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Folate Conjugates of Albumin-Binding Entities

Field of Invention

10 The present invention is directed towards new trifunctional folate-conjugates comprising a folate, an albumin binder and a radionuclide-based diagnostic or therapeutic moiety as well as pharmaceutical compositions thereof, their method of production and their use in diagnostic and therapeutic medical
15 applications, such as diagnostic nuclear imaging and radionuclide therapy.

Background

Cell-specific targeting for delivery of diagnostic or
20 therapeutic agents is a widely researched field and has led to the development of noninvasive diagnostic and/or therapeutic medical applications. In particular in the field of nuclear medicine procedures and treatments, which employ radioisotopes characterized by emission of gamma-rays or beta or alpha
25 particles or Auger electrons, selective localization of these radioactive compounds in targeted cells or tissues is required to achieve either high signal intensity and specificity for visualization (gamma- or positron radiation) of specific tissues, assessing a disease and/or monitoring effects of
30 therapeutic treatments, or to achieve high radiation dose

through particle radiation (beta- or alpha-radiation), for delivery of adequate doses of ionizing radiation to a specified diseased site, while preventing damage to healthy tissues.

The folate receptor (FR) is a high-affinity membrane-associated protein, which exhibits limited expression on healthy cells, but is frequently overexpressed on a wide variety of specific cell types, such as epithelial tumor cells (e.g. ovarian, endometrial, breast, colorectal, kidney, lung, nasopharyngeal) and activated (but not resting) macrophages, which are involved in inflammation and autoimmune diseases. This led to the use of folic acid and its derivatives as a targeting agent for the delivery of therapeutic and/or diagnostic agents to these specific cell populations to achieve a selective concentration of pharmaceutical and/or diagnostic agents in these specific cells relative to normal cells. Such folate-conjugates include folate radiopharmaceuticals (Leamon and Low, Drug Discov. Today 2001; 6:44-51; Ke et al., Adv Drug Deliv Rev 2004, 1143-1160, Müller and Schibli, J Nucl Med 2011; 52:1-4; Müller, Curr Pharm Design 2012; 18:1058-1083), folate-conjugates of chemotherapeutic agents (Leamon and Reddy, Adv. Drug Deliv. Rev. 2004; 56:1127-41; Leamon et al, Bioconjugate Chem. 2005; 16:803-11; Vlahov et al. Bioconjug Chem 2012; in press), proteins and protein toxins (Ward et al., J. Drug Target. 2000; 8:119-23; Leamon et al, J. Biol. Chem. 1993; 268:24847-54; Leamon and Low, J. Drug Target. 1994; 2:101-12), antisense oligonucleotides (Li et al, Pharm. Res. 1998; 15:1540-45; Zhao and Lee, Adv. Drug Deliv. Rev. 2004; 56:1193-204), liposomes (Lee and Low, Biochim. Biophys. Acta-Biomembr. 1995; 1233:134-44); Gabizon et al, Adv. Drug Deliv. Rev. 2004; 56:1177-92), hapten molecules (Paulos et al, Adv. Drug Deliv. Rev. 2004; 56:1205-17); MRI contrast agents (Konda et al, Magn. Reson. Mat. Phys. Biol. Med. 2001; 12:104-13) etc.

Known folate radiopharmaceuticals include for example conjugates with ¹²⁵I-labeled histamine (US 4,136,159), with small metal-chelates such as deferoxamine (US 5,688,488), acyclic or cyclic polyaminocarboxylates (e.g. DTPA, DTPA-BMA, DOTA and DO3A; US 6,221,334, Fani et al. Eur J Nucl Med Mol Imaging 2011;38: 108-119; Müller et al. Nucl Med Biol 2011; 38: 715-723), bisaminothiol (US 5,919,934), 6-hydrazinonicotinamido-hydrazido (Shuang Liu, Topics in Current Chemistry, vol 252 (2005), Springer Berlin/Heidelberg), and ethylenedicysteine (US 7,067,111), and small peptides (US 7,128,893).

However, there is still a need for alternative, highly selective radionuclide conjugates, which can be synthesized easily and which exhibit optimal target (i.e. tumor cell, activated macrophage, etc.) to non-target tissue ratios and are eliminated through the kidneys, for use as tumor imaging agents in highly selective and non-invasive procedures permitting early detection and treatment of tumor cells, activated macrophages (and other targeted cells exhibiting high ER expression, not yet identified).

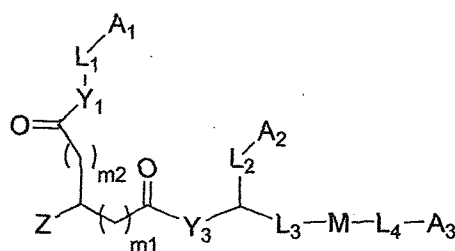
Applicants have now found novel trifunctional folate-conjugates that are able to overcome the drawbacks of known conjugates and meet the current needs by showing several advantages, such as stable complex formation, improved biodistribution and increased target tissue uptake. These novel trifunctional folate-conjugates comprise a folate, an albumin binder and a radionuclide-based therapeutic moiety or diagnostic moiety, e.g. a moiety suitable for diagnostic imaging or radiotherapeutic applications.

Summary of the Invention

The present invention is in a first aspect directed to new trifunctional folate-conjugates comprising a folate, an albumin

binder and a radionuclide-based therapeutic or diagnostic moiety (hereinafter also called compounds of the invention).

In one specific embodiment, the new folate conjugates are compounds of formula I,



I

wherein

Z is a pterooate or derivative thereof,

L₁, L₄ are independently of each other a covalent bond or a
10 straight-chain or branched C(1-8)alkyl, which is unsubstituted
or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂,
wherein R' represents H or C(1-8)alkyl,

L₂ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

L₃ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, and a five-

25 $\text{CO-NR}'-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$, $-\text{O}-\text{CO}-\text{O}-$, $-\text{S}-\text{R}'-$, $-\text{SO}_3\text{R}'-$, and a five-

membered azaheterocycle, wherein R' represents H or C(1-8)alkyl,

Y₁, Y₃ are independently of each other O, N or S,

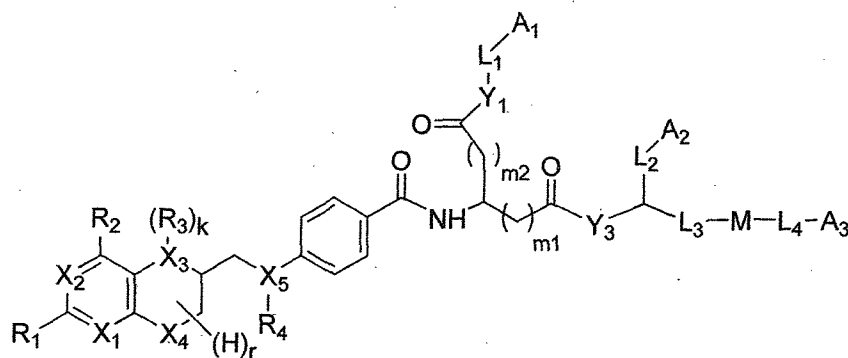
A₁, A₂, A₃ are independently of each other H, a capping group, or an albumin binder,

5 M is a radionuclide-based therapeutic or diagnostic moiety,

m₁, m₂ are independently of each other 0, 1, 2 or 3,

with the proviso that at least one, preferably one, of A₁, A₂ and A₃ is an albumin binder.

10 Preferably, the invention is directed towards compounds of formula II



II

15 wherein

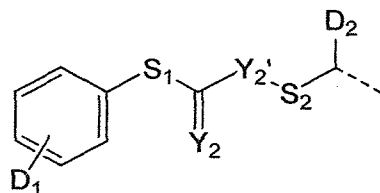
X₁ to X₅ are independently of each other C, N or O, preferably N or O,

R₁, R₂ are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, -CONHR₅, wherein R₅ represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -COR', -COOR', or -NHR', wherein R' is H or C(1-8)alkyl,

R_3 , R_4 are independently of each other H, nitroso, C(1-12)alkyl, -OR', -COR' or halosubstituted -COR', wherein R' represents H or C(1-8)alkyl,

and L_1 , L_2 , L_3 , L_4 , Y_1 , Y_3 , A_1 , A_2 , A_3 , M , m_1 , m_2 , k , and r are as defined above.

The albumin binder is preferably a linear or branched lipophilic group having 12-40 carbon atoms and a distal acidic group, such as a compound of formula III



III

wherein

Y_2 , Y_2' are independently of each other N, O or S,

S_1 , S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D_1 is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

D_2 is an acidic group,

and the broken line represents the linkage to L_1 , L_2 or L_4 .

The acidic group is preferably a group capable of ionizing to donate a hydrogen ion to a base to form a salt, preferably selected from the group consisting of $-\text{COOH}$, $-\text{SO}_3\text{H}$, $-\text{SO}_2\text{H}$, $-\text{NR}'\text{SO}_3\text{H}$, $-\text{P}(\text{O})(\text{OH})_2$, wherein R' represents H or C(1-8)alkyl.

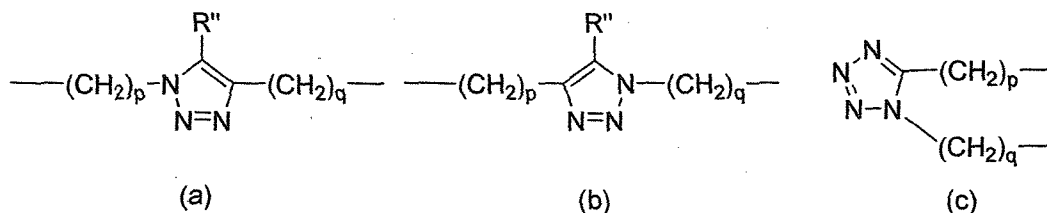
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In specific embodiments, either m_1 is 2 and m_2 is 0 or else m_1 is 0 and m_2 is 2.

Preferably, M is a radionuclide-based therapeutic or diagnostic moiety M_1 , which is a chelated metal radionuclide comprising a metal radionuclide and a metal chelator.

Alternatively, M is a radionuclide-based therapeutic or diagnostic moiety M_2 , which is a gamma- or positron-emitting non-metal radionuclide, optionally in combination with a prosthetic group.

15 In specific embodiments, L_3 is a group L_3' , which is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR' , NHR' , or $\text{CO}_2\text{R}'$, and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by a group selected from $-\text{O}-$, $-\text{CO}-$, $-\text{CO}-\text{O}-$, $-\text{NR}'-$, $-\text{NR}'-\text{CO}-$, or L_3 is a group of formulae (a), (b), or (c)



wherein

R'' is H or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, or NO_2 , and p and q are independently of each other 0, 1, 2, 3, 4, 5 or 6.

In a further aspect the present invention provides methods for

synthesizing the compounds of the invention.

In yet a further aspect the invention provides pharmaceutical compositions comprising a diagnostic imaging amount or a therapeutically effective amount of at least one compound of the present invention and a pharmaceutically acceptable carrier therefor.

In a further aspect the present invention provides uses of compounds and/or pharmaceutical compositions of the present invention for convenient and effective administration to a subject in need for diagnostic imaging or radionuclide therapy. The subject of the methods of the present invention is preferably a mammal, such as an animal or a human, preferably a human.

In a further aspect the present invention provides a single or multi-vial or multi-compartment kit containing all of the components needed to prepare the compounds of this invention.

Other features and advantages of the invention will be apparent from the following detailed description thereof and from the claims.

Brief Description of Figures

Figure 1: In vitro stability of ^{177}Lu -radiolabeled folate conjugates in PBS and human plasma (-■- represents ^{177}Lu -DOTA-folate in PBS, -▲- represents ^{177}Lu -DOTA-folate in plasma, -●- represents ^{177}Lu -DOTA-AB-folate in PBS, -◆- represents ^{177}Lu -DOTA-AB-folate in plasma).

Figure 2: Cell uptake and internalization of ^{177}Lu -DOTA-folate and ^{177}Lu -DOTA-AB-folate 16 in FR-positive KB-cells at 2 h and 4 h (vertically striped bars represent ^{177}Lu -DOTA-folate at 2h, horizontally striped bars represent ^{177}Lu -DOTA-AB-folate at 2h,

diagonally striped bars represent ^{177}Lu -DOTA-folate at 4h, filled bars represent ^{177}Lu -DOTA-AB-folate at 4h).

Figure 3: Biodistribution study in KB-tumor bearing mice 4 h and 24 h after injection of ^{177}Lu -DOTA-AB-folate and the control compound ^{177}Lu -DOTA-folate (dotted bars represent ^{177}Lu -DOTA-folate at 4h p.i., striped bars represent ^{177}Lu -DOTA-folate at 24h p.i., empty bars represent ^{177}Lu -DOTA-AB-folate at 4h p.i., filled bars represent ^{177}Lu -DOTA-AB-folate at 24h p.i.).

Figure 4(A), (B): SPECT/CT Images of tumor-bearing mice: Uptake of a ^{177}Lu -radiolabeled (A) and ^{161}Tb -radiolabeled DOTA-AB-folate (B) in a KB tumor xenograft (T) and in the kidneys (K) at 4h p.i. (left) and 24h p.i. (right).

Figure 5(A), (B): Effect of ^{177}Lu -radiolabeled DOTA-AB-folate (group A) or the control ^{177}Lu -radiolabeled DOTA-folate (group B) in KB tumor bearing nude mice over time, i.e. 19 days (-▲- represents group A, -◆- represents group B).

Detailed Description of the Invention

The present invention is in a first aspect directed to new trifunctional folate-conjugates comprising a folate, an albumin binder and a radionuclide-based therapeutic or diagnostic moiety (hereinafter also called compounds of the invention).

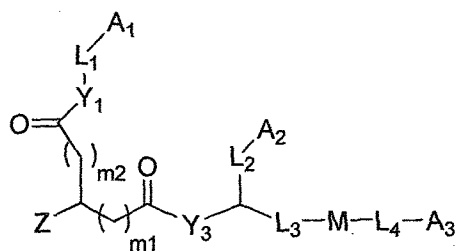
The radionuclide-based therapeutic or diagnostic moiety (hereinafter also called "radionuclide moiety") may be comprising any known metal or non-metal radionuclide such as a radioactive metal ion, a paramagnetic metal ion, a gamma-or positron emitting radiohalogen, a positron-emitting radioactive non-metal, a hyperpolarised NMR-active nucleus, a reporter suitable for in vivo optical imaging, or a beta-emitter suitable for intravascular detection.

Preferred radionuclide moieties for diagnostic purpose are any known moieties for use in the present invention which can be detected externally in a non-invasive manner following administration in vivo. Most preferred radionuclide moieties are
5 a chelated metal radionuclide (i.e. a radiometal) or a gamma- or positron-emitting non-metal radionuclide, i.e. particularly those suitable for detection using SPECT or PET. In some embodiments the folate, albumin binder and radionuclide moiety are in a radial arrangement, i.e. both the albumin binder and
10 radionuclide moiety are attached to the amino acid portion of the folate molecule. In other embodiments, the folate, albumin binder and radionuclide moiety are in a linear arrangement, i.e. only one of the albumin binder and radionuclide moiety is attached to the amino acid portion of the folate molecule. A
15 skilled person will know which arrangements are suitable, for example when A₂ is an albumin binder, M is preferably a chelated metal radionuclide (i.e. a radiometal).

Preferred radionuclide moieties for therapeutic purpose are any known moieties for use in the present invention, which can be
20 used for the treatment of any disease responsive to radionuclide treatment, e.g. cancer, following administration in vivo. Most preferred radionuclide moieties are a chelated metal radionuclide (i.e. a radiometal) or a gamma- or positron-emitting non-metal radionuclide. In some embodiments the folate,
25 albumin binder and radionuclide moiety are in a radial arrangement, i.e. both the albumin binder and radionuclide moiety are attached to the amino acid portion of the folate molecule. In other embodiments, the folate, albumin binder and radionuclide moiety are in a linear arrangement, i.e. only one
30 of the albumin binder and radionuclide moiety is attached to the amino acid portion of the folate molecule. A skilled person will know which arrangements are suitable, e.g. when A₂ is an albumin binder, M is preferably a chelated metal radionuclide (i.e. a

radiometal).

The present invention is specifically directed towards compounds of formula I,



I

wherein

Z is a pterate or derivative thereof,

L₁, L₄ are independently of each other a covalent bond or a
10 straight-chain or branched C(1-8)alkyl, which is unsubstituted
or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂,
wherein R' represents H or C(1-8)alkyl,

L₂ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

L₃ is a covalent bond or a linking group, such as straight-chain
or branched C(1-8)alkyl, which is unsubstituted or substituted
by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, and wherein one
or more of the non-adjacent CH₂ groups may independently be
replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-
, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -
C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, and a five-membered

azaheterocycle, wherein R' represents H or C(1-8)alkyl,

Y_1, Y_3 are independently of each other O, N or S,

A_1, A_2, A_3 are independently of each other H, a capping group, or an albumin binder,

5 M is a radionuclide-based therapeutic or diagnostic moiety, preferably M_1 or M_2 , wherein M_1 is a chelated metal radionuclide, and M_2 is a gamma- or positron-emitting non-metal radionuclide, optionally in combination with a prosthetic group,

m_1, m_2 are independently of each other 0, 1, 2 or 3,

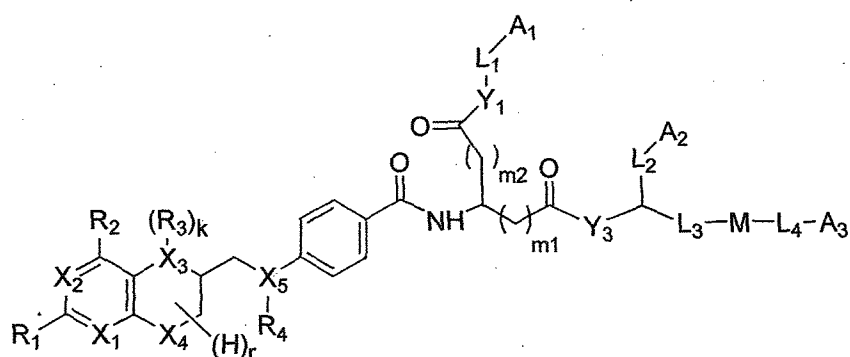
10 with the proviso that at least one, preferably one, of A_1, A_2 and A_3 is an albumin binder.

Unless specified otherwise all the definitions given hereinafter apply throughout the text (including all structural formulas).

The term "pteroate" ("pteroyl" or "pterotic") refers to compounds
15 based on a condensed pyrimidine heterocycle, which includes a pyrimidine fused with a further 5- or 6-membered heterocycle, such as for example a pteridine or a pyrrolopyrimidine bicycle. Such a pteroate may then be linked to an aminobenzoyl moiety, which may then be further derivatized in the para-position with
20 a linker of choice, such as a glutamic acid residue to give a folate. A folate is thus represented by a pteroyl-glutamic acid skeleton (more specifically N-[4(pteridin-6-ylmethylamino)benzoyl]-glutamic acid). Thus, as pteroate structures are precursors of folate structures, pteroate
25 derivatives include the analogous derivatives as those typically known for folate structures, which are for example optionally substituted folic acid, folinic acid, pteropolyglutamic acid, and folate receptor-binding pteridines such as tetrahydropterins, dihydrofolates, tetrahydrofolates, and their
30 deaza and dideaza analogs. The terms "deaza" and "dideaza"

analog refers to the art recognized analogs having a carbon atom substituted for one or two nitrogen atoms in the naturally occurring folic acid structure. For example, the deaza analogs include the 1-deaza, 3-deaza, 5-deaza, 8-deaza, and 10-deaza analogs. The dideaza analogs include, for example, 1,5-dideaza, 5,10-dideaza, 8,10-dideaza, and 5,8-dideaza analogs.

More specifically, the present invention is directed towards compounds of formula II



II

wherein

X_1 to X_5 are independently of each other C, N or O, preferably N or O,

R_1 , R_2 are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR_5$, $-COR_5$, $-COOR_5$, $-NHR_5$, $-CONHR_5$, $-CONHR_5$, wherein R_5 represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, or $-NHR'$, wherein R' is H or C(1-8)alkyl,

R_3 , R_4 are independently of each other H, nitroso, C(1-12)alkyl, $-OR'$, $-COR'$ or halosubstituted $-COR'$, wherein R' represents H or C(1-8)alkyl,

L_1, L_4 are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' , NHR' , CO_2R' or NO_2 ,

wherein R' represents H or C(1-8)alkyl,

L₂ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

L₃ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, and a five-membered

azaheterocycle, wherein R' represents H or C(1-8)alkyl,

Y₁, Y₃ are independently of each other O, N or S,

A₁, A₂, A₃ are independently of each other H, a capping group, or an albumin binder,

M is a radionuclide-based therapeutic or diagnostic moiety M₁ or M₂, wherein M₁ is a chelated metal radionuclide, and M₂ is a gamma- or positron-emitting non-metal radionuclide, optionally in combination with a prosthetic group,

m₁, m₂ are independently of each other 0, 1, 2 or 3,

k is 0 or 1, and

r has a value of 1 to 7,

with the proviso that at least one, preferably one, of A₁, A₂ and A₃ is an albumin binder.

It is understood, that the abbreviations "N" and "C" are

representative for all possible degrees of saturation, i.e. N includes -NH- and -N= linkages and C includes -CH₂- and -CH= linkages.

It is further understood, that (H)_r represents all hydrogen substituents on the indicated ring (i.e. on X₃, C6, C7 and X₄). For example r = 7 for a fully saturated 5,8-dideaza analog (X₃ = X₄ = C) and r = 1 for a fully unsaturated analog with X₃ = X₄ = N.

The term "alkyl", when used singly or in combination, refers to straight chain or branched alkyl groups containing the indicated number of carbon atoms. Thus, the term "C(1-12)alkyl" refers to a hydrocarbon radical whose carbon chain is straight-chain or branched and comprises 1 to 12 carbon atoms. Preferred alkyl groups include C(1-8)alkyl groups which refer to a hydrocarbon radical whose carbon chain is straight-chain or branched and comprises 1 to 8 carbon atoms, for example methyl, ethyl, propyl, isopropyl, butyl, sec-butyl, isobutyl, tertiary butyl, pentyl, isopentyl, neopentyl, hexyl, 2,3-dimethylbutane, neohexyl, heptyl, octyl. More preferred alkyl groups are C(1-6)alkyl groups containing one to six C-atoms, more preferably one to four carbon atoms. Optionally substituted alkylchains, such as designated by -(CHR)_x-, represent an alkyl chain having -CH₂- groups of the indicated value x, and wherein each of -CH₂- groups may independently of each other be substituted with the indicated group R. Thus, in case of multiple R-groups, the R groups may be the same or different.

The term "alkenyl", singly or in combination with other groups, refers to straight chain or branched alkyl groups as defined hereinabove having one or more carbon-carbon double bonds. Thus, the term "C(2-12)alkenyl" refers to a hydrocarbon radical whose carbon chain is straight-chain or branched and comprises 1 to 12

carbon atoms and one or more carbon-carbon double bonds. Preferred alkenyl groups include C(2-8)alkenyl groups, such as methylene, ethylene, propylene, isopropylene, butylene, t-butylene, sec-butylene, isobutylene, amylene, isoamylene, 5 pentylene, isopentylene, hexylene and the like. The preferred alkenyl groups contain two to six, more preferably two to four carbon atoms.

The term "alkynyl" as used herein refers to a linear or branched alkyl groups as defined hereinabove having one or more carbon-10 carbon triple bonds. The preferred alkynyl groups contain two to six, more preferably two to four carbon atoms.

As indicated above, the definition for "alkyl" applies both when used singly and in combination with other groups. Thus, alkoxy groups (or -O-alkyl) refer to alkyl groups as defined above, 15 substituted with an oxygen, such as methoxy, ethoxy, propoxy, isopropoxy, butoxy, tert-butoxy and the like. Alkanoyl groups as used herein include formyl and -CO-alkyl-groups, which refer to alkyl groups as defined above, terminally-substituted with a carbonyl such as acetyl, propanoyl, butanoyl, pentanoyl and the 20 like. Alkylamino groups (or -NHR-alkyl or -N(R)₂-alkyl) refer to alkyl groups as defined above, substituted with nitrogen, including both monoalkylamino such as methylamino, ethylamino, propylamino, tert-butylamino, and the like, and dialkylamino such as dimethylamino, diethylamino, methylpropylamino, and the 25 like.

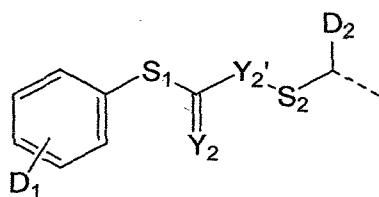
The term "halogen" or "halo" as used herein refers to any Group 7 element and includes fluoro, chloro, bromo, iodo.

The term "halosubstituted" as used herein refers to alkyl groups which have halogen moieties in the place of at least one 30 hydrogen.

In preferred embodiments, R₁ and R₂ may be independently of each other H, C(1-12)alkyl, -OR₅, -NHR₅, more preferably -OR₅, -NHR₅;

and/or R_3 is H, C(1-12)alkyl, or $-\text{CO}-\text{C}(1-8)\text{alkyl}$; and/or R_4 is H, nitroso, $-\text{O}-\text{C}(1-8)\text{alkyl}$, or $-\text{CO}-\text{C}(1-8)\text{alkyl}$.

The term "albumin binder" as used herein refers to a group, which binds noncovalently to human serum albumin (typically with
 5 a binding affinity less than about 10 μM). Albumin binding properties can be measured by surface plasmon resonance as described in J. Biol. Chem. 277(38), 35035-35042, (2002). Typical albumin binders suitable for use in the compounds of the present invention include linear and branched lipophilic groups
 10 having 12-40 carbon atoms and a distal acidic group. Suitable albumin binders for use in the compounds of the present invention are selected from compounds of formula III



III

15 wherein

Y_2, Y_2' are independently of each other N, O or S,

S_1, S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' ,
 20 NHR' , $\text{CO}_2\text{R}'$, SH, SO_3H or NO_2 , and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by $-\text{O}-$, $-\text{CO}-$, $-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-$, $-\text{NR}'-$, $-\text{NR}'-\text{CO}-$, $-\text{CO}-\text{NR}'-$, $-\text{NR}'-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-\text{NR}'-$, $-\text{NR}'-\text{CO}-\text{NR}'-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$, $-\text{O}-\text{CO}-\text{O}-$, $-\text{S}-\text{R}'-$, $-\text{SO}_3\text{R}'-$, wherein R' represents H or C(1-8)alkyl,

25 D_1 is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-\text{OR}_5$, $-\text{COR}_5$, $-\text{COOR}_5$, $-\text{NHR}_5$, $-\text{CONHR}_5$, wherein R_5 represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-

IVc

wherein

X₁ to X₅ are independently of each other C, N or O, preferably N or O,

5 Y₁, Y₃ are independently of each other N, O or S,

Y₂, Y_{2'} are independently of each other N, O or S,

R₁, R₂ are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -COR', -COOR', or -NHR',
10 wherein R' is H or C(1-8)alkyl,

R₃, R₄ are independently of each other H, nitroso, C(1-12)alkyl, -OR', -COR' or halosubstituted -COR', wherein R' represents H or C(1-8)alkyl,

15 A₁ is H or a capping group,

A₂ is H or a capping group,

A₃ is H or a capping group,

M is radionuclide-based therapeutic or diagnostic moiety M₁ or M₂, wherein M₁ is a chelated metal radionuclide, and M₂ is a
20 gamma- or positron-emitting non-metal radionuclide, optionally in combination with a prosthetic group,

m is 1, 2 or 3,

m₁, m₂ are independently of each other 0, 1, 2 or 3,

L₁, L₄ are independently of each other a covalent bond or a
25 straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl,

L₂ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

L₃ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, and a five-membered azaheterocycle, wherein R' represents H or C(1-8)alkyl,

S₁, S₂ are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D₁ is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

D₂ is an acidic group,

k is 0 or 1, and

r has a value of 1 to 7.

Groups L_1 and L_4 are independently of each other preferably a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl, preferably a covalent bond or a straight-chain or branched unsubstituted C(1-6)alkyl, most preferably a covalent bond.

Group L_2 is preferably a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

Preferably, m_1 and m_2 are not both 0. In specific embodiments m_1 is 2 and m_2 is 0, in other specific embodiments m_1 is 0 and m_2 is 2.

Y_1 is N, O, or S, preferably O; Y_3 is N, O, or S, preferably O; Y_2 is N, O, or S, preferably O; and Y_2' is N, O, or S, preferably N.

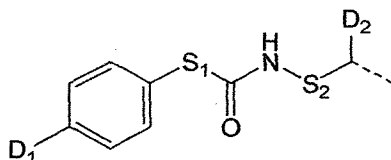
The acidic group D_2 is a group capable of ionizing to donate a hydrogen ion to a base to form a salt, and is preferably selected from the group consisting of -COOH, -SO₃H, -SO₂H, -NR'SO₃H, -P(O)(OH)₂, wherein R' represents H or C(1-8)alkyl.

Group D_1 may be in ortho-, meta- or para-position, preferably in para-position. D_1 is preferably a group selected from H, halogen, or C(1-12)alkyl, more preferably halogen, more preferably iodine (most preferably iodine in para-position).

Preferably, S_1 and S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-8)alkyl, wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -COO-, -NR'-, -NR'-CO-, -CO-NR'-, -CH=CH-, wherein R' represents H or C(1-8)alkyl.

In specific embodiments S_1 and S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-6)alkyl.

Thus, in specific embodiments the albumin binder for use in compounds of the present invention is a group of formula IIIa



IIIa

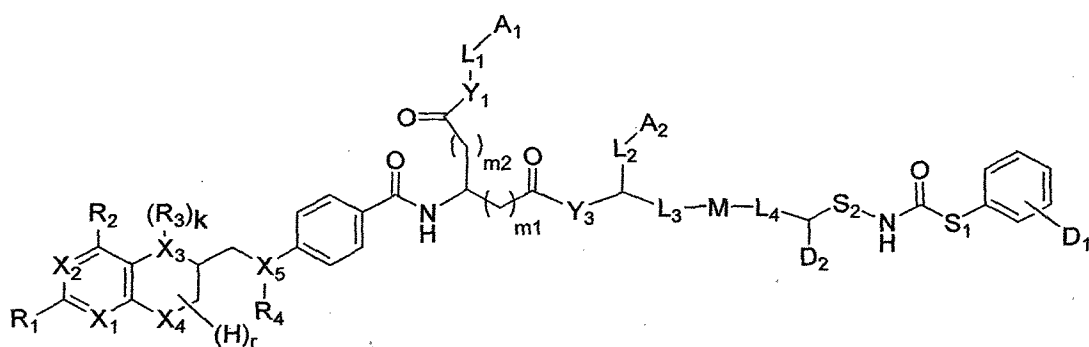
wherein

S_1 , S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-8)alkyl, wherein one or more of the non-adjacent CH_2 groups may independently be replaced by $-\text{O}-$, $-\text{CO}-$, $-\text{COO}-$, $-\text{NR}'-$, $-\text{NR}'-\text{CO}-$, $-\text{CO}-\text{NR}'-$, $-\text{CH}=\text{CH}-$, wherein R' represents H or C(1-8)alkyl, preferably a single bond or a spacer selected from a straight-chain or branched C(1-6)alkyl,

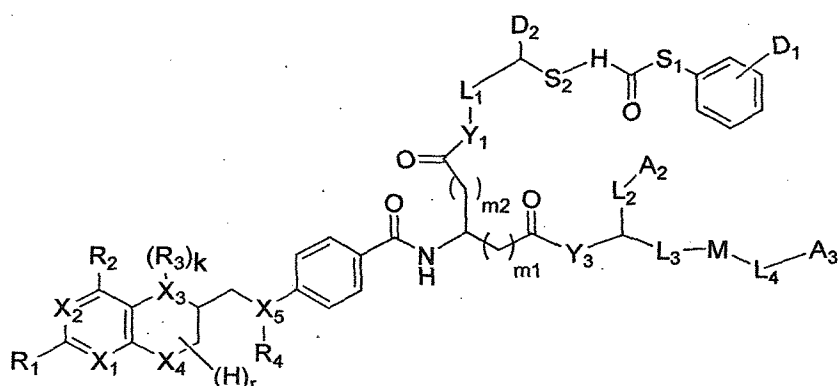
D_1 is a group selected from H, halogen, or C(1-12)alkyl, preferably halogen, more preferably iodine,

D_2 is $-\text{COOH}$, $-\text{SO}_3\text{H}$, $-\text{SO}_2\text{H}$, $-\text{NR}'\text{SO}_3\text{H}$, preferably $-\text{COOH}$, and the broken line represents the linkage to L_1 or L_2 (in compounds of formulae I or II).

