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(54) Title: SPLIT ANTIBODIES WHICH BIND TO CANCER CELLS AND TARGET RADIONUCLIDES TO SAID CELLS

(57) Abstract: 197 ABSTRACT OF THE DISCLOSURE The present invention relates to antibodies which bind to antigens on target cells and which target effector moieties to said cells, and to methods of using the same.

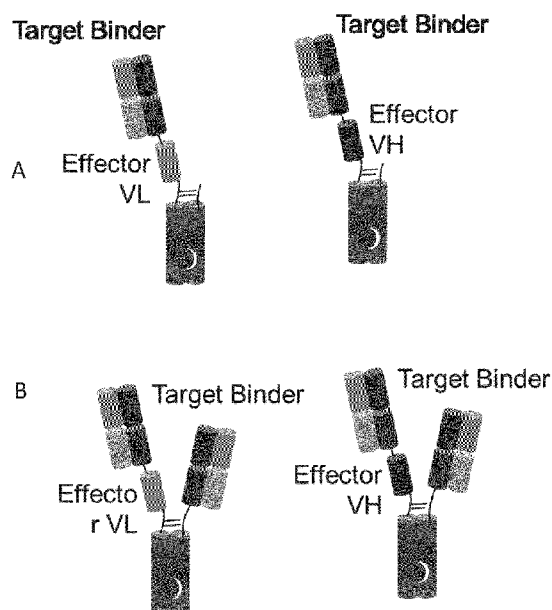


FIG. 29



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SPLIT ANTIBODIES WHICH BIND TO CANCER CELLS AND TARGET RADIONUCLIDES TO SAID CELLS

FIELD OF THE INVENTION

The present invention relates to novel formats for bi-specific antibodies. In certain embodiments the invention further relates to antibodies having this novel format which bind to antigens on target cells and which target radionuclides to said cells, and to methods of using the same.

BACKGROUND

10 The selective destruction of an individual cell or a specific cell type is often desirable in a variety of clinical settings. For example, it is a primary goal of cancer therapy to specifically destroy tumour cells, while leaving healthy cells and tissues intact and undamaged.

In this regard, bispecific antibodies have been designed which bind with one "arm" to a surface antigen on target cells, and with the second "arm" to an effector moiety such as a drug. A large variety of bispecific formats have been developed, but the task of developing bispecific antibodies is by no means trivial.

20 In pre-targeted radioimmunotherapy (PRIT), use is made of an antibody construct which has affinity for the tumour-associated antigen on the one hand and for a radiolabelled compound on the other. In a first step, the antibody is administered and localises to tumour. Subsequently, the radiolabelled compound is administered. Because the radiolabelled compound is small, it can be delivered quickly to the tumour and non-bound compound is fast-clearing, which reduces radiation exposure outside of the tumour (Goldenberg *et al* Theranostics 2012, 2(5), 523-540). A similar procedure can also be used for imaging. Pre-targeting can make use of a bispecific antibody or systems using avidin-biotin, although the latter has the disadvantage that avidin/streptavidin is immunogenic.

30 Methods of pre-targeted radioimmunotherapy or imaging commonly make use of a clearing or blocking agent, which is administered between the step of administering the antibody and the step of administering the radiolabelled compound. The purpose is to clear antibody from the blood and/or to block the binding site of the circulating antibody for the radiolabelled compound (see for instance Karacay et al, Bioconj. Chem., 13(5), 1054-1070

(2002)). The use of a clearing or blocking agent allows for sufficient levels of radioactivity to be administered for an efficient treatment while limiting adverse toxicity, but the timing and dosage must be chosen with care, and there is the possibility of the clearing agent introducing risks of adverse effects such as immune reactions. Thus, the use of a clearing phase is a complicating aspect in pre-targeting methods.

SUMMARY

The present invention provides novel formats for bi-specific antibodies in which the VH and VL domain for the effector moiety are split into two parts, and methods of using the same.

- 10 In particular, the present invention relates to a set of antibodies comprising
- i) a first antibody comprising:
 - a) an antigen binding moiety which binds to an antigen expressed on the surface of a target cell;
 - b) a polypeptide comprising or consisting of an antibody heavy chain variable domain (VH) of an antigen binding site for an effector moiety; and
 - c) an Fc domain comprising two subunits,wherein the polypeptide of (b) is fused by its N-terminus to the C-terminus of the antibody binding moiety of (a) and by its C-terminus to the N-terminus of one of the subunits of the Fc domain of (c);
- 20 and wherein the first antibody does not comprise a VL domain of an antigen binding site for the effector moiety; and
- ii) a second antibody comprising:
 - d) an antigen binding moiety which binds to an antigen expressed on the surface of a target cell;
 - e) a polypeptide comprising or consisting of an antibody light chain variable domain (VL) of an antigen binding site for the effector moiety; and
 - f) an Fc domain comprising two subunits,wherein the polypeptide of (e) is fused by its N-terminus to the C-terminus of the antigen binding moiety of (d) and by its C-terminus to the N-terminus of one of the subunits
- 30 of the Fc domain of (f);
- and wherein the second antibody does not comprise a VH domain of an antigen binding site for the effector moiety;

wherein said V_H domain of the first antibody and said V_L domain of the second antibody are together capable of forming a functional antigen binding site for the effector moiety.

Neither the first nor the second antibody comprise, on their own, a functional antigen binding site for the effector moiety. The first antibody has only a V_H domain from the functional binding site for the effector moiety, and not the V_L domain. The second antibody has only the V_L domain, and not the V_H domain.

10 A functional antigen binding site for the effector moiety is formed when the V_H and V_L domains of the first and second antibodies are associated. This may occur, for example, when the first and second antibodies are bound to the same individual target cell or to adjacent cells.

The first and second antibodies described herein may be referred to herein as “single domain split antibodies”, “split antibodies”, “SPLITs”, “hemibodies” or “demibodies”. The V_H and V_L domain which together form an antigen binding site capable of binding to the effector moiety are split between two antibodies, and not present as part of the same antibody.

20 The split domain format means that the effector moiety cannot bind to either the first antibody on its own or to the second antibody on its own. In the blood, there is little or no stable association between the first and the second antibody, and so little or no stable binding of the radiolabelled compound.

An antigen expressed on the surface of a target cell may be referred to herein as a “target antigen”, “target cell antigen”, or “TA”. According to the present invention, the first and the second antibody described above may have an antigen-binding moiety which binds to different target antigens, or for the same target antigen. (For the avoidance of doubt, where it stated that the antibodies bind the same target antigen, this means that they have a binding site capable of binding to the same target antigen and includes the possibility that the antibodies may bind to two individual antigen molecules that are the same as each other). For example, in one embodiment, both the first and the second antibody bind to CEA.

30 In some embodiments, the first and second antibody may bind to (have a binding site for) the same epitope of the same target antigen. In other embodiments, the first and second antibody may bind to (have a binding site for) different epitopes of the same target antigen.

In some embodiments, the first and second antibody may comprise the same antigen binding site for the target antigen. For instance, they may comprise an antigen binding site

capable of binding to the target antigen, comprising a V_L and V_H sequence, where the V_L and V_H sequences forming this antigen binding site are the same in the first and in the second antibodies.

The term “effector moiety” refers to a moiety which is responsible for the desired effect on the target cell. In one embodiment, the desired effect is cell killing. For instance, the effector moiety may be a radionuclide, drug or a toxin used for killing the target cell. In one embodiment, the effector moiety may be a radiolabelled compound suitable for radiotherapy. In another embodiment, the desired effect is labelling and the effector moiety may be a radiolabelled compound suitable for imaging.

10 In some embodiments, the antigen binding moiety of (a) and (d) may be an antibody fragment such as a Fv, Fab, cross-Fab, Fab', Fab'-SH, F(ab')₂; diabody; linear antibody; single-chain antibody molecule (e.g., scFv or scFab); or single domain antibody (dAbs) such as VHH; or a non-antibody binding scaffold such as a DARPIn (designed ankyrin repeat protein); affibody; Sso7d; monobody or anticalin.

In some embodiments, it may be preferred that the antigen binding moiety of (a) and the antigen binding moiety of (d) is a Fab. In such an embodiment, the polypeptide of (b) is fused by its N-terminus to the C-terminus of one of the chains of the Fab fragment of (a); and the polypeptide of (e) is fused by its N-terminus to the C-terminus of one of the chains of the Fab fragment of (d). Likewise, in other embodiments where the antigen-binding moiety
20 comprises more than one chain, the polypeptide may be fused by its N-terminus to the C-terminus of one of the chains.

The presence of an Fc region has benefits in the context of immunotherapy and imaging, e.g. prolonging the protein's circulating half-life and/or resulting in higher tumour uptake than may be observed with smaller fragments. The “split domain” format described herein may be particularly advantageous in this context, as it mitigates against the greater possibility of association with effector moiety/radiolabelled compound that would otherwise occur due to the prolonged presence of the circulating antibody. In some embodiments, the Fc domain is modified to reduce or eliminate Fc effector function.

30 The format described herein avoids significant off-target association of the V_L and V_H for the effector moiety. Moreover, antibodies according to the present invention may have advantages in terms of a reduced anti-drug antibody response and/or stability.

In another aspect, the present invention provides a pharmaceutical composition comprising the set of antibodies as described herein. In another aspect, the present invention

provides a kit comprising two separate pharmaceutical compositions, each comprising one of the antibodies described herein (i.e., the first and second antibody respectively).

In a further aspect, the present invention relates to a polynucleotide or set of polynucleotides encoding any of the antibodies or sets of antibodies described herein. In another aspect, the present invention relates to a vector or set of vectors comprising said polynucleotide or polynucleotides, optionally an expression vector or set of expression vectors. In a further object the present invention relates to a prokaryotic or eukaryotic host cell or a set of host cells comprising a vector or set of vectors of the present invention. In addition there is provided a method of producing an antibody comprising culturing the host
10 cell(s) so that the antibody is produced.

In some embodiments, where the effector molecule is a radiolabelled compound, antibodies as described herein find use in a method of pre-targeted radioimmunotherapy (PRIT) or in a method of pre-targeted radioimaging.

In one aspect, the present invention provides a method of pre-targeted radioimmunotherapy which comprises:

- i) administering to a subject a first antibody and a second antibody as described above; and
- ii) subsequently administering to said subject a radiolabelled compound.

In another aspect, the present invention provides a first and a second antibody
20 described above for use in a method of treatment comprising administering the first antibody and the second antibody to a subject, and subsequently administering to said subject a radiolabelled compound. In another aspect, the invention provides a first antibody as described above for use in a method of treatment comprising administering the first antibody and the second antibody to a subject, and subsequently administering to said subject a radiolabelled compound. In another aspect the invention provides the second antibody as described above for use in a method of treatment comprising administering the first antibody and the second antibody to a subject, and subsequently administering to said subject a radiolabelled compound.

In another aspect, the present invention provides a method of radioimaging which
30 comprises:

- i) administering to a subject a first and a second antibody as described herein, wherein the antibodies bind to the target antigen and localise to the surface of a cell expressing the target antigen;

- ii) subsequently administering a radiolabelled compound; and optionally
- iii) imaging the tissue or organ where the radionuclide has localised.

In another aspect, the present invention provides a first and a second antibody as described herein for use in a method of diagnosis carried out on the human or animal body, wherein the method comprises

- i) administering to subject a first and a second antibody as described herein, wherein the antibodies bind to the target antigen and localise to the surface of a cell expressing the target antigen;
- ii) subsequently administering a radiolabelled compound; and optionally
- iii) imaging the tissue or organ where the radionuclide has localised.

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The imaging step may be followed by a step of forming a diagnosis and optionally a step of delivering that diagnosis to the subject. In some embodiments the method may further comprise determining an appropriate treatment and optionally administering that treatment to the subject.

In each of the above methods/uses, binding of the first and the second antibody to the same or adjacent target cells results in association of the V_H and V_L domains of an antigen binding site for a radiolabelled compound and the formation of a functional antigen binding site for the radiolabelled compound. Thus, after administration of the radiolabelled compound, the radiolabelled compound binds to the functional antigen binding site formed by association of the V_H and V_L .

