE tissues.

Claims

1. A compound having the structure of Formula (Ia):

TM-L-AM (Ia),

wherein TM is an antibody or a functional fragment thereof that specifically binds to Her2/Neu, AM is an activating moiety that is represented by structure of formula (I):



(I)

wherein dashed line represents bond or absence of bond, ∿∿ is the point to be connected to the linker;

X is S or $-NR_1$, R_1 is $-W_0—W_1—W_2—W_3—W_4$,

W_0 is a bond, alkyl, alkenyl, alkynyl, alkoxy, or –alkyl-S-alkyl--,

W_1 is a bond, --O--, or $-NR_2$--, wherein R_2 is hydrogen, alkyl or alkenyl,

W_2 is a bond, --O--, --C(O)--, --C(S)--, or $-S(O)_2—$,

W_3 is a bond, $--NR_3$--, wherein R_3 is hydrogen, alkyl or alkenyl,

W_4 is hydrogen, alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, aryl, aryloxy, heteroaryl, or heterocyclyl, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, $--NH_2$, nitro, --alkyl-hydroxyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, $--O-R_4$, $--O-alkyl-R_4$, $--alkyl-O-R_4$, $--C(O)-R_4$, $--alkyl-C(O)-R_4$, $--alkyl-C(O)-O-R_4$, $--C(O)-O-R_4$, $--S-R_4$, $--S(O)_2-R_4$, $--NH-S(O)_2-R_4$, --alkyl-S-R4, $--alkyl-S(O)_2-R_4$, $--NHR_4$, $--NR_4R_4$,--NH-alkyl-R_4, halogen, --CN, $--NO_2$, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

Z is hydrogen, alkyl, alkenyl, alkynyl, alkoxy, aryl, haloalkyl, heteroaryl, heterocyclyl, each of which can be optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, aryl, heteroaryl, heterocyclyl, halogen, cyano, nitro, --$N(R_5)_2$, --alkoxy-alkyl, --alkoxy-alkenyl, --C(O)-alkyl, --C(O)-O-alkyl, --O-C(O)-alkyl, --C(O)-$N(R_5)_2$, aryl, heteroaryl, --CO-aryl, and –CO-heteroaryl, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl;

R is hydrogen, alkyl, alkoxy, haloalkyl, halogen, aryl, heteroaryl, heterocyclyl, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, aryl, heteroaryl, heterocyclyl, --NH_2, nitro, --alkyl-hydroxyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --alkyl-O-R_4, --C(O)-R_4, --C(O)-NH-R_4, --C(O)-NR_4R_4, --alkyl-C(O)-R_4, --alkyl-C(O)-O-R_4, --C(O)-O-R_4, --O-C(O)-R_4, --S-R_4, --C(O)-S-R_4, --S-C(O)-R_4, --$S(O)_2$-R_4, --NH-$S(O)_2$-R_4, --alkyl-S-R_4, --alkyl-$S(O)_2$-R_4, --NHR_4, --NR_4R_4,--NH-alkyl-R_4, halogen, --CN, and –SH, wherein R_4 is independently hydrogen, alkyl, alkenyl, alkoxy, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl;

n is 0, 1, 2, 3, or 4;

Y is –NR_6R_7, –$CR_6R_7R_8$, or –alkyl-NH_2, each of which can be optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, --NH_2, halogen, --$N(R_5)_2$, --alkoxy-alkyl, --alkoxy-alkenyl, --C(O)-alkyl, --C(O)-O-alkyl, --C(O)-$N(R_5)_2$, aryl, heteroaryl, --CO-aryl, and –CO-heteroaryl,

wherein R_6, R_7 and R_8 are independently hydrogen, alkyl, alkenyl, alkoxy, alkylamino, dialkylamino, alkylthio, arylthio, --alkyl-hydroxyl, --alkyl-C(O)-O-R_9, --alkyl-C(O)-R_9, or –alkyl-O-C(O)-R_9, wherein each R_5 is independently hydrogen, alkyl, haloalkyl, --alkyl-aryl, or –alkyl-heteroaryl, wherein R_9 is hydrogen, alkyl, alkenyl, halogen, or haloalkyl;

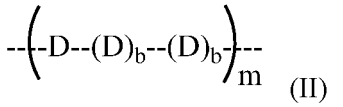
X and Z taken together may optionally form a (5-9)-membered ring;

or a pharmaceutically acceptable salt or solvate thereof.