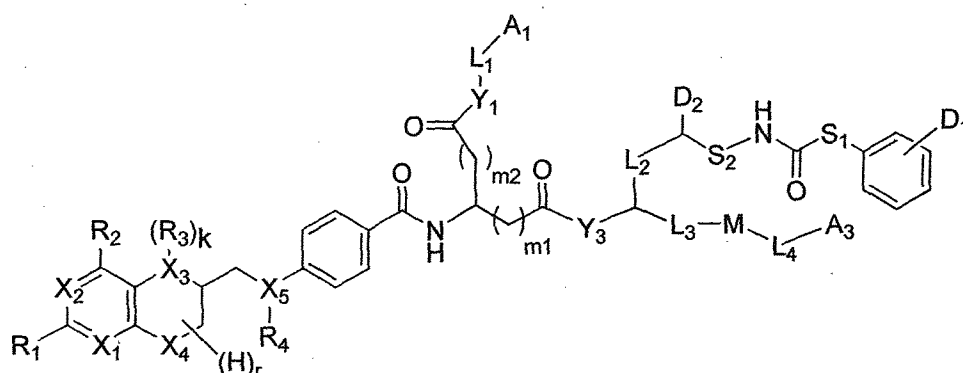
Thus, in preferred embodiments, compounds of formula IVa, IVb and IVc can be represented by compounds of formula Va, Vb and Vc



Va



Vb



Vc

wherein

X_1 to X_5 are independently of each other C, N or O, preferably N or O,

Y_1, Y_3 are independently of each other O, N or S,

R_1, R_2 are independently of each other H, halogen, C(1-12)alkyl,

C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -COR', -COOR', or -NHR', wherein R' is H or C(1-8)alkyl,

5 R₃, R₄ are independently of each other H, nitroso, C(1-12)alkyl, -OR', -COR' or halosubstituted -COR', wherein R' represents H or C(1-8)alkyl,

A₁ is H or a capping group,

A₂ is H or a capping group,

10 A₃ is H or a capping group,

M is a radionuclide-based therapeutic or diagnostic moiety M₁ or M₂, wherein M₁ is a chelated metal radionuclide, and M₂ is a gamma- or positron-emitting non-metal radionuclide, optionally in combination with a prosthetic group,

15 m is 1, 2 or 3,

m₁, m₂ are independently of each other 0, 1, 2 or 3,

L₁, L₄ are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂,

20 wherein R' represents H or C(1-8)alkyl,

L₂ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

L₃ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted

by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a five-membered azaheterocycle selected from a triazolyl or tetrazolyl group wherein R' represents H or C(1-8)alkyl,

5 S₁, S₂ are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-8)alkyl, wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -COO-, -NR'-, -NR'-CO-, -CO-NR'-, -CH=CH-
10 bond or a spacer selected from a straight-chain or branched C(1-6)alkyl,

D₁ is a group selected from H, halogen, or C(1-12)alkyl, preferably halogen, more preferably iodine

D₂ is -COOH, -SO₃H, -SO₂H, -NR'SO₃H, preferably -COOH, and the
15 broken line represents the linkage to L₁ or L₂ (in compounds of formulae I or II),

k is 0 or 1, and

r has a value of 1 to 7.

In a specific embodiment, the imaging moiety M is a chelated
20 radioimaging metal M₁ comprising a radioimaging metal ion and a metal chelator. The term "metal chelator" (or chelator) may be any of the metal chelators known in the art for complexing with a metal ion or radionuclide (and useful for the intended applications). The binding of a chelator to an ion/ radionuclide
25 may be determined by measuring the dissociation constant between chelator and ion/radionuclide. For the purposes of the present invention, the dissociation constant K_D between chelator and ion/radionuclide is from about 10⁻³ to about 10⁻¹⁵ M⁻¹. Preferably, the dissociation constant K_D between chelator and
30 ion/radionuclide is from about 10⁻⁶ to about 10⁻¹⁵ M⁻¹.

Examples of chelators are well known in the art, and include

bidentate, tridentate, and tetradentate ligands in linear, tripodal and macrocyclic form. Typical examples include bipyridyl (bipy); terpyridyl (terpy); crown ethers; aza-crown ethers; succinic acid; citric acid; salicylic acids; histidines;

5 imidazoles; ethyleneglycol-bis-(beta-aminoethyl ether) N,N'-tetraacetic acid (EGTA); nitroloacetic acid; acetylacetonate (acac); sulfate; dithiocarbamates; carboxylates; alkyldiamines; ethylenediamine (en); diethylenetriamine (dien); nitrate; nitro; nitroso; $(C_6H_5)_2PCH_2CH_2P(C_6H_5)_2$ (diphos); glyme; diglyme;

10 bis(acetylacetonate) ethylenediamine (acacen); ethylenediaminetetraacetic acid (EDTA), diethylenetriaminopentaacetic acid (DTPA); N-[2-[bis(carboxymethyl)amino]-3-(4-ethoxyphenyl)propyl]-N-[2-[bis(carboxymethyl)-amino]ethyl]-L-glycine (EOB-DTPA); N,N-

15 bis[2-[bis(carboxymethyl)amino]-ethyl]-L-glutamic acid (DTPA-Glu); N,N-bis[2-[bis(carboxymethyl)amino]-ethyl]-L-lysine (DTPA-Lys); mono- or bis-amide derivatives of DTPA such as N,N-bis[2-[carboxymethyl[(methylcarbamoyl)methyl]amino]-ethyl]glycine (DTPA-BMA); 4-carboxy-5,8,11-tris(carboxymethyl)-1-phenyl-2-oxa-

20 5,8,11-triazamidecan-13-oic acid (BOPTA); DTPA BOPTA, 1,4,7,10-tetraazacyclododecan-1,4,7-triacetic acid (DO3A); 1,4,7,10-tetraazacyclododecan-1,4,7,10-tetraactetic acid (DOTA); 10-(2-hydroxypropyl)-1,4,7,10-tetraazacyclododecan-1,4,7-triacetic acid (HPDO3A); 2-methyl-1,4,7,10-tetraazacyclododecan-1,4,7,10-

25 tetraacetic acid (MCTA); tetramethyl-1,4,7,10-tetraazacyclododecan-1,4,7,10-tetraacetic acid (DOTMA); 3,6,9,15-tetraazabicyclo[9.3.1]pentadeca-1(15); 11,13-triene-3,6,9-triacetic acid (PCTA); PCTA12; cyclo-PCTA12; N,N'-Bis(2-aminoethyl)-1,2-ethanediamine (TETA); 1,4,7,10-

30 tetraazacyclotridecane-N,N',N'',N'''-tetraacetic acid (TRITA); 1,12-dicarbonyl, 15-(4-isothiocyanatobenzyl) 1,4,7,10,13-pentaazacyclohexadecane-N,N',N'' triacetic acid (HETA); 1,4,7,10-tetraazacyclododecane-N,N',N'',N'''-tetraacetic acid mono-(N-

hydroxysuccinimidyl) ester (DOTA-NHS); N,N'-Bis(2-aminoethyl)-1,2-ethanediamine-N-hydroxy-succinimide ester (TETA-NHS); [(2S,5S,8S,11S)-4,7,10-tris-carboxymethyl-2,5,8,11-tetramethyl-1,4,7,10-tetraazacyclododecan-1-yl]acetic acid (M4DOTA);

5 [(2S,5S,8S,11S)-4,7-bis-carboxymethyl-2,5,8,11-tetramethyl-1; 4,7,10-tetraazacyclo-dodecan-1-yl]acetic acid; (M4DO3A); (R)-2-[(2S,5S,8S,11S)-4,7,10-tris-((R)-1-carboxyethyl)-2,5,8,11-tetramethyl-1,4,7,10-tetraazacyclododecan-1-yl]propionic acid (M4DOTMA); 10-phosphonomethyl-1,4,7,10-tetraazacyclododecane-

10 1,4,7-triacetic acid (MPDO3A); hydroxybenzyl-ethylenediamine-diacetic acid (HBED) and N,N'-ethylenebis-[2-(o-hydroxyphenolic)glycine] (EHPG).

Suitable metal chelators for use in the compounds of the present invention include bidentate, tridentate, and tetradentate,

15 ligands in linear, tripodal and macrocyclic form, as identified hereinabove. In specific embodiments, the metal chelators used for the present invention include linear or macrocyclic polyaminocarboxylates, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, EHPG, HBED, NOTA, DOTMA, TETMA, PDTA, TTHA, LICAM, MECAM,

20 preferably macrocyclic polyaminocarboxylates, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA.

The metal chelator may or may not be complexed with a radio imaging metal ion or radionuclide, and may include an optional spacer such as a single amino acid. Typical metal radionuclides

25 for scintigraphy or radionuclide therapy include ^{99}Tc , ^{51}Cr , ^{67}Ga , ^{68}Ga , ^{47}Sc , ^{51}Cr , ^{167}Tm , ^{141}Ce , ^{111}In , ^{168}Yb , ^{175}Yb , ^{140}La , ^{90}Y , ^{88}Y , ^{153}Sm , ^{166}Ho , ^{165}Dy , ^{166}Dy , ^{62}Cu , ^{64}Cu , ^{67}Cu , ^{97}Ru , ^{103}Ru , ^{186}Re , ^{188}Re , ^{203}Pb , ^{211}Bi , ^{212}Bi , ^{213}Bi , ^{214}Bi , ^{105}Rh , ^{109}Pd , ^{117}mSn , ^{149}Pm , ^{161}Tb , ^{155}Tb , ^{152}Tb , ^{149}Tb , ^{177}Lu , ^{198}Au and ^{199}Au . The choice of metal will

30 be determined based on the desired therapeutic or diagnostic application. For example, for diagnostic purposes radionuclides may include ^{64}Cu , ^{67}Ga , ^{68}Ga , $^{99\text{m}}\text{Tc}$, and ^{111}In , while for therapeutic purposes, radionuclides may include ^{64}Cu , ^{90}Y , ^{105}Rh

^{111}In , $^{117\text{m}}\text{Sn}$, ^{149}Pm , ^{153}Sm , ^{161}Tb , ^{166}Dy , ^{166}Ho , ^{175}Yb , ^{177}Lu , $^{186/188}\text{Re}$, and ^{199}Au . A skilled person will know which radionuclide to choose for the intended application. Preferred radionuclides include ^{177}Lu , ^{161}Tb , ^{213}Bi , and ^{111}In .

5 In another specific embodiment, the imaging moiety M is a gamma- or positron-emitting radioimaging non-metal M_2 selected from ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F , ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I . Preferred are positron-emitting radioimaging non-metal such as ^{11}C , ^{13}N , ^{18}F , more preferably ^{18}F . A radioimaging non-metal of choice may be in
10 combination with a prosthetic group, i.e. it may be linked directly or via a suitable prosthetic group, such as for example a benzoate derivative or a saccharide group, to its neighbouring group (i.e. groups L_3 and/or L_4).

The term "saccharide group" encompasses both cyclic
15 monosaccharides and cyclic oligosaccharides based on cyclic saccharide unit(s). The term "saccharide unit" as used herein refers to cyclic saccharide units which refer to intracellular cyclic hemiacetal or hemiketal forms of a linear (mono-/oligo-) saccharide. A monosaccharide comprises one saccharide unit,
20 whereas an oligosaccharide refers to a chain of saccharide units and comprises preferably 2 to 20 saccharide units, preferably 2 to 10 saccharide units, more preferably mono-, di-, and trisaccharides. An oligosaccharide may be linear or branched and the saccharide units within the oligosaccharide are linked to
25 each other by alpha- or beta (1-2), (1-4), or (1-6) linkages. Preferably the oligosaccharide of choice is linear, and more preferably the oligosaccharide is linear and the saccharide units within the oligosaccharide are linked by alpha - or beta (1-4) bonds. In the most preferred embodiment, the
30 oligosaccharide is linear and the saccharide units within the oligosaccharide are linked by alpha (1-4) bonds.

Preferably a saccharide unit is a pyranoside or a furanoside and

natural and synthetic derivatives thereof, preferably a pyranoside selected from allose, altrose, glucose, mannose, gulose, idose, galactose, talose and fucose, or a furanoside selected from ribose, arabinose, xylose, and lyxose. The term
5 derivative refers to any chemically or enzymatically modified monosaccharide unit, including those obtained by oxidation, deoxygenation, replacement of one or more hydroxyl groups by preferably a hydrogen atom, a halogen atom, an amino group or thiol group, etc., as well as alkylation, acylation, sulfation
10 or phosphorylation of hydroxy groups or amino groups. Preferred saccharide units of the present invention include for example glucose and galactose.

Thus in one specific embodiment, the saccharide group is a monosaccharide selected from ribose, arabinose, xylose, lyxose,
15 allose, altrose, glucose, mannose, gulose, idose, galactose, talose, fucose, preferably glucose and galactose.

In another specific embodiment, the saccharide group is an oligosaccharide comprising at least two, preferably 2 to 20 saccharide units which are identical or different and each
20 selected from the group consisting of ribose, arabinose, xylose, lyxose, allose, altrose, glucose, mannose, gulose, idose, galactose, talose, fucose, preferably glucose and galactose.

In more specific embodiments an oligosaccharide may be (a) a disaccharide, e.g. lactose, maltose, isomaltose, cellobiose, gentiobiose, melibiose, primeverose, rutinose; (b) a
25 disaccharide homologue, e.g. maltotriose, isomaltotriose, maltotetraose, isomaltotetraose, maltopentaose, maltohexaose, maltoheptaose, lactotriose, lactotetraose; (c) a uronic acid, e.g. glucuronic acid, galacturonic acid; (d) a branched
30 oligosaccharide, e.g. panose, isopanose; (e) an amino monosaccharide, e.g. galactosamine, glucosamine, mannosamine, fucosamine, quinovosamine, neuraminic acid, muramic acid, lactosdiamine, acosamine, bacillosamine, daunosamine,

desosamine, forosamine, garosamine, kanosamine, kansosamine, mycaminose, mycosamine, perosamine, pneumosamine, purpurosamine, rhodosamine; (f) a modified saccharide, e.g. abequose, amicetose, arcanose, ascarylose, boivinose, chacotriose, 5 chalcose, cladinose, colitose, cymarose, 2-deoxyribose, 2-deoxyglucose, diginose, digitalose, digitoxose, evalose, evernitrose, hamamelose, manninotriose, melibiose, mycarose, mycinose, nigerose, noviose, oleandrose, paratose, rhodinode, rutinose, sarmentose, sedoheptulose, solatriose, sophorose, 10 streptose, turanose, tyvelose.

In a more preferred embodiment, the saccharide group is a monosaccharide or an oligosaccharide, thus comprising one or more of the (same or different) saccharide unit(s) which is (are) selected from the group consisting of glucose, galactose, 15 glucosamine, galactosamine, glucuronic acid, gluconic acid, galacturonic acid, lactose, lactotetraose, maltose, maltotriose, maltotetraose, isomaltose, isomaltotriose, isomaltotetraose, and neuraminic acid.

It is understood that all isomers, including enantiomers, 20 diastereoisomers, rotamers, tautomers and racemates of the compounds of the invention are contemplated as being part of this invention. The invention includes stereoisomers in optically pure form and in admixture, including racemic mixtures. Isomers can be prepared using conventional techniques, 25 either by reacting optically pure or optically enriched starting materials or by separating isomers of a compound of formula I. This applies specifically to saccharide groups, or any other groups, e.g. amino acid groups present in a compound of formula I (and subsequent formulas), which may be present in the natural 30 L- or non-natural D-form, i.e. the glutamic acid portion (or derivatives thereof).

The term "capping group" (or terminal groups) as used herein

refers to a moiety attached to the functional groups L_1 , L_2 or L_4 , which is otherwise H or linked to an albumin binder. More specifically such capping groups represent suitable protecting groups for (i) Y_1 if L_1 represents a covalent bond, (ii) M if L_4 represents a covalent bond, and (iii) L_2 if L_2 is a functional group or carries a terminal functional group. These protecting groups depend on the nature of the functional group (typically an amino, carboxyl or hydroxy function) and thus are variable. Suitable protecting groups for amino functions include e.g. the t-butoxycarbonyl, the benzyloxycarbonyl, allyloxycarbonyl, methoxy- or ethoxycarbonyl, 2,2,2-trichloroethoxycarbonyl, acetyl or trifluoroacetyl, benzyl or 2,4,6-trimethoxybenzyl, the phthalolyl group, and the trityl or tosyl protecting group. Suitable protecting groups for the carboxyl function include e.g. silyl groups and alkyl, aryl or arylalkyl esters, more specifically alkyl esters such as methyl and t-butyl; alkoxyalkyl such as methoxymethyl; alkyl thioalkyl esters such as methyl, thiomethyl; haloalkyl esters such as 2,2,2-trichloroethyl and aralkyl ester, such as benzyl, p-methoxybenzyl, p-nitrobenzyl, diphenylmethyl. Suitable protecting groups for the hydroxy function include e.g. alkyl esters, t-butyl, benzyl or trityl groups, including methyl ethers, substituted methyl ethers (e.g., MOM (methoxymethyl ether), MTM (methylthiomethyl ether), BOM (benzyloxymethyl ether), PMBM or MPM (p-methoxybenzyloxymethyl ether)), substituted ethyl ethers, substituted benzyl ethers, silyl ethers (e.g., TMS (trimethylsilyl ether), TES (triethylsilyl ether), TIPS (triisopropylsilyl ether), TBDMS (tert-butyldimethylsilyl ether), tribenzyl silyl ether, TBDPS (tert-butyldiphenylsilyl ether)). The present invention is not intended to be limited to these protecting groups; rather, a variety of additional equivalent protecting groups can be readily identified and utilized in the present invention. The above and further

protecting groups as well as techniques to introduce and remove them are described in "Protective Groups in Organic Synthesis" Third Ed. Greene, T. W. and Wuts, P. G., Eds., John Wiley & Sons, New York: 1999, the entire contents of which are hereby
5 incorporated by reference.

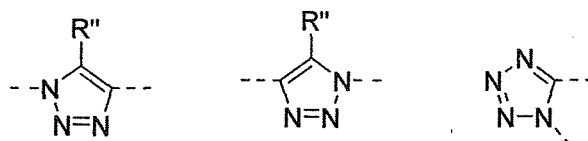
In specific embodiments Y_1 is O and L_1 is a covalent bond, and thus A_1 may be H or a carboxyl protecting group. Likewise, if M is a polyaminocarboxylate and L_4 is a covalent bond, then A_3 may be H or a carboxyl protecting group. If L_2 is a carboxyl group
10 (i.e. a Cl-alkyl, wherein the CH_2 group was replaced by $-COO-$), then A_2 may be H or a carboxyl protecting group. Likewise if M is a gamma- or positron-emitting non-metal radionuclide in combination with a prosthetic group such as a saccharide group, then A_3 may be a hydroxyl protecting group.

15 Group L_3 is preferably is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR' , NHR' , or $COOR'$, and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by a group selected from $-O-$, $-CO-$, $-$
20 $CO-O-$, $-O-CO-$, $-NR'-$, $-NR'-CO-$, $-CO-NR'-$, $-CH=CH-$, and a five-membered azaheterocycle, wherein R' represents H or C(1-8)alkyl.

More preferably, group L_3 is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR' , NHR' , or CO_2R' , and wherein one or more of the non-
25 adjacent CH_2 groups may independently be replaced by a five-membered azaheterocycle selected from a triazolyl or tetrazolyl group wherein R' represents H or C(1-8)alkyl.

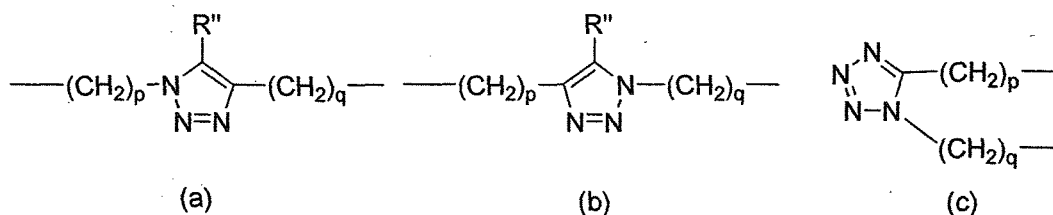
The term "azaheterocycle" refers to a heterocyclic group, which includes at least one nitrogen atom in a ring and may be
30 unsubstituted or substituted. The azaheterocyclic group may also be substituted as recognized in the art, e.g. by a C(1-6)alkyl.

For use in the present compound, the five-membered azaheterocyclic group is preferably a triazolyl or tetrazolyl group, more preferably a group of the following structures



- 5 wherein the dotted lines represent linking sites to the adjacent $-\text{CH}_2-$ groups within L_3 and R'' is H or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, or NO_2 .

Thus in preferred embodiments L_3 is a group L_3' , which is
 10 straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR' , NHR' , or $\text{CO}_2\text{R}'$, and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by a group selected from $-\text{O}-$, $-\text{CO}-$, $-\text{CO}-\text{O}-$, $-\text{NR}'-$, $-\text{NR}'-\text{CO}-$, or L_3 is a group of formulae (a), (b), or
 15 (c)



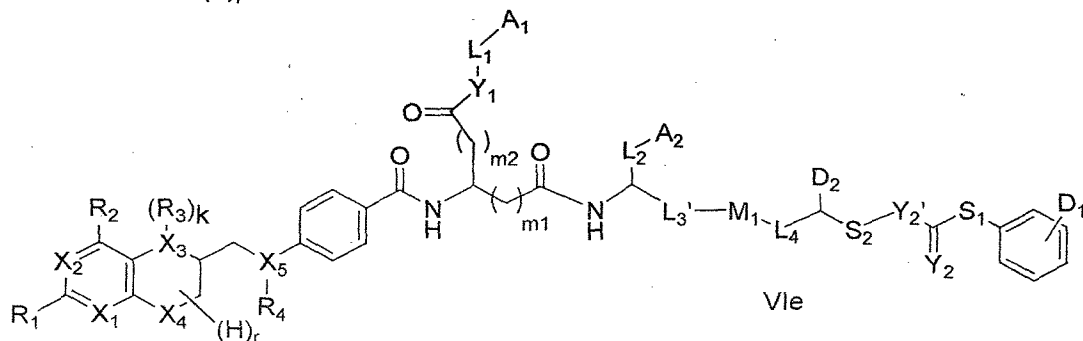
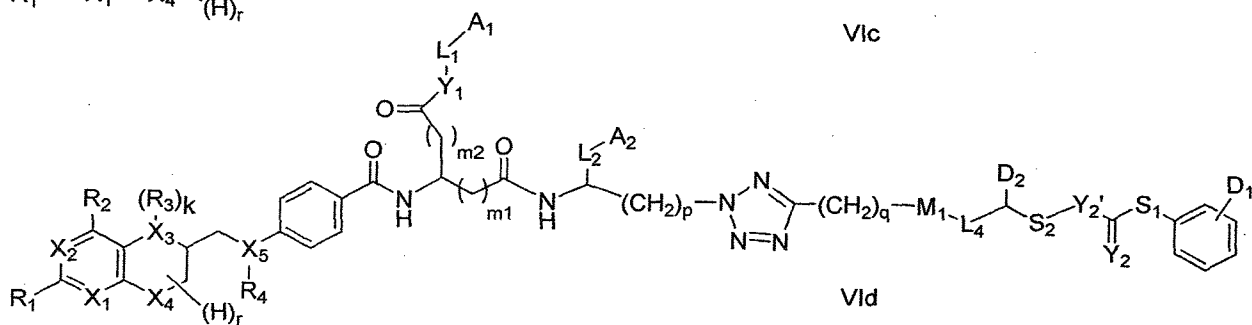
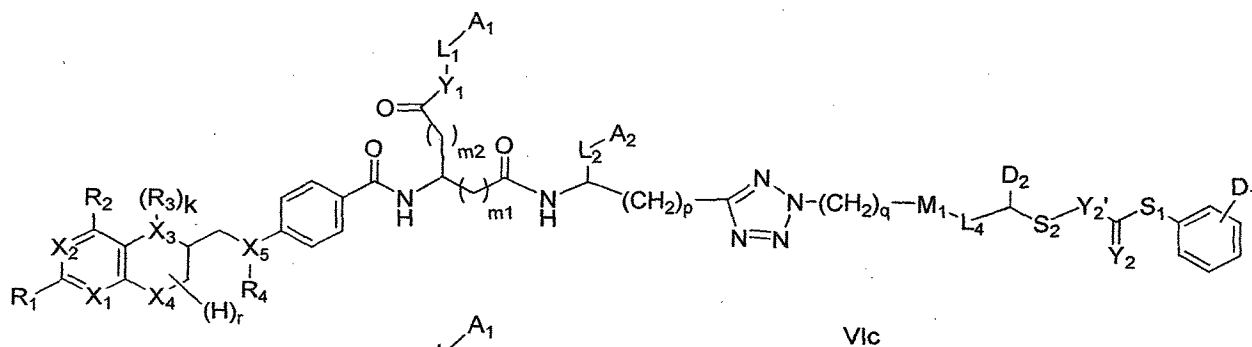
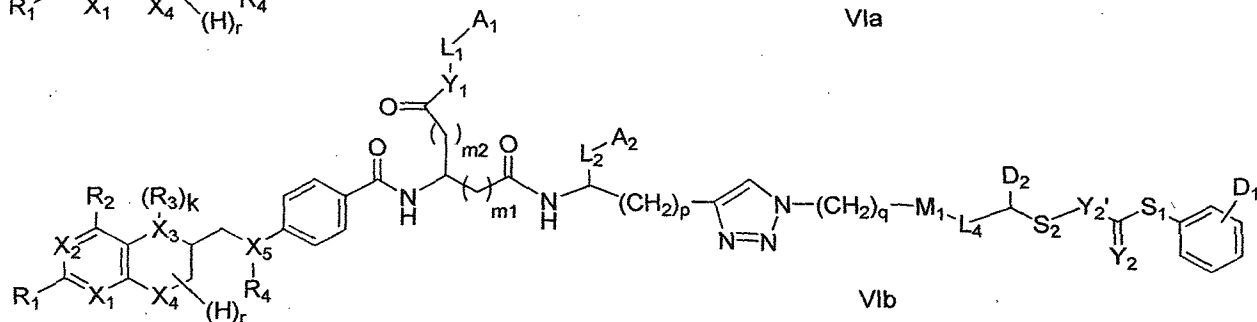
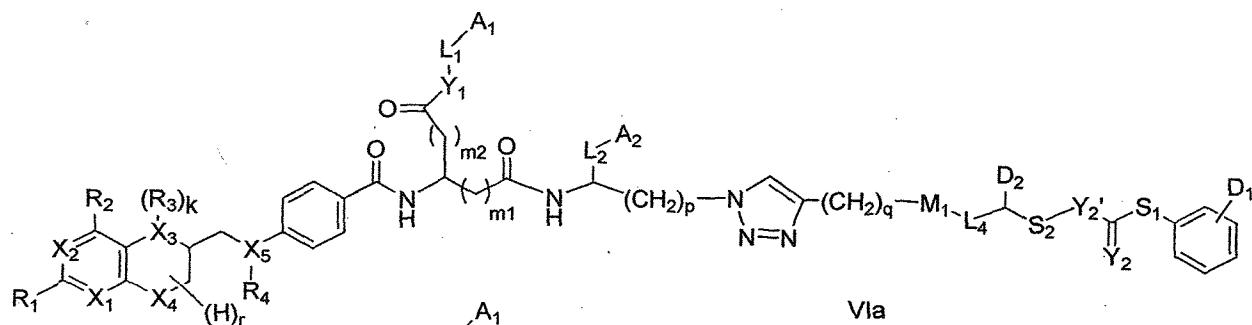
wherein

R'' is H or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, or NO_2 , and
 20 p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6.

In preferred embodiments, L_3' is straight-chain or branched C(1-6)alkyl, which is unsubstituted or substituted by at least one OH, NH_2 , or COOH, preferably COOH, and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by a group

selected from -CO-O- or -CO-NH-.