20

In any of the methods and uses described herein, the first and second antibodies can be administered simultaneously or sequentially, in either order.

Frequently in the art, methods of PRIT or radioimaging involve a clearing step. The clearing step comprises administering an agent between the administration of the antibody and the administration of the radiolabelled compound, wherein the agent increases the rate of removal of the antibody from blood and/or blocks binding of radiolabelled compound to the antibody.

In an embodiment of the methods and uses described herein, the method does not comprise a clearing step. That is, it does not comprise a step of administering a clearing agent or a blocking agent between the administration of the first and second antibodies and the administration of radiolabelled compound (i.e., after the administration of the antibodies but before administration of the radiolabelled compound). In another embodiment, no agent is administered between the administration of the first and second antibodies and the

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administration of radiolabelled compound, other than optionally a radiosensitizer, immunotherapeutic and/or a chemotherapeutic agent. In another embodiment, no agent is administered between the administration of the first and second antibodies and the administration of radiolabelled compound.

In some embodiments, the antibodies described herein may be administered as part of a combination therapy. For example, they may be administered in combination with one or more radiosensitizers, immunotherapeutics and/or chemotherapeutic agents: the radiosensitizer, immunotherapeutic or chemotherapeutic agent and the antibodies may be administered simultaneously or sequentially, in either order.

10 The methods of radioimaging and radioimmunotherapy described herein may optionally be combined as discussed further herein.

In a further aspect, the present invention provides a kit comprising:

- i) a first and a second antibody as described herein;
- ii) a radiolabelled compound which binds to the antigen binding site formed by association of the first and the second antibody.

Optionally the kit may exclude (i.e., does not comprise) a clearing agent or a blocking agent as described herein.

Optionally the kit may further comprise a radiosensitizer, immunotherapeutic or chemotherapeutic agent.

20 In some embodiments, the first and the second antibody may be present in the same pharmaceutical composition. In other embodiments, the first and second antibody may be present in separate pharmaceutical compositions. In some embodiments, the radiolabelled compound is present in a pharmaceutical composition separate from the antibodies.

In other embodiments, antibodies as described herein find use in a method of selective killing of a target cell, e.g., for cancer treatment.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 shows the schematic structure of a target antigen (TA)-DOTAM bispecific antibody (TA-DOTAM BsAb) belonging to the comparative examples, and exemplary TA-split-DOTAM-VH/VL antibodies.

30 Figure 2 is a schematic diagram showing the assembly of a split-VH/VL DOTAM binder on tumour cells. The TA-split-DOTAM-VH/VL antibodies will not significantly bind

²¹²Pb-DOTAM unless bound to tumour antigen (TA) on targeted cells, where the two domains of the DOTAM binder are assembled.

Figure 3 shows a schematic overview of an example of the Three-Step TA-PRIT concept, involving use of a clearing agent.

Figure 4 shows a schematic overview of an example of the Two-Step TA-PRIT concept, in which a clearing agent is not used.

Figure 5 shows binding of split antibodies to MKN45 cells to demonstrate CEA binding competence. Detection of antibodies is done using human IgG specific secondary antibodies

10 Figure 6 shows binding of split antibodies to MKN45 cells to demonstrate DOTAM binding competence. Detection of antibodies is done using Pb-DOTAM-FITC.

Figure 7A shows an exemplary protocol for two-step PRIT with a CEA-split-DOTAM-VH/VL, carried out in SCID mice carrying SC BxPC3 tumours (h = hours, d = days, w = weeks).

Figure 7B shows an exemplary protocol for a three-step PRIT control, carried out in SCID mice carrying SC BxPC3 tumours (h=hours, d=days, w=weeks).

Figure 8 shows the biodistribution of pretargeted ²¹²Pb-DOTAM in SCID mice carrying SC BxPC3 tumors, 6 hours after injection of ²¹²Pb -DOTAM, pretargeted either by CEA-split-DOTAM-VH alone, CEA-split-DOTAM-VL alone, or the two complementary
20 antibodies combined, or using standard three-step PRIT (%ID/g ± SD, n = 4).

Figure 9 shows CEA-Split-DOTAM-VH/VL pharmacokinetics after IV injection in SCID mice.

Figure 10 shows the experimental design of protocol 158, comprising CEA-PRIT in 2 (top) or 3 steps (bottom) in SCID mice carrying SC BxPC3 tumors. *CEA split DOTAM BsAb dose adjusted to compensate for hole/hole impurities in 2/4 constructs.

Figure 11 shows the biodistribution of pretargeted ²¹²Pb -DOTAM in SCID mice carrying SC BxPC3 tumors (6 h p.i.). The distribution is of ²¹²Pb in tumour-bearing SCID mice, 6 hours after injection of ²¹²Pb -DOTAM, pretargeted by CEA-DOTAM BsAb or bi-paratopic combinations of CEA-split-DOTAM antibodies. The radioactive content in organs
30 and tissues is expressed as average % ID/g ± SD (n = 4).

Figure 12 shows the experimental schedule of protocol 160, comprising one cycle of 3-step CEA-PRIT (top), 2-step CEA-PRIT (middle), or 1-step CEA-RIT in SCID mice carrying SC BxPC3 tumors. Biodistribution (BD) scouts were euthanized 24 hours after the

radioactive injection, whereas mice in the efficacy groups were maintained and monitored carefully until the termination criteria were reached.

Figure 13 shows biodistribution of pretargeted ^{212}Pb -DOTAM and ^{212}Pb -DOTAM-CEA-DOTAM in SCID mice carrying SC BxPC3 tumors (24 h p.i.). The distribution is of ^{212}Pb in tumor-bearing SCID mice 24 hours after injection of CEA-DOTAM-pretargeted ^{212}Pb -DOTAM or pre-incubated ^{212}Pb -DOTAM-CEA-DOTAM. The radioactive content in organs and tissues is expressed as average % ID/g $\pm$ SD (n = 3).

Figure 14 shows tumor growth averages with standard error for PRIT-treated groups and control (groups A-E) in the BxPC3 model (n=10). Curves were truncated at n<5. Dotted vertical lines indicate ^{212}Pb -DOTAM administration (20 μCi) for some or all groups, according to the study design.

Figure 15 shows individual tumor growth curves for PRIT-treated groups and control (groups A-E) in the BxPC3 model (n=10). Dotted vertical lines indicate administration of ^{212}Pb -labeled compounds (20 μCi).

Figure 16 shows average body weight loss in mice treated with CEA-PRIT and CEA-RIT (groups A-E, n=10) in the BxPC3 model. Curves were truncated at n<5. Dotted vertical lines indicate administration of ^{212}Pb -labeled compounds for some or all groups, according to the study design.

Figure 17 shows the experimental design of protocol 175, comprising two-step CEA-PRIT in SCID mice carrying SC BxPC3 tumors, with sacrifice and necropsy 24 hours after the ^{212}Pb -DOTAM injection. The CEA-split-DOTAM-VH-AST dose was adjusted to compensate for hole/hole impurities.

Figure 18 shows distribution of ^{212}Pb in tumor-bearing SCID mice 24 hours after injection of ^{212}Pb -DOTAM, pretargeted by CEA-split-DOTAM-VH/VL antibodies (protocol 175). The radioactive content in organs and tissues is expressed as average % ID/g $\pm$ SD (n = 4).

Figure 19 shows the experimental design of protocol 185, comprising two-step CEA-PRIT in SCID mice carrying SC BxPC3 tumors, with sacrifice and necropsy 6 hours after the ^{212}Pb -DOTAM injection. The CEA-split-DOTAM-VH-AST (CH1A1A) dose was adjusted to compensate for hole/hole impurities.

Figure 20 shows distribution of ^{212}Pb in tumor-bearing SCID mice 6 hours after injection of ^{212}Pb -DOTAM, pretargeted by CEA-split-DOTAM-VH/VL antibodies (protocol

185). The radioactive content in organs and tissues is expressed as average % ID/g $\pm$ SD (n = 5).

Figure 21 shows distribution of CEA-split-DOTAM-VH/VL pairs (VH and VL antibodies combined) in two selected SC BxPC3 tumors 7 days after injection. A and B show sections of a tumor from mouse A3, injected with CEA-split-DOTAM-VH/VL targeting T84.66, where A shows the CEA expression, and B shows the corresponding CEA-split-DOTAM-VH/VL distribution. C and D show tumor sections from mouse C5, injected with CEA-split-DOTAM-VH/VL targeting CH1A1A: C showing the CEA expression and D the corresponding CEA-split-DOTAM-VH/VL distribution.

10 Figure 22 shows the experimental design of protocol 189, comprising two-step CEA-PRIT in SCID mice carrying SC BxPC3 tumors, with sacrifice and necropsy 6 hours after the ^{212}Pb -DOTAM injection. The CEA-split-DOTAM-VH-AST (CH1A1A) dose was adjusted to compensate for hole/hole impurities.

Figure 23 shows distribution of ^{212}Pb in tumor-bearing SCID mice 6 hours after injection of ^{212}Pb -DOTAM, pretargeted by bi-paratopic pairs of CEA-split-DOTAM-VH/VL antibodies (T84.66 and CH1A1A), compared with the positive control (CH1A1A only). The radioactive content in organs and tissues is expressed as average % ID/g $\pm$ SD.

Figure 24 shows mean Fluorescence Intensity (MFI) as determined by FACS for SPLIT antibodies. Binding of Pb-DOTA-FITC determined by FACS can only be shown for a
20 co-incubation of both SPLIT antibodies with Pb-DOTA-FITC. Single SPLIT antibodies did not give rise to a significant signal.

Figure 25A-C shows exemplary formats of split antibodies.

Figure 26 shows results from example 8, experiment 1, assessing binding of individual TA-split-DOTAM-VH and TA-split-DOTAM-VL antibodies to biotinylated DOTAM captured on a chip.

Figure 27 shows results from example 8, experiment 2, assessing binding of DOTAM to individual TA-split-DOTAM-VH and TA-split-DOTAM-VL antibodies captured on a chip.

30 Figure 28 shows results from example 8, experiment 3, assessing binding of DOTAM to TA-split-DOTAM-VH/VL antibodies (antibody pairs), captured on a chip.

Figure 29 shows the schematic structure of further exemplary TA-split-DOTAM-VH/VL antibodies. Figure 29A shows a format which is monovalent for the target antigen. Figure 29B shows a format which is bivalent for the target antigen.

Figure 30 shows the experimental design of protocol 193, comprising a 2-step SPLIT PRIT regimen in SCID mice carrying SC BxPC3 tumors.

Figure 31 shows distribution of ^{212}Pb in tumor-bearing SCID mice 6 hours after injection of ^{212}Pb -DOTAM, pretargeted by N-terminal SPLITs (results from example 9).

DETAILED DESCRIPTION OF THE INVENTION

I. DEFINITIONS

10 An “acceptor human framework” for the purposes herein is a framework comprising the amino acid sequence of a light chain variable domain (VL) framework or a heavy chain variable domain (VH) framework derived from a human immunoglobulin framework or a human consensus framework, as defined below. An acceptor human framework “derived from” a human immunoglobulin framework or a human consensus framework may comprise the same amino acid sequence thereof, or it may contain amino acid sequence changes. In some aspects, the number of amino acid changes are 10 or less, 9 or less, 8 or less, 7 or less, 6 or less, 5 or less, 4 or less, 3 or less, or 2 or less. In some aspects, the VL acceptor human framework is identical in sequence to the VL human immunoglobulin framework sequence or human consensus framework sequence.

20 “Affinity” refers to the strength of the sum total of noncovalent interactions between a single binding site of a molecule (e.g., an antibody) and its binding partner (e.g., an antigen). Unless indicated otherwise, as used herein, “binding affinity” refers to intrinsic binding affinity which reflects a 1:1 interaction between members of a binding pair (e.g., antibody and antigen). The affinity of a molecule X for its partner Y can generally be represented by the dissociation constant (K_D). Affinity can be measured by common methods known in the art, including those described herein. Specific illustrative and exemplary methods for measuring binding affinity are described in the following.