2. The compound of claim **1**, wherein AM is a compound selected from the group consisting of: 2-propylthiazolo[4,5-c]quinolin-4-amine, 1-(2-methylpropyl)-1H-imidazo[4,5-c]quinolin-4-amine, 4-amino-2-(ethoxymethyl)-a,a-di-methyl-1H-imidazo[4,5-c]quinoline-1-ethanol, 1-(4-amino-2-ethylaminomethylimidazo-[4,5-c]quinolin-1-yl)-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl-]methanesulfonamide, 4-amino-2-ethoxymethyl-aa-dimethyl-6,7,8,9-tetrahydro-1h-imidazo[4,5-c]quinoline-1-ethanol, 4-amino-aa-dimethyl-2-methoxyethyl-1h-imidazo[4,5-c]quinoline-1-ethanol, 1-{2-[3-(benzyloxy)propoxy]ethyl}-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-

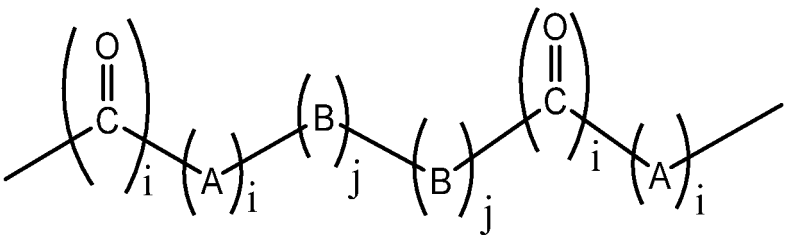
4-amine, N-[4-(4-amino-2-butyl-1H-imidazo[4,5- c][1,5]naphthyridin-1-yl)butyl]-n'-butylurea, N1-[2-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5] naphthyridin-1-yl)ethyl]-2-amino-4- methylpentanamide, N-(2-{2-[4-amino-2-(2-methoxyethyl)-1H- imidazo[4,5-c]quinolin-1-yl]ethoxy}ethyl)-n'- phenylurea, 1-(2-amino-2-methylpropyl)-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, 1-{4-[(3,5-dichlorophenyl)sulfonyl]butyl}-2-ethyl- 1H-imidazo[4,5-c]quinolin-4-amine, N-(2-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]ethoxy}ethyl)-n'-cyclohexylurea, N-{3-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]propyl}-n'-(3-cyanophenyl)thiourea, N-[3-(4-amino-2-butyl-1H-imidazo[4,5-c]quinolin-1- yl)-2,2-dimethylpropyl]benzamide, 2-butyl-1-[3-(methylsulfonyl)propyl]-1H- imidazo[4,5-c]quinolin-4-amine, N-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]-1,1-dimethylethyl}-2- ethoxyacetamide, 1-[4-amino-2-ethoxymethyl-7-(pyridin-4-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 1-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, N-{3-[4-amino-1-(2-hydroxy-2-methylpropyl)-2- (methoxyethyl)-1H-imidazo[4,5-c]quinolin-7-yl]phenyl}methanesulfonamide, 1-[4-amino-7-(5-hydroxymethylpyridin-3-yl)-2-(2- methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 3-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]propane-1,2-diol, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-propylurea, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-cyclopentylurea, 1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methyl]-2- (ethoxymethyl)-7-(4-hydroxymethylphenyl)-1H- imidazo[4,5-c]quinolin-4-amine, 4-[4-amino-2-ethoxymethyl-1-(2-hydroxy-2-methylpropyl)-1H-imidazo[4,5-c]quinolin-7-yl]-N- methoxy-N-methylbenzamide, 2-ethoxymethyl-N1-isopropyl-6,7,8,9-tetrahydro-1H-imidazo[4,5-c]quinoline-1,4-diamine, 1-[4-amino-2-ethyl-7-(pyridin-4-yl)-1H-imidazo[4,5- c]quinolin-1-yl]-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl]methanesulfonamide, and N-[4-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)butyl]-n'-cyclohexylurea.