In some embodiments the present invention provides compounds of formula I having formulae VI a-e



wherein

X₁ to X₅ are independently of each other C, N or O, preferably N or O,

Y₁, Y₂, Y_{2'} are independently of each other N, O or S,

5 R₁, R₂ are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -COR', -COOR', or -NHR', wherein R' is H or C(1-8)alkyl,

10 R₃, R₄ are independently of each other H, nitroso, C(1-12)alkyl, -OR', -COR' or halosubstituted -COR', wherein R' represents H or C(1-8)alkyl,

A₁ is H or a capping group,

A₂ is H or a capping group,

15 M₁ is a linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a radioimaging metal ion,

m₁, m₂ are independently of each other 0, 1, 2 or 3,

20 L₁, L₄ are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl,

L₂ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted
25 by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

S₁, S₂ are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl

D₁ is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

D₂ is an acidic group,

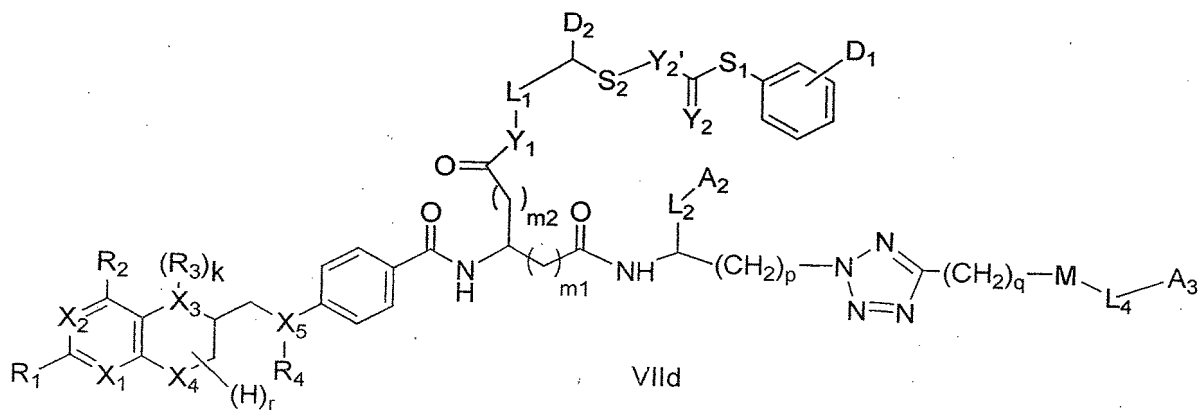
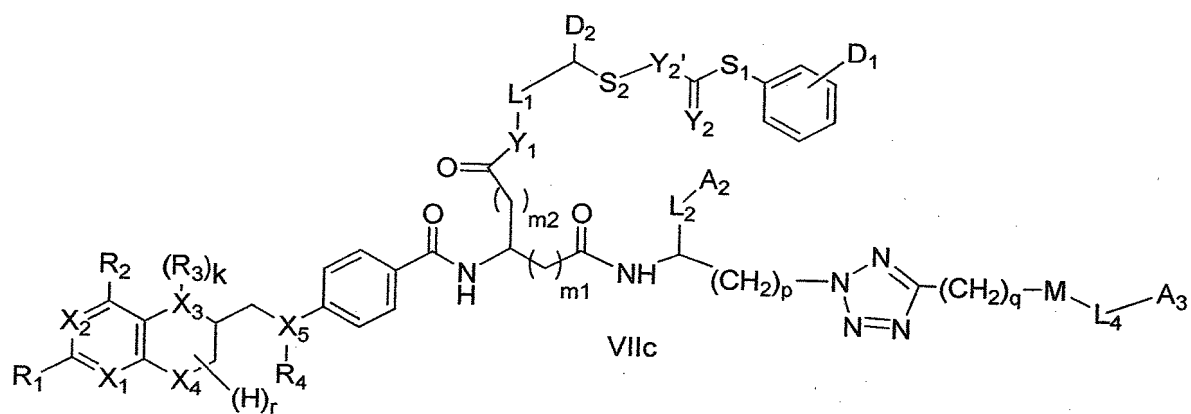
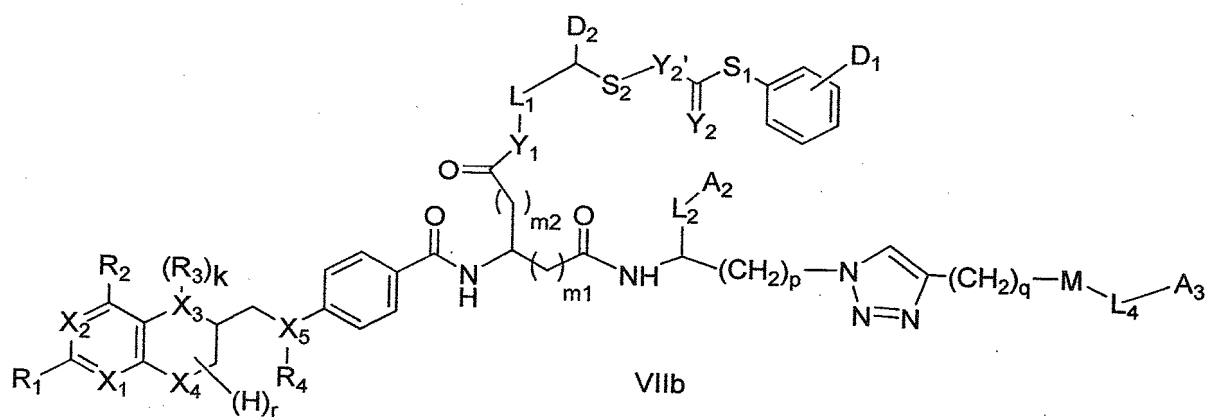
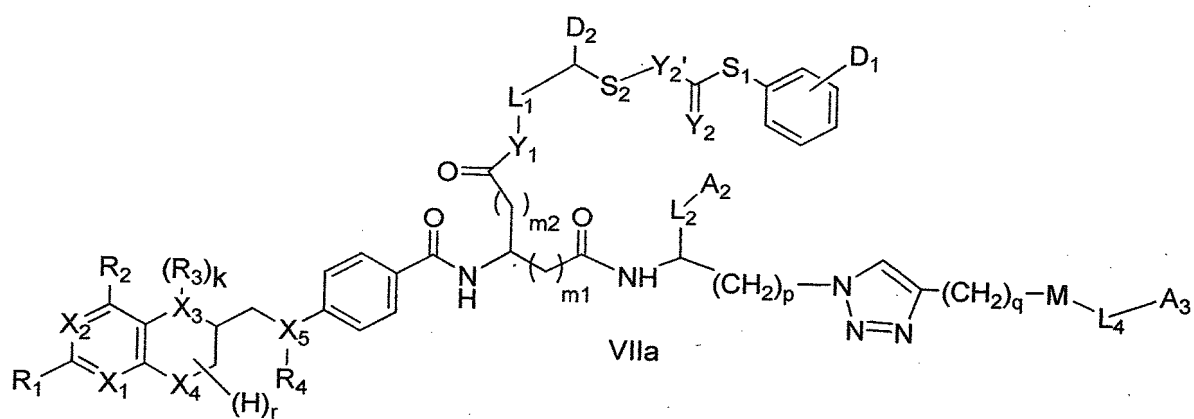
L₃ is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-, wherein R' represents H or C(1-8)alkyl,

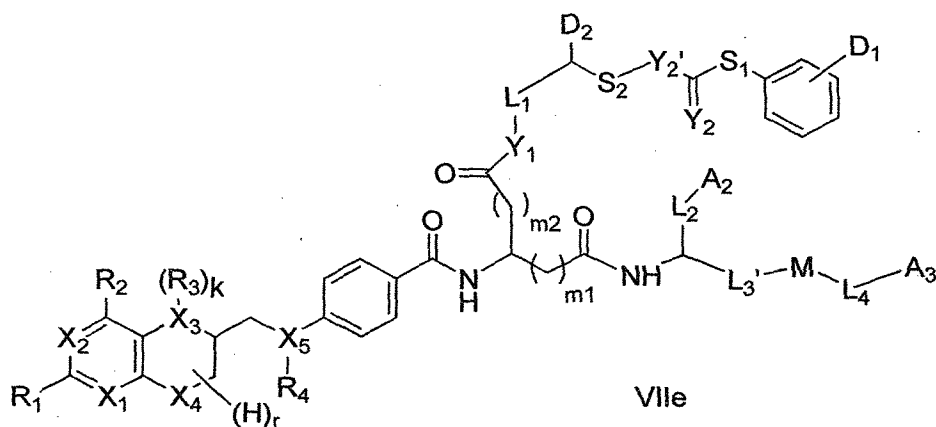
p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6,

k is 0 or 1, and

r has a value of 1 to 7.

In yet further embodiments the present invention provides compounds of formula I having formulae VII a-e





wherein

X₁ to X₅ are independently of each other C, N or O, preferably N or O.

5 Y_1, Y_2, Y_2' are independently of each other $N, 0$ or S ,

R₁, R₂ are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -COR', -COOR', or -NHR', wherein R' is H or C(1-8)alkyl,

R₃, R₄ are independently of each other H, nitroso, C(1-12)alkyl, -OR', -COR' or halosubstituted -COR', wherein R' represents H or C(1-8)alkyl.

A_2 is H or a capping group,

15 A_3 is H or a capping group,

M is a radionuclide-based therapeutic or diagnostic moiety M_1 or M_2 , wherein M_1 is a chelated metal radionuclide selected from a linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a metal radionuclide, and wherein M_2 is a gamma- or positron-emitting radioimaging non-metal radionuclide, preferably selected from ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F , ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I , optionally

in combination with a prosthetic group,

m_1 , m_2 are independently of each other 0, 1, 2 or 3,

L_1 , L_4 are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl,

L_2 is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

S_1 , S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D_1 is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

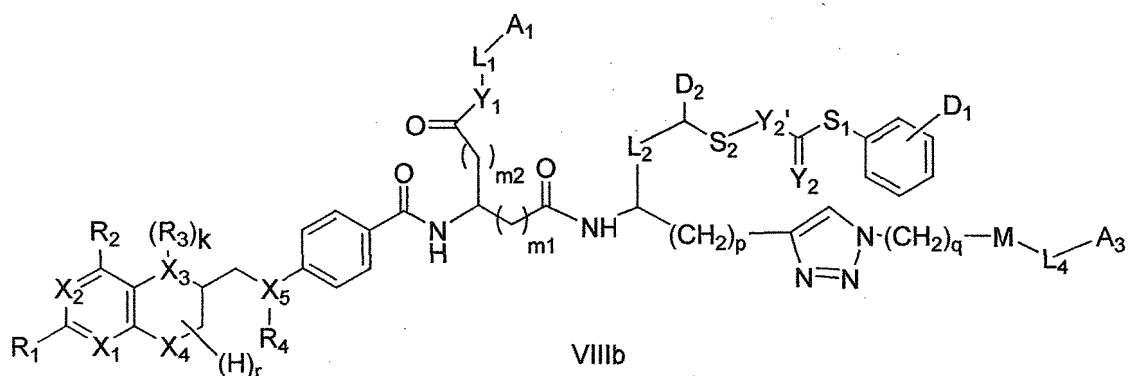
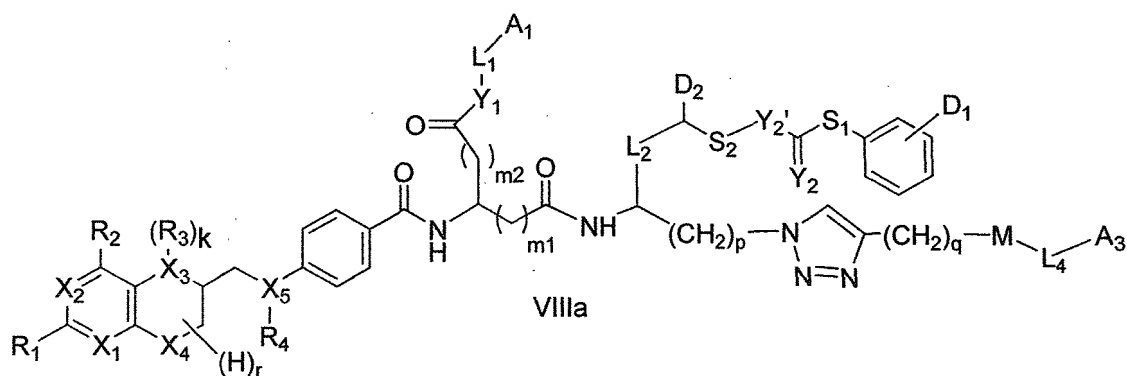
D_2 is an acidic group,

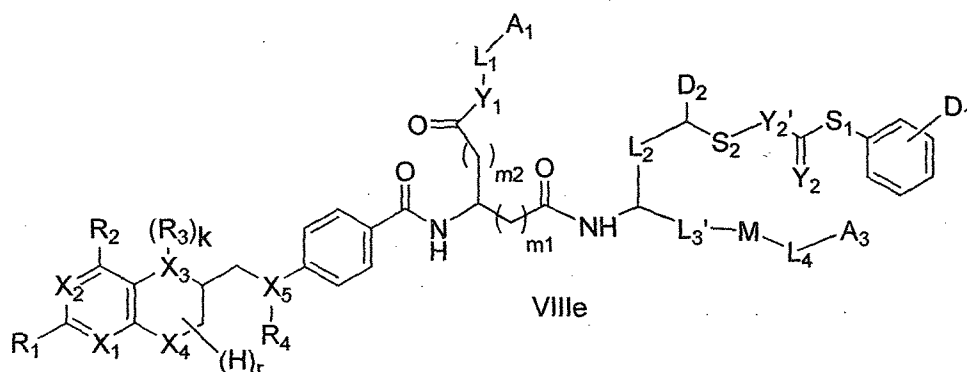
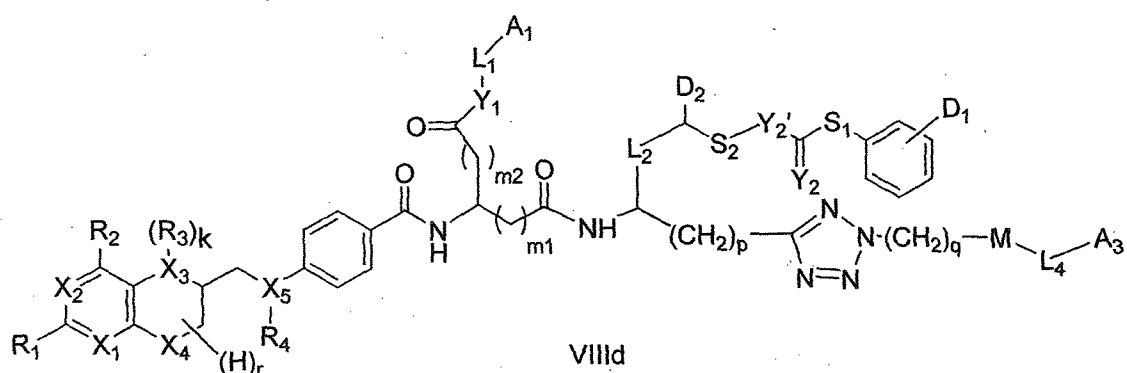
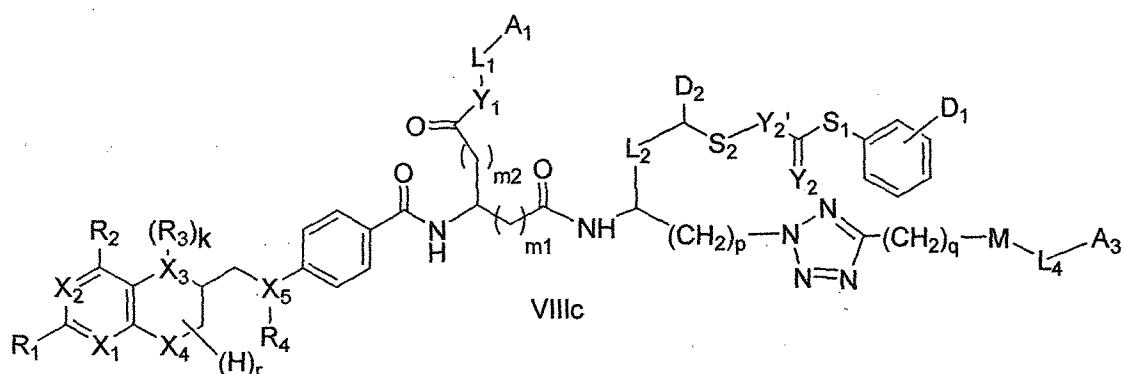
L_3 is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -

CO-O-, -NR'-, -NR'-CO-, wherein R' represents H or C(1-8)alkyl,
p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6,
k is 0 or 1,
r has a value of 1 to 7.

5

In yet further embodiments the present invention provides compounds of formula I having formulae VIII a-e





wherein

X_1 to X_5 are independently of each other C, N or O, preferably N or O,

Y_1, Y_2, Y_2' are independently of each other N, O or S,

R_1, R_2 are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR_5$, $-COR_5$, $-COOR_5$, $-NHR_5$, $-CONHR_5$, wherein R_5 represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, or $-NHR'$, wherein R' is H or C(1-8)alkyl,

R_3 , R_4 are independently of each other H, nitroso, C(1-12)alkyl, -OR', -COR' or halosubstituted -COR', wherein R' represents H or C(1-8)alkyl,

A_1 is H or a capping group,

5 A_3 is H or a capping group,

M is a radionuclide-based therapeutic or diagnostic M_1 or M_2 , wherein M_1 is a chelated metal radiometal selected from a linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a radionuclide
10 metal, and wherein M_2 is a gamma- or positron-emitting non-metal radionuclide, preferably selected from ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F , ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I , optionally in combination with a prosthetic group,

m_1 , m_2 are independently of each other 0, 1, 2 or 3,

15 L_1 , L_4 are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl,

L_2 is a covalent bond or a linking group, such as straight-chain
20 or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

25 S_1 , S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -

CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D₁ is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

D₂ is an acidic group,

L₃ is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-, wherein R' represents H or C(1-8)alkyl,

p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6,

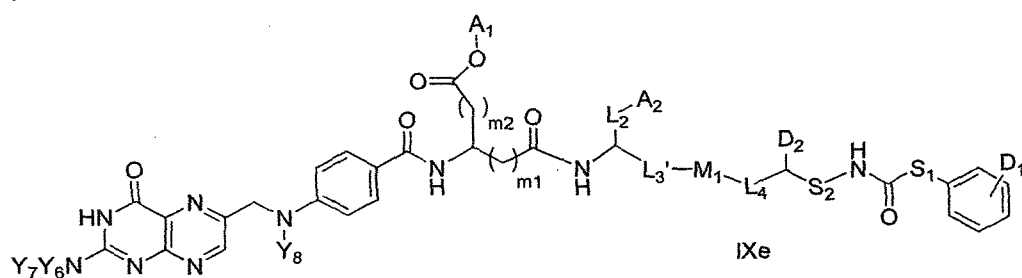
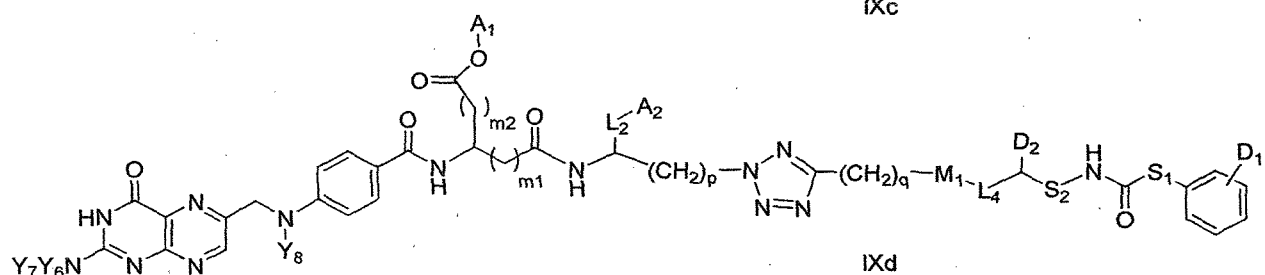
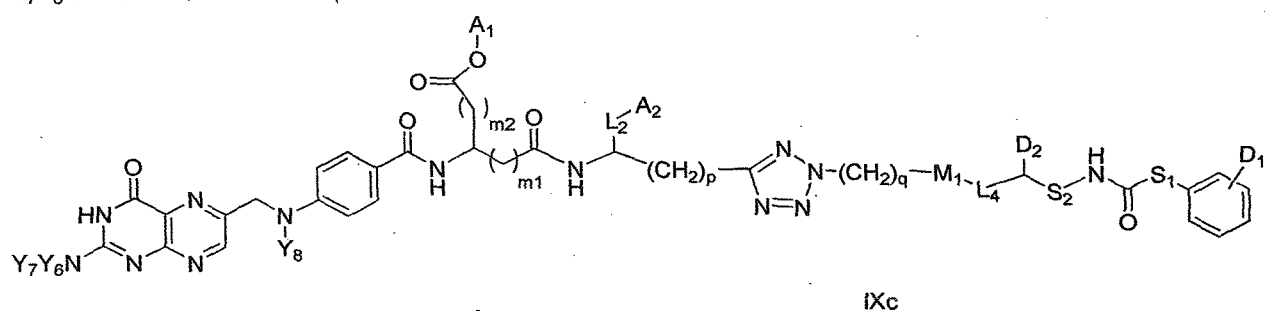
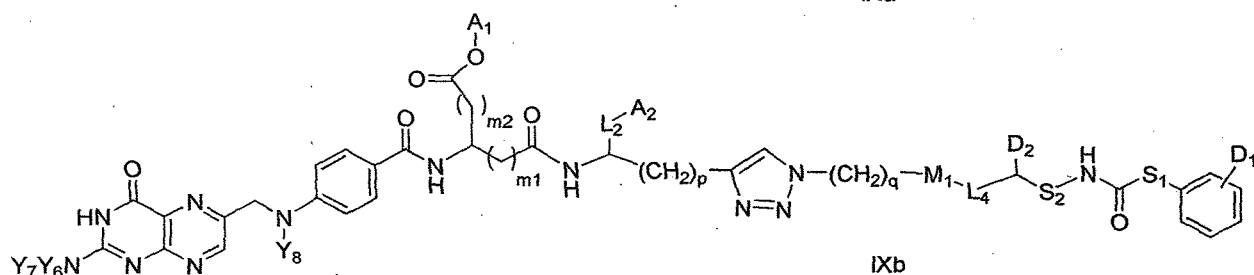
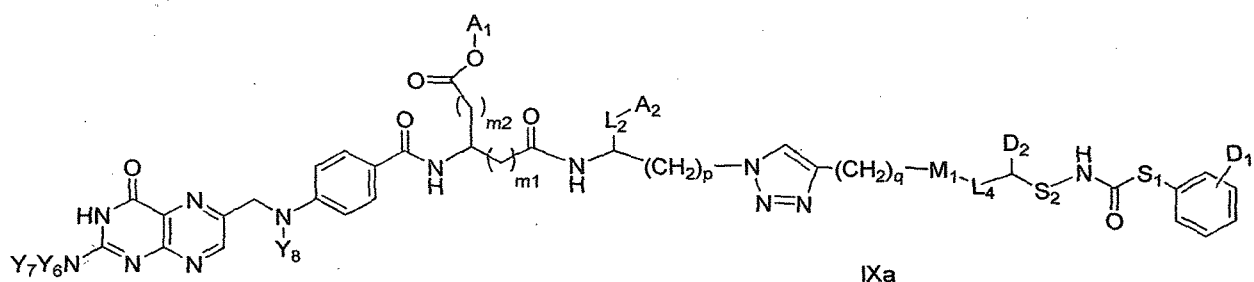
k is 0 or 1,

r has a value of 1 to 7.

In preferred embodiments, Y₂ is O and Y₂' is NH.

Further preferred embodiments of the compounds of the invention include for example compounds wherein X₁ to X₅ are N, R₁ is NY₆Y₇, R₂ is O, R₄ is Y₈, k is 0 and r is 1.

Thus the invention contemplates more specifically compounds of formulas IX a-e, wherein Y₁, Y₂ are O, Y₃, Y₂' are NH and L₁ is preferably a covalent bond,



5 wherein,

Y_6 , Y_7 are independently of each other selected from H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -COR', -COOR', and -NHR', wherein R' is H or C(1-8)alkyl,

Y_8 is selected from H, nitroso, C(1-12)alkyl, -OR', -COR', and

halosubstituted $-\text{COR}'$, wherein R' is H or C(1-12)alkyl,

A_1 is H or a carboxy protecting group,

A_2 is H or capping group

M_1 is a linear or macrocyclic polyaminocarboxylate, such as DTPA,

5 DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a radioimaging metal ion,

m_1 , m_2 are independently of each other 0, 1, 2 or 3, preferably

m_1 is 0 and m_2 is 2 or m_1 is 2 and m_2 is 0,

10 L_2 is a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR' , NHR' , or $\text{CO}_2\text{R}'$, and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by a group selected from $-\text{O}-$, $-\text{CO}-$, $-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-$, $-\text{NR}'-$, $-\text{NR}'-\text{CO}-$, $-\text{CO}-\text{NR}'$, wherein R' represents H or C(1-8)alkyl,

15 L_4 is a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' , NHR' , $\text{CO}_2\text{R}'$ or NO_2 , wherein R' represents H or C(1-8)alkyl,

20 S_1 , S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' , NHR' , $\text{CO}_2\text{R}'$, SH, SO_3H or NO_2 , and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by $-\text{O}-$, $-\text{CO}-$, $-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-$, $-\text{NR}'-$, $-\text{NR}'-\text{CO}-$, $-\text{CO}-\text{NR}'-$, $-\text{NR}'-\text{CO}-\text{O}-$, $-\text{O}-\text{CO}-\text{NR}'-$, $-\text{NR}'-\text{CO}-\text{NR}'-$, $-\text{CH}=\text{CH}-$, $-\text{C}\equiv\text{C}-$, $-\text{O}-\text{CO}-\text{O}-$, $-\text{S}-\text{R}'-$, $-\text{SO}_3\text{R}'-$, wherein R' represents H or C(1-8)alkyl,

D_1 is a group selected from H, halogen, C(1-12)alkyl, preferably halogen,

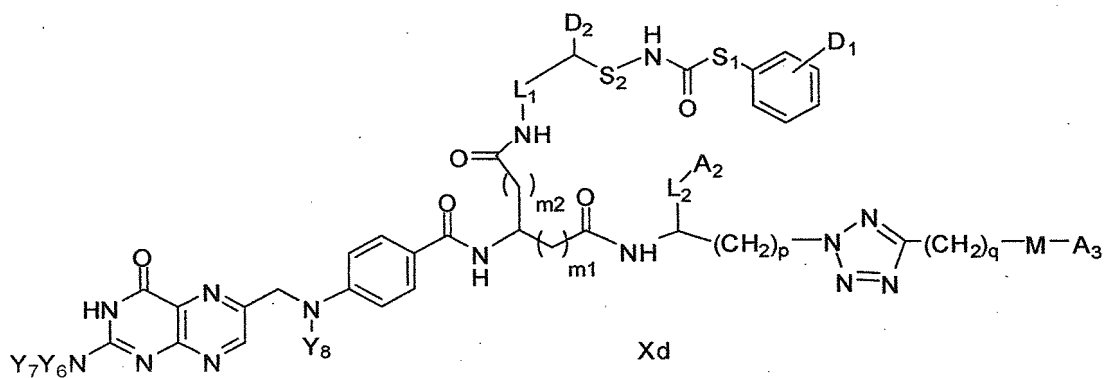
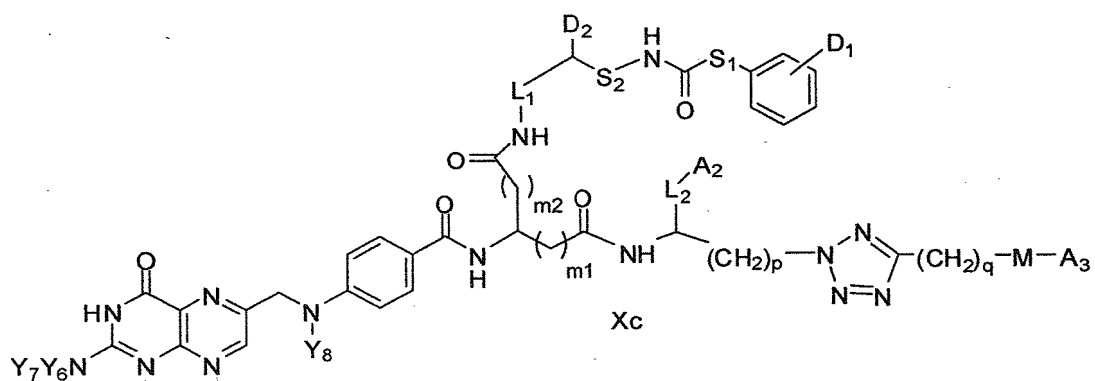
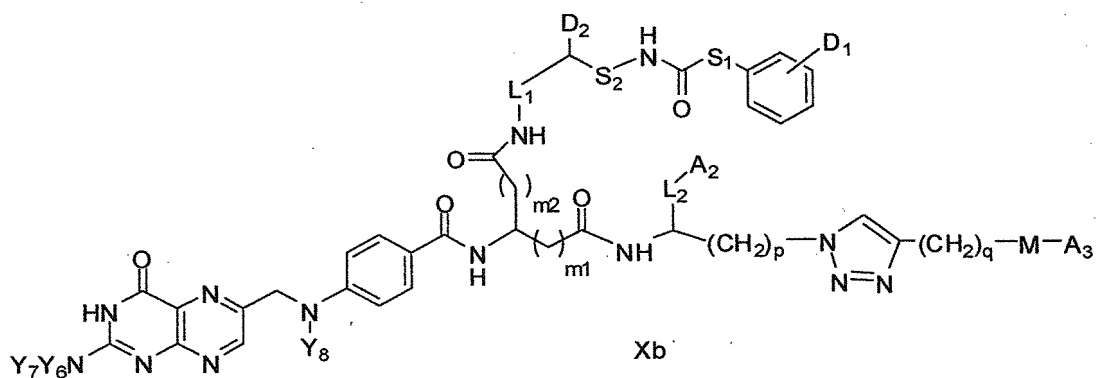
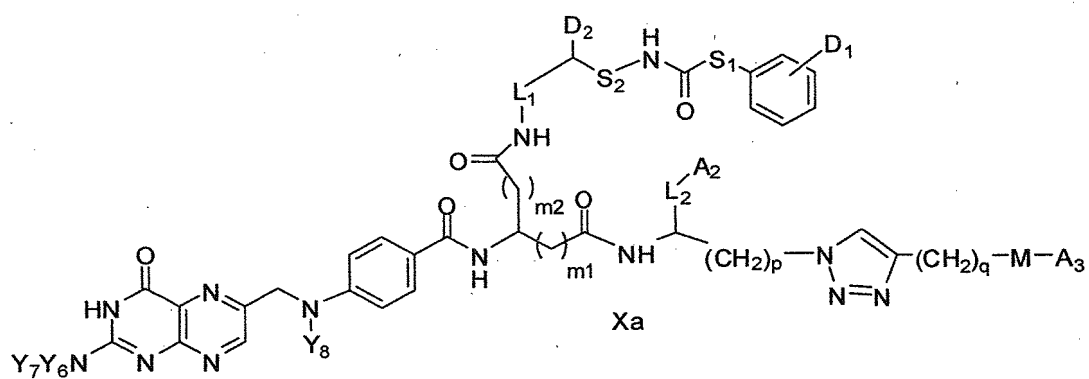
D_2 is an acidic group selected from $-\text{COOH}$, $-\text{SO}_3\text{H}$, $-\text{SO}_2\text{H}$, $-\text{NR}'\text{SO}_3\text{H}$,

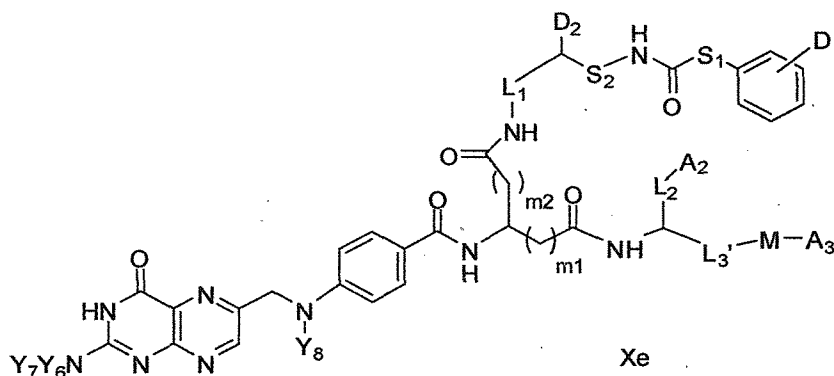
-P(O)(OH)₂,

L₃' is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-, wherein R' represents H or C(1-8)alkyl, p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6.

In preferred embodiments, L₂ is a covalent bond or a straight-chain or branched unsubstituted C(1-6)alkyl, most preferably a covalent bond; and/or (i) m₁ is 0 and m₂ is 2 or (ii) m₁ is 2 and m₂ is 0; and/or S₁ and S₂ are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-8)alkyl, wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -COO-, -NR'-, -NR'-CO-, -CO-NR'-, -CH=CH-, wherein R' represents H or C(1-8)alkyl, more preferably a single bond or a spacer selected from a straight-chain or branched C(1-6)alkyl.