An “affinity matured” antibody refers to an antibody with one or more alterations in one or more complementary determining regions (CDRs), compared to a parent antibody which does not possess such alterations, such alterations resulting in an improvement in the affinity of the antibody for antigen.

30 The term “an antibody that binds to an antigen expressed on the surface of a target cell” refers to an antibody that is capable of binding said antigen with sufficient affinity such that the antibody is useful as a diagnostic and/or therapeutic agent in targeting said antigen. In one aspect, the extent of binding of the antibody to an unrelated, non antigen protein is less

than about 10% of the binding of the antibody to the antigen as measured, e.g., by surface plasmon resonance (SPR). In certain aspects, an antibody that binds to an antigen expressed on the surface of a target cell has a dissociation constant (K_D) of $\leq 1\mu\text{M}$, $\leq 100\text{ nM}$, $\leq 10\text{ nM}$, $\leq 1\text{ nM}$, $\leq 0.1\text{ nM}$, $\leq 0.01\text{ nM}$, or $\leq 0.001\text{ nM}$ (e.g., 10^{-8} M or less, e.g., from 10^{-8} M to 10^{-13} M , e.g., from 10^{-9} M to 10^{-13} M). An antibody is said to “specifically bind” to an antigen expressed on the surface of a target cell when the antibody has a K_D of $1\mu\text{M}$ or less. In certain aspects, the antibody binds to an epitope of said antigen that is conserved among said antigen from different species.

The terms “an antigen binding site for an effector moiety” or “a functional antigen binding site for an effector moiety” refer to an antigen binding site comprising V_H and a V_L domain, capable of binding to the effector moiety with sufficient affinity such that the antibody is useful as a diagnostic and/or therapeutic agent to associate the effector moiety with the antibody. In one aspect, the extent of binding of the antigen binding site to an unrelated, non antigen -compound is less than about 10% of the binding of the antibody to the effector moiety as measured, e.g., by surface plasmon resonance (SPR). In certain aspects, an antigen binding site that binds to an effector moiety has a dissociation constant (K_D) of $\leq 1\mu\text{M}$, $\leq 100\text{ nM}$, $\leq 10\text{ nM}$, $\leq 1\text{ nM}$, $\leq 0.1\text{ nM}$, $\leq 0.01\text{ nM}$, or $\leq 0.001\text{ nM}$ (e.g., 10^{-8} M or less, e.g., from 10^{-8} M to 10^{-13} M , e.g., from 10^{-9} M to 10^{-13} M). It may be preferred that it has a K_D of 100 pM , 50 pM , 20 pM , 10 pM , 5 pM , 1 pM or less, e.g., 0.9 pM or less, 0.8 pM or less, 0.7 pM or less, 0.6 pM or less or 0.5 pM or less. For instance, the functional binding site may bind the effector moiety with a K_D of about 1 pM - 1 nM , e.g., about 1 - 10 pM , 1 - 100 pM , 5 - 50 pM , 100 - 500 pM or 500 pM - 1 nM . An antigen binding site is said to “specifically bind” to an effector moiety when the antigen binding site has a K_D of $1\mu\text{M}$ or less.

The term “antibody” herein is used in the broadest sense and encompasses various antibody structures, including but not limited to monoclonal antibodies, polyclonal antibodies, multispecific antibodies (e.g., bispecific antibodies), and antibody fragments so long as they exhibit the desired antigen-binding activity. As used herein, the term “antibody” also encompasses individual hemibodies, which comprise either a V_H domain or a V_L domain of a functional antigen binding site.

An “antibody fragment” refers to a molecule other than an intact antibody that comprises a portion of an intact antibody that binds the antigen to which the intact antibody binds. Examples of antibody fragments include but are not limited to F_v , Fab, cross-Fab, Fab', Fab'-SH, $F(ab')_2$; diabodies; linear antibodies; single-chain antibody molecules (e.g.,

scFv, and scFab); single domain antibodies (dAbs); and multispecific antibodies formed from antibody fragments. For a review of certain antibody fragments, see Holliger and Hudson, Nature Biotechnology 23:1126-1136 (2005). The term "Fab fragment" refers to a protein consisting of the VH and CH1 domain of the heavy chain and the VL and CL domain of the light chain of an immunoglobulin. "Fab' fragments" differ from Fab fragments by the addition of residues at the carboxy terminus of the CH1 domain including one or more cysteines from the antibody hinge region. For discussion of Fab and F(ab')₂ fragments comprising salvage receptor binding epitope residues and having increased *in vivo* half-life, see U.S. Patent No. 5,869,046.

10 As used herein, a reference to a "Fab fragment" is intended to include a cross-Fab fragment or a scFab as well as a conventional Fab fragment (i.e., one comprising a light chain comprising a VL domain and a CL domain, and a heavy chain fragment comprising a VH domain and a CH1 domain).

The term "cross-Fab fragment" or "xFab fragment" or "crossover Fab fragment" refers to a Fab fragment, wherein either the variable regions or the constant regions of the heavy and light chain are exchanged. A cross-Fab fragment comprises a polypeptide chain composed of the light chain variable region (VL) and the heavy chain constant region 1 (CH1), and a polypeptide chain composed of the heavy chain variable region (VH) and the light chain constant region (CL). For clarity, in a crossover Fab molecule wherein the
20 variable regions of the Fab light chain and the Fab heavy chain are exchanged, the peptide chain comprising the heavy chain constant region is referred to herein as the "heavy chain" of the crossover Fab molecule. Conversely, in a crossover Fab molecule wherein the constant regions of the Fab light chain and the Fab heavy chain are exchanged, the peptide chain comprising the heavy chain variable region is referred to herein as the "heavy chain" of the crossover Fab molecule.

As used herein, the term "single-chain" refers to a molecule comprising amino acid monomers linearly linked by peptide bonds. A single-chain Fab molecule is a Fab molecule wherein the Fab light chain and the Fab heavy chain are connected by a peptide linker to form a single peptide chain. In a particular such embodiment, the C-terminus of the Fab light
30 chain is connected to the N-terminus of the Fab heavy chain in the single-chain Fab molecule.

Asymmetrical Fab arms can also be engineered by introducing charged or non-charged amino acid mutations into domain interfaces to direct correct Fab pairing. See e.g., WO 2016/172485.

A “single-chain variable fragment” or “scFv” is a fusion protein of the variable domains of the heavy (VH) and light chains (VL) of an antibody, connected by a peptide linker. In particular, the linker is a short polypeptide of 10 to 25 amino acids and is usually rich in glycine for flexibility, as well as serine or threonine for solubility, and can either connect the N-terminus of the VH with the C-terminus of the VL, or *vice versa*. This protein retains the specificity of the original antibody, despite removal of the constant regions and the introduction of the linker. For a review of scFv fragments, see, e.g., Plückthun, in *The Pharmacology of Monoclonal Antibodies*, vol. 113, Rosenberg and Moore eds., (Springer-Verlag, New York), pp. 269-315 (1994); see also WO 93/16185; and U.S. Patent Nos. 5,571,894 and 5,587,458.

The term “blocking agent” refers to an agent which blocks the binding of an effector molecule, in particular the radiolabelled compound, to a functional binding site for that effector molecule. Generally said blocking agent binds to the functional binding site for the effector molecule, e.g., specifically binds to the said functional binding site.

The term “clearing agent” refers to an agent which increases the rate of clearance of an antibody from the circulation of the subject. Generally the clearing agent binds to the antibody, e.g., specifically binds to the antibody.

The term “clearing step” or “clearing phase” as used herein encompasses the use of either a blocking agent or a clearing agent. Some agents can function as both a clearing and as a blocking agent.

The term “epitope” denotes the site on an antigen, either proteinaceous or non-proteinaceous, to which an antibody binds. Epitopes can be formed both from contiguous amino acid stretches (linear epitope) or comprise non-contiguous amino acids (conformational epitope), e.g., coming in spatial proximity due to the folding of the antigen, i.e. by the tertiary folding of a proteinaceous antigen. Linear epitopes are typically still bound by an antibody after exposure of the proteinaceous antigen to denaturing agents, whereas conformational epitopes are typically destroyed upon treatment with denaturing agents. An epitope comprises at least 3, at least 4, at least 5, at least 6, at least 7, or 8-10 amino acids in a unique spatial conformation.

Screening for antibodies binding to a particular epitope (i.e., those binding to the same epitope) can be done using methods routine in the art such as, e.g., without limitation, alanine scanning, peptide blots (see *Meth. Mol. Biol.* 248 (2004) 443-463), peptide cleavage analysis, epitope excision, epitope extraction, chemical modification of antigens (see *Prot. Sci.* 9

(2000) 487-496), and cross-blocking (see “Antibodies”, Harlow and Lane (Cold Spring Harbor Press, Cold Spring Harb., NY).

Antigen Structure-based Antibody Profiling (ASAP), also known as Modification-Assisted Profiling (MAP), allows to bin a multitude of monoclonal antibodies specifically binding to an antigen based on the binding profile of each of the antibodies from the multitude to chemically or enzymatically modified antigen surfaces (see, e.g., US 2004/0101920). The antibodies in each bin bind to the same epitope which may be a unique epitope either distinctly different from or partially overlapping with epitope represented by another bin.

10 Also competitive binding can be used to easily determine whether an antibody binds to the same epitope as, or competes for binding with, a reference antibody. For example, an “antibody that binds to the same epitope” as a reference antibody refers to an antibody that blocks binding of the reference antibody to its antigen in a competition assay by 50% or more, and conversely, the reference antibody blocks binding of the antibody to its antigen in a competition assay by 50% or more. Also for example, to determine if an antibody binds to the same epitope as a reference antibody, the reference antibody is allowed to bind to the antigen under saturating conditions. After removal of the excess of the reference antibody, the ability of an antibody in question to bind to the antigen is assessed. If the antibody in question is able to bind to the antigen after saturation binding of the reference antibody, it can
20 be concluded that the antibody in question binds to a different epitope than the reference antibody. But, if the antibody in question is not able to bind to the antigen after saturation binding of the reference antibody, then the antibody in question may bind to the same epitope as the epitope bound by the reference antibody. To confirm whether the antibody in question binds to the same epitope or is just hampered from binding by steric reasons routine experimentation can be used (e.g., peptide mutation and binding analyses using ELISA, RIA, surface plasmon resonance, flow cytometry or any other quantitative or qualitative antibody-binding assay available in the art). This assay should be carried out in two set-ups, i.e. with both of the antibodies being the saturating antibody. If, in both set-ups, only the first (saturating) antibody is capable of binding to the antigen, then it can be concluded that the
30 antibody in question and the reference antibody compete for binding to the antigen.

In some aspects, two antibodies are deemed to bind to the same or an overlapping epitope if a 1-, 5-, 10-, 20- or 100-fold excess of one antibody inhibits binding of the other by

at least 50%, at least 75%, at least 90% or even 99% or more as measured in a competitive binding assay (see, e.g., Junghans et al., Cancer Res. 50 (1990) 1495-1502).

In some aspects, two antibodies are deemed to bind to the same epitope if essentially all amino acid mutations in the antigen that reduce or eliminate binding of one antibody also reduce or eliminate binding of the other. Two antibodies are deemed to have “overlapping epitopes” if only a subset of the amino acid mutations that reduce or eliminate binding of one antibody reduce or eliminate binding of the other.

10 The term “chimeric” antibody refers to an antibody in which a portion of the heavy and/or light chain is derived from a particular source or species, while the remainder of the heavy and/or light chain is derived from a different source or species.

The “class” of an antibody refers to the type of constant domain or constant region possessed by its heavy chain. There are five major classes of antibodies: IgA, IgD, IgE, IgG, and IgM, and several of these may be further divided into subclasses (isotypes), e.g., IgG₁, IgG₂, IgG₃, IgG₄, IgA₁, and IgA₂. In certain aspects, the antibody is of the IgG₁ isotype. In certain aspects, the antibody is of the IgG₁ isotype with the P329G, L234A and L235A mutation to reduce Fc-region effector function. In other aspects, the antibody is of the IgG₂ isotype. In certain aspects, the antibody is of the IgG₄ isotype with the S228P mutation in the hinge region to improve stability of IgG₄ antibody. The heavy chain constant domains that correspond to the different classes of immunoglobulins are called α , δ , ϵ , γ , and μ ,
20 respectively. The light chain of an antibody may be assigned to one of two types, called kappa (κ) and lambda (λ), based on the amino acid sequence of its constant domain.