3. The compound of claims **1-2**, wherein L is represented by structure of formula (II):



(II)

m is 1, 2, 3, 4, 5, or 6, each b independently is 0 or 1, and D is independently represented by structure of formula (III):



(III)

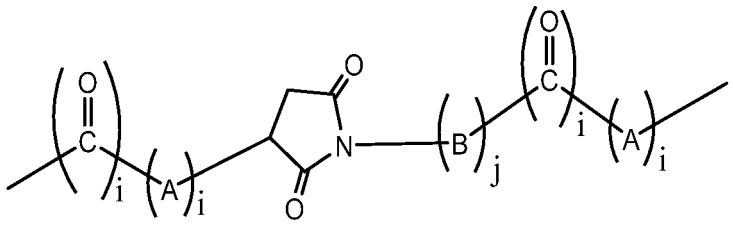
wherein each i independently is 0 or 1;

each j independently is 0, 1, 2, 3, 4, 5, or 6;

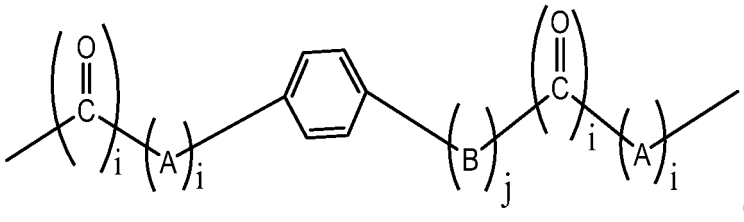
each A independently is S, O, or N-Ra, wherein Ra is hydrogen, alkyl, alkenyl, or alkoxy;

each B independently is alkyl, alkenyl, --O-alkyl--, --alkyl-O--, --S-alkyl--, --alkyl-S--, aryl, heteroaryl, heterocyclyl, or peptide, each of which is optionally substituted by one or more substituents selected from the group consisting of hydroxyl, alkoxy, alkyl, alkenyl, alkynyl, cycloalkyl, --alkyl-aryl, --alkyl-heteroaryl, --alkyl-heterocyclyl, --O-R_4, --O-alkyl-R_4, --C(O)-R_4, --C(O)-O-R_4, --S-R_4, --$S(O)_2$-R_4, --NHR_4, --NH-alkyl-R_4, halogen, --CN, --NO_2, and –SH, wherein R_4 is alkyl, alkenyl, --alkyl-hydroxyl, aryl, heteroaryl, heterocyclyl, or haloalkyl.

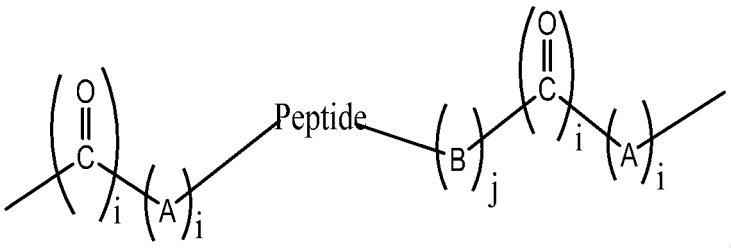
4. The compound of claim **3**, wherein the linker is presented by the following strucutres of formula (V)-(VII):



(V)