Further compounds include compounds of formulas X a-e, wherein Y₂ is O; Y₁, Y₃, Y₂' are NH; and L₄ is preferably a covalent bond,





wherein

Y₆, Y₇ are independently of each other selected from H, halo,
 5 C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -COR', -
 COOR', and -NHR', wherein R' is H or C(1-8)alkyl,

Y₈ is selected from H, nitroso, C(1-12)alkyl, -OR', -COR', and
 halosubstituted -COR', wherein R' is H or C(1-12)alkyl,

A₂ is H or a capping group,

10 A₃ is H or a capping group, e.g. a protecting group,

M is a radionuclide-based therapeutic or diagnostic moiety M₁ or
 M₂, wherein M₁ is a chelated metal radionuclide selected from a
 linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA,
 DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a metal
 15 radionuclide, and wherein M₂ is a gamma- or positron-emitting
 non-metal radionuclide, preferably selected from ¹¹C, ¹³N, ¹⁵O,
¹⁷F, ¹⁸F, ⁷⁵Br, ⁷⁶Br, ⁷⁷Br, ¹²³I, ¹²⁴I, ¹³¹I, optionally in combination
 with a prosthetic group,

m₁, m₂ are independently of each other 0, 1, 2 or 3, preferably
 20 m₁ is 0 and m₂ is 2 or m₁ is 2 and m₂ is 0,

S₁, S₂ are independently of each other a single bond or a spacer
 selected from a straight-chain or branched C(1-12)alkyl, which
 is unsubstituted or substituted by at least one CN, Hal, OR',

NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D₁ is a group selected from H, halogen, C(1-12)alkyl, preferably halogen,

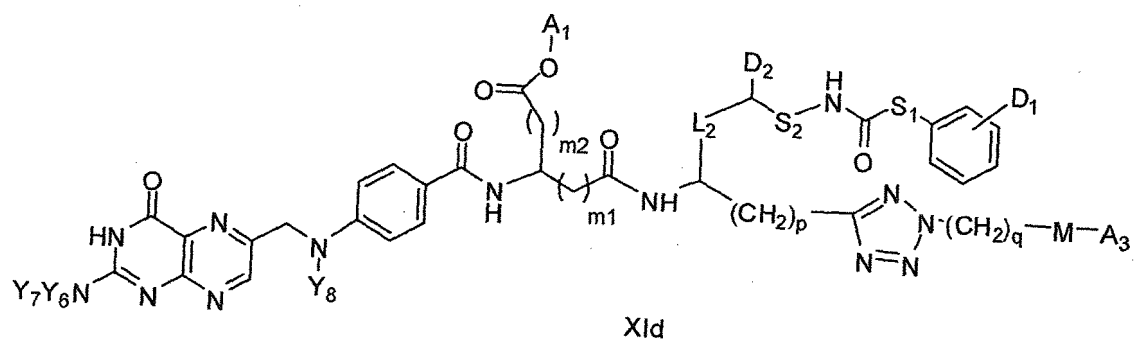
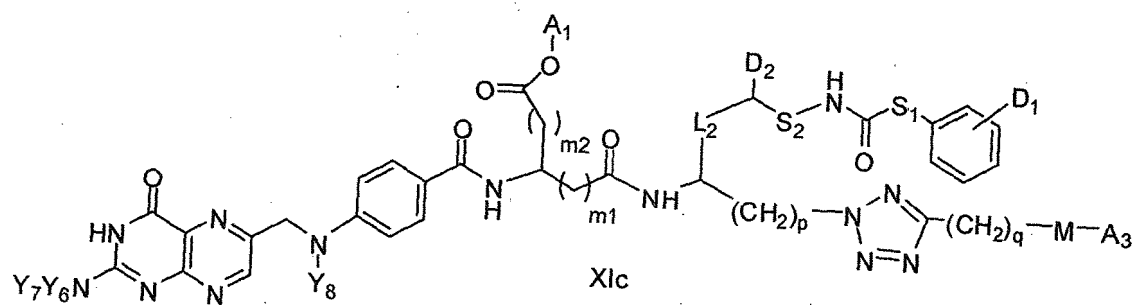
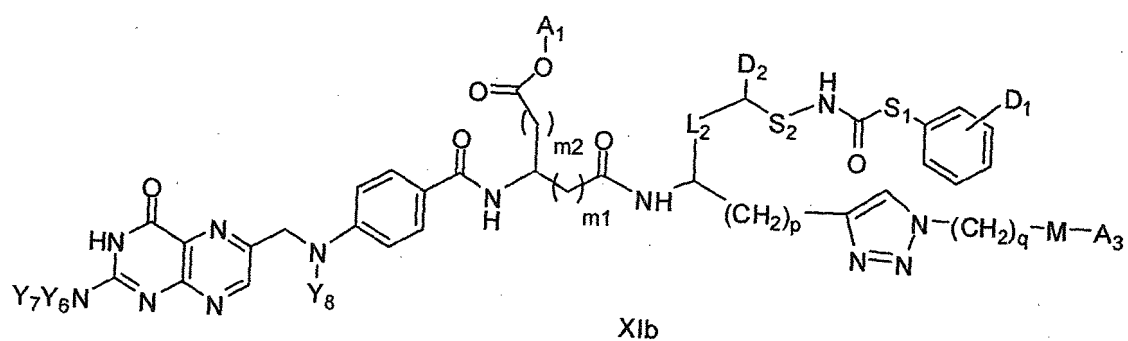
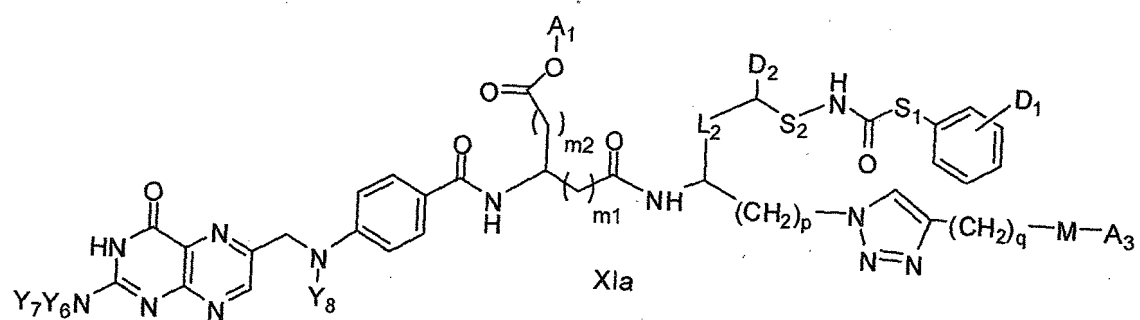
D₂ is an acidic group selected from -COOH, -SO₃H, -SO₂H, -NR'SO₃H, -P(O)(OH)₂,

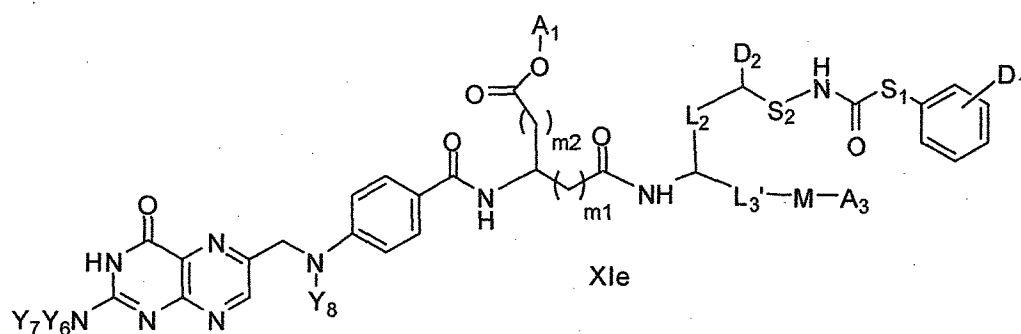
L₁ is a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl,

L₂ is a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl, L₃ is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-, wherein R' represents H or C(1-8)alkyl,

p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6.

The invention also contemplates compounds of formulas XI a-e, wherein Y₁, Y₂ are O, Y₃, Y₂ are NH and L₁ and L₄ are preferably a covalent bond,





wherein

5 Y_6, Y_7 are independently of each other selected from H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, and $-NHR'$, wherein R' is H or C(1-8)alkyl,

Y_8 is selected from H, nitroso, C(1-12)alkyl, $-OR'$, $-COR'$, and halosubstituted $-COR'$, wherein R' is H or C(1-12)alkyl,

A_1 is H or carboxy protecting group,

10 A_3 is H or a carboxy protecting group or a hydroxyl protecting group,

M is a radionuclide-based therapeutic or diagnostic moiety M_1 or M_2 , wherein M_1 is a chelated metal radionuclide selected from a linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a metal radionuclide, and wherein M_2 is a gamma- or positron-emitting non-metal radionuclide, preferably selected from ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F , ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I , optionally in combination with a prosthetic group,

20 m_1, m_2 are independently of each other 0, 1, 2 or 3, preferably m_1 is 0 and m_2 is 2 or m_1 is 2 and m_2 is 0,

S_1, S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' ,

NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D₁ is a group selected from H, halogen, C(1-12)alkyl, preferably halogen,

D₂ is an acidic group selected from -COOH, -SO₃H, -SO₂H, -NR'SO₃H, -P(O)(OH)₂,

L₂ is a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

L₃ is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-, wherein R' represents H or C(1-8)alkyl, p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6.

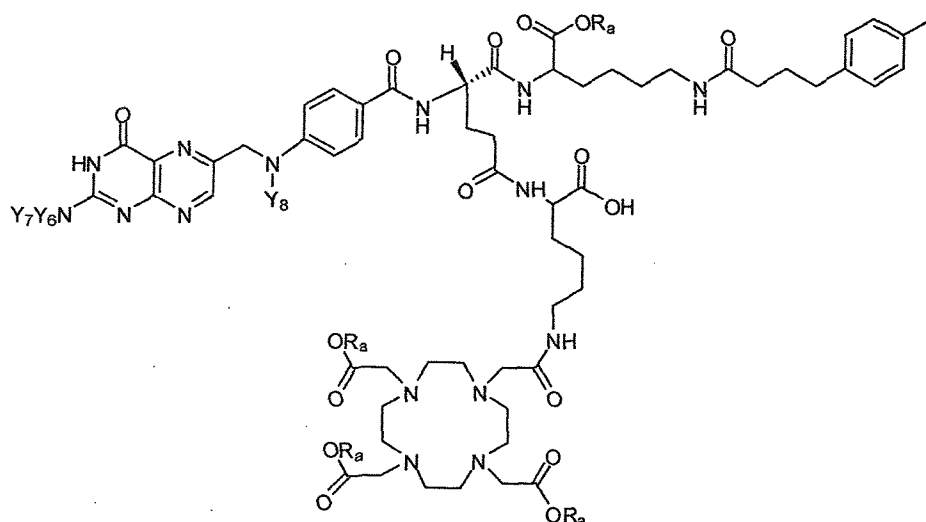
In preferred embodiments, L₁ is a covalent bond or a straight-chain or branched unsubstituted C(1-6)alkyl, most preferably a covalent bond; and/or (i) m₁ is 0 and m₂ is 2 or (ii) m₁ is 2 and m₂ is 0; and/or S₁ and S₂ are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-8)alkyl, wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -COO-, -NR'-, -NR'-CO-, -CO-NR'-, -CH=CH-, wherein R' represents H or C(1-8)alkyl, more preferably a single bond or a spacer selected

from a straight-chain or branched C(1-6)alkyl.

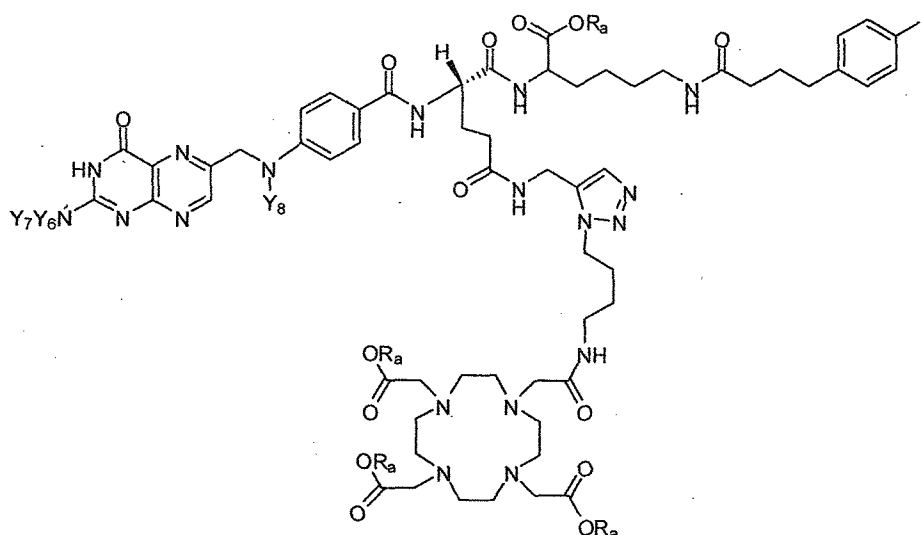
L₂ is preferably a covalent bond or a straight-chain or branched C1-, C2-, C3- or C4-alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-.

L₃ is preferably straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -CO-O- or -NR'-CO-, wherein R' represents H or C(1-8)alkyl.

Most preferred compounds of the invention are e.g.



and



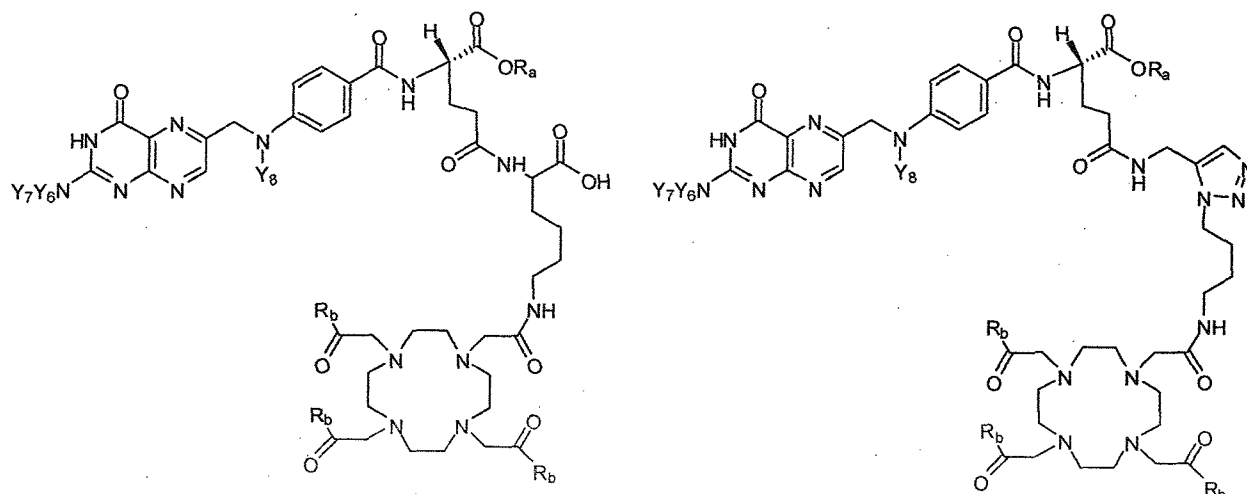
wherein

5 Y_6, Y_7 are independently of each other selected from H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, and $-NHR'$, wherein R' is H or C(1-8)alkyl,

Y_8 is selected from H, nitroso, C(1-12)alkyl, $-OR'$, $-COR'$, and halosubstituted $-COR'$, wherein R' is H or C(1-12)alkyl,

R_a is H or C(1-8)alkyl;

10 and compounds



wherein

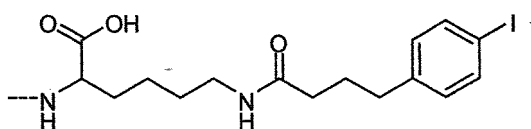
Y_6, Y_7 are independently of each other selected from H, halo, C(1-

12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -COR', -COOR', and -NHR', wherein R' is H or C(1-8)alkyl,

Y₈ is selected from H, nitroso, C(1-12)alkyl, -OR', -COR', and halosubstituted -COR', wherein R' is H or C(1-12)alkyl,

5 R_a is H or C(1-8)alkyl,

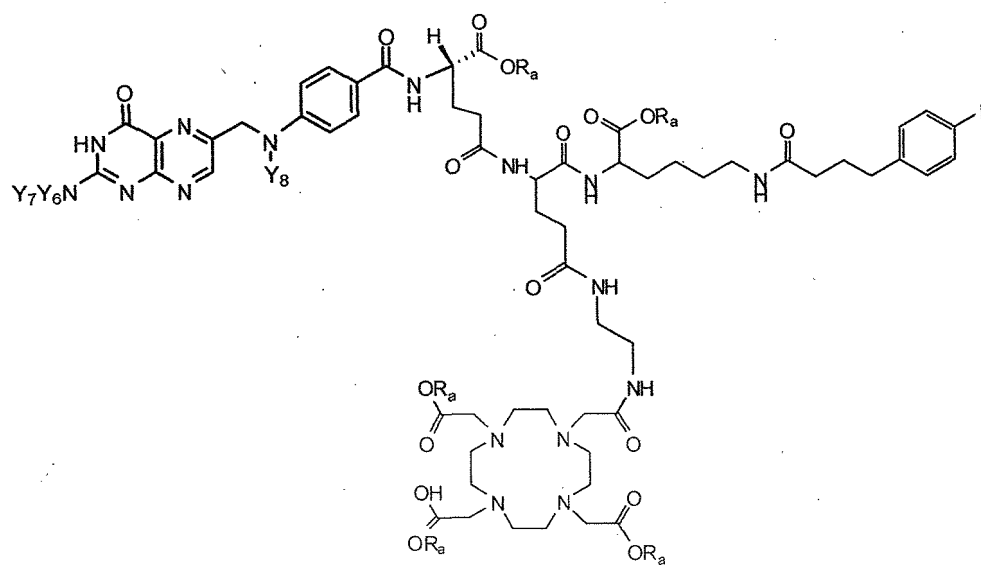
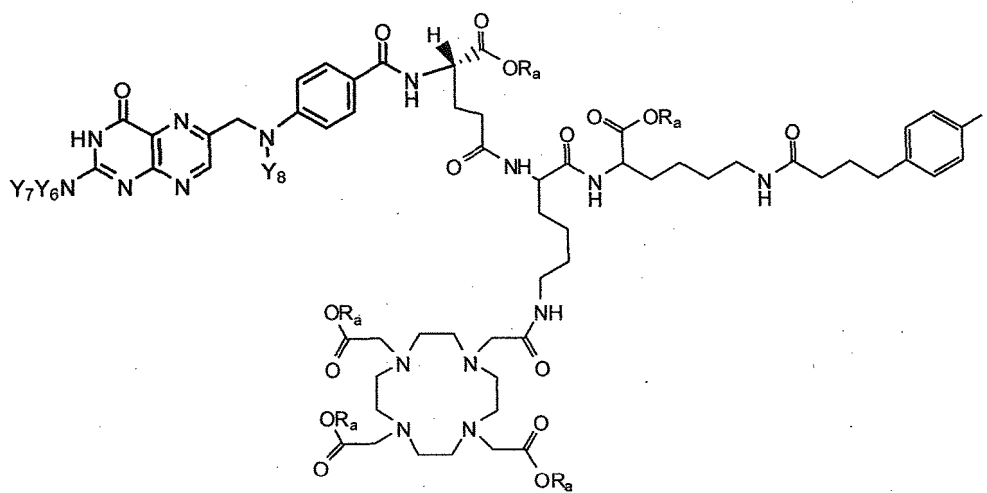
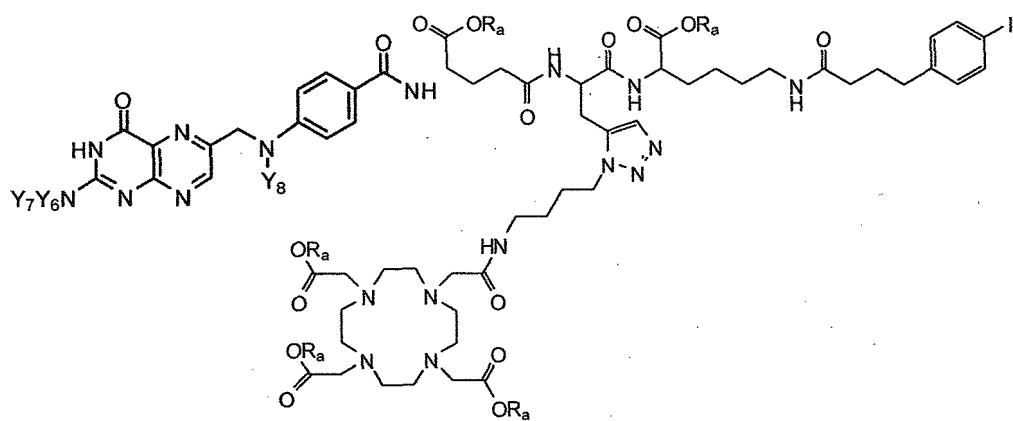
R_b is independently of each other selected from -OH, -OC(1-8)alkyl, or a group of formula (i)



(i)

10 with the proviso that at least one of groups R is a group of formula (i);

and compounds



wherein

Y_6, Y_7 are independently of each other selected from H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, and $-NHR'$, wherein R' is H or C(1-8)alkyl,

Y_8 is selected from H, nitroso, C(1-12)alkyl, $-OR'$, $-COR'$, and
5 halosubstituted $-COR'$, wherein R' is H or C(1-12)alkyl,

R_a is H or C(1-8)alkyl.

In another aspect the present invention also provides methods of synthesizing a compound of the invention. The synthesis is
10 preferably based on a modular approach (using appropriately derivatized functionalities, i.e., folate group, albumin binder group and chelating group) and is based on standard coupling chemistries known in the art, including esterification, amidation, and the click-reaction. The latter reaction has been
15 proven to be particularly useful and is based on the coupling of an azide and an alkyne group in a cycloaddition under thermal conditions or in the presence of a catalyst to obtain the final compound of choice (Kolb and Sharpless, Drug Discovery Today 2003, 8, 1128; Kolb et al. Angew. Chem. Int. Ed. 2001, 40, 2004; Rostovtsev, V. V. et al. Angew. Chem. Int. Ed. 2002, 41, 2596; US
20 2005/02222427; WO 06/116629). These reactions are known as Huisgen 1,3-dipolar cycloaddition (thermal conditions) and Click-Reaction (catalytic conditions) and have been described in the art (Kolb and Sharpless, Drug Discovery Today 2003, 8, 1128; Kolb et al. Angew. Chem. Int. Ed. 2001, 40, 2004; Rostovtsev et al. Angew. Chem. Int. Ed. 2002, 41, 2596; US 2005/02222427; WO 06/116629). More specifically compounds of formula I wherein the
25 fivemembered heterocycle is a triazole are obtained by cycloaddition of an azide R_a-N_3 with an alkyne $R_b-C\equiv C-R_c$ and
30 compounds of formula I wherein the five-membered heterocycle is a tetrazole are obtained by cycloaddition of an azide R_a-N_3 with a cyanide $R_b-C\equiv N$. All possible combinations are contemplated

herein, i.e. R_a being the folate derivative and R_b being a chelating moiety or precursor thereof as well as R_b being the folate derivative and R_a being a chelating moiety or precursor thereof. Thus the modular and versatile nature of the reaction
5 allows employing a wide variety of linkers to couple the imaging moiety to folic acid.

It will be obvious for a skilled person to select appropriate conditions for the various coupling steps and choose appropriate protecting groups PG (e.g. see Greene & Wuts, Eds., Protective
10 Groups in Organic Synthesis, 2nd Ed., 1991, John Wiley & Sons, NY.) and leaving groups LG (e.g. a halogen, tosylate, mesylate, triflate, carbonate group).

It will also be obvious for a skilled person that the last step in the synthesis of the compounds of the invention preferably
15 includes introducing the metal radionuclide (if the imaging moiety is a chelated metal radionuclide M_1) or the gamma- or positron-emitting non-metal radionuclide (if the imaging moiety is a gamma- or positron-emitting non-metal radionuclide M_2 .)

Thus, if the imaging moiety is a chelated metal radionuclide M_1 ,
20 such as selected from a linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, the present invention further provides a method of synthesizing a complex of the invention, which comprises labeling such a compound of the invention, which
25 includes the steps of first obtaining a compound of the invention, and reacting the compound with a radionuclide as specified hereinabove, generally in the presence of a reducing agent to form a metal chelate complex between the compound of the invention and the radionuclide. The reducing agent may be
30 any known reducing agent, but will preferably be a dithionite ion or a stannous ion.

Alternatively, if the imaging moiety is a gamma- or positron-

emitting non-metal radionuclide M_2 , preferably selected from ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F , ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I , the present invention further provides a method of synthesizing the respective compound of the invention, which comprises
5 introducing non-metal radionuclide M_2 from the corresponding precursor according to known methods in the art.

In a further aspect the invention provides pharmaceutical compositions comprising a diagnostic imaging amount or a therapeutically effective amount of at least one compound of the
10 present invention and a pharmaceutically acceptable carrier therefore. In a preferred embodiment, the pharmaceutical compositions contain at least one compound wherein M is a chelated metal radionuclide M_1 selected from DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a metal
15 radionuclide, selected from e.g. ^{177}Lu , ^{161}Tb , ^{213}Bi , or ^{111}In . In another preferred embodiment, the pharmaceutical compositions contain at least one compound wherein M is a non-metal radionuclide M_2 selected from ^{18}F , ^{123}I , ^{124}I , ^{131}I .

As used herein, a pharmaceutically acceptable carrier, which is
20 present in an appropriate dosage, includes solvents, dispersion media, antibacterial and antifungal agents, isotonic agents, and the like which are physiologically acceptable. The use of such media and agents are well-known in the art.

In a further aspect the present invention provides uses of
25 compounds and/or pharmaceutical compositions of the present invention for convenient and effective administration to a subject in need for diagnostic imaging or radionuclide therapy. The subject of the methods of the present invention is preferably a mammal, such as an animal or a human, preferably a
30 human.

Thus in a particular embodiment the present invention provides a method for diagnostic imaging of a cell or population of cells

expressing a folate-receptor, said method comprising the steps of administering at least one compound or composition of the present invention in a diagnostic imaging amount, and obtaining a diagnostic image of said cell or population of cells.

- 5 The compounds and/or compositions of the present invention may also be used for radionuclide therapy agents useful for the treatment of a subject in need thereof.

Thus in another particular embodiment the present invention provides a method for radionuclide therapy comprising the steps
10 of administering to a subject in need thereof at least one compound or composition of the present invention in therapeutically effective amounts, and after localization of said at least one compound or composition in the desired tissues, subjecting the tissues to irradiation to achieve the
15 desired therapeutic effect.

In yet another embodiment the present invention provides a method for simultaneous diagnosis and radionuclide therapy comprising the steps of administering to a subject in need thereof at least one compound or composition of the present
20 invention in a diagnostically and therapeutically effective amount, and after localization of said at least one compound or composition in the desired tissues, subjecting the tissues to irradiation, and obtaining a diagnostic image of said tissues to follow the course of treatment.

- 25 An image of a cell or tissue expressing the folate receptor, i.e. a tumor cell or tissue, labeled with one or more of the compounds or compositions of the present invention can be detected using a radiation detector, e.g. a γ -radiation detector. One such procedure utilizes scintigraphy. Tomographic imaging
30 procedures such as single photon emission computed tomography (SPECT) or positron emission tomography (PET) can also be used to improve visualization. Selection and use of such radiation

detectors is within the skill of one of ordinary skill in the art. Thus, a diagnostic imaging amount of a compound or composition of the present invention to be administered may be selected in an amount sufficient such as to produce a diagnostic
5 image of an organ or other site of the subject as described hereinabove. A therapeutically effective amount of a compound or composition of the present invention to be administered may be selected in an amount sufficient such as to produce a desired radiotherapeutic effect. More specifically a therapeutically
10 effective amount is an amount of at least one of the compounds of the present invention, which will permit sufficient tumor localization of the complex to stop and/or diminish tumor growth or size. As provided herein tumor growth or size can be monitored using the methods of the present invention or any
15 other known diagnostic imaging procedure.

Clearly the specific activity of the metal radionuclide of choice, e.g. ^{99m}Tc , $^{186/188}\text{Re}$, $^{111}\text{In}^{+3}$, $^{67/68}\text{Ga}^{+3}$, $^{90}\text{Y}^{+3}$, $^{109}\text{Pd}^{+2}$, $^{105}\text{Rh}^{+3}$, ^{177}Lu , $^{64/67}\text{Cu}$, ^{166}Ho , ^{213}Bi , preferably ^{99m}Tc or $^{186/188}\text{Re}$, or the non-metal radionuclide of choice, e.g. ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F ,
20 ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I , preferably ^{18}F , ^{123}I , ^{124}I , ^{131}I , will be taken into consideration in determining a dosage for diagnostic imaging or radionuclide therapy.

Generally, the unit dose to be administered has a radioactivity of about 0.01 mCi to about 300 mCi, preferably 10 mCi to about
25 200 mCi. For a solution to be injected a preferred unit dosage is from about 0.01 mL to about 10 mL. After e.g. intravenous administration, imaging of the organ or tumor in vivo can take place, if desired, from within minutes to hours or even longer, after the radiolabeled reagent has been administered to a
30 subject. Typically, a sufficient amount of the administered dose will accumulate in the targeted area to be imaged within about 1-4 hours.

The compounds and/or compositions of the present invention may be administered by an appropriate route such as parentally (for example, intravenously), intramuscularly or intraperitoneally or by any other suitable method. For example, the compounds and/or compositions of this invention may be administered to a subject by bolus or slow infusion intravenous injection. The suitable forms for injection include sterile aqueous solutions or dispersions and sterile powders of the above mentioned compounds and/or compositions of the present invention.

10 The compounds or pharmaceutical compositions are preferably sterile. Sterilization can be accomplished by any art recognized technique, including but not limited to, addition of antibacterial or antifungal agents, for example, paraben, chlorobutanol, phenol, sorbic acid, thimerosal, and the like.

15 The compounds and/or compositions of the invention may also be used for in vitro detection of a cell expressing the folate receptor in a tissue biopsy taken from a subject. Thus in a further embodiment the present invention provides a method for in vitro detection of a cell expressing the folate receptor, e.g. a tumor cell, in a tissue sample which includes contacting
20 said tissue sample with a compound or composition of the present invention in effective amounts and for sufficient time and conditions to allow binding to occur and detecting such binding by imaging techniques.

25 Samples can be collected by procedures known to the skilled person, e.g., by collecting a tissue biopsy or a body fluid, by aspirating for tracheal or pulmonary samples and the like.

Tissue samples to be tested include any tissue suspected to contain a cell expressing a folate receptor, such as tumor cells, epithelial cells, kidneys, gastrointestinal or the
30 hepatobiliary system, and others. Samples can be sectioned, e.g., with a microtome, to facilitate microscopic examination

and observation of bound complex. Samples can also be fixed with an appropriate fixative either before or after incubation with one of the compounds or compositions of the present invention to improve the histological quality of sample tissues.

- 5 Time and conditions sufficient for binding of a complex of the present invention to a folate receptor on the cell include standard tissue culture conditions, i.e. samples can be cultured in vitro and incubated with one of the compounds or compositions of the present invention in physiological media. Such conditions
10 are well known to the skilled person. Alternatively, samples can be fixed and then incubated with a complex or composition of the present invention in an isotonic or physiological buffer.

A typical amount of said complex of the present invention for in vitro detection of a tumor cell can range from about 1 ng/l to
15 about 1000 µg/l. A preferred amount is about 1 µg/l to about 100 µg/l. Preferred compounds of the invention, wherein the imaging moiety is a chelated metal radionuclide M_1 to be used for in vitro diagnosis of a tumor cell are the same as for in vivo applications and include ^{99m}Tc , $^{186/188}\text{Re}$, $^{111}\text{In}^{+3}$, $^{67/68}\text{Ga}^{+3}$, $^{90}\text{Y}^{+3}$,
20 $^{109}\text{Pd}^{+2}$, $^{105}\text{Rh}^{+3}$, ^{177}Lu , $^{64/67}\text{Cu}$, ^{161}Tb , ^{213}Bi , preferably ^{177}Lu , ^{161}Tb , ^{213}Bi , or ^{111}In .

Likewise, preferred compounds of the invention, wherein the imaging moiety is a non-metal radionuclide M_2 to be used for in vitro diagnosis of a tumor cell are the same as for in vivo
25 applications and include ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F , ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I , preferably ^{18}F , ^{123}I , ^{124}I , ^{131}I .

For detection of cellular binding of one of the present compounds, samples can be incubated in the presence of a selected compound, then washed and counted in a standard
30 scintillation counter. Alternative methods apply and are known to the skilled person.

It is understood that the above methods of the invention may be performed in combination with any other methods of cancer diagnosis or therapy including methods using other already developed diagnostic and/or therapeutic agents and utilizing x-ray computed tomography (CT), magnetic resonance imaging (MRI), functional magnetic resonance imaging (fMRI), single photon emission computed tomography (SPECT), optical imaging, and ultrasound.

For diagnostic or radionuclide therapy applications it may be convenient to prepare the compounds of the present invention at, or near, the site where they are to be used. Thus for the compounds of the invention, wherein M is a chelated metal radionuclide M_1 , the present invention provides in a further aspect a single or multi-vial kit containing all of the components needed to prepare those compounds or compositions of this invention, other than the radioimaging metal ion itself. Thus a preferred single-vial kit of the present invention comprises a compound of the present invention, wherein M is a chelated metal radionuclide M_1 , and a source of a pharmaceutically acceptable reducing agent such as a stannous salt. In addition, the kit comprises optionally further additives, for example the kit is buffered with a pharmaceutically acceptable acid or base to adjust the pH to a desired value for complex formation. Such a single vial kit may optionally contain exchange ligands such as glucoheptonate, gluconate, mannitol, maleate, citric or tartaric acid and may also contain reaction modifiers, such as diethylenetriaminepentaacetic acid or ethylenediamine tetraacetic acid. Additional additives, such as solubilizers (for example a cyclodextrin), antioxidants (for example ascorbic acid) and/or fillers (for example, NaCl) may be employed to improve the radiochemical purity and stability of the final product, or to aid in the production of the kit. The

radionuclide, e.g. Lu, Tb, Bi, or In, will preferably be added separately in the form of a solution.