“Fc effector functions” refer to those biological activities attributable to the Fc region of an antibody, which vary with the antibody isotype. Examples of antibody effector functions include: C1q binding and complement dependent cytotoxicity (CDC); Fc receptor binding; antibody-dependent cell-mediated cytotoxicity (ADCC); phagocytosis; down regulation of cell surface receptors (e.g., B cell receptor); and B cell activation.

An “effective amount” of an agent, e.g., a pharmaceutical composition, refers to an amount effective, at dosages and for periods of time necessary, to achieve the desired therapeutic or prophylactic result.

30 The term “tandem Fab” refers to an antibody comprising two Fab fragments connected via a peptide linker/tether. In some embodiments, a tandem Fab may comprise one Fab fragment and one cross-Fab fragment, connected by a peptide linker/tether.

The term "Fc region" herein is used to define a C-terminal region of an immunoglobulin heavy chain that contains at least a portion of the constant region. The term "Fc domain" herein is used to define a C-terminal region of an immunoglobulin that contains the constant regions of two heavy chains, excluding the first constant region. Thus, Fc domain refers to the last two constant region immunoglobulin domains of IgA, IgD, and IgG, and the last three constant region immunoglobulin domains of IgE and IgM. The term includes native sequence Fc regions and variant Fc regions. In one aspect, a human IgG heavy chain Fc region extends from Cys226, or from Pro230, to the carboxyl-terminus of the heavy chain. However, antibodies produced by host cells may undergo post-translational cleavage of one or more, particularly one or two, amino acids from the C-terminus of the heavy chain. Therefore an antibody produced by a host cell by expression of a specific nucleic acid molecule encoding a full-length heavy chain may include the full-length heavy chain, or it may include a cleaved variant of the full-length heavy chain. This may be the case where the final two C-terminal amino acids of the heavy chain are glycine (G446) and lysine (K447, numbering according to EU index). Therefore, the C-terminal lysine (Lys447), or the C-terminal glycine (Gly446) and lysine (Lys447), of the Fc region may or may not be present. In one aspect, a heavy chain including an Fc region as specified herein, comprised in an antibody according to the invention, comprises an additional C-terminal glycine-lysine dipeptide (G446 and K447, numbering according to EU index). In one aspect, a heavy chain including an Fc region as specified herein, comprised in an antibody according to the invention, comprises an additional C-terminal glycine residue (G446, numbering according to EU index). Unless otherwise specified herein, numbering of amino acid residues in the Fc region or constant region is according to the EU numbering system, also called the EU index, as described in Kabat et al., *Sequences of Proteins of Immunological Interest*, 5th Ed. Public Health Service, National Institutes of Health, Bethesda, MD, 1991. A "subunit" of an Fc domain as used herein refers to one of the two polypeptides forming the dimeric Fc domain, i.e. a polypeptide comprising C-terminal constant regions of an immunoglobulin heavy chain, capable of stable association with the other of the two polypeptides forming the dimeric Fc domain. For example, a subunit of an IgG Fc domain comprises an IgG CH2 and an IgG CH3 constant domain.

"Framework" or "FR" refers to variable domain residues other than complementary determining regions (CDRs). The FR of a variable domain generally consists of four FR domains: FR1, FR2, FR3, and FR4. Accordingly, the CDR and FR sequences generally

appear in the following sequence in VH (or VL): FR1-CDR-H1(CDR-L1)-FR2- CDR-H2(CDR-L2)-FR3- CDR-H3(CDR-L3)-FR4.

The terms “full length antibody”, “intact antibody”, and “whole antibody” are used herein interchangeably to refer to an antibody having a structure substantially similar to a native antibody structure or having heavy chains that contain an Fc region as defined herein.

By "fused" is meant that the components are linked by peptide bonds, either directly or via one or more peptide linkers.

The terms “host cell”, “host cell line”, and “host cell culture” are used interchangeably and refer to cells into which exogenous nucleic acid has been introduced, including the progeny of such cells. Host cells include “transformants” and “transformed cells”, which include the primary transformed cell and progeny derived therefrom without regard to the number of passages. Progeny may not be completely identical in nucleic acid content to a parent cell, but may contain mutations. Mutant progeny that have the same function or biological activity as screened or selected for in the originally transformed cell are included herein.

A “human antibody” is one which possesses an amino acid sequence which corresponds to that of an antibody produced by a human or a human cell or derived from a non-human source that utilizes human antibody repertoires or other human antibody-encoding sequences. This definition of a human antibody specifically excludes a humanized antibody comprising non-human antigen-binding residues.

A “human consensus framework” is a framework which represents the most commonly occurring amino acid residues in a selection of human immunoglobulin VL or VH framework sequences. Generally, the selection of human immunoglobulin VL or VH sequences is from a subgroup of variable domain sequences. Generally, the subgroup of sequences is a subgroup as in Kabat et al., *Sequences of Proteins of Immunological Interest*, Fifth Edition, NIH Publication 91-3242, Bethesda MD (1991), vols. 1-3. In one aspect, for the VL, the subgroup is subgroup kappa I as in Kabat et al., *supra*. In one aspect, for the VH, the subgroup is subgroup III as in Kabat et al., *supra*.

A “humanized” antibody refers to a chimeric antibody comprising amino acid residues from non-human CDRs and amino acid residues from human FRs. In certain aspects, a humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the CDRs correspond to those of a non-human antibody, and all or substantially all of the FRs correspond to those of a human antibody. A

humanized antibody optionally may comprise at least a portion of an antibody constant region derived from a human antibody. A “humanized form” of an antibody, e.g., a non-human antibody, refers to an antibody that has undergone humanization.

The term “hypervariable region” or “HVR” as used herein refers to each of the regions of an antibody variable domain which are hypervariable in sequence and which determine antigen binding specificity, for example “complementarity determining regions” (“CDRs”).

Generally, antibodies comprise six CDRs: three in the VH (CDR-H1, CDR-H2, CDR-H3), and three in the VL (CDR-L1, CDR-L2, CDR-L3). Exemplary CDRs herein include:

10 (a) hypervariable loops occurring at amino acid residues 26-32 (L1), 50-52 (L2), 91-96 (L3), 26-32 (H1), 53-55 (H2), and 96-101 (H3) (Chothia and Lesk, *J. Mol. Biol.* 196:901-917 (1987));

(b) CDRs occurring at amino acid residues 24-34 (L1), 50-56 (L2), 89-97 (L3), 31-35b (H1), 50-65 (H2), and 95-102 (H3) (Kabat et al., *Sequences of Proteins of Immunological Interest*, 5th Ed. Public Health Service, National Institutes of Health, Bethesda, MD (1991)); and

(c) antigen contacts occurring at amino acid residues 27c-36 (L1), 46-55 (L2), 89-96 (L3), 30-35b (H1), 47-58 (H2), and 93-101 (H3) (MacCallum et al. *J. Mol. Biol.* 262: 732-745 (1996)).

20 Unless otherwise indicated, the CDRs are determined according to Kabat et al., *supra*. One of skill in the art will understand that the CDR designations can also be determined according to Chothia, *supra*, McCallum, *supra*, or any other scientifically accepted nomenclature system. Instead of the above, the sequence of CDR-H1 as described herein may extend from Kabat26 to Kabat35, e.g., for the Pb-DOTAM binding variable domain.

In one aspect, CDR residues comprise those identified in the sequence tables or elsewhere in the specification.

Unless otherwise indicated, HVR/CDR residues and other residues in the variable domain (e.g., FR residues) are numbered herein according to Kabat et al., *supra*.

An “immunoconjugate” is an antibody conjugated to one or more heterologous molecule(s), including but not limited to a cytotoxic agent.

30 An “individual” or “subject” is a mammal. Mammals include, but are not limited to, domesticated animals (e.g., cows, sheep, cats, dogs, and horses), primates (e.g., humans and non-human primates such as monkeys), rabbits, and rodents (e.g., mice and rats). In certain aspects, the individual or subject is a human.

Molecules as described herein may be “isolated”. An “isolated” antibody is one which has been separated from a component of its natural environment. In some aspects, an antibody is purified to greater than 95% or 99% purity as determined by, for example, electrophoretic (e.g., SDS-PAGE, isoelectric focusing (IEF), capillary electrophoresis) or chromatographic (e.g., ion exchange or reverse phase HPLC) methods. For a review of methods for assessment of antibody purity, see, e.g., Flatman et al., *J. Chromatogr. B* 848:79-87 (2007).

The term “nucleic acid molecule” or “polynucleotide” includes any compound and/or substance that comprises a polymer of nucleotides. Each nucleotide is composed of a base, specifically a purine- or pyrimidine base (i.e. cytosine (C), guanine (G), adenine (A), thymine (T) or uracil (U)), a sugar (i.e. deoxyribose or ribose), and a phosphate group. Often, the nucleic acid molecule is described by the sequence of bases, whereby said bases represent the primary structure (linear structure) of a nucleic acid molecule. The sequence of bases is typically represented from 5' to 3'. Herein, the term nucleic acid molecule encompasses deoxyribonucleic acid (DNA) including e.g., complementary DNA (cDNA) and genomic DNA, ribonucleic acid (RNA), in particular messenger RNA (mRNA), synthetic forms of DNA or RNA, and mixed polymers comprising two or more of these molecules. The nucleic acid molecule may be linear or circular. In addition, the term nucleic acid molecule includes both, sense and antisense strands, as well as single stranded and double stranded forms.

Moreover, the herein described nucleic acid molecule can contain naturally occurring or non-naturally occurring nucleotides. Examples of non-naturally occurring nucleotides include modified nucleotide bases with derivatized sugars or phosphate backbone linkages or chemically modified residues. Nucleic acid molecules also encompass DNA and RNA molecules which are suitable as a vector for direct expression of an antibody of the invention *in vitro* and/or *in vivo*, e.g., in a host or patient. Such DNA (e.g., cDNA) or RNA (e.g., mRNA) vectors, can be unmodified or modified. For example, mRNA can be chemically modified to enhance the stability of the RNA vector and/or expression of the encoded molecule so that mRNA can be injected into a subject to generate the antibody *in vivo* (see e.g., Stadler et al, Nature Medicine 2017, published online 12 June 2017, doi:10.1038/nm.4356 or EP 2 101 823 B1).

An “isolated” nucleic acid refers to a nucleic acid molecule that has been separated from a component of its natural environment. An isolated nucleic acid includes a nucleic acid molecule contained in cells that ordinarily contain the nucleic acid molecule, but the

nucleic acid molecule is present extrachromosomally or at a chromosomal location that is different from its natural chromosomal location.

“Isolated nucleic acid encoding an antibody” refers to one or more nucleic acid molecules encoding antibody heavy and light chains (or fragments thereof), including such nucleic acid molecule(s) in a single vector or separate vectors, and such nucleic acid molecule(s) present at one or more locations in a host cell.

The term “monoclonal antibody” as used herein refers to an antibody obtained from a population of substantially homogeneous antibodies, i.e., the individual antibodies comprising the population are identical and/or bind the same epitope, except for possible variant antibodies, e.g., containing naturally occurring mutations or arising during production of a monoclonal antibody preparation, such variants generally being present in minor amounts. In contrast to polyclonal antibody preparations, which typically include different antibodies directed against different determinants (epitopes), each monoclonal antibody of a monoclonal antibody preparation is directed against a single determinant on an antigen. Thus, the modifier “monoclonal” indicates the character of the antibody as being obtained from a substantially homogeneous population of antibodies, and is not to be construed as requiring production of the antibody by any particular method. For example, the monoclonal antibodies in accordance with the present invention may be made by a variety of techniques, including but not limited to the hybridoma method, recombinant DNA methods, phage-
display methods, and methods utilizing transgenic animals containing all or part of the human immunoglobulin loci, such methods and other exemplary methods for making monoclonal antibodies being described herein.

A “naked antibody” refers to an antibody that is not conjugated to a heterologous moiety (e.g., a cytotoxic moiety) or radiolabel. The naked antibody may be present in a pharmaceutical composition.