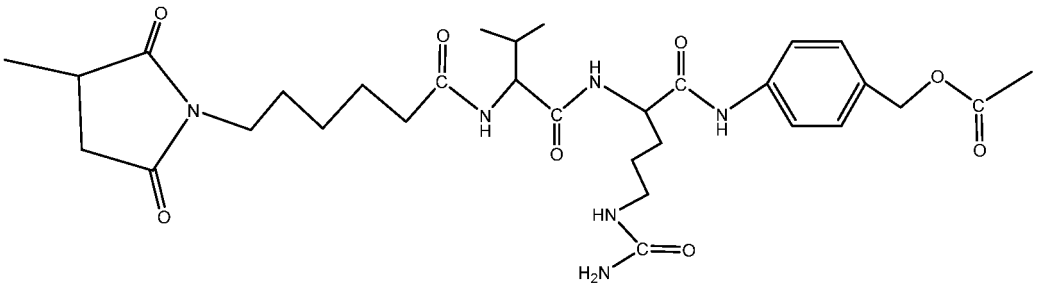
(VI)



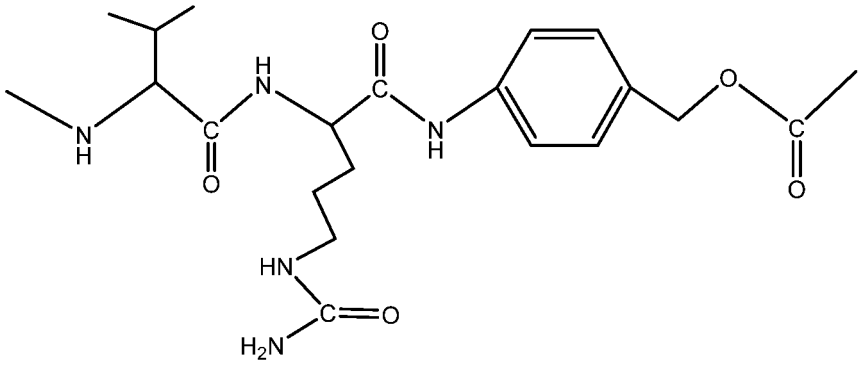
(VII)

A, B, i and j are defined above.

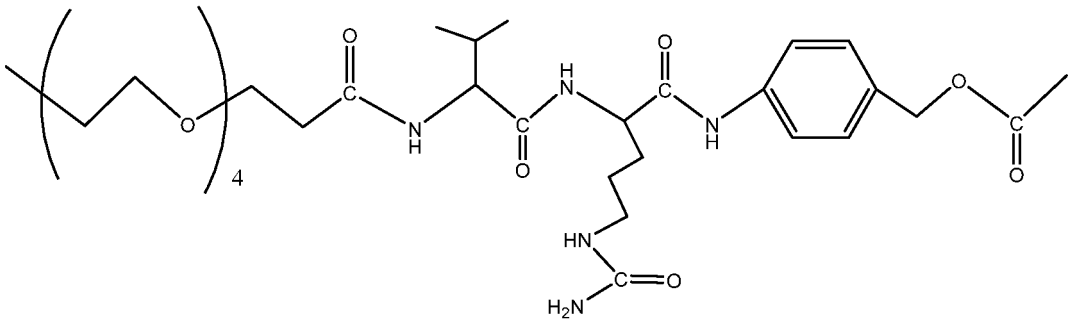
5. The compound of claim **4**, wherein the linker is selected from S1, S2, S3, S4, S5, S6, S7, –Gly-Phe-Leu-Gly-, -Ala-Leu-Ala-Leu-, -Phe-Arg-, -Phe-Lys-, -Val-Lys-, -Val-Ala-, or Val-Cit-, wherein S1-S7 are represented by the following structures:



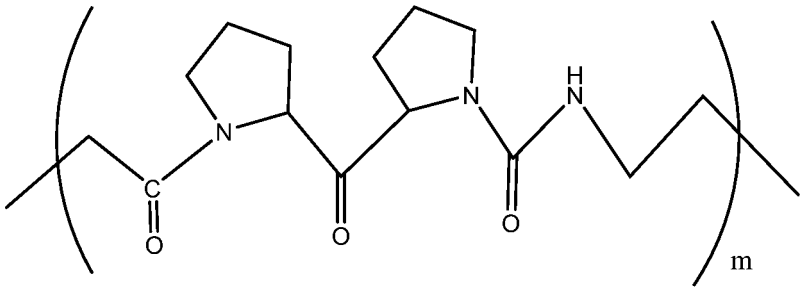
S1



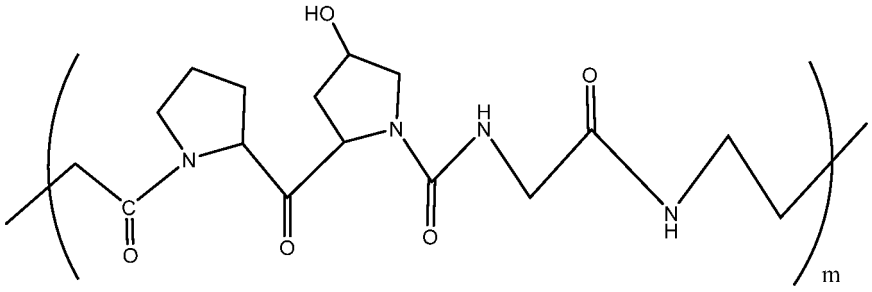
S2



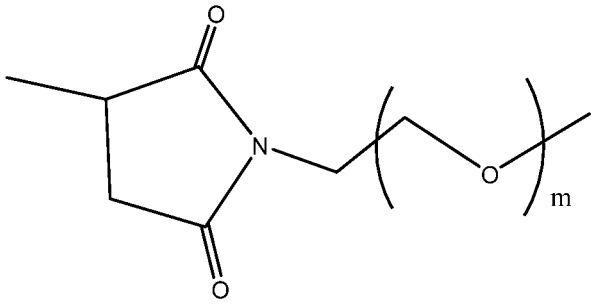
S3



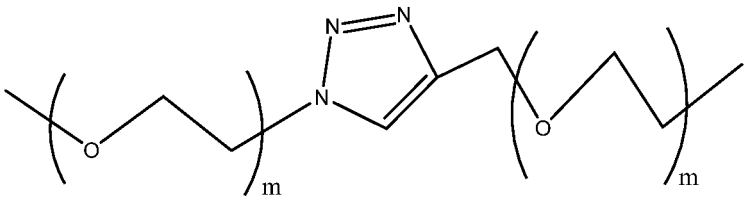
S4



S5



S6



S7,

wherein each m is independently 1 to 20.