Likewise, a preferred multi-vial kit of the present invention comprises, in one vial, the components, other than the radionuclide itself, required to form a labile radionuclide complex, that is, an exchange ligand and a pharmaceutically acceptable reducing agent such as a stannous salt. A compound of the present invention, wherein M is a chelated metal radionuclide M_1 , is contained in a second vial, as well as optional additives such as buffers appropriate to adjust the pH to its optimal value. Optionally the radionuclide will be provided in form of a solution to be added.

All components of a kit may be in liquid, frozen or dry form. In a preferred embodiment, kit components are provided in lyophilized form.

All of the compounds, compositions and/or methods disclosed and claimed herein can be made and executed without undue experimentation in light of the present disclosure. It will be apparent to those of skill in the art that variations may be applied to the present invention without departing from the scope of the invention. The Examples provided herein are intended to be illustrative and are not exhaustive; therefore the illustrated Examples should not be viewed as limiting the invention in any way.

Examples

Materials and Methods

All chemicals were purchased from Sigma-Aldrich or Fluka, Buchs, Switzerland. All chemicals and solvents were of reagent grade and were used without further purification unless otherwise stated. The Boc-protected amino acids were purchased from Bachem

AG, Bubendorf, Switzerland. The pterioic acid precursor was a kind gift from Merck Eprova AG, Schaffhausen, Switzerland and DOTA-azide and DOTA-NHS-Ester was received from Macrocyclics, Dallas, USA. $[\text{Na}][^{99\text{m}}\text{TcO}_4]$ was eluted from a $^{99}\text{Mo}/^{99\text{m}}\text{Tc}$ -generator (Mallinckrodt-Tyco, Petten, the Netherlands) with a 0.9% saline solution. $^{67}\text{GaCl}_3$ was received from Mallinckrodt-Tyco, Petten, the Netherlands and $^{177}\text{LuCl}_3$ was received from Nuclear Research and Consultancy Group, NRG, Petten, the Netherlands.

Reactions were monitored by HPLC or by thin-layer chromatography (TLC) using precoated silica gel 60 F254 aluminium sheets (Merck), and visualized by UV absorption or stained with a solution of ninhydrin in EtOH (0.2 g in 100 ml). Column chromatography was performed using silica gel 60 (Fluka; particle size 0.04-0.063 mm). Analytical HPLC was performed using a Merck-Hitachi L-7000 system equipped with an L-7400 tunable absorption detector and an XBridge™ C-18 reverse phase column (5 μM , 4.6 x 150 mm, Waters). HPLC solvents were either water with 0.1% TFA (solvent A) and MeCN (solvent B), or 0.1% TFA in water (solvent A) and MeOH (solvent B) with a flow rate of 1 mL/min. The gradient was as follows: 0-15 min: gradient from 95% A to 20% A. Sep-PakR columns (Waters) were washed with methanol and water prior to use. Analytical radio-HPLC was performed on a Merck-Hitachi L-2130 system equipped with a L-2450 diode array detector and a Berthold radiodetector with a reversed-phase column (Gemini C18, 5 μM , 4.6 x 250 mm, Phenomenex) using 0.05 M NH_4HCO_3 (solvent A) and acetonitrile (solvent B) as solvent system with a gradient from 0 - 15 min 80% A, 15 - 20 min 80 - 30% A, 20 - 25 min 30% A and a flow rate of 1 mL/min. Semipreparative radio-HPLC was performed on a HPLC system equipped with a Smartline pump 1000, Smartline Manager 5000, a Smartline UV detector 2500 (Knauer) and a GabiStar radiodetector (Raytest) using a reversed-phase semipreparative column (Gemini C18, 5 μM , 250 x 10 mm, Phenomenex) at a flow

rate of 4 mL/min with 0.05 M NH_4HCO_3 (solvent A) and acetonitrile (solvent B) as solvent system and a gradient as follows: 0 - 20 min 80% A, 20 - 25 min 80 - 30% A, 25 - 30 min 30% A.

5 Nuclear magnetic resonance spectra were recorded on a 400 MHz Bruker spectrometer. ^1H and ^{13}C chemical shifts were reported relative to residual solvent peaks or water as a reference. Values of the coupling constant J were given in Hertz (Hz). The following abbreviations are used for the description of ^1H -NMR spectra: singlet (s), doublet (d), triplet (t), quartet (q),
10 multiplet (m). The chemical shifts of complex multiplets are given as the range of their occurrence. Low resolution mass spectra were recorded with a Micromass Quattro microTM API LC-ESI using either the negative or positive ionization mode. High resolution mass spectra were recorded with a Bruker FTMS 4.7T
15 BioAPEXII.

Cell culture: KB cells (human nasopharyngeal carcinoma cell line) were purchased from the American Type Culture Collection (CCL-17). JK24-FBP cells were a kind gift from Dr. Patrick Hwu (National Cancer Institute, Bethesda, MD). The cells were
20 cultured as monolayers at 37°C in a humidified atmosphere containing 5% CO_2 . Importantly, the cells were cultured in a special RPMI cell culture medium herein referred as FFRPMI (modified RPMI 1640 medium, without folic acid, vitamin B12, and phenol red; Cell Culture Technologies GmbH). FFRPMI medium was
25 supplemented with 10% heat-inactivated fetal calf serum (FCS, as the only source of folate), L-glutamine, and antibiotic (penicillin 100 IU/ml, streptomycin 100 $\mu\text{g}/\text{ml}$, fungizone 0.25 $\mu\text{l}/\text{ml}$). Eighteen to twenty hours prior to each experiment, the cells were seeded in 12-well plates (7×10^5 cells/2 ml) to form
30 confluent monolayers over night.

Uptake and internalization studies: Receptor binding studies were performed according to the following general procedure: The

monolayers of the cells were rinsed with PBS pH 7.4. Pure FFRPMI medium (without FCS, L-glutamine, antibiotics, 975 μ l) only, FFRPMI medium (475 μ l) and a folic acid solution for receptor blocking studies (500 μ l, 200 μ M in medium) or FFRPMI medium containing 4% BSA were added into the corresponding wells. The well plates were pre-incubated at 37°C for 15 min. Solutions of the ^{67}Ga -labelled compounds 16 or 6 (25 μ l, 1.5 MBq/ml) were added to each well. Plates were incubated at 37°C for up to 4 h. Then, each well was rinsed twice with PBS buffer pH 7.4 or once with PBS pH 7.4 and once with an acidic stripping buffer (aqueous solution of 0.1 M acetic acid and 0.15 M NaCl), respectively, to remove bound complex from the FR on the cell surface. The monolayers were dissolved in 1 ml NaOH (1M) and transferred to 4 ml tubes. Samples as well as 4 standard-samples (25 μ l labelled ^{67}Ga -16 or ^{67}Ga -6) were counted for radioactivity using a cobraTM II γ -counter. The concentration of proteins was determined for each sample in order to normalize measured radioactivity of the samples with respect to the averaged sample protein content that accounted for 0.5 mg or 0.3 mg (as indicated under results) protein per well (Thermo Scientific Pierce BCA Protein Assay, Product #23225).

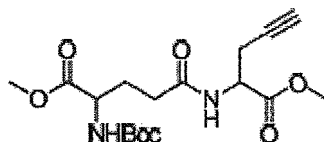
Albumin binding studies: Albumin binding studies using HPLC and a SuperoseTM 12 size exclusion column were performed according to the following general procedure: Compound 16 and 6 were radiolabelled according to the above described protocol. Samples of 1MBq were injected onto the column, using 0.05% tweenR 20 in PBS pH 7.4 as eluent. The radioactive compounds ^{67}Ga -16 and ^{67}Ga -6 (12 μ l, approximately 1 MBq) were incubated for 30 min at 37°C with either HSA solution 20% (100 μ l) or human plasma (100 μ l). A volume of 100 μ l of the incubated solution was injected on the SuperoseTM 12 size exclusion column. Elution of the plasma proteins and of the radiolabelled compound was monitored by HPLC in the UV trace (220 nm) and the radioactive γ -trace,

respectively. Albumin binding studies using ultracentrifugation devices were performed according to the following general procedure⁸³: The HPLC-purified radiolabelled compounds 16, 6, FA2, FA4 and 22 (2.5-3.0 MBq) were incubated for 10 min at 37°C with 350 µl of human plasma or 350 µl of 1x PBS, pH 7.4. The samples were loaded onto the ultrafilter (CentrifreeR, Ultracel YMT devices from Millian, Geneva) and centrifugated for 35 min at 2000 rpm. The filtrate (3 x 25 µl) and initially incubated solution (3 x 25 µl, standard) were counted for radioactivity using a Cobra™ II γ-counter. For standardization, the filtrate of the compounds in PBS was indicated being 100%. This 100% fraction was compared to the fraction of active compound found in the filtrate of the plasma sample and the activity of the plasma sample were expressed in % of the activity in the PBS filtrate.

In vitro autoradiography: The slides with mouse tissue sections (KB tumour xenografts and kidneys) were preincubated in Tris-HCl buffer (170 mM, pH 7.6, with 5 mM MgCl₂) with 0.25% (w/v) BSA for 10 min at room temperature. Then, the sections were incubated with a solution of ⁶⁷Ga-16 and ⁶⁷Ga-6 (0.5 MBq/ml in Tris-HCl buffer containing 1% BSA) for 60 min at room temperature. After incubation, the sections were rinsed twice for 5 min in cold Tris-HCl buffer (with 25% BSA), then washed for 5 min in pure Tris-HCl buffer, and finally rinsed with cold distilled water. The sections were air-dried and exposed to phosphor imaging screens.

Example 1: Synthesis of trifunctional conjugate 16

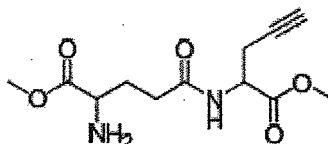
(a)



1

Boc-Glu-OMe (181.5 mg, 0.65 mmol) was dissolved in DMF (1.5 ml), HBTU (314.2 mg, 0.83 mmol, 1.2 eq.) and NEt₃ (2 eq.) were added. The solution was stirred at 0 °C for 1 h. The solution was added
5 to H-Pra-OMe·HCl (120.2 mg, 0.74 mmol, 1.06 eq.) in DMF (2 ml) and NEt₃ (4 eq.). The suspension was stirred over night, 0 °C → RT. MeOH was added to dissolve remaining particles. The solution was stirred at 0 °C for 3h. The product was extracted with citric acid (1 M) and ethyl acetate. The organic phase was
10 rinsed with brine, dried over Na₂SO₄ and concentrated under reduced pressure. Purification was achieved by CC (SiO₂, MeOH/CH₂Cl₂ 1:50) to give a clear oil. 233 mg (0.63 mmol, 90%). LC-MS [M-C₅H₇O₂(Boc)]H⁺= 270.87 (calc. for C₁₇H₂₆N₂O₇ , 370.17). ¹H-NMR (DMSO-d₆) δ 8.3, 7.17, 4.36-3.31, 3.92-3.87, 3.57, 3.55,
15 3.24, 2.8, 2.52, 2.15, 1.87-1.79/1.72-1.62, 1.3 ppm.

(b)

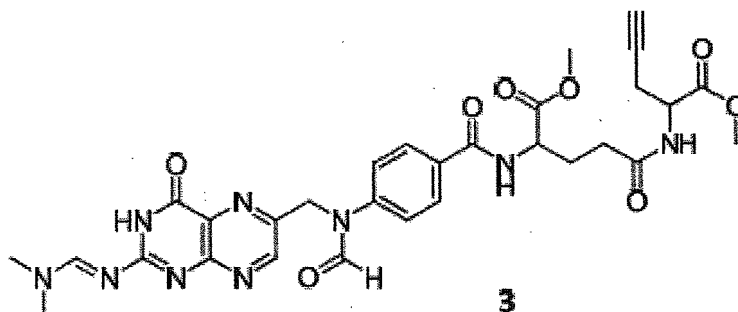


2

1 (233 mg, 0.63 mmol) was dissolved in TFA (10% in CH₂Cl₂). The solution was stirred at RT for 3 h. The solvent was evaporated
20 under reduced pressure. Purification was achieved by CC (SiO₂, MeOH/CH₂Cl₂ 1:30) to give the TFA salt as a clear solid. 224 mg (0.58 mmol, 92%). LC-MS [M+H]⁺ = 270.84 (calc. for C₁₂H₁₈N₂O₅,

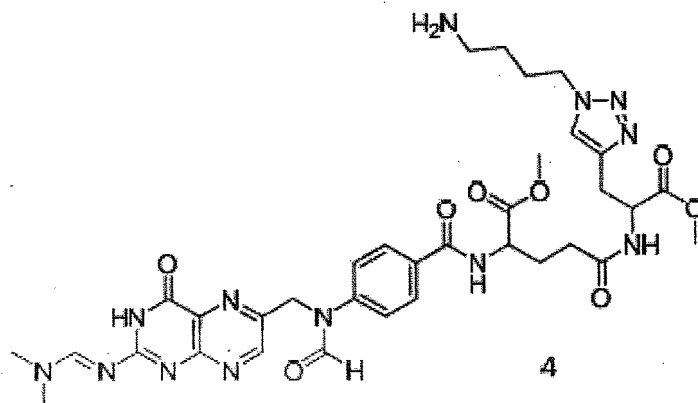
270.12). ¹H-NMR (MeOH-d₄) δ 7.98, 4.58, 4.10, 3.85, 3.74, 2.82, 2.71, 2.54, 2.19.

(c)



5. Pterioic acid (166.9 mg, 0.41 mmol, 1.04 eq.) was dissolved in DMF (3 ml). HBTU (208.9 mg, 0.55 mmol, 1.4 eq.) and NEt₃ (2 eq.) were added. The suspension was stirred at 0 °C for 1.5 h. The suspension was added to 2 (105.2 mg, 0.39 mmol, 1 eq.) in DMF (2 ml) and NEt₃ (2 eq.). The solution was stirred at 0 °C for 4 h.
- 10 The solvent was evaporated under reduced pressure. Purification was achieved by CC (SiO₂, MeOH/CH₂Cl₂ 1:20) to give a yellow powder. 90 mg (0.14 mmol, 36%). LC-MS [M+H]⁺ = 647.93 (calc. for C₃₀H₃₃N₉O₈, 647.25).

(d)



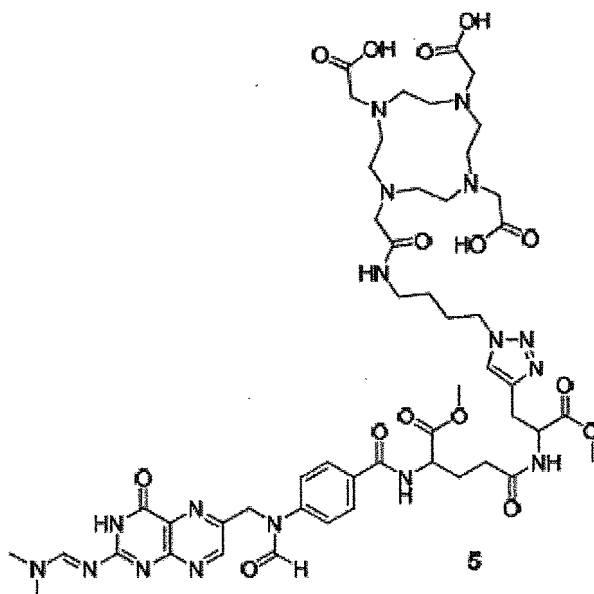
15

3 (22.88 mg, 35 μmol) and 4-azidobutyl-1-amine·TFA (27.7 mg, 0.12 mmol, 3.4 eq.) were mixed in 3:1 tBuOH/H₂O (1.1 ml).

Cu(OAc)₂·H₂O (31 µl, from a 100 mM stem solution, 0.1 eq.) and Na ascorbate (1.5 mg, from a 100 mM stem solution, 0.2 eq.) were added drop wise. The solution was stirred at RT for several h. Some of the solvent was evaporated under reduced pressure.

5 Purification was achieved by semi-preparative HPLC using 0.1% TFA in H₂O and acetonitrile as the eluent to give a yellow powder. 8.64 mg (11.3 µmol, 32%). LC-MS [M-C₃H₇N (dimethylformamide protecting group)]⁺ = 706.95 (calc. for C₃₄H₄₃N₁₃O₈, 761.34).

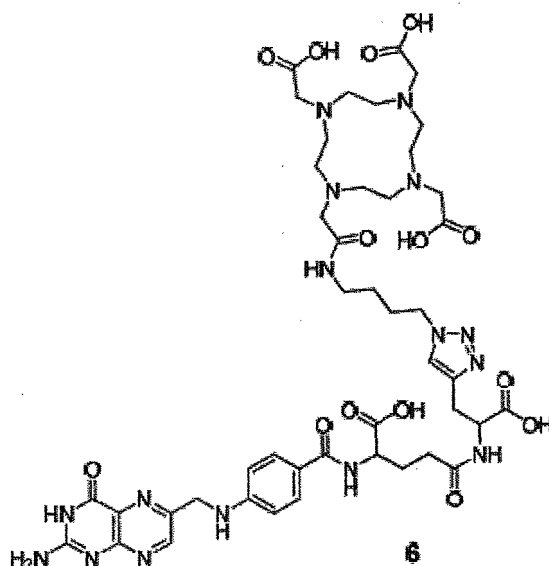
10 (e)



4 (8.64 mg, 11.3 µmol) and NHS-DOTA·PF₆ (8.05 mg, 12.4 µmol, 1.1 eq.) were dissolved in a mixture of DMF (0.4 ml) and DIPEA (4 eq.). The reaction was stirred at 40 °C for several h. The

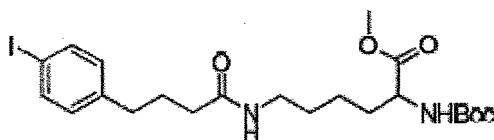
15 reaction was forced to completion by adding more NHS-DOTA PF₆. The solvent was removed under reduced pressure and the product was used without further purification. LC-MS [M-C₃H₇N (dimethylformamide protecting group)]⁺ = 1093.02 (calc. for C₅₀H₆₉N₁₇O₁₅, 1147.52).

20 (f)



5 was dissolved in a 1:1 mixture of MeOH /H₂O and adjusted to pH 13 using NaOH (1M). The solution was stirred at RT for 2 h. The solvent was evaporated under reduced pressure. The resulting solid was redissolved in H₂O and the solution was neutralized using HCl (1M). Final purification was achieved by solid phase extraction using a reverse phase SepPak column with H₂O and increasing concentrations of MeOH as the eluent. Some of the solvent was evaporated under reduced pressure. The compound was lyophilized to give a bright yellow powder. 4.1 mg (3.95 μmol, 35%). LC-MS [M+Na]⁺ = 1059.44, (calc. for C₄₄H₆₀N₁₆O₁₄, 1036.45).

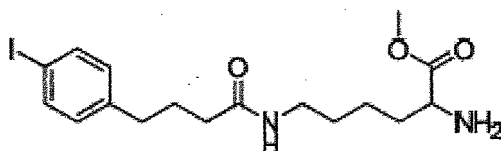
(g)



4-p-iodophenyl-butyric acid (502 mg, 1.7 mmol, 1.07 eq.) was dissolved in DMF (1.5 ml) and NEt₃ (2 eq.). HBTU (751 mg, 1.98 mmol, 1.23 eq.) was added. The solution was stirred at 0 °C for 1 h. The solution was added drop wise to Boc-Lys-OMe (428 mg,

1.6 mmol) in DMF (1.5 ml) and NEt_3 (2 eq.). The solution was stirred at 0 °C for 4 h. The product was extracted with citric acid (1 M) and ethyl acetate/n-Hexane (9:1). The organic phase was rinsed with brine, dried over Na_2SO_4 and concentrated under reduced pressure. Purification was achieved by CC (SiO_2 , MeOH/ CH_2Cl_2 1:50) to give a clear oil. 653.5 mg (1.23 mmol, 75%). LC-MS $[\text{M}+\text{H}]^+ = 532.77$ (calc. for $\text{C}_{22}\text{H}_{33}\text{IN}_2\text{O}_5$, 532.14). $^1\text{H-NMR}$ (MeOH- d_4) δ 7.59, 6.98, 4.09, 3.69, 3.15, 2.59, 2.29, 2.18, 1.88, 1.78/1.64, 1.51, 1.42 ppm.

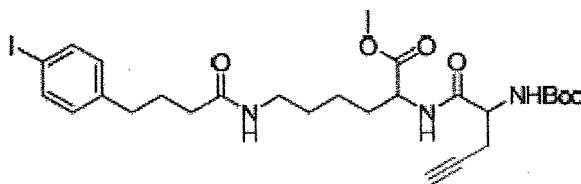
10 (h)



8

7 (568.5 mg, 1.07 mmol) was deprotected with TFA 10% in CH_2Cl_2 at RT for 2h. The solvent was evaporated under reduced pressure. Purification was achieved by CC (SiO_2 , MeOH/ CH_2Cl_2 1:30) to give the TFA salt as a slightly yellow tar. 478.9 mg (0.88 mmol, 82%). LC-MS $[\text{M}+\text{H}]^+ = 432.94$, (calc. for $\text{C}_{17}\text{H}_{25}\text{IN}_2\text{O}_3$, 432.09). $^1\text{H-NMR}$ (MeOH- d_4) δ 7.60, 6.99, 3.96, 3.80, 3.17, 2.58, 2.19, 1.88, 1.53, 1.4 ppm.

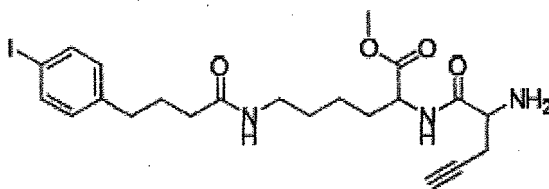
(i)



9

Boc-Pra-OH (50.7 mg, 0.24 mmol, 1.06 eq.) was dissolved in DMF (1 ml) and NEt_3 (1.8 eq.). HBTU (101.4 mg, 0.27 mmol, 1.2 eq.) was added. The solution was stirred at 0 °C for 1h. The solution was added drop wise to 8-TFA (122.6 mg, 0.224 mmol) in DMF (1.5 ml) and NEt_3 (1.8 eq.). The solution was stirred at 0 °C for 4 h. The solvent was evaporated under reduced pressure. Purification was achieved by CC (SiO_2 , first CH_2Cl_2 , then $\text{MeOH}/\text{CH}_2\text{Cl}_2$ 1:50) to give a yellow oil. 125 mg (0.2 mmol, 89%). LC-MS $[\text{M}+\text{H}]^+ = 627.86$ (calc. for $\text{C}_{27}\text{H}_{38}\text{IN}_3\text{O}_6$, 627.18). $^1\text{H-NMR}$ (MeOH-d_4) δ 7.98, 6.99, 4.42, 4.25, 3.69, 3.15, 2.68/2.63, 2.58, 2.38, 2.18, 1.87, 1.72, 1.45, 1.45 ppm.

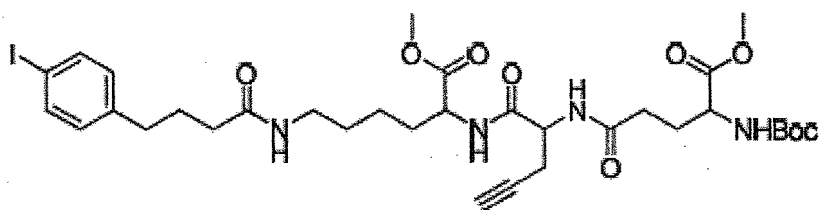
(j)



10

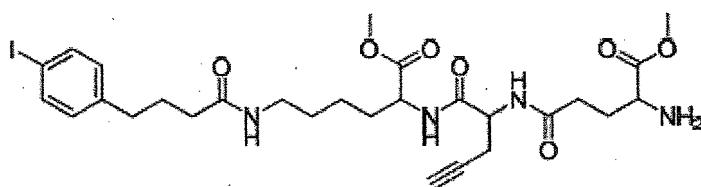
9 (122.2 mg, 0.195 mmol) was deprotected with 10% TFA in CH_2Cl_2 (10 ml). The solvent was evaporated under reduced pressure. Purification was achieved by CC (SiO_2 , first $\text{MeOH}/\text{CH}_2\text{Cl}_2$ 1:30, than $\text{MeOH}/\text{CH}_2\text{Cl}_2$ 1:20) to give the TFA salt as a yellow oil. 92.5 mg (0.144 mmol, 74%). LC-MS $[\text{M}+\text{H}]^+ = 527.87$ (calc. for $\text{C}_{22}\text{H}_{30}\text{IN}_3\text{O}_4$, 527.13). $^1\text{H-NMR}$ (MeOH-d_4) δ 7.6, 6.99, 4.45, 3.7, 3.59, 3.15, 2.59, 2.44, 1.28, 1.88, 1.76, 1.51, 1.41 ppm.

(k)

**11**

Boc-Glu-OMe (38.9 mg, 0.149 mmol, 1.01 eq.) was dissolved in DMF (1 ml) and NEt_3 (2 eq.). HBTU (60.2 mg, 0.159 mmol, 1.08 eq.) was added. The solution was stirred at 0 °C for 1 h. The solution
 5 was added to 10·TFA (90 mg, 0.147 mmol) in DMF (1.5 ml) and NEt_3 (2 eq.). The reaction was stirred at 0 °C for 3 h and stored 48 h at 5 °C. The solvent was evaporated under reduced pressure. Purification was achieved by CC (SiO_2 , $\text{MeOH}/\text{CH}_2\text{Cl}_2$ 1:20) to give
 10 a clear oil. 53.9 mg (0.07 mmol, 48%). LC-MS $[\text{M}+\text{H}]^+ = 771.02$ (calc. for $\text{C}_{33}\text{H}_{47}\text{IN}_4\text{O}_9$, 770.24). $^1\text{H-NMR}$ ($\text{CDCl}_3/\text{MeOH-d}_4$) δ 7.55, 7.45, 6.92, 4.49, 4.42, 4.17, 3.70, 3.68, 3.12, 2.63, 2.55, 2.14, 1.84, 1.68, 1.45, 1.40, 1.32 ppm.

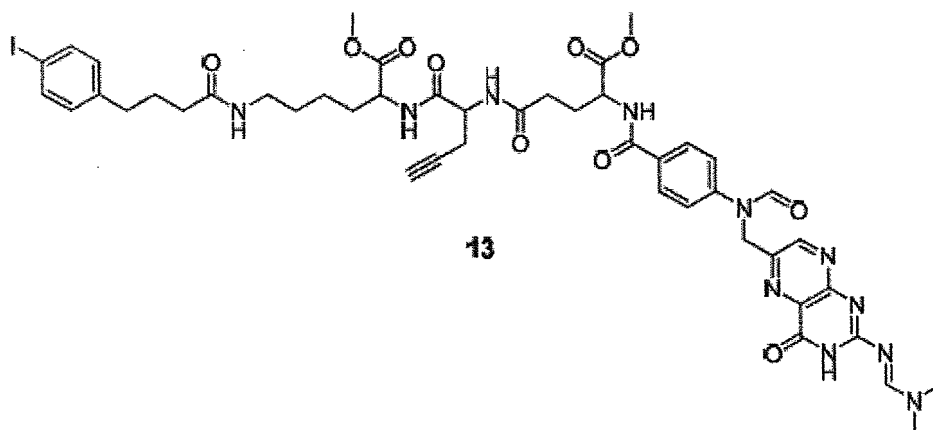
(1)

**12**

15 11 (319 mg, 0.414 mmol) was deprotected with 10% TFA in CH_2Cl_2 (10 ml). The solvent was evaporated under reduced pressure. Purification was achieved by CC (SiO_2 , first CH_2Cl_2 , then $\text{MeOH}/\text{CH}_2\text{Cl}_2$ 1:20) to give the multiple TFA salt as a clear oil. 331.5 mg. LC-MS $[\text{M}+\text{H}]^+ = 670.96$ (calc. for $\text{C}_{28}\text{H}_{39}\text{IN}_4\text{O}_7$, 670.19).
 20 $^1\text{H-NMR}$ (MeOH-d_4) δ 7.6, 6.99, 4.58-4.50, 4.4, 3.86, 3.79, 3.69,

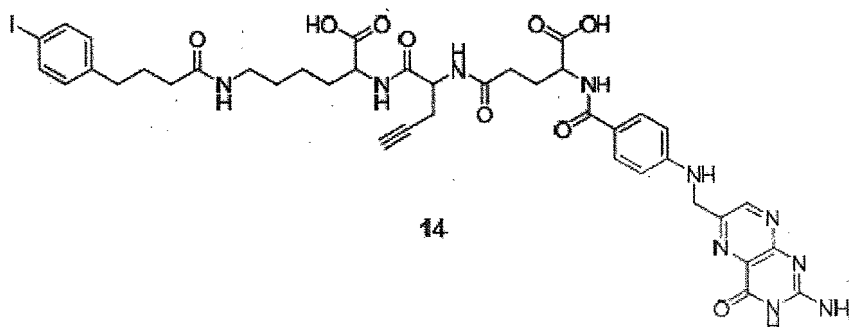
3.20-3.09, 2.73- 2.62, 2.47, 2.43-2.41, 2.18, 2.14-2.0, 1.93-1.81, 1.76-1.67, 1.53-1.46, 1.43-1.34, 1.33-1.25, 1.18 ppm.

(m)



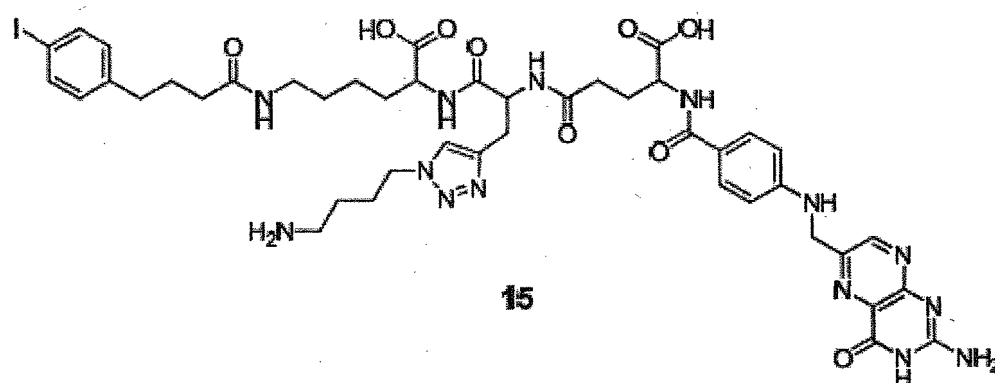
5 Pterioic acid (167.6 mg, 0.41 mmol, 1.05 eq.) was dissolved in DMF (1 ml) and NEt_3 (2 eq.). HBTU (177.9 mg, 0.47 mmol, 1.2 eq.) was added. The solution was stirred at 0 °C for 1 h. The solution was added to 12-TFA (305.5 mg, 0.39 mmol, 1.0 eq.) in DMF (1.5 ml) and NEt_3 (2 eq.). The reaction was stirred at 0 °C
10 for 4 h. The solvent was evaporated under reduced pressure. Purification was achieved by CC (SiO_2 , first CH_2Cl_2 , then $\text{MeOH}/\text{CH}_2\text{Cl}_2$ 1:10) to give a yellow solid. 178.4 mg (0.17 mmol, 44%).