“Native antibodies” refer to naturally occurring immunoglobulin molecules with varying structures. For example, native IgG antibodies are heterotetrameric glycoproteins of about 150,000 daltons, composed of two identical light chains and two identical heavy chains that are disulfide-bonded. From N- to C-terminus, each heavy chain has a variable domain (VH), also called a variable heavy domain or a heavy chain variable region, followed by three constant heavy domains (CH1, CH2, and CH3). Similarly, from N- to C-terminus, each light chain has a variable domain (VL), also called a variable light domain or a light chain variable region, followed by a constant light (CL) domain.

The term “package insert” is used to refer to instructions customarily included in commercial packages of therapeutic products, that contain information about the indications, usage, dosage, administration, combination therapy, contraindications and/or warnings concerning the use of such therapeutic products.

“Percent (%) amino acid sequence identity” with respect to a reference polypeptide sequence is defined as the percentage of amino acid residues in a candidate sequence that are identical with the amino acid residues in the reference polypeptide sequence, after aligning the sequences and introducing gaps, if necessary, to achieve the maximum percent sequence identity, and not considering any conservative substitutions as part of the sequence identity for the purposes of the alignment. Alignment for purposes of determining percent amino acid sequence identity can be achieved in various ways that are within the skill in the art, for instance, using publicly available computer software such as BLAST, BLAST-2, Clustal W, Megalign (DNASTAR) software or the FASTA program package. Those skilled in the art can determine appropriate parameters for aligning sequences, including any algorithms needed to achieve maximal alignment over the full length of the sequences being compared. Alternatively, the percent identity values can be generated using the sequence comparison computer program ALIGN-2. The ALIGN-2 sequence comparison computer program was authored by Genentech, Inc., and the source code has been filed with user documentation in the U.S. Copyright Office, Washington D.C., 20559, where it is registered under U.S. Copyright Registration No. TXU510087 and is described in WO 2001/007611.

Unless otherwise indicated, for purposes herein, percent amino acid sequence identity values are generated using the ggsearch program of the FASTA package version 36.3.8c or later with a BLOSUM50 comparison matrix. The FASTA program package was authored by W. R. Pearson and D. J. Lipman (1988), “Improved Tools for Biological Sequence Analysis”, PNAS 85:2444-2448; W. R. Pearson (1996) “Effective protein sequence comparison” Meth. Enzymol. 266:227- 258; and Pearson et. al. (1997) Genomics 46:24-36 and is publicly available from www.fasta.bioch.virginia.edu/fasta_www2/fasta_down.shtml or www.ebi.ac.uk/Tools/sss/fasta. Alternatively, a public server accessible at fasta.bioch.virginia.edu/fasta_www2/index.cgi can be used to compare the sequences, using the ggsearch (global protein:protein) program and default options (BLOSUM50; open: -10; ext: -2; Ktup = 2) to ensure a global, rather than local, alignment is performed. Percent amino acid identity is given in the output alignment header.

The term “pharmaceutical composition” or “pharmaceutical formulation” refers to a preparation which is in such form as to permit the biological activity of an active ingredient contained therein to be effective, and which contains no additional components which are unacceptably toxic to a subject to which the pharmaceutical composition would be administered.

A “pharmaceutically acceptable carrier” refers to an ingredient in a pharmaceutical composition or formulation, other than an active ingredient, which is nontoxic to a subject. A pharmaceutically acceptable carrier includes, but is not limited to, a buffer, excipient, stabilizer, or preservative.

10 A reference to a target antigen as used herein, refers to any native target antigen from any vertebrate source, including mammals such as primates (e.g., humans) and rodents (e.g., mice and rats), unless otherwise indicated. The term encompasses “full-length”, unprocessed target antigen as well as any form of target antigen that results from processing in the cell. The term also encompasses naturally occurring variants of the target antigen, e.g., splice variants or allelic variants. For instance, the target antigen CEA may have the amino acid sequence of human CEA, in particular Carcinoembryonic antigen-related cell adhesion molecule 5 (CEACAM5), which is shown in UniProt (www.uniprot.org) accession no. P06731 (version 119), or NCBI (www.ncbi.nlm.nih.gov/) RefSeq NP_004354.2. Another example of a target antigen is Fibroblast Activation Protein (FAP). The amino acid sequence
20 of human FAP is shown in UniProt (www.uniprot.org) accession no. Q12884 (version 149), or NCBI (www.ncbi.nlm.nih.gov/) RefSeq NP_004451.2. Another example of a target antigen is GPRC5D (see UniProt no. Q9NZD1 (version 115); NCBI RefSeq no. NP_061124.1 for the human sequence).

The terms “split antibody”, “split antibodies”, “single domain split antibodies” or “SPLIT PRIT” as referred to herein mean that the VH and VL domain which together form an antigen binding site capable of binding to the effector moiety are split between two antibodies, and not present as part of the same antibody (before assembly in vivo). “CEA-targeted SPLIT PRIT” refers to a split antibody targeting CEA. The term “SPLIT PRIT” may also be used interchangeably with the term “TA-split-DOTAM-VH/VL” (e.g., where “TA” or
30 target antigen is CEA, FAP or GPRC5D). The term “CEA-targeted SPLIT PRIT” may be used interchangeably with the term “CEA-split-DOTAM-VH/VL”.

As used herein, “treatment” (and grammatical variations thereof such as “treat” or “treating”) refers to clinical intervention in an attempt to alter the natural course of a disease

in the individual being treated, and can be performed either for prophylaxis or during the course of clinical pathology. Desirable effects of treatment include, but are not limited to, preventing occurrence or recurrence of disease, alleviation of symptoms, diminishment of any direct or indirect pathological consequences of the disease, preventing metastasis, decreasing the rate of disease progression, amelioration or palliation of the disease state, and remission or improved prognosis. In some aspects, antibodies of the invention are used to delay development of a disease or to slow the progression of a disease.

10 The term “variable region” or “variable domain” refers to the domain of an antibody heavy or light chain that is involved in binding the antibody to antigen. The variable domains of the heavy chain and light chain (VH and VL, respectively) of a native antibody generally have similar structures, with each domain comprising four conserved framework regions (FRs) and three complementary determining regions (CDRs). (See, e.g., Kindt et al. *Kuby Immunology*, 6th ed., W.H. Freeman and Co., page 91 (2007).) A single VH or VL domain may be sufficient to confer antigen-binding specificity. Furthermore, antibodies that bind a particular antigen may be isolated using a VH or VL domain from an antibody that binds the antigen to screen a library of complementary VL or VH domains, respectively. See, e.g., Portolano et al., *J. Immunol.* 150:880-887 (1993); Clarkson et al., *Nature* 352:624-628 (1991).

20 The term “vector”, as used herein, refers to a nucleic acid molecule capable of propagating another nucleic acid to which it is linked. The term includes the vector as a self-replicating nucleic acid structure as well as the vector incorporated into the genome of a host cell into which it has been introduced. Certain vectors are capable of directing the expression of nucleic acids to which they are operatively linked. Such vectors are referred to herein as “expression vectors”.

The terms “Pb” or “lead” as used herein include ions thereof, e.g., Pb(II). References to other metals also include ions thereof. Thus, the skilled reader understands that, for example, the terms lead, Pb, ²¹²Pb or ²⁰³Pb are intended to encompass ionic forms of the element, in particular, Pb(II).

II. COMPOSITIONS AND METHODS

30 In one aspect, the invention is based, in part, on a set of antibodies comprising a first and a second antibody, wherein each antibody can bind to an antigen on a target cell, but wherein a functional antigen binding site for an effector moiety is formed only when the first and second antibodies are associated with each other. In one embodiment, sets of antibodies

according to the invention are useful in cell killing/cancer treatment. In another embodiment, sets of antibodies of the invention are useful, e.g., for methods of pre-targeted immunotherapy and/or for pre-targeted imaging. In preferred aspects such methods eliminate a step of administering a clearing agent or blocking agent.

The split format has advantages for reducing off-target effects. In the context of PRIT, it avoids the need for a clearing agent, as demonstrated herein. Moreover, the particular format as set out herein avoids a free C-terminus of the VH domain of the split antigen-binding site, and thus reduces the potential for an anti-drug antibody response involving pre-existing human anti-VH (HAVH) autoantibody. Further, and without wishing to be bound by theory, the inventors believe that the format as set out herein assists in protecting the hydrophobic interfaces of the DOTAM VH/VL, and thus assists with stability.

A. Antibody formats

As described above, the present invention provides novel formats for bi-specific antibodies in which the VH and VL domain for the effector antigen are split between two parts, and methods of using the same.

In particular, the present invention relates to a set of antibodies comprising

i) a first antibody comprising:

a) an antigen binding moiety, wherein the antigen binding moiety binds to an antigen expressed on the surface of a target cell;

b) a polypeptide comprising or consisting of an antibody heavy chain variable domain (VH) of an antigen binding site for an effector moiety; and

c) an Fc domain comprising two subunits,

wherein the polypeptide of (b) is fused by its N-terminus to the C-terminus of the antigen binding moiety of (a) and by its C-terminus to the N-terminus of one of the subunits of the Fc domain of (c);

and wherein the first antibody does not comprise a VL domain of an antigen binding site for the effector moiety; and

ii) a second antibody comprising:

d) an antigen binding moiety, wherein the antigen binding moiety binds to an antigen expressed on the surface of a target cell;

e) a polypeptide comprising or consisting of an antibody light chain variable domain (VL) of an antigen binding site for the effector moiety; and

f) an Fc domain comprising two subunits,

wherein the polypeptide of (e) is fused by its N-terminus to the C-terminus of the antigen binding moiety and by its C-terminus to the N-terminus of one of the subunits of the Fc domain of (f);

and wherein the second antibody does not comprise a VH domain of an antigen binding site for the effector moiety;

wherein said VH domain of the first antibody and said VL domain of the second antibody are together capable of forming a functional antigen binding site for the effector moiety.

The fusion may be direct or indirect, e.g., via a peptide linker.

10 In some embodiments, the antigen binding moiety of (a) and/or (d) may be a Fab.

It may be preferred that the polypeptide of (b) is fused by its N-terminus to the C-terminus of the heavy chain of the Fab fragment of (a). In some embodiments, the Fab fragment of (a) comprises a light chain comprising a VL domain and a CL domain and a heavy chain fragment comprising a VH domain and a CH1 domain and the polypeptide of (b) is fused by its N-terminus to the C-terminus of the CH1 domain.

It may similarly be preferred that the polypeptide of (e) is fused by its N-terminus to the C-terminus of the heavy chain of the Fab fragment of (d). In some embodiments, the Fab fragment of (d) comprises a light chain comprising a VL domain and a CL domain and a heavy chain fragment comprising a VH domain and a CH1 domain and the polypeptide of (e)
20 is fused by its N-terminus to the C-terminus of the CH1 domain.

It may be preferred that the polypeptides (b) and (e) do not comprise a constant region (e.g., CH1 or CL). In some embodiments, it may be preferred that the polypeptide of (b) consists of an antibody heavy chain variable domain (VH) of an antigen binding site for an effector moiety and/or that the polypeptide of (e) consists of an antibody light chain variable domain (VL) of an antigen binding site for the effector moiety. This may assist correct assembly of the light chains forming part of the Fab fragments of (a) and (d) and/or reduce the tendency of the two parts to form a binding competent moiety in the circulation.

It may be preferred that the association of the first and second antibody results in the formation of only one functional antigen binding for the effector moiety – i.e., the two
30 associated antibodies provide monovalent binding for the effector moiety. Thus, the first antibody may comprise only one VH domain of an antigen binding site for the effector moiety, and the second antibody may comprise only one VL domain of an antigen binding

site for the effector moiety, so that together they form only one complete functional binding site for the effector moiety.

In some embodiments, the first and/or second antibodies further comprise another antigen binding moiety binding to a target antigen, e.g., another antibody fragment such as another Fab fragment binding to a target antigen. Thus, in some embodiments the first and/or second antibodies (generally both) each comprise two antigen binding moieties capable of binding to a target antigen. The two antigen binding moieties are preferably capable of binding to the same target antigen. Optionally, the first and second antibodies each comprise not more than two antigen binding moieties capable of binding to a target antigen. In other
 10 embodiments, the first and second antibodies may each comprise more than two antigen binding moieties capable of binding to a target antigen.