6. The compound of claims **1-5**, wherein the antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22).

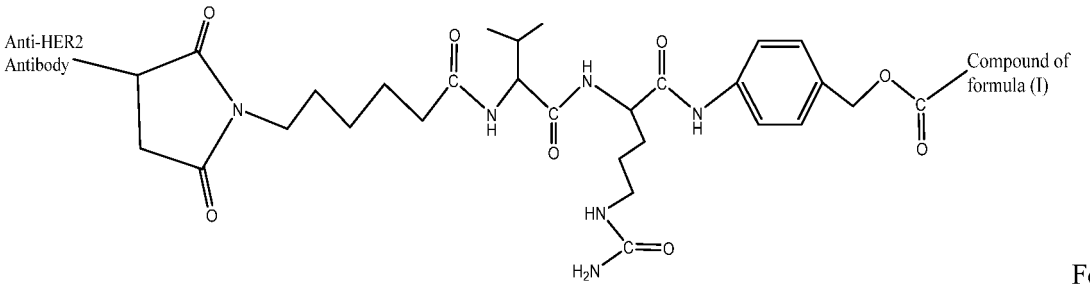
7. The compound of claim **1**, wherein TM is an anti-HER2 antibody; L is selected from S1, S2, S3, S4, S5, S6, S7, –Gly-Phe-Leu-Gly-, -Ala-Leu-Ala-Leu-, -Phe-Arg-, -Phe-Lys-, -Val-Lys-, -Val-Ala-, or Val-Cit-; AM is a compound of formula (I) selected from selected from 2-propylthiazolo[4,5-c]quinolin-4-amine, 1-(2-methylpropyl)-1H-imidazo[4,5-c]quinolin-4-amine, 4-amino-2-(ethoxymethyl)-a,a-dimethyl-1H-imidazo[4,5-c]quinoline-1-ethanol, 1-(4-amino-2-ethylaminomethylimidazo-[4,5-c]quinolin-1-yl)-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl-]methanesulfonamide, 4-amino-2-ethoxymethyl-aa-dimethyl-6,7,8,9-tetrahydro-1h-imidazo[4,5-c]quinoline-1-ethanol, 4-amino-aa-dimethyl-2-methoxyethyl-1h-imidazo[4,5-c]quinoline-1-ethanol, 1-{2-[3-(benzyloxy)propoxy]ethyl}-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, N-[4-(4-amino-2-butyl-1H-imidazo[4,5- c][1,5]naphthyridin-1-yl)butyl]-n'-butylurea, N1-[2-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5] naphthyridin-1-yl)ethyl]-2-amino-4- methylpentanamide, N-(2-{2-[4-amino-2-(2-methoxyethyl)-1H- imidazo[4,5-c]quinolin-1-yl]ethoxy}ethyl)-n'- phenylurea, 1-(2-amino-2-methylpropyl)-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, 1-{4-[(3,5-dichlorophenyl)sulfonyl]butyl}-2-ethyl- 1H-imidazo[4,5-c]quinolin-4-amine, N-(2-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]ethoxy}ethyl)-n'-cyclohexylurea, N-{3-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]propyl}-n'-(3-cyanophenyl)thiourea, N-[3-(4-amino-2-butyl-1H-imidazo[4,5-c]quinolin-1- yl)-2,2-dimethylpropyl]benzamide, 2-butyl-1-[3-(methylsulfonyl)propyl]-1H- imidazo[4,5-c]quinolin-4-amine, N-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]-1,1-dimethylethyl}-2- ethoxyacetamide, 1-[4-amino-2-ethoxymethyl-7-(pyridin-4-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 1-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, N-{3-[4-amino-1-(2-hydroxy-2-methylpropyl)-2- (methoxyethyl)-1H-imidazo[4,5-c]quinolin-7-yl]phenyl}methanesulfonamide, 1-[4-amino-7-(5-hydroxymethylpyridin-3-yl)-2-(2- methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 3-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]propane-1,2-diol, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-

3-propylurea, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-cyclopentylurea, 1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methyl]-2- (ethoxymethyl)-7-(4-hydroxymethylphenyl)-1H- imidazo[4,5-c]quinolin-4-amine, 4-[4-amino-2-ethoxymethyl-1-(2-hydroxy-2-methylpropyl)-1H-imidazo[4,5-c]quinolin-7-yl]-N- methoxy-N-methylbenzamide, 2-ethoxymethyl-N1-isopropyl-6,7,8,9-tetrahydro-1H-imidazo[4,5-c]quinoline-1,4-diamine, 1-[4-amino-2-ethyl-7-(pyridin-4-yl)-1H-imidazo[4,5- c]quinolin-1-yl]-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl]methanesulfonamide, or N-[4-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)butyl]-n'-cyclohexylurea, wherein the amine group on the quinoline ring is the point of connection with the linker.

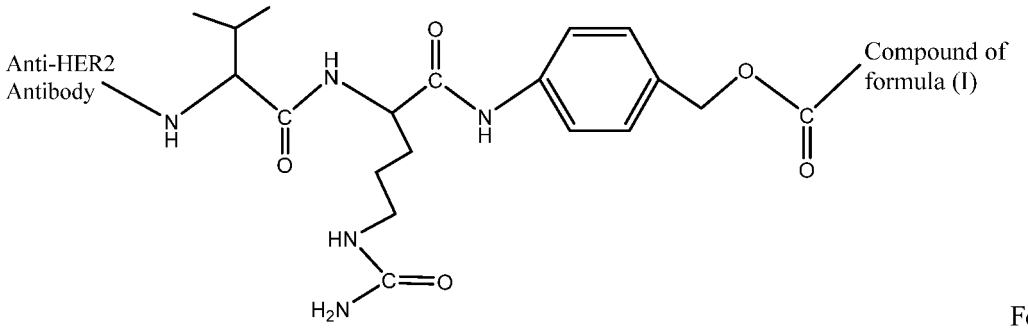
8. The compound of claim **7**, wherein the anti-HER2 antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); L is S1, S2 or S3.