(n)



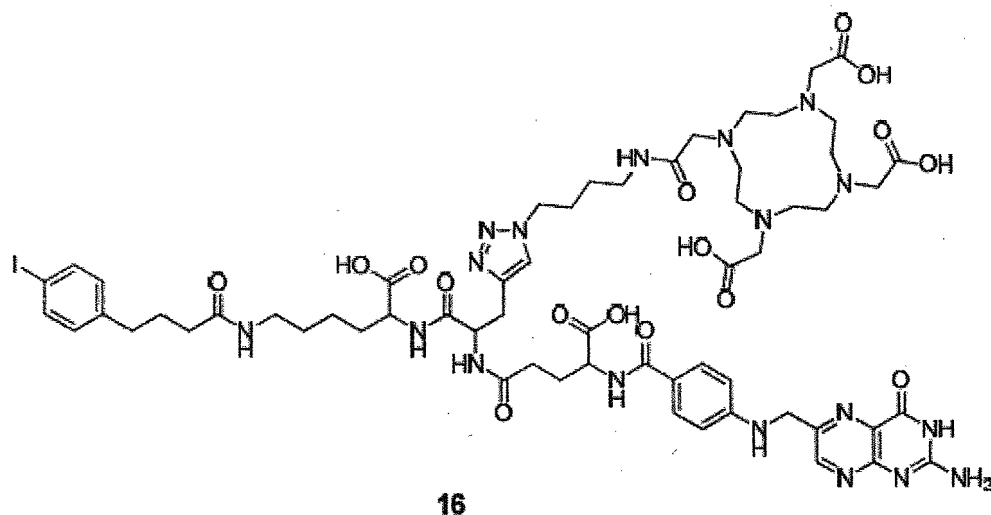
13 (178.4 mg, 0.17 mmol) was deprotected in MeOH/NaOH (aq.) pH 12.8 (10 ml) over night at RT. Purification was achieved by solid phase extraction using a reverse phase SepPak column, elution with H₂O and increasing amounts of MeOH and NaOH (aq.).
5 211.36 mg (0.23 mmol). LC-MS [M+H]⁺ = 936.93 (calc. for C₄₀HtIN₁₀O₉, 936.24).

(o)

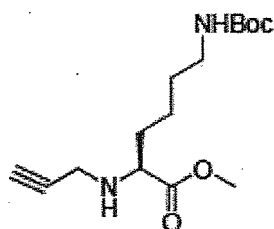


14 (21.9 mg, 23.4 μmol) and 4-azidobutyl-1-amine·TFA (5.24 mg, 4.7 μmol, 2 eq.) were mixed in 1:1 tBuOH/H₂O (1.2 ml). Cu(OAc)₂·H₂O (0.43 mg, from a 100 mM stem solution, 0.1 eq.) and Na ascorbate (0.93 mg, from a 100 mM stem solution, 0.2 eq.) were added drop wise. The solution was stirred over night at RT and at 50 °C for some h. The solvent was evaporated under reduced pressure. Purification was achieved by solid phase extraction using a reverse phase SepPak column, elution with H₂O, slightly basic EDTA and increasing concentrations of MeOH and NaOH (aq.). 10.58 mg (10.1 μmol, 43%). LC-MS [M+H]⁺ = 1050.95 (calc. for C₄₄H₅₅IN₁₄O₉, 1050.33).

20 (p)

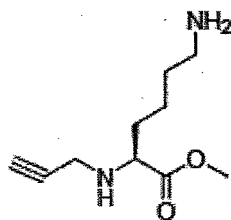


15 (10.58 mg, 10.1 μmol) and NHS-DOTA $\cdot\text{PF}_6$ (9.7 mg, 15.0 μmol , 1.5 eq.) were dissolved in DMF (1 ml) and DIPEA (2.03 eq.). The reaction was stirred at 40 °C for several h. The solvent was removed under reduced pressure. The solid was redissolved in basic water and precipitated in acid. The precipitate was centrifuged and dried under high vacuum to give a yellow powder. 6.7 mg (4.7 μmol , 46%). LC-MS $[\text{M}+\text{H}]^+ = 1437.76$ (calc. for $\text{C}_{60}\text{H}_{81}\text{IN}_{18}\text{O}_{16}$, 1436.51).



H-Lys-(Boc)OMe $\cdot\text{HCl}$ (291.9 mg, 0.98 mmol), 3-Bromo-1-propyne (80% in Toluene, 110.85 μl , 1.16 mmol, 1.2 eq.) and Cs_2CO_3 (643 mg, 2.0 mmol, 2.05 eq.) were stirred in DMF (4.5 ml) at RT and 60 °C for several h. The suspension was filtered and the solvent was evaporated under reduced pressure. Purification was achieved by

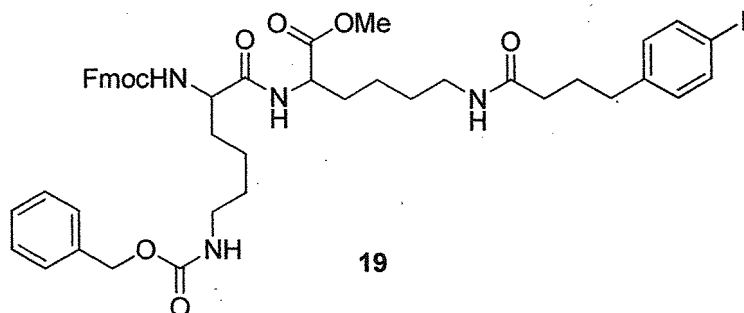
CC (SiO₂, first pure CH₂Cl₂, then MeOH/CH₂Cl₂ 1:50) to give a clear oil. 164.1 mg (0.55 mmol, 56%). LC-MS [M+H]⁺ = 298.89 (calc. for C₁₅H₂₆N₂O₄, 298.19). ¹H-NMR (CDCl₃-d) δ 7.97, 3.68, 3.38-3.36, 3.35-3.33, 3.05, 2.17, 1.81, 1.72-1.53, 1.48-1.28, 1.39 ppm.

**18**

17 (164.1 mg, 0.55 mmol) was deprotected in 10% TFA in CH₂Cl₂ at RT for 3 h. Purification was achieved by solid phase extraction using a reverse phase SepPak column with H₂O as eluent to give a clear oil. 144 mg (0.46 mmol, 84%). LC-MS [M+H]⁺ = 198.92 (calc. for C₁₀H₁₈N₂O₂, 198.14).

Example 2: Synthesis of trifunctional DOTA-Lys-conjugate 26

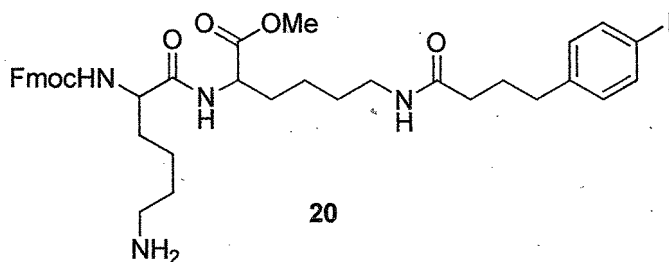
15 (a)

**19**

To 5 g of Fmoc-Lys(Z)-OH (Z = benzyloxycarbonyl) in 50 ml abs. DMF were added 2.2 ml triethylamine and 4.1 g HBTU (N,N,N',N'-tetramethyl-O-(1H-benzotriazol-1-yl)-uronium-hexafluorophosphate). The mixture was stirred for 15 min. at 25°C. Then a

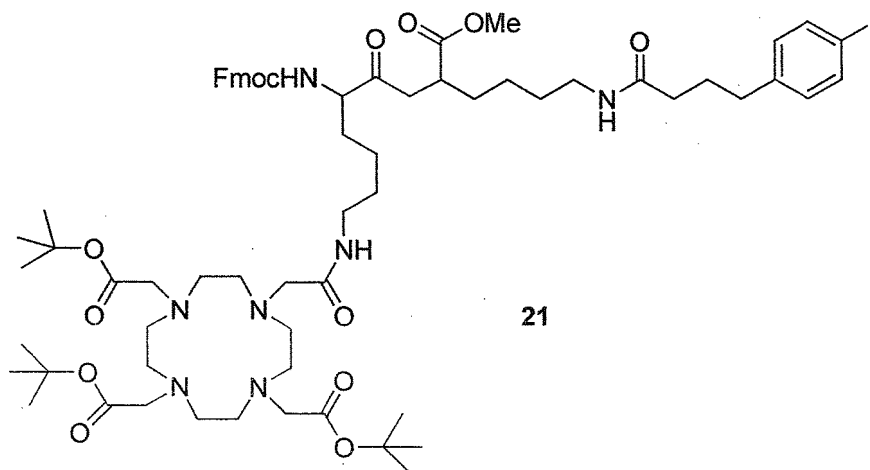
5 mixture of 4.9 g of compound 8 and 2.2 ml triethylamine in 50 ml DMF was added. After stirring for 20 hours, the solvent was removed in vacuum and the residue was dissolved in 50 ml methylene chloride. The methylene chloride solution was washed
10 twice with 30 ml 5% aqueous citric acid solution, twice with 30 ml aqueous 5% sodium bicarbonate solution and twice with 30 ml of water. The organic layer was dried over magnesium sulphate and then evaporated to dryness to give 8.4 g of a waxy residue which was purified by chromatography on silica gel using
10 $\text{CH}_2\text{Cl}_2/\text{MeOH}$ 95:5 as eluent to give 7.3 g of pure 19.

(b)



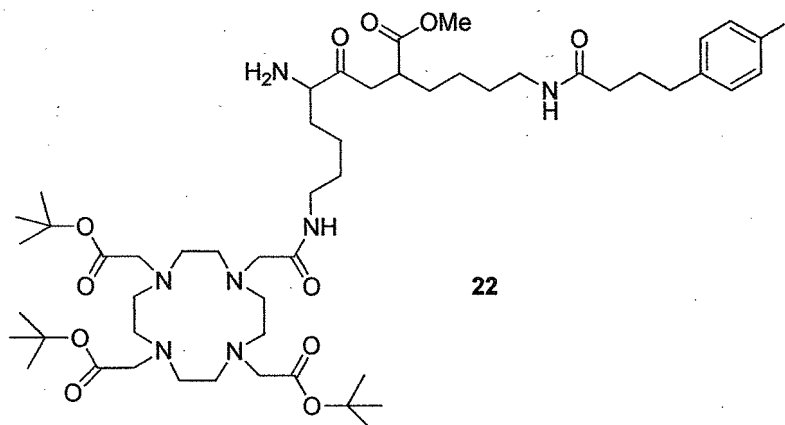
5 g of compound 19 were dissolved in 100 ml of methanol and 200 mg of Pd/C were added. The mixture was hydrogenated at 25°C
15 using 3 bar hydrogen for 24 hours. Solids were removed by filtration and the filtrate was evaporated to dryness to give compound 20.

(c)



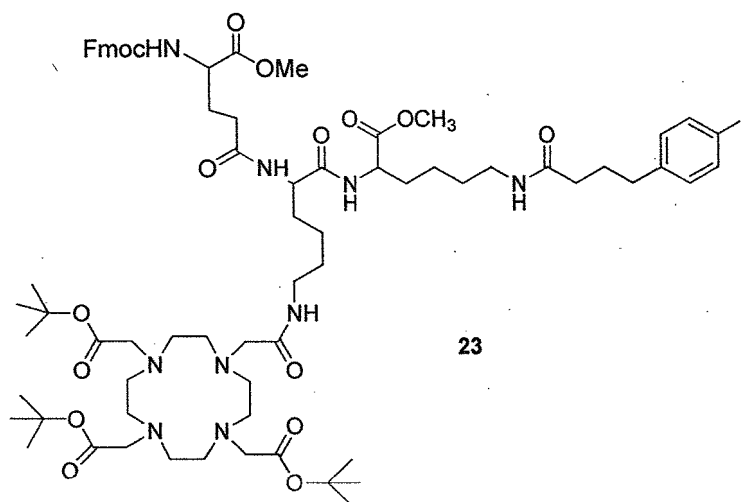
Compound 20 (1 equivalent) was dissolved in CH_2Cl_2 and an equimolar amount of DOTA-NHS-ester (purchased from Macrocyclics) was added. The mixture was stirred for 20 hours at 25°C . The solvent was evaporated and the residue was purified by chromatography on silica gel to give pure 21.

(d)



Compound 21 (1 equivalent) was dissolved in CH_2Cl_2 . After addition of 30 equivalents of diethylamine the mixture was stirred for 20 hours. After evaporation to dryness the residue was purified by chromatography on silica gel to give pure 22.

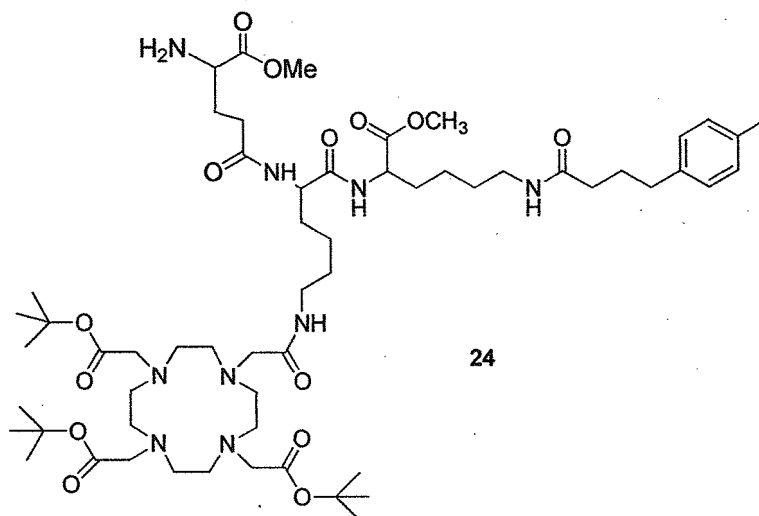
(e)



Fmoc-Glu-OMe (1 equivalent) was dissolved in abs. DMF. After addition of 2 equivalents of diisopropylethylamine and 1.1 equivalents of HBTU (N,N,N',N' -tetramethyl- O -(1H-benzotriazol-1-

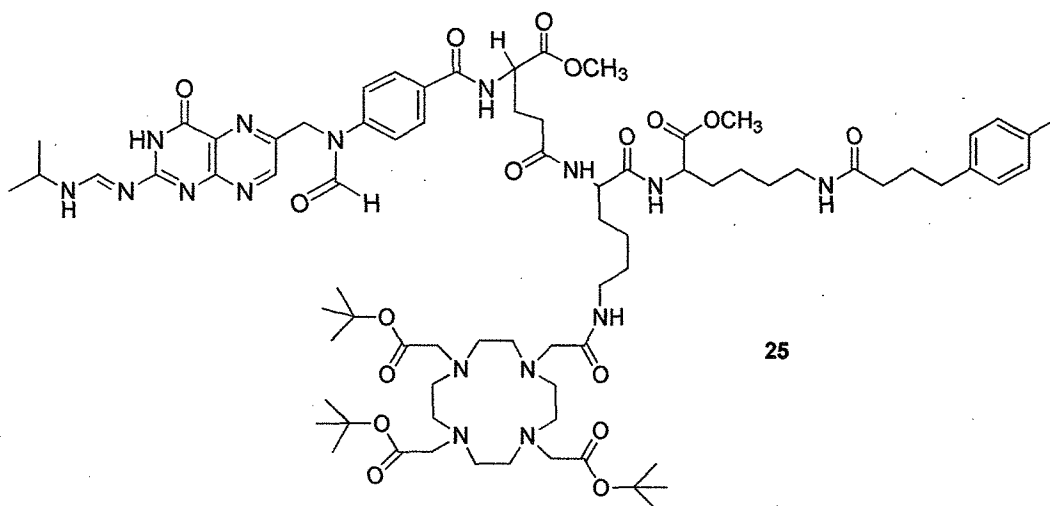
yl)-uronium-hexafluoro-phosphate) the mixture was stirred for 15 min. at 25°C. Then a solution of 1 equivalent of compound 22 in DMF was added and the mixture was stirred for 20 hours. The solvent was removed in vacuum and the residue was dissolved in methylene chloride. The methylene chloride solution was washed twice with 5% aqueous citric acid solution, twice with aqueous 5% sodium bicarbonate solution and twice with water. The organic layer was dried over magnesium sulphate and then evaporated to dryness to give a residue, which was purified by chromatography on silica gel to give pure 23.

(f)



Compound 23 (1 equivalent) was dissolved in CH₂Cl₂. After addition of 30 equivalents of diethylamine the mixture was stirred for 20 hours. After evaporation to dryness the residue was purified by chromatography on silica gel to give pure 24.

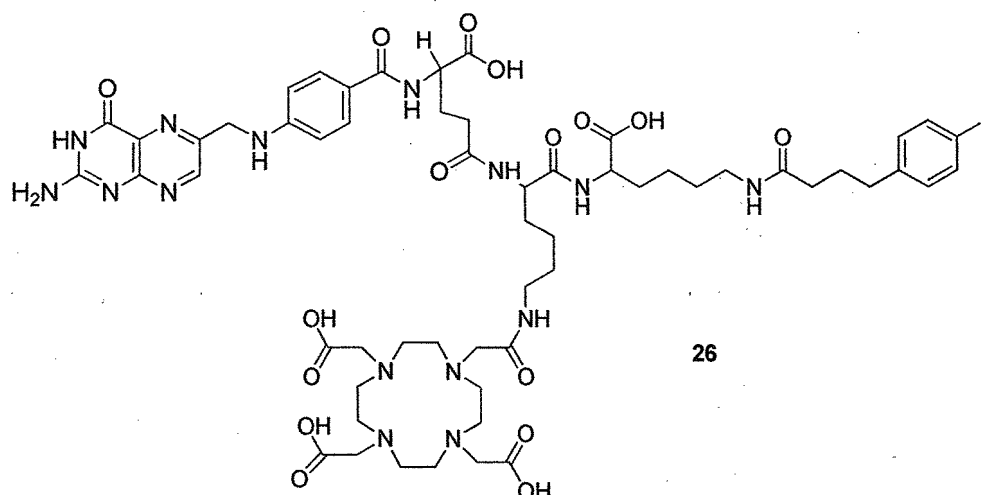
(g)



25

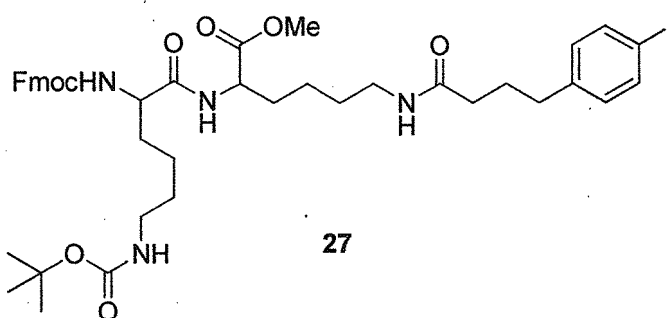
N²-(N,N-dimethylaminomethylene)-10-formylpteroic acid (1 equivalent) was dissolved in abs. DMF. 2 Equivalents of triethylamine and 1.2 equivalents of HBTU (N,N,N',N'-tetramethyl-O-(1H-benzotriazol-1-yl)-uronium-hexafluoro-phosphate) were added and the mixture was stirred for 30 min. Then a solution of compound 24 in abs. DMF was added and the mixture was stirred for 20 hours at 25°C. After removal of solvent in vacuum the residue was dissolved in methylene chloride. The methylene chloride solution was washed twice with 5% aqueous citric acid solution, twice with aqueous 5% sodium bicarbonate solution and twice with water. The organic layer was dried over magnesium sulphate and then evaporated to dryness to give a residue, which was purified by chromatography on silica gel to give pure 25.

(h)



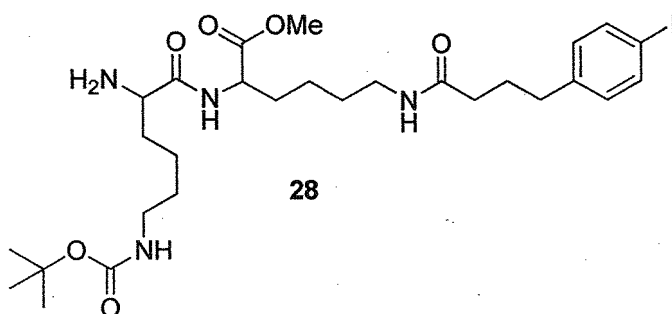
Compound 25 (1 equivalent) was dissolved in tetrahydrofuran and an aqueous solution of lithium hydroxide (3 equivalents) in water was added. After 4 hours at 25°C the mixture was acidified to pH=1 by addition of aqueous hydrochloric acid. After heating to 50°C for 2 hours the mixture was cooled to 25°C and the precipitated product was isolated by centrifugation.

Example 3: Alternative synthesis of trifunctional DOTA-Lys-conjugate 26

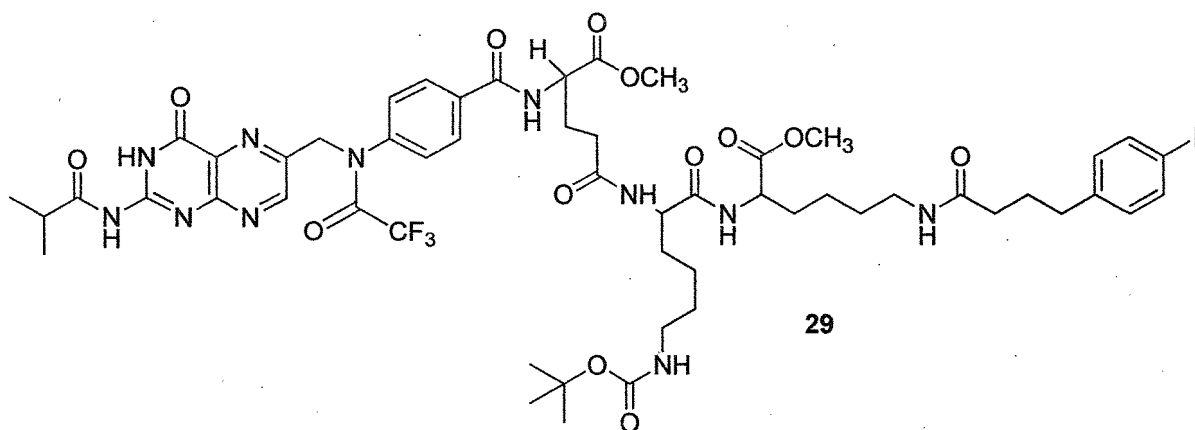


Fmoc-Lys(Boc)-OH (6.4g, 13.7 mmol) was dissolved in dry N,N-dimethylformamide (53 ml). After addition of triethylamine (3.75 ml) and HBTU (N,N,N',N'-tetramethyl-O-(1H-benzotriazol-1-yl)-uronium-hexafluoro-phosphate, 5.2 g) the solution was stirred at room temperature for 15 min. A mixture of compound 8 (6.4 g, 13.7 mmol) in dry N,N-dimethylformamide (64 ml) and

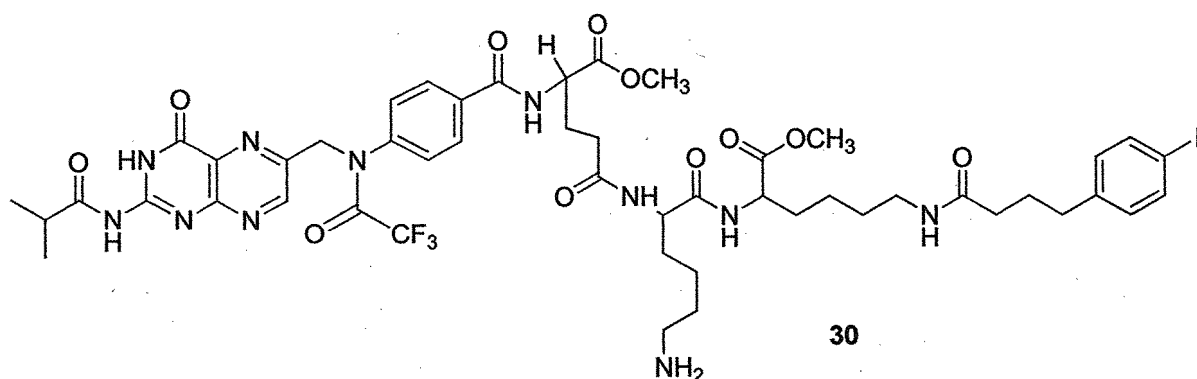
triethylamine (3.75 ml) was added. The reaction mixture was stirred for 17 hours at room temperature. Then the solids were drawn off and the filtrate was evaporated in vacuum at 40°C to give an oily residue. The residue was dissolved in methylene chloride (100 ml) and washed with 5% aqueous citric acid (3 x 50 ml), with 5% aqueous sodium hydrogen carbonate (3 x 50 ml) and water (50 ml). The organic layer was dried over magnesium sulphate and evaporated to dryness in vacuum at 40°C to give compound 27 (7.2 g, 60% yield) which was used without further purification for the synthesis of compound 28.



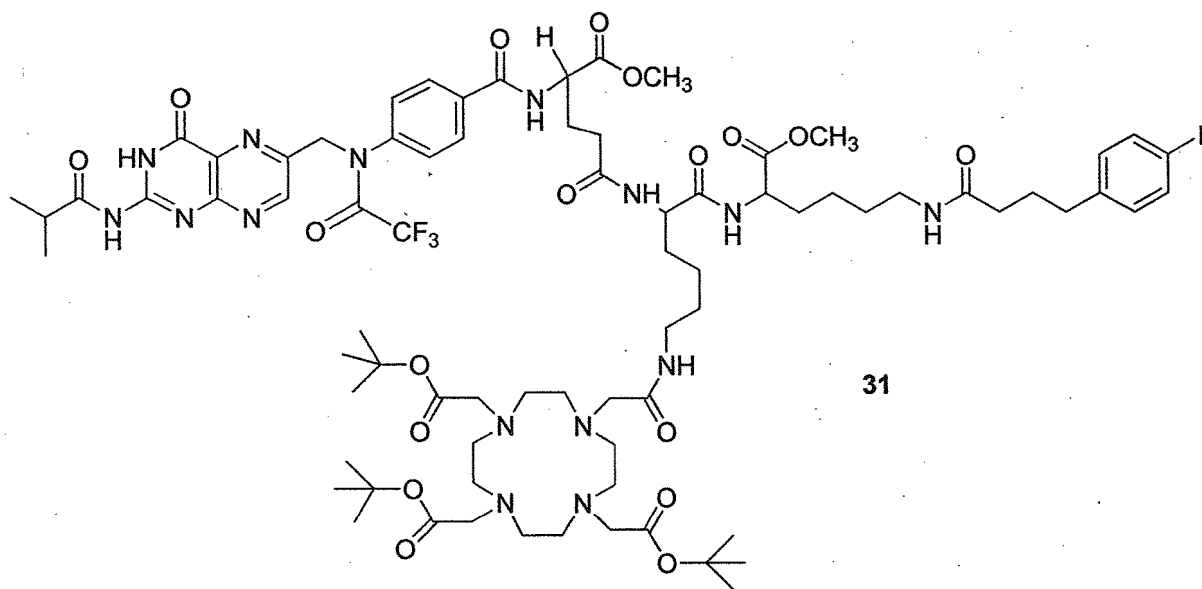
To compound 27 (7.2 g crude product) in dry methylene chloride (36 ml) was added diethylamine (36 ml). The mixture was stirred 2 hours at room temperature and then evaporated to dryness in vacuum at 40°C. Then a mixture of methylene chloride/methanol 90:10 (50 ml) was added to the residue and solids were removed. The filtrate was evaporated to dryness and the residue was purified by column chromatography using SiO₂ (70 g) and methylene chloride / methanol 90:10 as eluent to give compound 28 (4.1g, 76% yield) as a white solid (SiO₂, R_f=0.34 (CH₂Cl₂/MeOH/90:10)).



To a solution of N²-isobutyryl-10-trifluoroacetyl-folic acid- α -methylester (3.4 g, 5.54 mmol, prepared according to WO 2009/082606) in dry N,N-dimethylformamide (34 ml) was added triethylamine (1.5 ml) and HBTU (N,N,N',N'-tetramethyl-O-(1H-benzotriazol-1-yl)-uronium-hexafluoro-phosphate, 2.1 g) and the mixture was stirred for 10 min. at room temperature. Then a solution of compound 28 (3.8 g, 5.8 mmol) in dry N,N-dimethylformamide (38 ml) was added and the mixture was stirred for 20 hours at room temperature. The reaction mixture was evaporated to dryness in vacuum at 40°C. The residue was dissolved in methylene chloride (128 ml) and washed with 5% aqueous citric acid (50 ml), with 5% aqueous sodium hydrogen carbonate (50 ml) and water (50 ml). The organic layer was dried over magnesium sulphate and evaporated to dryness in vacuum at 40°C to give an off-white foam (7.2 g) which was purified by column chromatography using SiO₂ (210 g) and methylene chloride/methanol 93:7 as eluent to give an off-white foam, which was purified a second time by column chromatography using SiO₂ (250 g) and methylene chloride/ methanol 95:5 as eluent to give compound 29 (3.7g, 53% yield) as an off-white foam, R_f=0.33 (SiO₂, CH₂Cl₂/MeOH/90:10).



To a solution of compound 29 (3.5 g, mmol) in methylene chloride (88 ml) was added trifluoroacetic acid (8.8 ml) and triisopropylsilane (0.88 ml). The mixture was stirred at room temperature for 30 min. and then evaporated to dryness in vacuum at 40°C. The residue was purified by column chromatography using SiO₂ (235 g) and methylene chloride/ methanol 90:10 as eluent to give compound 30 as an off-white foam (2.8 g, 79% yield), R_f=0.16 (SiO₂, CH₂Cl₂/MeOH/90:10).



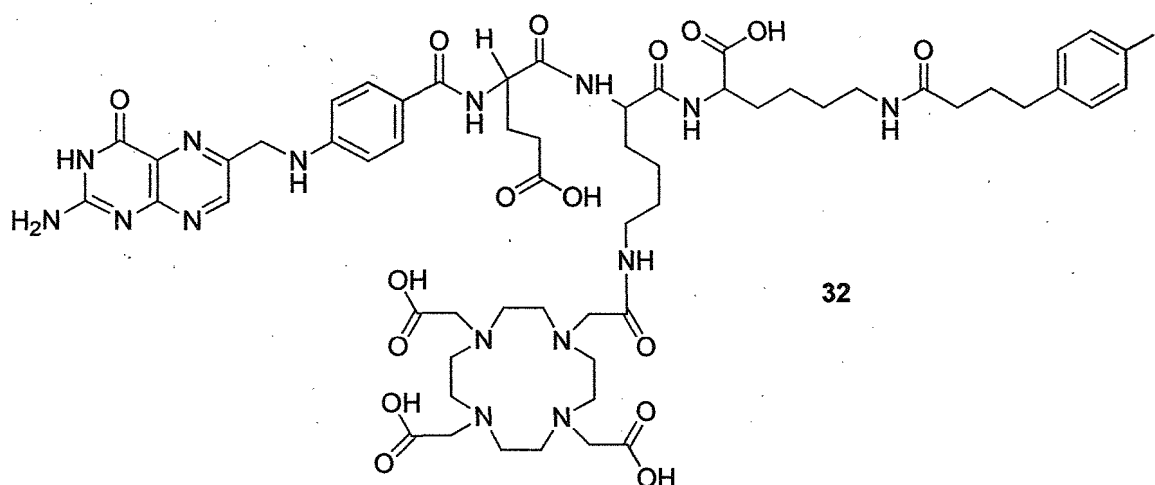
To a solution of DOTA -tris (tBu)ester (224 mg, 0.39 mmol, purchased from Macrocyclics, No. B-260) in dry N,N-dimethylformamide (12.5 ml) was added HBTU (N,N,N',N'-tetramethyl-O-(1H-benzotriazol-1-yl)-uronium-hexafluoro-phosphate, 148 mg) and triethylamine (54.2 μl). After stirring for 5 min. at room temperature a solution of compound 30 (500

mg, 0.39 mmol) in dry N,N-dimethylformamide (5 ml) and triethylamine (54.2 μ l) was added. The reaction mixture was stirred for 20 hours at room temperature and then evaporated to dryness in vacuum at 40°C. The residue was dissolved in 5 methylene chloride (25 ml) and washed with 5% aqueous citric acid (2x5 ml), with 5% aqueous sodium hydrogen carbonate (2x5 ml) and water (2x5ml). The organic layer was dried over magnesium sulphate and evaporated to dryness in vacuum at 40°C to give a yellow foam (0.5 g) which was purified by column 10 chromatography using SiO₂ (17 g) and methylene chloride/ methanol 93:7 as eluent to give compound 31 as an off-white foam (0.43 g, 64% yield), R_f=0.45 (SiO₂, CH₂Cl₂/MeOH/90:10).