In one embodiment, this further antigen binding moiety, e.g., Fab fragment, is fused by its C-terminus to the N-terminus of the other subunit of the Fc domain. (In the case of a Fab or other moiety composed of more than one chain, it may be fused by the C-terminus of one of its chains, e.g., its heavy chain, to the N-terminus of the other subunit of the Fc domain). Thus, in one embodiment, the first and/or second antibodies may be a two-armed antibody, wherein each arm bears a binding site for a target antigen.

Thus, in one embodiment, the present invention relates to a set of antibodies comprising:

- 20 i) a first antibody comprising:
- a) a first Fab fragment, wherein the Fab fragment binds to an antigen expressed on the surface of a target cell;
- b) a polypeptide comprising or consisting of an antibody heavy chain variable domain (VH) of an antigen binding site for an effector moiety; and
- c) an Fc domain comprising a first and a second subunit,
- wherein the polypeptide of (b) is fused by its N-terminus to the C-terminus of one of the chains of the Fab fragment of (a) and by its C-terminus to the N-terminus of the first subunit of the Fc domain of (c);
- and further comprising a second Fab fragment which binds to an antigen expressed on
 30 the surface of a target cell, wherein the second Fab is fused by the C-terminus of one of its chains to the N-terminus of the second subunit of the Fc domain of (c);
- wherein the first antibody does not comprise a VL domain of an antigen binding site for the effector moiety; and

ii) a second antibody comprising:

d) a first Fab fragment, wherein the Fab fragment binds to an antigen expressed on the surface of a target cell;

e) a polypeptide comprising or consisting of an antibody light chain variable domain (VL) of an antigen binding site for the effector moiety; and

f) an Fc domain comprising a first and a second subunit,

wherein the polypeptide of (e) is fused by its N-terminus to the C-terminus of one of the chains of the Fab fragment of (d) and by its C-terminus to the N-terminus of the first subunit of the Fc domain of (f);

10 and further comprising a second Fab fragment which binds to an antigen expressed on the surface of a target cell, wherein the second Fab is fused by the C-terminus of one of its chains to the N-terminus of the second subunit of the Fc domain of (f);

wherein the second antibody does not comprise a VH domain of an antigen binding site for the effector moiety; and

wherein said VH domain of the first antibody and said VL domain of the second antibody are together capable of forming a functional antigen binding site for the effector moiety.

It may be preferred that second Fab fragment of the first antibody is fused by the C-terminus of its heavy chain to the second subunit of the Fc domain. It may be preferred that
 20 the second Fab fragment of the first antibody comprises a light chain comprising a VL domain and a CL domain and a heavy chain fragment comprising a VH domain and a CH1 domain and that the second Fab fragment is fused by the C-terminus of its CH1 domain to the second subunit of the Fc domain. It may similarly be preferred that the second Fab fragment of the second antibody is fused by the C-terminus of its heavy chain to the second subunit of the Fc domain. It may be preferred that the second Fab fragment of the second antibody comprises a light chain comprising a VL domain and a CL domain and a heavy chain fragment comprising a VH domain and a CH1 domain and that the second Fab fragment is fused by the C-terminus of its CH1 domain to the second subunit of the Fc domain.

It may be preferred that the first and second antigen binding moiety (e.g., Fab
 30 fragment) of the first antibody bind to the same target antigen as each other, i.e., the first antibody is bivalent for the target antigen. It may similarly be preferred that the first and second antigen binding moiety (e.g., Fab fragment) of the second antibody bind to the same

target antigen as each other, i.e., the second antibody is bivalent for a target antigen. In some embodiments the first and second antibody also bind to the same target antigen as each other.

In some embodiments the first and/or second antibody are multivalent (e.g., bivalent) and monospecific for an epitope of the target antigen. Thus, in some embodiments, the first and second antigen binding moiety (e.g., Fab fragment) of the first antibody bind to the same epitope of the target antigen as each other; and/or the first and second antigen binding moiety (e.g., Fab fragment) of the second antibody bind to the same epitope on the target antigen as each other. Preferably the target antigen bound by the first and second antibody is the same. In some embodiments, the first and second antibody also bind to the same epitope in that target antibody as each other. Thus, the variable domain sequences (VH and VL) of the first and second Fab of the first antibody, or the first and second Fab of the second antibody, or all four Fabs, may in some embodiments be the same.

In other embodiments, the first antibody and second antigen bind to the same target antigen but each bind to a different epitope on that target antigen – e.g., the first and second Fab fragment of the first antibody bind to epitope A of the target antigen and the first and second Fab fragment of the second antibody bind to epitope B of that target antigen.

In some embodiments, the first antibody may comprise the following peptides:

- i) a first heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VH domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);
- ii) a Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (i) to form a binding site for a target antigen;
- iii) a second heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; and an Fc subunit (e.g., CH2-CH3); and
- iv) a further Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (iii) to form a binding site for a target antigen.

Optionally the Fab heavy chain in (i) has the same sequence as the Fab heavy chain in (iii) and the Fab light chains of (ii) and (iv) have the same sequence as each other.

The second antibody may comprise the following peptides:

- v) a first heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VL domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);

vi) a Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (v) to form a binding site for a target antigen;

vii) a second heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; and an Fc subunit (e.g., CH2-CH3); and

viii) a further Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (vii) to form a binding site for a target antigen.

Optionally the Fab heavy chain in (v) has the same sequence as the Fab heavy chain in (vii) and the Fab light chains of (vi) and (viii) have the same sequence as each other.

Optionally the Fab heavy chains in (i), (iii), (v) and (vii) have the same sequence as each other and the Fab light chains of (ii), (iv) (vi) and (viii) have the same sequence as each other.

In other embodiments, which may in some instances be preferred, the first and/or the second antibody each have a single antigen binding moiety capable of specific binding to a target antigen. Thus, the first antibody and/or second antibody may be monospecific and monovalent for a target antigen. Preferably the first and second antibody bind to the same target antigen as each other, at the same or at different epitopes.

In one embodiment, the first and/or second antibody is a one-armed antibody. In such embodiments, the Fc subunit of the first antibody which is not fused to the polypeptide of (b) is also not fused to any other antigen binding moiety; and/or the Fc subunit of the second antibody which is not fused to the polypeptide of (e) is also not fused to any other antigen binding moiety. Thus, the Fc domain may comprise a subunit which is lacking Fd. In some embodiments, one of the polypeptides making up the antibody may consist or consist essentially of the Fc subunit.

Thus, in some embodiments, the first antibody may comprise the following polypeptides:

i) a polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VH domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);

ii) a Fab light chain polypeptide (e.g., VL-CL); and

iii) an Fc subunit polypeptide (e.g., CH2-CH3);

wherein the Fab heavy chain of (i) and the Fab light chain of (ii) form a Fab fragment capable of binding to a target antigen.

The second antibody may comprise the following polypeptides:

iv) a polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VL domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);

v) a Fab light chain polypeptide (e.g., VL-CL), and

vi) an Fc subunit polypeptide (e.g., CH2-CH3);

wherein the Fab heavy chain of (iv) and the Fab light chain of (v) form a Fab fragment capable of binding to a target antigen.

10 In some embodiments of these one-armed antibodies, the Fab heavy chain of (i) and of (iv) may have the same sequence as each other; and the Fab light chain polypeptide of ii) and (v) may have the same sequence as each other.

In any embodiments where an antibody comprises two Fab fragments with different paratopes (e.g., binding different antigens and/or epitopes), it may be preferred that one of the Fabs is a conventional Fab (comprising a heavy chain VH-CH1 and a light chain VL-CL) and the other is a cross-Fab or scFab. Fabs having a first specificity/variable domain sequence may be conventional Fabs, and Fabs having a second specificity/variable domain sequence may be selected from a cross-Fab or a scFab. This reduces the potential for mispairing of the light chains.

20 In any of the above, the antigen binding moieties may be fused to the Fc domain or to each other directly or through a peptide linker, comprising one or more amino acids. Peptide linkers are known in the art and are described herein. The linker (e.g., the linker between the Fab fragment and the VH/VL for the effector moiety and/or between the VH/VL for the effector moiety and the Fc domain) may be a peptide of at least 5 amino acids or at least 10 amino acids, preferably 5 to 100, e.g., 10 to 70, 10 to 60, or 10 to 50 amino acids. In some embodiments, it may be preferred that the linker is 15-30 amino acids in length, e.g., 15-25, e.g., 16, 17, 18, 19, 20, 21, 22, 23 or 24 amino acids in length. The linker may be a rigid linker or a flexible linker. In some embodiments, it is a flexible linker comprising or consisting of Thr, Ser, Gly and/or Ala residues. For example, it may comprise or consist of
30 Gly and Ser residues. In some embodiments it may have a repeating motif such as (Gly-Gly-Gly-Ser)_n, where n is for instance 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10. Suitable, non-immunogenic peptide linkers include, for example, (G₄S)_n, (SG₄)_n, (G₄S)_n or G₄(SG₄)_n peptide linkers, where "n" is generally a number between 1 and 10, typically between 2 and

4. In another embodiment said peptide linker is $(GxS)_n$ or $(GxS)_nG_m$ with G = glycine, S = serine, and $(x = 3, n = 3, 4, 5 \text{ or } 6, \text{ and } m = 0, 1, 2 \text{ or } 3)$ or $(x = 4, n = 2, 3, 4 \text{ or } 5 \text{ and } m = 0, 1, 2 \text{ or } 3)$, e.g., $x = 4$ and $n = 2$ or 3 , e.g., with $x = 4, n = 2$. In some embodiments, the linker may be or may comprise the sequence GGGGSGGGGSGGGGSGGGGS (SEQ ID NO.: 31). In another embodiment the linker may be or comprise GGGGSGGGGSGGGGSGGSGG (SEQ ID NO: 148) or GGGGSGGGGSGGGGSGGSGGS (SEQ ID NO: 149) or GGGGSGGGGSGGGGSGGSGGG (SEQ ID NO: 150). Another exemplary peptide linker is EPKSC(D)-(G₄S)₂. (SEQ ID NO: 151) Additionally, where an antigen binding moiety is fused to the N-terminus of an Fc domain subunit, it may be fused via an immunoglobulin hinge region or a portion thereof, with or without an additional peptide linker.

The present inventors have determined that in a peptide linker consisting of y amino acids, a Ser in the y position (i.e., a Ser as the last/C-terminal amino acid of the linker) may induce glycosylation of the y +2 amino acid (i.e., of the amino acid positioned 2 residues in the C-terminal direction from the last amino acid in the linker), depending on the nature of this y+2 amino acid. Therefore it may be preferred that the last serine residue of the linker is placed in the y-2 or y-3 position (i.e., that the last serine residue of the linker is at a position 2 or 3 amino acids in the N-terminal direction from the last amino acid in the linker). In some embodiments, the linker may consist of y consecutive amino acid residues selected from the group consisting of Gly and Ser, e.g., wherein y=at least 5 or at least 10 and less than or equal to 100, e.g., 5 to 100, 10 to 70, 10 to 60 or 10 to 50, e.g., 15 to 31 or 15 to 30, e.g., 15, 16, 17, 18, 19, 20, 21, 22, 23, 24 or 25, and wherein the last serine is in the y-2 or y-3 position. (Thus, there may be a serine in the y-2 position and a glycine in the y-1 and y position; or there may be a serine in the y-3 position and a glycine in the y-2, y-1 and y positions). In some embodiments it may be preferred that y=20 or 21. In some embodiments, it may be preferred that the linker is $(GxS)_n(GGSGG)$ or $(GxS)_n(GGSGGG)$ with G = glycine, S = serine, $x = 4$ and $n = 1$ to 20, e.g., 1 to 10, e.g., 2, 3, 4, 5, 6, 7, 8, or 9, e.g., $n = 2$ to 4. For instance, the linker may be GGGGSGGGGSGGGGSGGSGG (SEQ ID NO: 148) or GGGGSGGGGSGGGGSGGSGGG (SEQ ID NO: 150).