9. The compound of claim **8**, wherein the anti-HER2 antibody is trastuzumab (Herceptin); AM is Resiquimod or Imiquimod, wherein the amine group on the quinoline ring is the point of connection with the linker.

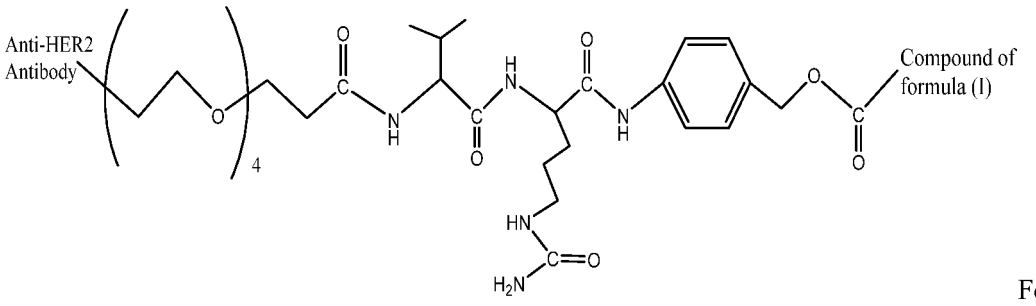
10. A compopund having the structures of Formula A-C:



Formula A,



Formula B,



Formula C;

or a pharmaceutically acceptable salt or solvate thereof.

11. The compound of claim **10**, the anti-HER2 antibody is trastuzumab (Herceptin), pertuzumab or margetuximab (MGAH22); the compound of formula (I) is selected from 2-propylthiazolo[4,5-c]quinolin-4-amine, 1-(2-methylpropyl)-1H-imidazo[4,5-c]quinolin-4-amine, 4-amino-2-(ethoxymethyl)-a,a-di-methyl-1H-imidazo[4,5-c]quinoline-1-ethanol, 1-(4-amino-2-ethylaminomethylimidazo-[4,5-c]quinolin-1-yl)-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl-]methanesulfonamide, 4-amino-2-ethoxymethyl-aa-dimethyl-6,7,8,9-tetrahydro-1h-imidazo[4,5-c]quinoline-1-ethanol, 4-amino-aa-dimethyl-2-methoxyethyl-1h-imidazo[4,5-c]quinoline-1-ethanol, 1-{2-[3-(benzyloxy)propoxy]ethyl}-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, N-[4-(4-amino-2-butyl-1H-imidazo[4,5- c][1,5]naphthyridin-1-yl)butyl]-n'-butylurea, N1-[2-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5] naphthyridin-1-yl)ethyl]-2-amino-4- methylpentanamide, N-(2-{2-[4-amino-2-(2-methoxyethyl)-1H- imidazo[4,5-c]quinolin-1-yl]ethoxy}ethyl)-n'- phenylurea, 1-(2-amino-2-methylpropyl)-2-(ethoxymethyl)-1H-imidazo[4,5-c]quinolin-4-amine, 1-{4-[(3,5-dichlorophenyl)sulfonyl]butyl}-2-ethyl- 1H-imidazo[4,5-c]quinolin-4-amine, N-(2-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]ethoxy}ethyl)-n'-cyclohexylurea, N-{3-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]propyl}-n'-(3-cyanophenyl)thiourea, N-[3-(4-amino-2-butyl-1H-imidazo[4,5-c]quinolin-1- yl)-2,2-dimethylpropyl]benzamide, 2-butyl-1-[3-(methylsulfonyl)propyl]-1H- imidazo[4,5-c]quinolin-4-amine, N-{2-[4-amino-2-(ethoxymethyl)-1H-imidazo[4,5- c]quinolin-1-yl]-1,1-dimethylethyl}-2- ethoxyacetamide, 1-[4-amino-2-ethoxymethyl-7-(pyridin-4-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 1-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, N-{3-[4-amino-1-(2-hydroxy-2-methylpropyl)-2- (methoxyethyl)-1H-imidazo[4,5-c]quinolin-7-yl]phenyl}methanesulfonamide, 1-[4-amino-7-(5-hydroxymethylpyridin-3-yl)-2-(2- methoxyethyl)-1H-imidazo[4,5-c]quinolin-1-yl]-2-methylpropan-2-ol, 3-[4-amino-2-(ethoxymethyl)-7-(pyridin-3-yl)-1H- imidazo[4,5-c]quinolin-1-yl]propane-1,2-diol, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-propylurea, 1-[2-(4-amino-2-ethoxymethyl-1H-imidazo[4,5- c]quinolin-1-yl)-1,1-dimethylethyl]-3-cyclopentylurea, 1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methyl]-2- (ethoxymethyl)-7-(4-