Compound 26 was obtained from compound 31 according to the following procedure: To a solution of compound 31 (0.43 g, 0.25 15 mmol) in methylene chloride (20 ml) was added trifluoroacetic acid (20 ml) and the resulting solution was stirred for 20 hours at room temperature. The reaction mixture was evaporated to dryness in vacuum at 40°C and the residue was dissolved in tetrahydrofuran (20 ml). Then a solution of lithium hydroxide 20 (120 mg) in water (20 ml) was added and the reaction mixture was stirred for 20 hours. The tetrahydrofuran was removed under reduced pressure at 40°C and to the residual aqueous solution was added aqueous 1M hydrochloric acid (2.6 ml) until pH=3. The precipitated product was isolated by centrifugation (5000 rounds 25 per min.) and washed with water (4 x 3ml). The product was dried at 40°C in vacuum for 48 hours to give compound 26 as an orange-yellow solid (279 mg, 82% yield). HPLC: 97.0% area. (5.9 mg product were dissolved in 5 ml water and 3 drops buffer (4g Na₂CO₃ and 4 g KHCO₃ dissolved in 200 ml water), Injection 30 volume: 2 μ l, column: Phenomenex, XB-C18, 75x4.6 mm, 2.6 μ m, eluent A: 0.1% trifluoroacetic acid in water, eluent B: 0.1% trifluoroacetic acid acetonitrile in water /90:10, flow: 2.0 ml/min., pressure: 230 bar, gradient: in 20 min. from 0% eluent

B to 100% eluent B. UV-detector, wavelength 230 nm.). LC-MS: $[M+H]^+ = 1356.8$ (calc. for $C_{57}H_{78}IN_{18}O_{16}$, 1355.5).

Example 4: Synthesis of trifunctional DOTA-Lys- α -conjugate 32

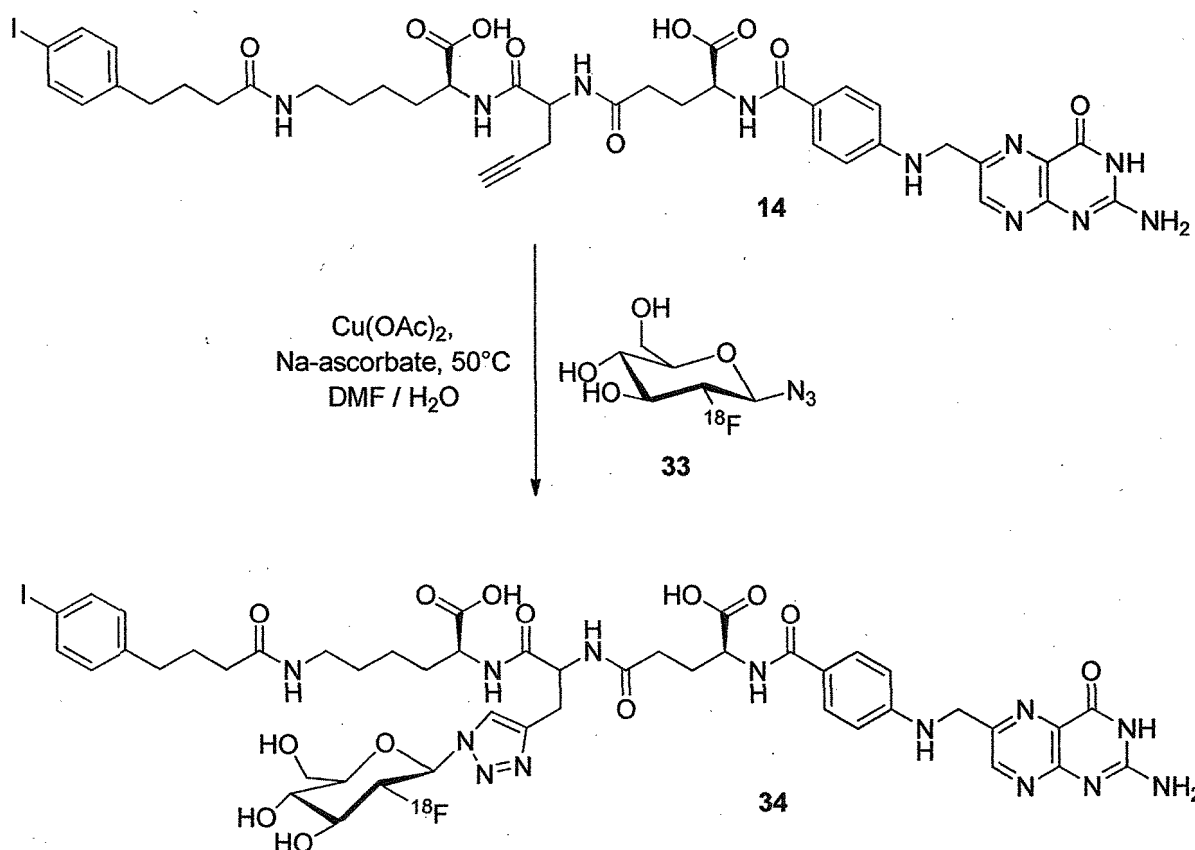


The synthesis was achieved in analogy to example 4 using N²-isobutyryl-10-trifluoroacetyl-folic acid- γ -methylester instead of N²-isobutyryl-10-trifluoroacetyl-folic acid- α -methylester (the N²-isobutyryl-10-trifluoroacetyl-folic acid- γ -methylester was prepared in analogy to the N²-isobutyryl-10-trifluoroacetyl-folic acid- α -methylester according to WO 2009/082606.)

Compound 32 was obtained as an orange-yellow solid. HPLC: 95.0% area (same method as described for analysis of compound 26). LC-MS: $[M+H]^+ = 1356.6$ (calc. for $C_{57}H_{78}IN_{18}O_{16}$, 1355.5).

Example 5: Synthesis of trifunctional conjugate [¹⁸F]FDG-AB-folate 34

(a)



The 3,4,6-tri-*O*-acetyl-2-*O*-trifluoromethanesulfonyl- β -D-mannopyranosyl azide precursor used for coupling the ^{18}F -substituted glucose via click reaction to the folate, was obtained according to literature procedures (e.g. Maschauer and Prante, Carbohydr. Res. 2009, 753; Takatani et al Carbohydr. Res. 2003, 1073).

(b) Radiosynthesis of 2- ^{18}F fluoroglucopyranosyl azide 33

To the dry ^{18}F -fluoride-cryptate complex the precursor, 3,4,6-tri-*O*-acetyl-2-*O*-trifluoromethanesulfonyl- β -D-mannopyranosyl azide (3.0 mg, 6.5 μmol), in 0.30 mL of anhydrous acetonitrile was added. The mixture was stirred for 5 min at 80°C to afford a ^{18}F -incorporation of maximum 75% according to radio-UPLC analysis. After 5 min of cooling and addition of 8 mL of water, the mixture was passed through a reversed-phase cartridge (Sep-Pak C18 Plus; Waters; preconditioned with MeOH and H_2O). The cartridge was washed with 5 mL of water. The ^{18}F -labelled

protected intermediate, 3,4,6-tri-O-acetyl-2-deoxy-2-
[¹⁸F]fluoroglucopyranosyl azide, was eluted with 2.0 mL of
acetonitrile into another 10 ml sealed reaction vessel and dried
under reduced pressure and a nitrogen stream at 80°C. For
5 hydrolysis, 0.25 ml of 60 mM sodium hydroxide solution was added
and the mixture was heated to 65°C for 5 min to complete the
deacetylation. After cooling, the mixture was neutralized with
0.25 ml of 60 mM hydrogen chloride solution and directly used
for the click reaction without further purification.

10

(c) Coupling to folate alkyne precursor 14

The deprotected 2-deoxy-2-[¹⁸F]fluoroglucopyranosyl azide 33
obtained in step (b) was transferred into another reaction
vessel containing the folate alkyne 14 in 300 µl DMF, followed
15 by addition of 10 µl 0.1 M Cu(OAc)₂ solution and 20 µl 0.1 M
sodium ascorbate solution. The reaction mixture was stirred at
50°C for 15 min. After addition of 3 ml of 0.05 M NH₄HCO₃
solution the mixture was submitted to the semipreparative radio-
HPLC system. The product fraction [¹⁸F]FDG-albuminbinder folate
20 (R_T = 19.3 min) was collected into 20 ml of water and the mixture
was passed through a reversed-phase cartridge (Sep-Pak C18
light; Waters; preconditioned with EtOH (5 ml) and H₂O (5 ml)).
The cartridge was washed with 10 ml of water and the ¹⁸F-labeled
product [¹⁸F]FDG-albuminbinder folate 34 was eluted with 1.0 ml
25 of ethanol into a sterile, pyrogen-free vial. After evaporation
of the ethanol, the final product solution was diluted with 2 ml
PBS for injection.

The overall decay-corrected yield of the isolated product
reached 1-2% after a total synthesis time of 3 h with a
30 radiochemical purity always greater than 95%. The specific
activity of [¹⁸F]FDG-albuminbinder folate 34 was around 40-
50 GBq/µmol.

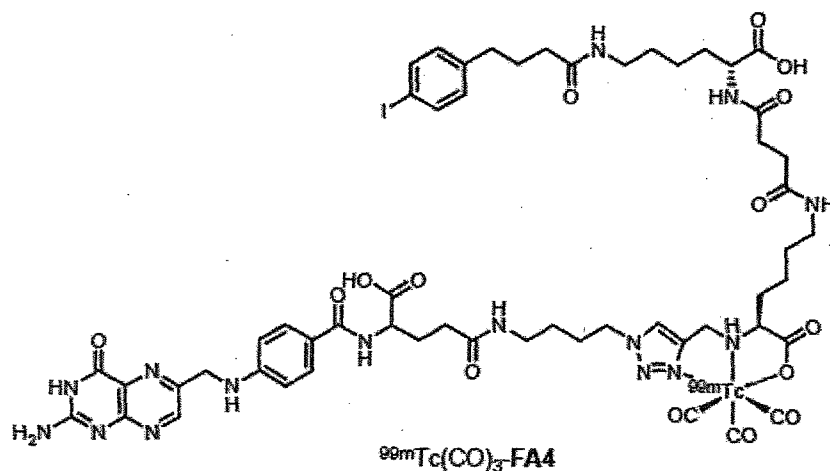
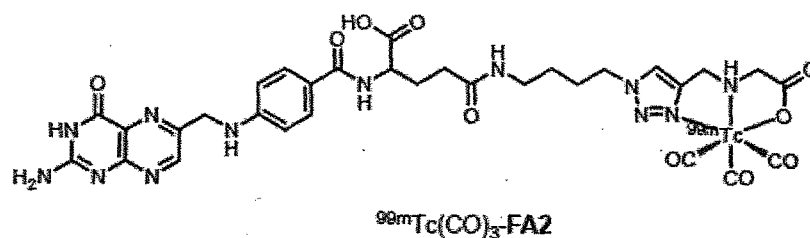
The logD_{7.4} value of [¹⁸F]FDG-albuminbinder folate was found to be

-3.2 $\pm$ 0.4 by the shake flask method.

Example 6: Radiolabelling of compounds of Example 1(f) and 1(p)

(a) with ^{67}Ga and ^{177}Lu

- 5 A volume of 15 μl of compound 16 and 6 (1 mM) were mixed with 250 μl of Na-acetate buffer (0.4 M, pH 5) and $^{67}\text{GaCl}_3$ solution (40 MBq, 2.7×10^{-11} mol) or $^{177}\text{LuCl}_3$ solution (50 MBq, 0.7×10^{-11} mol for 30 nmol compound). The mixture was incubated for 30 min at 90°C, then cooled to RT. DTPA solution (10 μl , 5 mM, pH 5)
- 10 was added for complexation of potential traces of unreacted ^{67}Ga or ^{177}Lu respectively. Quality control was performed by HPLC. The HPLC system comprised a Waters system with a tunable absorbance detector and a Unispec multichannel analyser γ -detector and a Waters XTerraR MS C18 (5 μm , 4.6 mm x 150 mm) with an eluent
- 15 that consisted of aqueous 0.05 M triethylammonium phosphate buffer, at pH 2.25 or pH 7.5, and methanol with a linear gradient to 80% methanol over 30 min. For *in vitro* cell experiments and the albumin binding assay compounds ^{67}Ga -16 and ^{67}Ga -6 or ^{177}Lu -16 and ^{177}Lu -6 respectively were purified by HPLC.
- 20 For all the *in vitro* experiments the radioactive compounds were purified via HPLC. The radioactive peak was collected and diluted with PBS. MeOH was removed under a flow of N_2 .



(b) with $^{99m}\text{Tc}(\text{CO})_3$

Eluted $[\text{Na}][^{99m}\text{TcO}_4]$ in 0.9% saline solution (1 ml, 2-10 GBq) was added to an Isolink Kit (4.5 mg BC, 2.9 Borax, 7.8 mg $\text{Na}_2(\text{CO})_3$, 9 mg Na-K-Tartrate under N_2 atmosphere. The solution was heated for 20 min to 100°C , then cooled to RT and neutralized using an acidic HCl/phosphate buffer. PBS (250 μl of 1 x, pH 7.4), 50 nmol of compound FA2, FA4 or 22 and 200 μl of $^{99m}\text{Tc}(\text{CO})_3(\text{H}_2\text{O})_3$ (around 0.8 GBq) were mixed in a penicilline vial and heated for 45 min to 80°C . Quality of the radiolabelling was analyzed via HPLC according to above mentioned protocol.

Example 7: In vitro stability of ^{177}Lu -radiolabeled DOTA-AB-folate 16

The ^{177}Lu -radiolabeled DOTA-AB-folate 16 displayed higher stability in human plasma and PBS than the control compound (Figure 1, filled squares: ^{177}Lu -DOTA-folate in PBS, filled triangles: ^{177}Lu -DOTA-folate in plasma, filled circles: ^{177}Lu -DOTA-AB-folate in PBS, filled diamonds: ^{177}Lu -DOTA-AB-folate in plasma). The reason for the relatively quick degradation of the control compound ^{177}Lu -DOTA-folate in PBS is still unknown. Both of the radiofolates showed FR-specific binding to KB tumor cells (human cervical carcinoma cell line overexpressing the FR). The uptake was high and comparable for both compounds. About 75 % of total added radiotracer per 0.5 mg cell protein was bound at steady state (after 2 h incubation at 37 °C) whereof about 30 % were internalized in the case of ^{177}Lu -DOTA-AB-folate and 15 % in the case of the control compound.

Example 8: Cell uptake and internalization of ^{177}Lu -radiolabeled DOTA-AB-folate 16 in FR-positive KB-cells

Determination octanol/PBS pH 7.4 distribution coefficient (logD value) showed high negative values (-4.04 ± 0.01 for ^{177}Lu -DOTA-AB-folate 16 and -4.44 ± 0.29 for ^{177}Lu -DOTA-folate). Filter tests performed for the determination of the albumin bound fraction of the radiotracers revealed about 90% binding of ^{177}Lu -DOTA-AB-folate whereas the control compound did not show binding to albumin (Figure 2 (from left to right): columns 1 (vertically striped) and 3 (diagonally striped): ^{177}Lu -DOTA-folate at 2h and 4h, respectively; columns 2 (horizontally striped) and 4 (filled): ^{177}Lu -DOTA-AB-folate at 2h and 4h respectively).

Example 9: Biodistribution study of ^{177}Lu -radiolabeled DOTA-AB-folate 16 in KB-tumor bearing mice

In vivo experiments were performed in female nude mice bearing KB tumor xenografts. Mice have been euthanized at different time

points after injection of either ^{177}Lu -radiolabeled DOTA-AB-folate 16 or the control compound ^{177}Lu -DOTA-folate, respectively. As expected, we found an increased circulation time of the ^{177}Lu -DOTA-AB-folate 16 in the blood ($> 4\%$ ID/g and $> 1\%$ ID/g at 4 h and 24 h p.i.) compared to the control compound ($\sim 0.2\%$ ID/g and $\sim 0.05\%$ ID/g at 4 h and 24 h p.i.). The tumor uptake of ^{177}Lu -DOTA-AB-folate 16 was very high already 4 h after injection ($18.12 \pm 1.80\%$ ID/g) and retained over time. This was about 4-fold higher the uptake achieved with the control compound ($4.98 \pm 1.21\%$ ID/g, 4 h p.i.). In contrast, kidney uptake was only about 30% ID/g at 4 h and 24 h p.i. of ^{177}Lu -DOTA-AB-folate compared to more than 70% ID/g for the control compound. Thanks to the albumin binding moiety the tissue distribution of the DOTA-folate could be significantly improved in that the tumor-to-kidney ratio was increased severalfold (see Figure 3: Biodistribution in KB-tumor bearing mice 4 h and 24 h after injection of ^{177}Lu -DOTA-AB-folate (white and black, columns 3 and 4 from left to right) and the control compound ^{177}Lu -DOTA-folate (dotted and striped, columns 1 and 2 from left to right)). Figure 4(A) illustrates the uptake in a KB tumor xenograft (T) and in the kidneys (K) of a ^{177}Lu -radiolabeled DOTA-AB-folate 16 at 3 h p.i. (left image) and at 24 h p.i. (right image) at a dosage of 59 MBq. Figure 4(B) illustrates the uptake in a KB tumor xenograft (T) and in the kidneys (K) of a ^{161}Tb -radiolabeled DOTA-AB-folate 16 at 2 h p.i. (left image) and at 48 h p.i. (right image) at a dosage of 58 MBq.

Figure 5 illustrates the efficacy of therapy in KB tumor bearing nude mice over time (19 days). The mice were inoculated with 4.5 Mio KB tumor cells on each shoulder 5 days before therapy. Group A represents the mice injected at $t = 0$ days with ^{177}Lu -radiolabeled DOTA-AB-folate 16. Group B represents the control group, i.e. mice injected at $t = 0$ with PBS pH 7.4. Figure 5(A)

illustrates the relative tumor size (y-axis) of group A (filled diamonds) and group B (filled triangles) over time (x-axis). Figure 5(B) shows the results on $t = 17$ days.

5 Example 10: Biodistribution study of ^{18}F -FDG-AB-folate 34 in KB-tumor bearing mice

In vivo experiments were performed in female nude mice bearing KB tumor xenografts.

10 Animals were injected with ~ 3 MBq of $[^{18}\text{F}]$ FDG-AB-folate 34 via a lateral tail vein. Animals were sacrificed 2 h after radiotracer injection. Organs- and tissues were dissected and measured in the γ -counter (Wizard, PerkinElmer). The incorporated radioactivity was expressed as percentage injected dose (%ID) per gram of tissue. The biodistribution data is summarized in
15 Table 1.

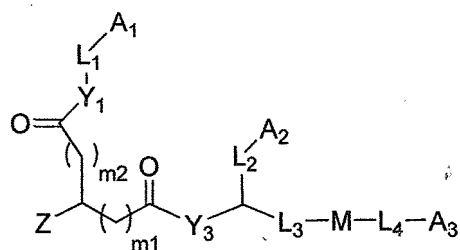
Table 1: Ex vivo biodistribution studies with $[^{18}\text{F}]$ FDG-AB-folate in nude mice bearing KB tumor xenografts 2 h after injection

Organ or tissue	2 h p.i. ($n = 3$)
%ID/g in:	
Blood	2.93 ± 0.55
Brain	0.89 ± 0.17
Heart	3.20 ± 0.53
Lungs	2.44 ± 0.58
Spleen	0.81 ± 0.03
Liver	7.14 ± 0.78
Gallbladder	83.78 ± 16.78
Kidneys	14.98 ± 1.47
Stomach	1.93 ± 0.51

Intestine	8.84 ± 7.07
Feces	53.02 ± 26.52
Salivary glands	7.55 ± 0.08
Bone	1.50 ± 0.24
Muscle	1.62 ± 0.26
Tumor	9.51 ± 0.08
Ratios:	
Tumor / Liver	1.34 ± 0.16
Tumor / Kidneys	0.64 ± 0.07
Tumor / Blood	3.33 ± 0.59

Claims

1. Compounds of formula I,



I

wherein

Z is a pterate or derivative thereof,

L₁, L₄ are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl,

L₂ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

L₃ is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-,

-S-R'-, -SO₃R'-, and a five-membered azaheterocycle,
wherein R' represents H or C(1-8)alkyl,

Y₁, Y₃ are independently of each other O, N or S,

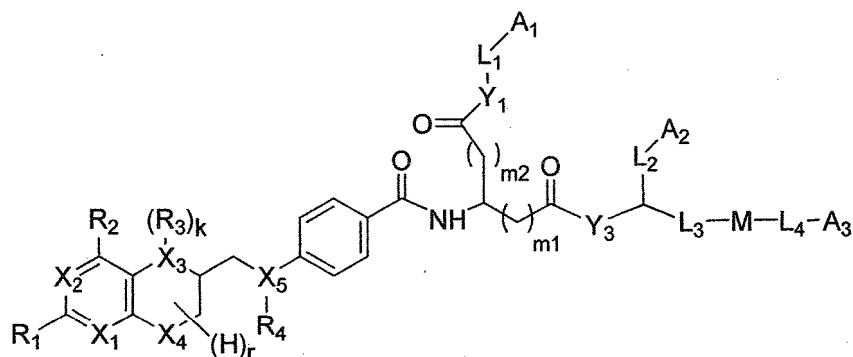
A₁, A₂, A₃ are independently of each other H, a capping group, or an
5 albumin binder,

M is a radionuclide-based therapeutic or diagnostic moiety,

m₁, m₂ are independently of each other 0, 1, 2 or 3,

with the proviso that at least one, preferably one, of A₁, A₂ and
A₃ is an albumin binder.

2. Compound according to claim 1 having formula II



II

wherein

X₁ to X₅ are independently of each other C, N or O, preferably N
or O,

R₁, R₂ are independently of each other H, halogen, C(1-12)alkyl,
C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -
20 NHR₅, -CONHR₅, -CONHR₅, wherein R₅ represents H, halo,
C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR', -
COR', -COOR', or -NHR', wherein R' is H or C(1-8)alkyl,

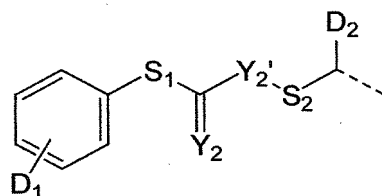
R₃, R₄ are independently of each other H, nitroso, C(1-12)alkyl,
-OR', -COR' or halosubstituted -COR', wherein R'
25 represents H or C(1-8)alkyl,

- L_1, L_4 are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl,
- L_2 is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,
- L_3 is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, and a five-membered azaheterocycle, wherein R' represents H or C(1-8)alkyl,
- Y_1, Y_3 are independently of each other O, N or S,
- A_1, A_2, A_3 are independently of each other H, a capping group, or an albumin binder,
- M is a radionuclide-based therapeutic or diagnostic moiety, preferably an imaging moiety M_1 or M_2 , wherein M_1 is a chelated metal radionuclide, and M_2 is a gamma- or positron-emitting non-metal radionuclide optionally in combination with a prosthetic group,
- m_1, m_2 are independently of each other 0, 1, 2 or 3,
- k is 0 or 1,

r has a value of 1 to 7.

with the proviso that at least one, preferably one, of A₁, A₂ and A₃ is an albumin binder.

3. Compound according to claims 1 or 2 wherein the albumin binder is a linear or branched lipophilic group having 12-40 carbon atoms and a distal acidic group, such as a compound of formula III



III

wherein

Y₂, Y₂' are independently of each other N, O or S,

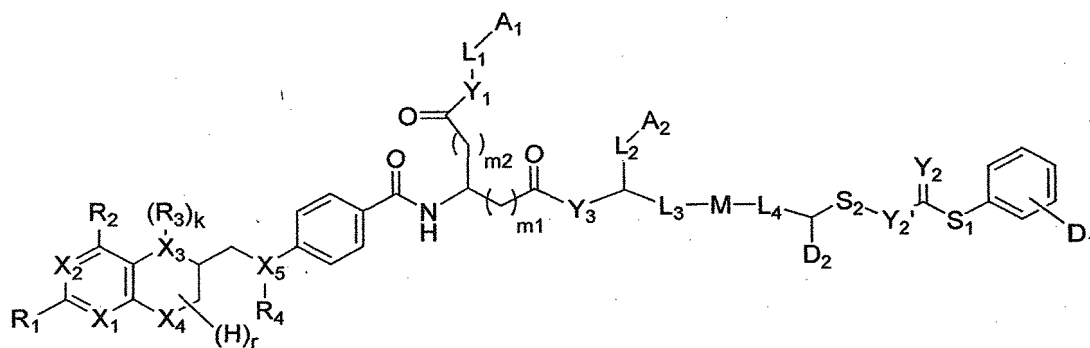
S₁, S₂ are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D₁ is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

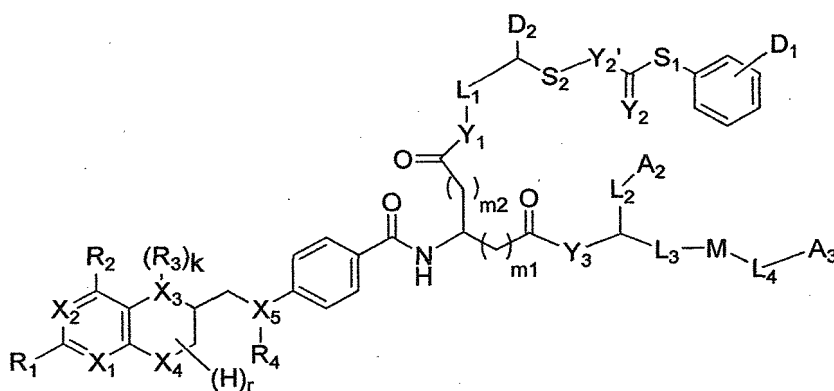
D₂ is an acidic group,

and the broken line represents the linkage to L₁, L₂ or L₄.

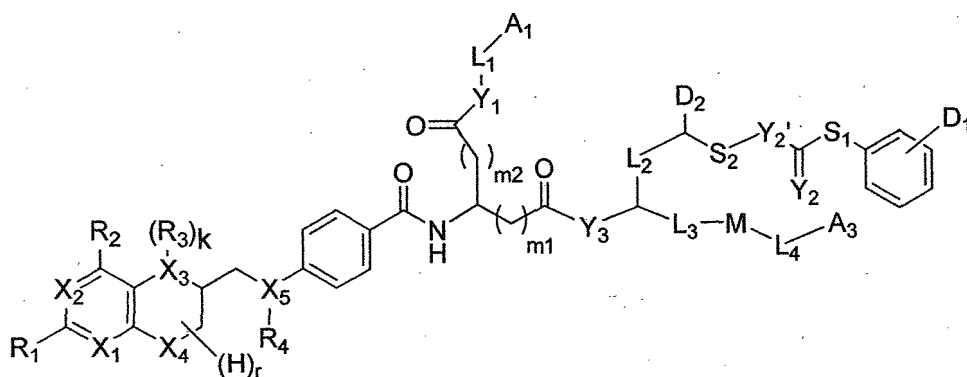
4. A compound according to any preceding claim, having formula IVa, IVb and IVc



IVa



IVb



IVc

wherein

- 10 X_1 to X_5 are independently of each other C, N or O, preferably N or O,

Y_1, Y_3 are independently of each other N, O or S,

Y_2, Y_2' are independently of each other N, O or S,

- R_1, R_2 are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR_5$, $-COR_5$, $-COOR_5$, $-NHR_5$, $-CONHR_5$, wherein R_5 represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, or $-NHR'$, wherein R' is H or C(1-8)alkyl,
- R_3, R_4 are independently of each other H, nitroso, C(1-12)alkyl, $-OR'$, $-COR'$ or halosubstituted $-COR'$, wherein R' represents H or C(1-8)alkyl,
- A_1 is H or a capping group,
- A_2 is H or a capping group,
- A_3 is H or a capping group,
- M is a radionuclide-based therapeutic or diagnostic moiety, preferably an imaging moiety M_1 or M_2 , wherein M_1 is a chelated metal radionuclide, and M_2 is a gamma- or positron-emitting non-metal radionuclide optionally in combination with a prosthetic group,
- m_1, m_2 are independently of each other 0, 1, 2 or 3,
- L_1, L_4 are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' , NHR' , CO_2R' or NO_2 , wherein R' represents H or C(1-8)alkyl,
- L_2 is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR' , NHR' , or CO_2R' , and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$, $-NR'-$, $-NR'-CO-$, $-CO-NR'$, wherein R' represents H or C(1-8)alkyl,
- L_3 is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or

substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, and a five-membered azaheterocycle, wherein R' represents H or C(1-8)alkyl,

S₁, S₂ are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D₁ is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

D₂ is an acidic group,

k is 0 or 1,

r has a value of 1 to 7.

5. A compound according to claim 4, wherein the acidic group is a group capable of ionizing to donate a hydrogen ion to a base to form a salt, preferably selected from the group consisting of -COOH, -SO₃H, -SO₂H, -NR'SO₃H, -P(O)(OH)₂, wherein R' represents H or C(1-8)alkyl.

6. A compound according to claim..., wherein S₁ and S₂ are independently of each other a single bond or a spacer

selected from a straight-chain or branched C(1-8)alkyl, wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -COO-, -NR'-, -NR'-CO-, -CO-NR'-, -CH=CH-, wherein R' represents H or C(1-8)alkyl.

7. A compound according to any preceding claim, wherein m₁ is 2 and m₂ is 0 or wherein m₁ is 0 and m₂ is 2.

8. A compound according to any preceding claim, wherein M is a radionuclide-based therapeutic or diagnostic moiety M₁ or M₂, wherein M₁ is a chelated metal radionuclide, and M₂ is a gamma- or positron-emitting non-metal radionuclide optionally in combination with a prosthetic group.

9. A compound according to any preceding claim, with the proviso that when A₂ is an albumin binder, M is a chelated metal radionuclide M₁.

10. A compound according to any preceding claim, wherein the imaging moiety M is a chelated metal radionuclide M₁ comprising a metal radionuclide and a metal chelator.

11. A compound according to claim 9, wherein the metal radionuclide is selected from ⁹⁹Tc, ⁵¹Cr, ⁶⁷Ga, ⁶⁸Ga, ⁴⁷Sc, ⁵¹Cr, ¹⁶⁷Tm, ¹⁴¹Ce, ¹¹¹In, ¹⁶⁸Yb, ¹⁷⁵Yb, ¹⁴⁰La, ⁹⁰Y, ⁸⁸Y, ¹⁵³Sm, ¹⁶⁶Ho, ¹⁶⁵Dy, ¹⁶⁶Dy, ⁶²Cu, ⁶⁴Cu, ⁶⁷Cu, ⁹⁷Ru, ¹⁰³Ru, ¹⁸⁶Re, ¹⁸⁸Re, ²⁰³Pb, ²¹¹Bi, ²¹²Bi, ²¹³Bi, ²¹⁴Bi, ¹⁰⁵Rh, ¹⁰⁹Pd, ¹¹⁷mSn, ¹⁴⁹Pm, ¹⁶¹Tb, ¹⁷⁷Lu, ¹⁹⁸Au and ¹⁹⁹Au.

12. A compound according to claim 9, wherein the metal

chelator is selected from a bidentate, tridentate, or tetradentate, linear, tripodal or macrocyclic ligand, preferably a linear or macrocyclic polyaminocarboxylate chelator, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, EHPG, HBED, NOTA, DOTMA, TETMA, PDTA, TTHA, LICAM, MECAM, more preferably a macrocyclic polyaminocarboxylate.