In a particular embodiment of any of the above sets of antibodies, the Fc domain is an IgG Fc domain. In a specific embodiment, the Fc domain is an IgG1 Fc domain. In another specific embodiment, the Fc domain is an IgG4 Fc domain. In an even more specific embodiment, the Fc domain is an IgG4 Fc domain comprising the amino acid substitution S228P (Kabat numbering). In particular embodiments the Fc domain is a human Fc domain.

It may be preferred that the Fc region is engineered to reduce or eliminate Fc effector function. This may include substitution of one or more of Fc region residues 234, 235, 238, 265, 269, 270, 297, 327 and/or 329, e.g., one or more of 234, 235 and/or 329. In some embodiments, the Fc region may be engineered to include the substitution of Pro 329 to Gly, Leu 234 to Ala and/or Leu 235 to Ala (numbering according to EU index). Modifications to reduce Fc effector function are discussed further below.

Techniques which are known for making multispecific antibodies can also be used to make any of the heterodimers described herein. These include, but are not limited to, 10 recombinant co-expression of two immunoglobulin heavy chain-light chain pairs having different specificities (see Milstein and Cuello, *Nature* 305: 537 (1983)) and “knob-in-hole” engineering (see, e.g., U.S. Patent No. 5,731,168, and Atwell et al., *J. Mol. Biol.* 270:26 (1997)). Other methods include engineering electrostatic steering effects for making antibody Fc-heterodimeric molecule (see, e.g., WO 2009/089004); cross-linking two or more antibodies or fragments (see, e.g., US Patent No. 4,676,980, and Brennan et al., *Science*, 229: 81 (1985)); using leucine zippers (see, e.g., Kostelny et al., *J. Immunol.*, 148(5):1547-1553 (1992) and WO 2011/034605); and using the common light chain technology for circumventing the light chain mis-pairing problem (see, e.g., WO 98/50431).

The CH3 domains of the full length antibody as described above can be altered by the 20 “knob-into-holes” technology which is described in detail with several examples in e.g. WO 96/027011, Ridgway, J.B., et al., *Protein Eng* 9 (1996) 617-621; and Merchant, A.M., et al., *Nat Biotechnol* 16 (1998) 677-681. In this method the interaction surfaces of the two CH3 domains are altered to increase the heterodimerisation of both heavy chains containing these two CH3 domains. Each of the two CH3 domains (of the two heavy chains) can be the “knob”, while the other is the “hole”. For instance one comprises called “knob mutations” (T366W and optionally one of S354C or Y349C) and the other comprises the so-called “hole mutations” (T366S, L368A and Y407V and optionally Y349C or S354C) (see, e.g., Carter, P. et al., *Immunotechnol.* 2 (1996) 73) according to EU index numbering.

The introduction of a disulfide bridge may additionally or alternatively be used to 30 stabilize the heterodimers (Merchant, A.M., et al., *Nature Biotech* 16 (1998) 677-681; Atwell, S., et al., *J. Mol. Biol.* 270 (1997) 26-35) and increase the yield.

Thus in some embodiments the first and/or second antibody is further characterized in that: the CH3 domain of one heavy chain of the full length antibody and the CH3 domain of

the other heavy chain of the full length antibody each meet at an interface which comprises an original interface between the antibody CH3 domains; wherein said interface is altered to promote the formation of the antibody, wherein the alteration is characterized in that:

a) the CH3 domain of one heavy chain is altered, so that within the original interface the CH3 domain of one heavy chain that meets the original interface of the CH3 domain of the other heavy chain within the antibody, an amino acid residue is replaced with an amino acid residue having a larger side chain volume, thereby generating a protuberance within the interface of the CH3 domain of one heavy chain which is positionable in a cavity within the interface of the CH3 domain of the other heavy chain

10 and

b) the CH3 domain of the other heavy chain is altered, so that within the original interface of the second CH3 domain that meets the original interface of the first CH3 domain within the antibody an amino acid residue is replaced with an amino acid residue having a smaller side chain volume, thereby generating a cavity within the interface of the second CH3 domain within which a protuberance within the interface of the first CH3 domain is positionable.

Said amino acid residue having a larger side chain volume may optionally be selected from the group consisting of arginine (R), phenylalanine (F), tyrosine (Y), tryptophan (W).

20 Said amino acid residue having a smaller side chain volume may optionally be selected from the group consisting of alanine (A), serine (S), threonine (T), valine (V).

Optionally, in some embodiments, both CH3 domains are further altered by the introduction of cysteine (C) as amino acid in the corresponding positions of each CH3 domain such that a disulfide bridge between both CH3 domains can be formed.

30 In some embodiments, multispecific (e.g., biparatopic) antibodies may also comprise amino acid substitutions in Fab molecules (including cross-Fab molecules) comprised therein which are particularly efficient in reducing mispairing of light chains with non-matching heavy chains (Bence-Jones-type side products), which can occur in the production of Fab-based bi-/multispecific antigen binding molecules with a VH/VL exchange in one (or more, in case of molecules comprising more than two antigen-binding Fab molecules) of their binding arms (see also PCT publication no. WO 2015/150447, particularly the examples therein, incorporated herein by reference in its entirety). The ratio of a desired multispecific antibodies compared to undesired side products, in particular Bence Jones-type side products

occurring in one of their binding arms, can be improved by the introduction of charged amino acids with opposite charges at specific amino acid positions in the CH1 and CL domains of a Fab molecule (sometimes referred to herein as “charge modifications”).

Therefore, in some embodiments, an antibody of the present invention comprising Fab molecules, comprises at least one Fab with a heavy chain constant domain CH1 domain comprising charge modifications as described herein, and a light chain constant CL domain comprising charge modifications as described herein.

Charge modifications can be made either in the conventional Fab molecule(s) comprised in the antibodies of the present invention, or in the crossover Fab molecule(s) comprised in the antibodies of the present invention (but not in both). In particular
10 embodiments, the charge modifications are made in the conventional Fab molecule(s) comprised in the antibodies of the present invention.

In some embodiments, in a Fab or cross-Fab comprising a light chain constant domain CL comprising charge modifications and a heavy chain constant domain CH1 comprising charge modifications, charge modifications in the light chain constant domain CL are at position 124 and optionally at position 123 (numbering according to Kabat), and charge modifications in the heavy chain constant domain CH1 are at position 147 and/or 213 (numbering according to Kabat EU Index). In some embodiments, in the light chain constant domain CL the amino acid at position 124 is substituted independently by lysine (K), arginine
20 (R) or histidine (H) (numbering according to Kabat) (in one preferred embodiment independently by lysine (K)), and in the heavy chain constant domain CH1 the amino acid at position 147 and/or the amino acid at position 213 is substituted independently by glutamic acid (E) or aspartic acid (D) (numbering according to Kabat EU index).

B. Target Antigens

The antigen expressed on the surface of the target cell is also termed herein the “target antigen” or “target cell antigen”. These terms are used interchangeably herein.

Insofar as the invention relates to treatment methods and to products for use therein, it is applicable to any condition that is treatable by cytotoxic activity targeted to cells of the
30 patient, e.g., diseased cells. Thus, the target cell is any cell against which it is desired to target cytotoxicity, e.g., any diseased cell. The treatment is preferably of a tumour or cancer. However, the applicability of the invention is not limited to tumours and cancers. For example, the treatment may also be of viral infection (by targeting infected cells) or T-cell

driven autoimmune disease (by targeting T cells). Immunotoxins directed against viral antigens expressed on the surface of infected cells have been investigated for a variety of viral infections such as HIV, rabies and EBV. Cai and Berger 2011 Antiviral Research 90(3):143-50 used an immunotoxin containing PE38 for targeted killing of cells infected with Kaposi's sarcoma-associated herpesvirus. In addition, Resimmune® (A-dmDT390-bisFv(UCHT1)) selectively kills human malignant T cells and transiently depletes normal T cell and is considered to have potential for the treatment of T-cell driven autoimmune diseases such as multiple sclerosis and graft-versus-host disease, as well as T cell blood cancers for which it is undergoing clinical trials. Likewise, methods of the invention may be applicable to any cell type for which imaging is desirable, including but not limited to cancer or tumour cells.

Thus, suitable target antigens may include cancer cell antigens, viral antigens or microbial antigens.

The antigens are usually normal cell surface antigens which are either over-expressed or expressed at abnormal times. Ideally the target antigen is expressed only on diseased cells (such as tumour cells), however this is rarely observed in practice. As a result, target antigens are usually selected on the basis of differential expression between diseased and healthy tissue.

The cell surface marker or target antigen can be, for example, a tumour-associated antigen.

The term "tumour-associated antigen" or "tumour specific antigen" as used herein refers to any molecule (e.g., protein, peptide, lipid, carbohydrate, etc.) solely or predominantly expressed or over-expressed by tumour cells and/or cancer cells, or by other cells of the stroma of the tumour such as cancer-associated fibroblasts, such that the antigen is associated with the tumour(s) and/or cancer(s). The tumour-associated antigen can additionally be expressed by normal, non-tumour, or non-cancerous cells. However, in such cases, the expression of the tumour-associated antigen by normal, non-tumour, or non-cancerous cells is not as robust as the expression by tumour or cancer cells. In this regard, the tumour or cancer cells can over-express the antigen or express the antigen at a significantly higher level, as compared to the expression of the antigen by normal, non-tumour, or non-cancerous cells. Also, the tumour-associated antigen can additionally be expressed by cells of a different state of development or maturation. For instance, the tumour-associated antigen can be additionally expressed by cells of the embryonic or foetal stage, which cells are not

normally found in an adult host. Alternatively, the tumour-associated antigen can be additionally expressed by stem cells or precursor cells, which cells are not normally found in an adult host.

The tumour-associated antigen can be an antigen expressed by any cell of any cancer or tumour, including the cancers and tumours described herein. The tumour-associated antigen may be a tumour-associated antigen of only one type of cancer or tumour, such that the tumour-associated antigen is associated with or characteristic of only one type of cancer or tumour. Alternatively, the tumour-associated antigen may be a tumour-associated antigen (e.g., may be characteristic) of more than one type of cancer or tumour. For example, the
 10 tumour-associated antigen may be expressed by both breast and prostate cancer cells and not expressed at all by normal, non-tumour, or non-cancer cells.

Exemplary tumour-associated antigens to which the antibodies of the invention may bind include, but are not limited to, Melanoma-associated Chondroitin Sulfate Proteoglycan (MCSP), mucin 1 (MUC1; tumour-associated epithelial mucin), preferentially expressed antigen of melanoma (PRAME), carcinoembryonic antigen (CEA), prostate specific membrane antigen (PSMA), PSCA, EpCAM, Trop2 (trophoblast-2, also known as EGP-1), granulocyte-macrophage colony-stimulating factor receptor (GM-CSFR), CD56, human epidermal growth factor receptor 2 (HER2/neu) (also known as erbB-2), CDS, CD7, tyrosinase related protein (TRP) I, and TRP2. In another embodiment, the tumour antigen
 20 may be selected from the group consisting of cluster of differentiation (CD) 19, CD20, CD21, CD22, CD25, CD30, CD33 (sialic acid binding Ig-like lectin 3, myeloid cell surface antigen), CD79b, CD123 (interleukin 3 receptor alpha), transferrin receptor, EGF receptor, mesothelin, cadherin, Lewis Y, Glypican-3, FAP (fibroblast activation protein alpha), GPRC5D (G Protein-Coupled Receptor Class C Group 5 Member D), PSMA (prostate specific membrane antigen), CA9 = CAIX (carbonic anhydrase IX), L1 CAM (neural cell adhesion molecule L 1), endosialin, HER3 (activated conformation of epidermal growth factor receptor family member 3), Alkl/BMP9 complex (anaplastic lymphoma kinase 1/bone morphogenetic protein 9), TPBG = 5T4 (trophoblast glycoprotein), ROR1 (receptor tyrosine kinase-like surface antigen), HER1 (activated conformation of epidermal growth factor receptor), and CLL1 (C-
 30 type lectin domain family 12, member A). Mesothelin is expressed in, e.g., ovarian cancer, mesothelioma, non-small cell lung cancer, lung adenocarcinoma, fallopian tube cancer, head and neck cancer, cervical cancer, and pancreatic cancer. CD22 is expressed in, e.g., hairy cell leukaemia, chronic lymphocytic leukaemia (CLL), prolymphocytic leukaemia (PLL), non-

Hodgkin's lymphoma, small lymphocytic lymphoma (SLL), and acute lymphatic leukaemia (ALL). CD25 is expressed in, e.g., leukemias and lymphomas, including hairy cell leukaemia and Hodgkin's lymphoma. Lewis Y antigen is expressed in, e.g., bladder cancer, breast cancer, ovarian cancer, colorectal cancer, esophageal cancer, gastric cancer, lung cancer, and pancreatic cancer. CD33 is expressed in, e.g., acute myeloid leukaemia (AML), chronic myelomonocytic leukaemia (CML), and myeloproliferative disorders.