hydroxymethylphenyl)-1H- imidazo[4,5-c]quinolin-4-amine, 4-[4-amino-2-ethoxymethyl-1-(2-hydroxy-2-methylpropyl)-1H-imidazo[4,5-c]quinolin-7-yl]-N- methoxy-N-methylbenzamide, 2-ethoxymethyl-N1-isopropyl-6,7,8,9-tetrahydro-1H-imidazo[4,5-c]quinoline-1,4-diamine, 1-[4-amino-2-ethyl-7-(pyridin-4-yl)-1H-imidazo[4,5- c]quinolin-1-yl]-2-methylpropan-2-ol, N-[4-(4-amino-2-ethyl-1H-imidazo[4,5-c]quinolin-1-yl)butyl]methanesulfonamide, or N-[4-(4-amino-2-butyl-1H-imidazo[4,5-c][1,5]naphthyridin-1-yl)butyl]-n'-cyclohexylurea, wherein the amine group on the quinoline ring is the point of connection with the linker.

12. The compound of claim **11**, wherein the compound of formula (I) is Resiquimod or Imiquimod, wherein the amine group on the quinoline ring is the point of connection with the linker.

13. The comound of claims **10-12**, wherein the anti-HER2 antibody is trastuzumab (Herceptin).

14. A pharmaceutical composition comprising an effective amount of the compound according to claims 1-13 and one or more pharmaceutically acceptable carriers.

15. The pharmaceutical composition of claim **14**, further comprising an effective amount of an additional therapeutic agent.

16. The pharmaceutical composition of claim **15**, wherein the additional therapeutic agent is an anticancer agent.

17. The pharmaceutical composition of claim **16**, wherein the anticancer agent is an antimetabolite, an inhibitor of topoisomerase I and II, an alkylating agent, a microtubule inhibitor, an antiandrogen agent, a GNRh modulator or mixtures thereof.

18. A method of inhibiting proliferation of a HER2 positive tumor cell comprising administering to said tumor cell the compound of claims **1-13**.

19. A method of treating a HER2 positive tumor/cancer in a subject comprising administering to the subject the compound of claims **1-13**.

20. The method of claims **18** or **19**, wherein the the tumor/cell is selected from esophageal, stomach, colon, rectal, , pancreatic, lung, breast, cervix uteri, corpus uteri, ovary, bladder, head and neck,endometrial, osteosarcoma, prostate, neuroblastoma,

21. Use of the compound according to claims **1-13** in the manufacturer of a medicament for the treatment of a HER2 positive tumor/cancer in a subject.

22. Use of the pharmaceutical composition according to claims **14-17** in the manufacturer of a medicament for the treatment of a HER2 positive tumor/cancer in a subject.