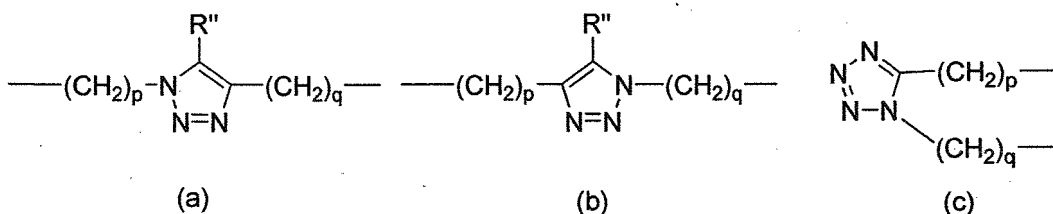
- 5
13. A compound according to any preceding claim, wherein the imaging moiety M is a gamma- or positron-emitting non-metal radionuclide M₂ selected from ¹¹C, ¹³N, ¹⁵O, ¹⁷F, ¹⁸F, ⁷⁵Br, ⁷⁶Br, ⁷⁷Br, ¹²³I, ¹²⁴I, ¹³¹I, optionally in combination with a prosthetic group.
- 10
14. Compound according to any preceding claim, wherein L₁ is a covalent bond or straight-chain or branched C(1-6)alkyl.
- 15
15. Compound according to any one of claims 4 to 14, wherein L₂ is a covalent bond or straight-chain or branched C(1-6)alkyl.
- 20
16. Compound according to any one of claims 3 to 14, wherein Y₂ is O and/or Y₂' is N.
- 25
17. Compound according to any preceding claim, wherein Y₁ is O or N and/or Y₃ is N.
- 30
18. Compound according to any preceding claim, wherein L₃ is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a

group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-, and a five-membered azaheterocycle selected from a triazolyl or tetrazolyl group wherein R' represents H or C(1-8)alkyl.

5

19. A compound according to any preceding claim, wherein L₃ is a group L₃', which is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-, or L₃ is a group of formulae (a), (b), or (c)

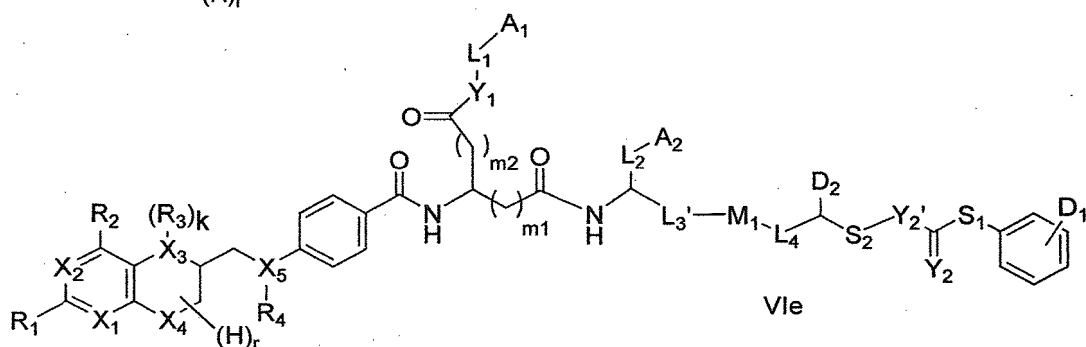
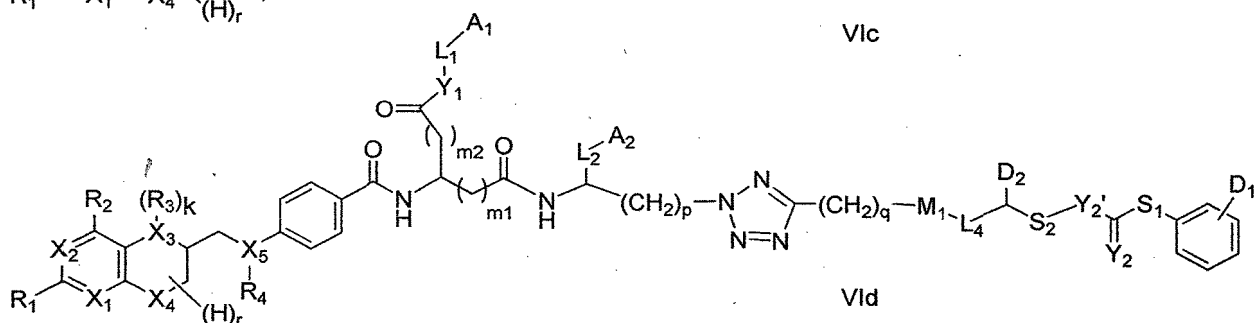
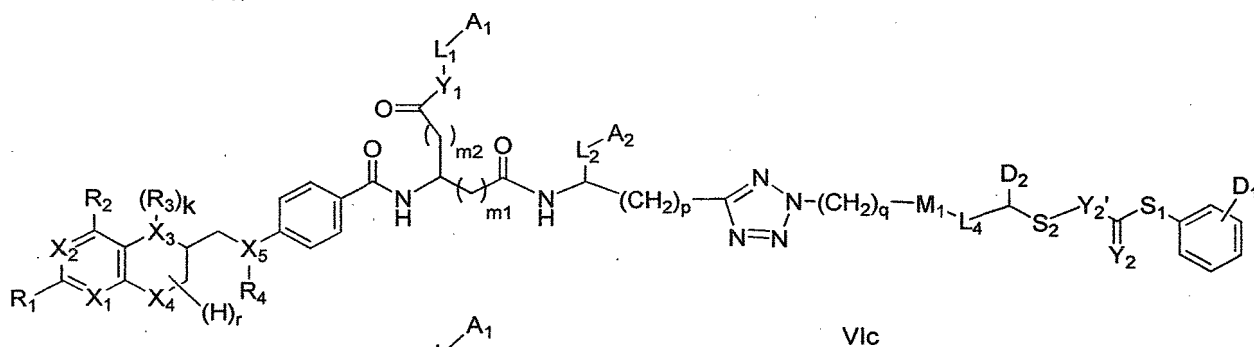
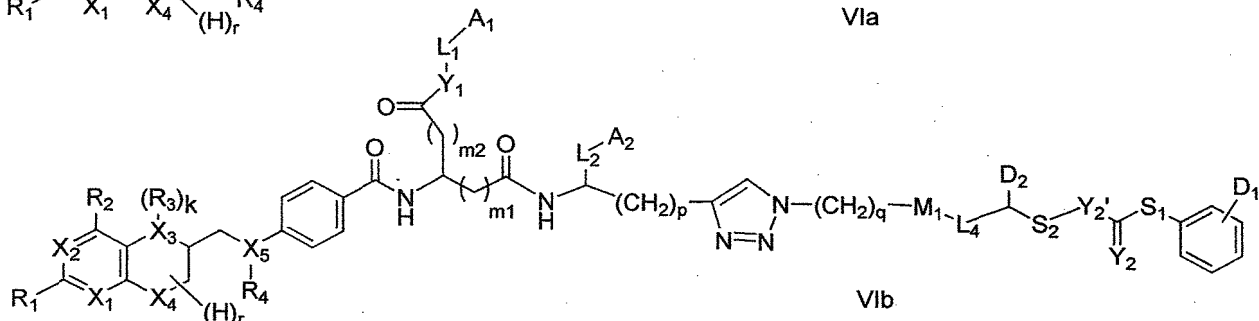
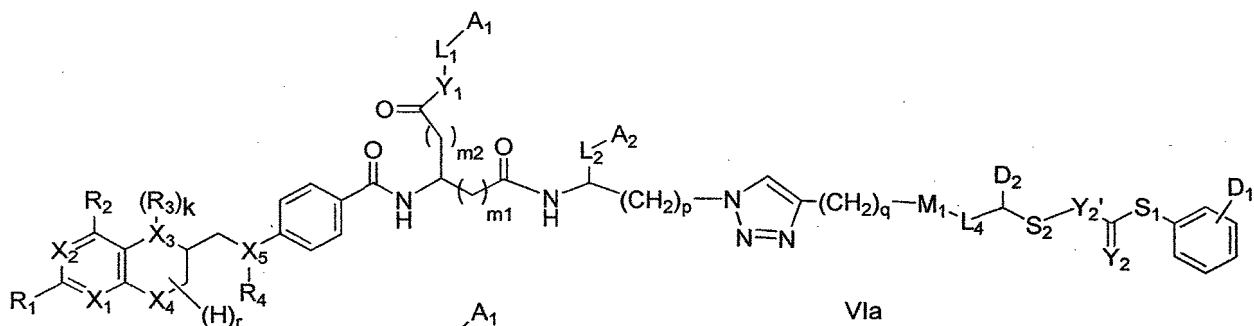
10



wherein

- 15 R'' is H or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, or NO₂,
- p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6.

- 20 20. A compound according to any preceding claim having formulae VI a-e



5 wherein

X_1 to X_5 are independently of each other C, N or O, preferably N or O,

Y_1, Y_2, Y_2' are independently of each other N, O or S,

R_1, R_2 are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR_5$, $-COR_5$, $-COOR_5$, $-NHR_5$, $-CONHR_5$, wherein R_5 represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, or $-NHR'$, wherein R' is H or C(1-8)alkyl,

R_3, R_4 are independently of each other H, nitroso, C(1-12)alkyl, $-OR'$, $-COR'$ or halosubstituted $-COR'$, wherein R' represents H or C(1-8)alkyl,

R_5 is H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

A_1 is H or a capping group,

A_2 is H or a capping group,

M_1 is a linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a radioimaging metal ion,

m_1, m_2 are independently of each other 0, 1, 2 or 3,

L_1, L_4 are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' , NHR' , CO_2R' or NO_2 , wherein R' represents H or C(1-8)alkyl,

L_2 is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR' , NHR' , or CO_2R' , and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$, $-NR'-$, $-NR'-CO-$, $-CO-NR'$, wherein R' represents H or C(1-8)alkyl,

S_1, S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN,

Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-,
 5 , -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl

D₁ is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,
 10

D₂ is an acidic group,

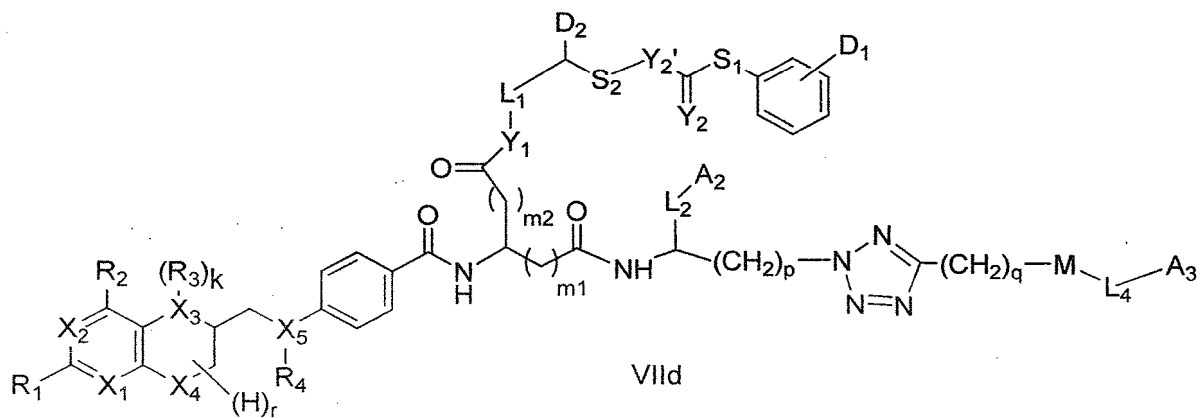
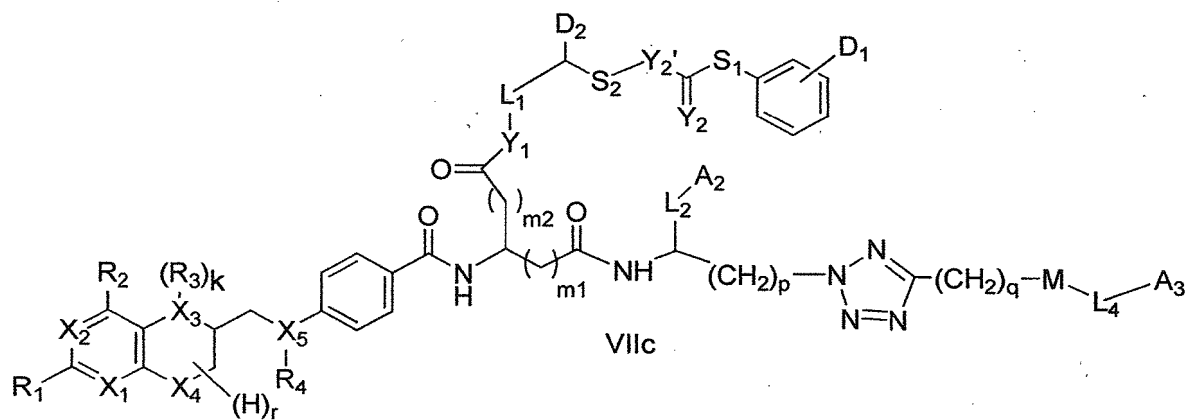
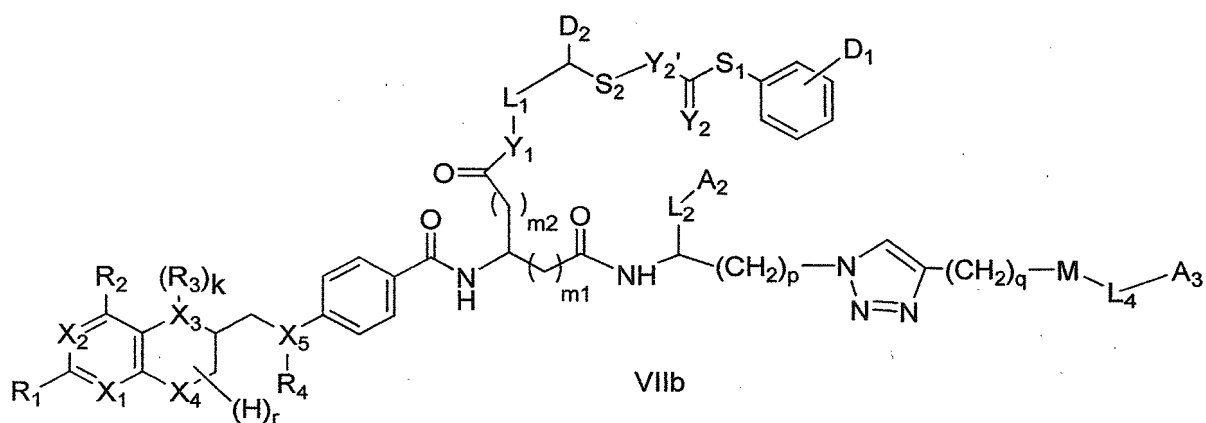
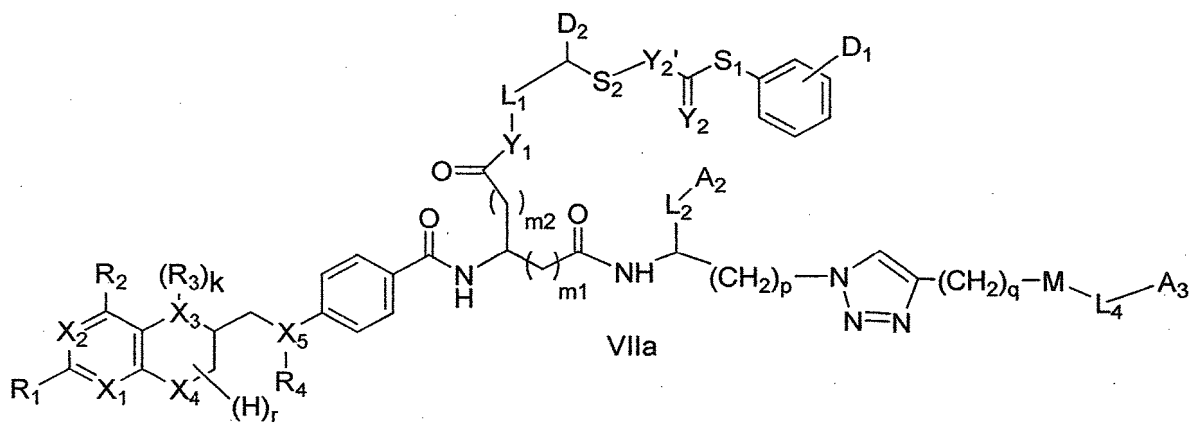
L₃ is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-,
 15 wherein R' represents H or C(1-8)alkyl,

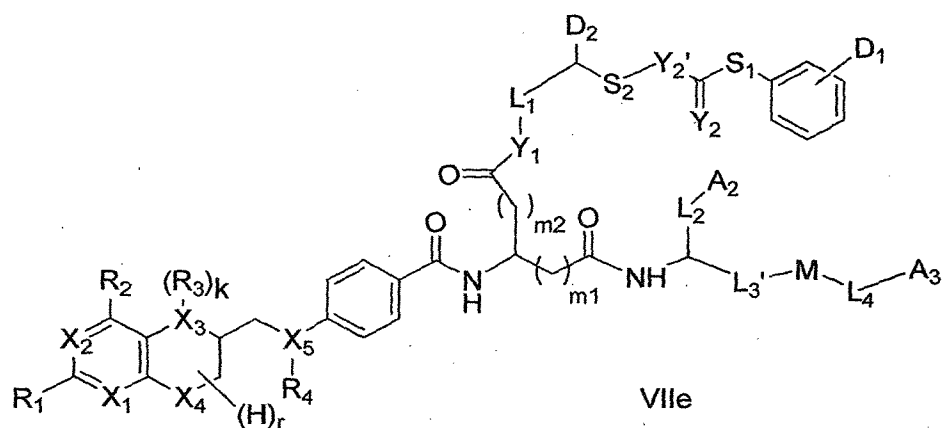
p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6,

k is 0 or 1,

20 r has a value of 1 to 7.

21. A compound according to any preceding claim having formulae VII a-e





wherein

X_1 to X_5 are independently of each other C, N or O, preferably N or O,

5 Y_1, Y_2, Y_2' are independently of each other N, O or S,

R_1, R_2 are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR_5$, $-COR_5$, $-COOR_5$, $-NHR_5$, $-CONHR_5$, wherein R_5 represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, or $-NHR'$, wherein R' is H or C(1-8)alkyl,

10 R_3, R_4 are independently of each other H, nitroso, C(1-12)alkyl, $-OR'$, $-COR'$ or halosubstituted $-COR'$, wherein R' represents H or C(1-8)alkyl,

A_2 is H or a capping group,

15 A_3 is H or a capping group,

M is an imaging moiety M_1 or M_2 , wherein M_1 is a chelated metal radionuclide selected from a linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a metal radionuclide ion, and wherein M_2 is a gamma- or positron-emitting non-metal radionuclide, preferably selected from ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F , ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I optionally in combination with a prosthetic group,

m_1, m_2 are independently of each other 0, 1, 2 or 3,

L_1, L_4 are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R' or NO₂, wherein R' represents H or C(1-8)alkyl,

L_2 is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a group selected from -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR', wherein R' represents H or C(1-8)alkyl,

S_1, S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR', NHR', CO₂R', SH, SO₃H or NO₂, and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein R' represents H or C(1-8)alkyl,

D_1 is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

D_2 is an acidic group,

L_3 is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a

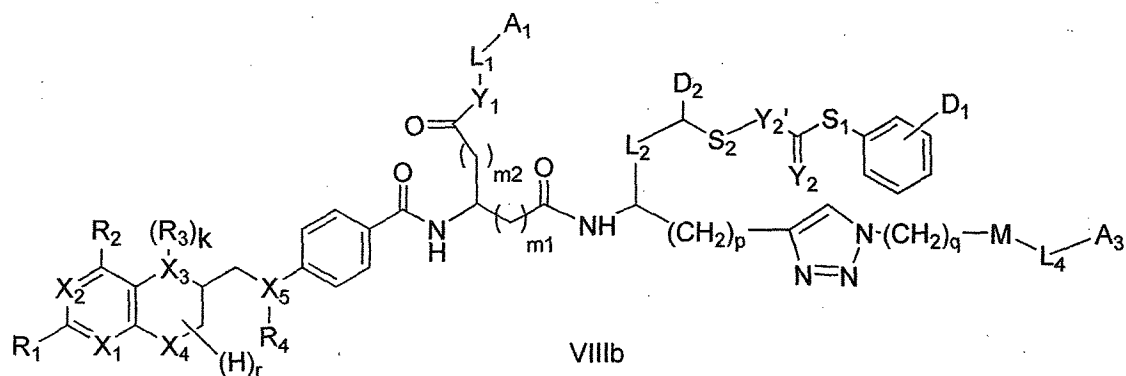
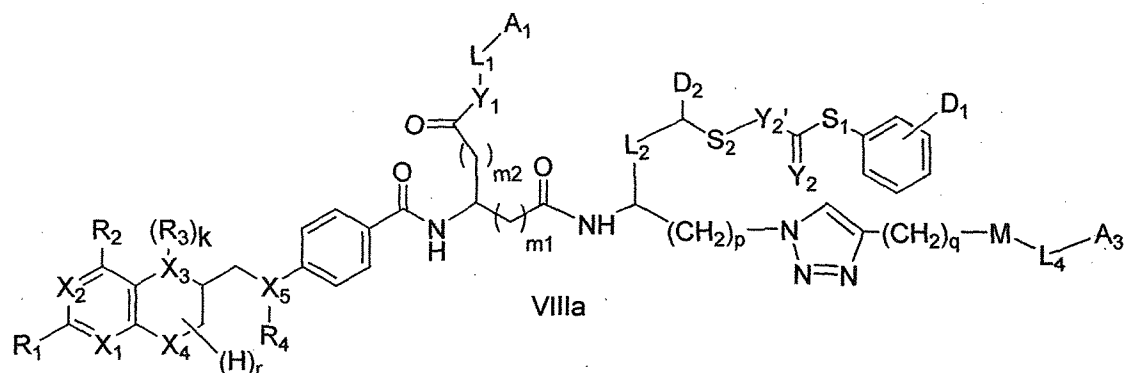
group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-,
wherein R' represents H or C(1-8)alkyl,

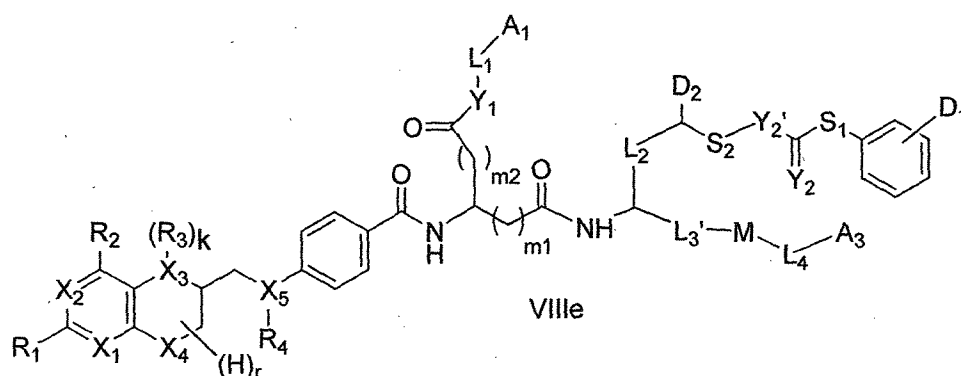
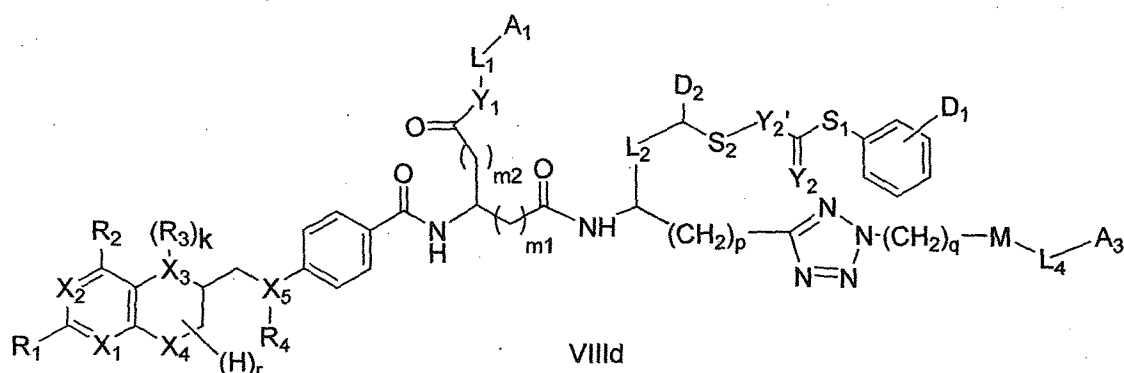
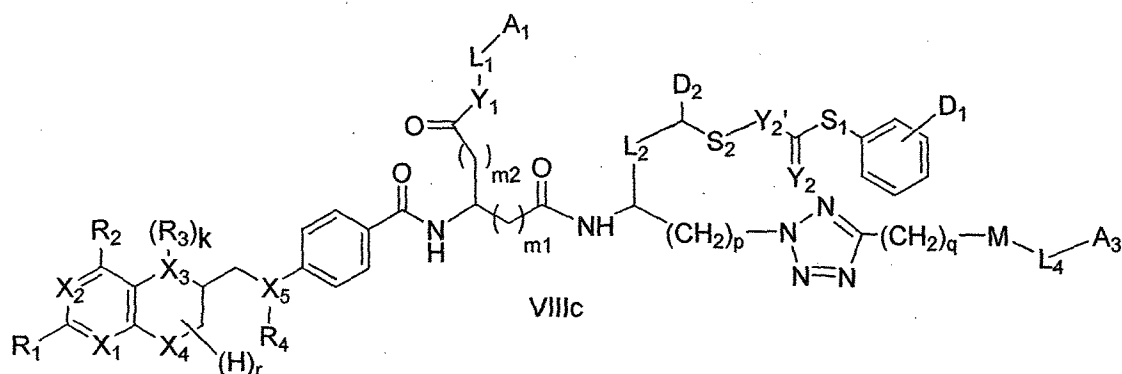
p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6,

k is 0 or 1,

5 r has a value of 1 to 7.

22. A compound according to any preceding claim having formulae
VIII a-e





wherein

X_1 to X_5 are independently of each other C, N or O, preferably N or O,

Y_1, Y_2, Y_2' are independently of each other N, O or S,

R_1, R_2 are independently of each other H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR_5$, $-COR_5$, $-COOR_5$, $-NHR_5$, $-CONHR_5$, wherein R_5 represents H, halo, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, $-OR'$, $-COR'$, $-COOR'$, or $-NHR'$, wherein R' is H or C(1-8)alkyl,

- R_3, R_4 are independently of each other H, nitroso, C(1-12)alkyl, $-OR'$, $-COR'$ or halosubstituted $-COR'$, wherein R' represents H or C(1-8)alkyl,
- A_1 is H or a capping group,
- 5 A_3 is H or a capping group,
- M is an imaging moiety M_1 or M_2 , wherein M_1 is a chelated metal radionuclide selected from a linear or macrocyclic polyaminocarboxylate, such as DTPA, DOTA, DO3A, HP-DO3A, EDTA, TETA, NOTA, DOTMA, complexed with a radioimaging metal ion, and wherein M_2 is a gamma- or positron-emitting non-metal radionuclide, preferably selected from ^{11}C , ^{13}N , ^{15}O , ^{17}F , ^{18}F , ^{75}Br , ^{76}Br , ^{77}Br , ^{123}I , ^{124}I , ^{131}I , optionally in combination with a prosthetic group,
- 10 m_1, m_2 are independently of each other 0, 1, 2 or 3,
- 15 L_1, L_4 are independently of each other a covalent bond or a straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' , NHR' , CO_2R' or NO_2 , wherein R' represents H or C(1-8)alkyl,
- 20 L_2 is a covalent bond or a linking group, such as straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one OR' , NHR' , or CO_2R' , and wherein one or more of the non-adjacent CH_2 groups may independently be replaced by a group selected from $-O-$, $-CO-$, $-CO-O-$, $-O-CO-$, $-NR'-$, $-NR'-CO-$, $-CO-NR'$, wherein R' represents H or C(1-8)alkyl,
- 25 S_1, S_2 are independently of each other a single bond or a spacer selected from a straight-chain or branched C(1-12)alkyl, which is unsubstituted or substituted by at least one CN, Hal, OR' , NHR' , CO_2R' , SH, SO_3H or NO_2 ,
- 30

and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by -O-, -CO-, -CO-O-, -O-CO-, -NR'-, -NR'-CO-, -CO-NR'-, -NR'-CO-O-, -O-CO-NR'-, -NR'-CO-NR'-, -CH=CH-, -C≡C-, -O-CO-O-, -S-R'-, -SO₃R'-, wherein
5 R' represents H or C(1-8)alkyl,

D₁ is a group selected from H, halogen, C(1-12)alkyl, C(2-12)alkenyl, C(2-12)alkynyl, -OR₅, -COR₅, -COOR₅, -NHR₅, -CONHR₅, wherein R₅ represents H, C(1-12)alkyl, C(2-12)alkenyl, or C(2-12)alkynyl,

10 D₂ is an acidic group,

L₃ is straight-chain or branched C(1-8)alkyl, which is unsubstituted or substituted by at least one Hal, OR', NHR', or CO₂R', and wherein one or more of the non-adjacent CH₂ groups may independently be replaced by a
15 group selected from -O-, -CO-, -CO-O-, -NR'-, -NR'-CO-, wherein R' represents H or C(1-8)alkyl,

p, q are independently of each other 0, 1, 2, 3, 4, 5 or 6,

k is 0 or 1,

r has a value of 1 to 7.

20 23. A compound according to any one of claims 2 to 22, wherein R₃ is H, C(1-12)alkyl, or -CO-C(1-8)alkyl.

24. A compound according to any one of claims 2 to 23,
25 wherein R₄ is H, nitroso, -O-C(1-8)alkyl, or -CO-C(1-8)alkyl.

25. A compound according to any one of claims 2 to 24,
wherein R₁ and R₂ are independently of each other H, C(1-12)alkyl, -OR₅, -NHR₅, wherein R₅ is H, C(1-12)alkyl,
30

C(2-12)alkenyl, or C(2-12)alkynyl.

26. Pharmaceutical composition comprising at least one compound according to any preceding claim.
- 5 27. Use of a compound according to any one of claims 1 to 25 for diagnostic imaging of a cell or population of cells expressing a folate-receptor in vitro or in vivo.
- 10 28. Use of a compound according to any one of claims 1 to 25 for convenient and effective administration to a subject in need for diagnostic imaging.
- 15 29. Method for diagnostic imaging of a cell or population of cells expressing a folate-receptor, said method comprising the steps of administering at least one compound according to any one of claims 1 to 25 in a diagnostic imaging amount, and obtaining a diagnostic image of said cell or population of cells.
- 20 30. Method according to claim 29, wherein the diagnostic imaging is performed of a cell or population of cells expressing a folate-receptor in vitro or in vivo.
- 25 31. Method for in vitro detection of a cell expressing the folate receptor in a tissue sample which includes contacting said tissue sample with a compound according to any one of claims 1 to 25 in effective amounts and for sufficient time and conditions to allow binding to occur and detecting such binding by PET imaging.
- 30 32. Method of diagnostic imaging or monitoring a subject comprising the steps of (i) administering at least one

compound according to any one of claims 1 to 25 in a diagnostic imaging amount, and (ii) performing diagnostic imaging using PET by detecting a signal from said at least one compound.

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33. Method of monitoring cancer therapy in a subject comprising the steps of (i) administering to a subject in need thereof at least one compound according to any one of claims 1 to 25 in a diagnostic imaging amount in combination with a therapeutically active, and (ii) performing diagnostic imaging using PET by detecting a signal from said at least one compound to follow the course of cancer therapy.

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15 34. Method of claims 32 or 33 used in combination with any other methods of cancer diagnosis or therapy.