Exemplary antibodies that specifically bind to tumour-associated antigens include, but are not limited to, antibodies against the transferrin receptor (e.g., HB21 and variants thereof), antibodies against CD22 (e.g., RFB4 and variants thereof), antibodies against CD25 (e.g., anti-Tac and variants thereof), antibodies against mesothelin (e.g., SS 1, MORAb-009, SS, HN1, HN2, MN, MB, and variants thereof) and antibodies against Lewis Y antigen (e.g., B3 and variants thereof). In this regard, the targeting moiety (cell-binding agent) may be an antibody selected from the group consisting of B3, RFB4, SS, SS1, MN, MB, HN1, HN2, HB21, and MORAb-009, and antigen binding portions thereof. Further exemplary targeting moieties suitable for use in the inventive chimeric molecules are disclosed e.g., in U.S. Patents 5,242,824 (anti-transferrin receptor); 5,846,535 (anti-CD25); 5,889,157 (anti-Lewis Y); 5,981,726 (anti-Lewis Y); 5,990,296 (anti-Lewis Y); 7,081,518 (anti-mesothelin); 7,355,012 (anti-CD22 and anti-CD25); 7,368,110 (anti-mesothelin); 7,470,775 (anti-CD30); 7,521,054 (anti-CD25); and 7,541,034 (anti-CD22); U.S. Patent Application Publication 2007/0189962 (anti-CD22); Frankel et al., Clin. Cancer Res., 6: 326-334 (2000), and Kreitman et al., AAPS Journal, 8(3): E532-E551 (2006), each of which is incorporated herein by reference.

Further antibodies have been raised to target specific tumour related antigens including: Cripto, CD30, CD19, CD33, Glycoprotein NMB, CanAg, Her2 (ErbB2/Neu), CD56 (NCAM), CD22 (Siglec2), CD33 (Siglec3), CD79, CD138, PSCA, PSMA (prostate specific membrane antigen), BCMA, CD20, CD70, E-selectin, EphB2, Melanotransferin, Muc16 and TMEFF2. Any of these, or antigen-binding fragments thereof, may be useful in the present invention, i.e., may be incorporated into the antibodies described herein.

In some embodiments of the present invention, it may be preferred that the tumour-associated antigen is carcinoembryonic antigen (CEA).

CEA is advantageous in the context of the present invention because it is relatively slowly internalized, and thus a high percentage of the antibody will remain available on the surface of the cell after initial treatment, for binding to the radionuclide. Other low

internalizing targets/tumour associated antigens may also be preferred. Other examples of tumour-associated antigen include CD20 or HER2. In still further embodiments, the target may be EGP-1 (epithelial glycoprotein-1, also known as trophoblast-2), colon-specific antigen-p (CSAp) or a pancreatic mucin MUC1. See for instance Goldenberg et al 2012 (Theranostics 2(5)), which is incorporated herein by reference. This reference also describes antibodies such as Mu-9 binding to CSAp (see also Sharkey et al Cancer Res. 2003; 63: 354-63), hPAM4 binding to MUC1 (see also Gold et al Cancer Res. 2008; 68: 4819-26), valtuzumab binding to CD20 (see also Sharkey et al Cancer Res. 2008; 68: 5282-90) and hRS7 which binds to EGP-1 (see also Cubas et al Biochim Biophys Acta 2009; 1796: 309-14). Any of these or antigen-binding portions thereof may be useful in the present invention, i.e., may be incorporated into the antibodies described herein. One example of an antibody that has been raised against CEA is T84.66 (as shown in NCBI Acc No: CAA36980 for the heavy chain and CAA36979 for the light chain, or as shown in SEQ ID NO 317 and 318 of WO2016/075278) and humanized and chimeric versions thereof, such as T84.66-LCHA as described in WO2016/075278 A1 and/or WO2017/055389. Another example is CH1A1a, an anti-CEA antibody as described in WO2012/117002 and WO2014/131712, and CEA hMN-14 (see also US 6 676 924 and US 5 874 540). Another anti-CEA antibody is A5B7 as described in M.J. Banfield et al, Proteins 1997, 29(2), 161-171. Humanized antibodies derived from murine antibody A5B7 have been disclosed in WO 92/01059 and WO 2007/071422. See also co-pending application PCT/EP2020/067582. An example of a humanized version of A5B7 is A5H1EL1(G54A). A further exemplary antibody against CEA is MFE23 and the humanized versions thereof described in US7626011 and/or co-pending application PCT/EP2020/067582. A still further example of an antibody against CEA is 28A9. Any of these or an antigen binding fragment thereof may be useful to form a CEA-binding moiety in the present invention.

FAP (fibroblast activation protein alpha) or GPRC5D (G Protein-Coupled Receptor Class C Group 5 Member D) may also be preferred in some embodiments. FAP is an established target for imaging and therapy, due to its broad expression in the microenvironment of a number of tumor types, e.g. pancreas, breast, and lung cancer (Lindner, T., Loktev, A., Giesel, F. et al. Targeting of activated fibroblasts for imaging and therapy. EJNMMI radiopharm. chem. 4, 16 (2019)). In one embodiment of the invention, SPLIT PRIT using FAP as the target antigen would thus be expected to generate specific accumulation of ²¹²Pb-DOTAM on activated cancer-associated fibroblasts. Consequently,

the emitted alpha radiation would be expected to negatively affect the immune suppression of FAP-expressing malignant tumors, in addition to a limited direct tumor-killing effect on adjacent tumor cells. G-protein coupled receptor family C group 5 member D (GPRC5D) is overexpressed on multiple myeloma plasma cells (Atamaniuk J, Gleiss A, Porpaczy E, Kainz B, Grunt TW, Raderer M, et al. Overexpression of G protein-coupled receptor 5D in the bone marrow is associated with poor prognosis in patients with multiple myeloma. *Eur J Clin Invest.* 2012;42:953-60.), with established SC (subcutaneous) *in vivo* models reflecting the expression found in multiple myeloma patients, e.g. OPM-2 and NCI-H929 (Kodama T, Kochi Y, Nakai W, Mizuno H, Baba T, Habu K, et al. Anti-GPRC5D/CD3 bispecific T-cell-
10 redirecting antibody for the treatment of multiple myeloma. *Mol Cancer Ther.* (2019) 18:1555–64). In one embodiment of the invention, we therefore expect SPLIT PRIT using GPRC5D as the target antigen - to generate tumor-specific accumulation of ^{212}Pb -DOTAM followed by radiation-induced tumor cell death.

In some embodiments, the antibodies of the invention may bind specifically to the target antigen (e.g., any of the target antigens discussed herein). In some embodiments, they may bind with a dissociation constant (K_D) of $\leq 1\mu\text{M}$, $\leq 100\text{ nM}$, $\leq 10\text{ nM}$, $\leq 1\text{ nM}$, $\leq 0.1\text{ nM}$, $\leq 0.01\text{ nM}$, or $\leq 0.001\text{ nM}$ (e.g. 10^{-7}M or less, e.g. from 10^{-7} to 10^{-13} , 10^{-8} M or less, e.g. from 10^{-8} M to 10^{-13} M , e.g., from 10^{-9} M to 10^{-13} M).

In one embodiment, the first and second antibody may each bind to the same target
20 antigen, which can be termed “antigen A” (i.e., they have binding specificity for the same target antigen). They may each having binding specificity for the same epitope on antigen A. Alternatively, the first antibody may bind to a first epitope on antigen A and the second antibody may bind to a different, second epitope on antigen A. For instance, in one embodiment, one of the antibodies may bind to the T84.66 epitope of CEA and the other may bind to the A5B7 epitope of CEA. In some embodiments, one or both of the first and/or second antibodies may be biparatopic for antigen A – i.e., each of the individual antibodies may bind to two different epitopes of antigen A. The first antibody may comprise a first and a second binding site, which bind to a first and a second epitope of antigen A respectively, wherein the first and second epitopes are different from each other. Alternatively or
30 additionally, the second antibody may comprise a first and a second binding site, which bind to a first and a second epitope of antigen A, wherein the first and second epitopes are different from each other. In some embodiments, one or both of the epitopes bound by the first antibody may be different from one or both of the epitopes bound by the second

antibody. In other embodiments, the two epitopes bound by the first antibody may be the same as the two epitopes bound by the second antibody.

In another embodiment, the first antibody and second antibody may respectively bind to different target antigens, which may be termed antigen A and antigen B respectively.

C. Effector moieties

According to the present invention, association of the first and second antibodies forms a functional binding site for an effector moiety.

In one particular embodiment, an effector moiety according to the present invention is selected from a drug, a cytotoxin, an imaging agent, and a radiolabelled compound.

In a particular embodiment, an effector moieties according to the present invention is a radiolabelled compounds which comprise a radioisotope, e.g., are a radiolabelled hapten.

In some embodiments, the effector molecule may comprise a chelated radioisotope.

In some embodiments, the functional binding site for the effector molecule may bind to a chelate comprising the chelator and the radioisotope. In other embodiments, the antibody may bind to a moiety which is conjugated to the chelated radioisotope, for instance, histamine-succinyl-glycine (HSG), digoxigenin, biotin or caffeine

The chelator may be, for example, a multidentate molecule such as an aminopolycarboxylic acid or an aminopolythiocarboxylic acid, or a salt or functional variant thereof. The chelator may be, for example, bidentate or tridentate or tetradentate. Examples of suitable metal chelators include molecules comprising EDTA (Ethylenediaminetetraacetic acid, or a salt form such as CaNa_2EDTA), DTPA (Diethylenetriamine Pentaacetic Acid), DOTA (1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid), NOTA (2,2',2''-(1,4,7-Triazanonane-1,4,7-triyl)triacetic acid), IDA (Iminodiacetic acid), MIDA ((Methylimino)diacetic acid), TTHA (3,6,9,12-Tetrakis(carboxymethyl)-3,6,9,12-tetraazatetradecanedioic acid), TETA (2,2',2'',2'''-(1,4,8,11-Tetraazacyclotetradecane-1,4,8,11-tetrayl)tetraacetic acid), DOTAM (1,4,7,10-Tetrakis(carbamoylmethyl)-1,4,7,10-tetraazacyclododecane), HEHA (1,4,7,10,13,16-hexaazacyclohexadecane-1,4,7,10,13,16-hexaacetic acid, available from Macrocyclics, Inc., Plano, Texas), NTA (nitrilotriacetic acid) EDDHA (ethylenediamine-N, N'-bis(2-hydroxyphenylacetic acid), BAL (2,3,-dimercaptopropanol), DMSA (2,3-dimercaptosuccinic acid), DMPS (2,3-dimercapto-1-propanesulfonic acid), D-penicillamine (B-dimethylcysteine), MAG_3 (mercaptoacetyltriglycine), Hynic (6-hydrazinopridine-3-carboxylic acid), p-

isothiocyanatobenzyl-desferrioxamine (e.g., labelled with zirconium for imaging), and salts or functional variants/derivatives thereof capable of chelating the metal. In some embodiments, it may be preferred that the chelator is DOTA or DOTAM or a salt or functional variant/derivative thereof capable of chelating the metal. Thus, the chelator may be or may comprise DOTA or DOTAM with a radioisotope chelated thereto.

The effector molecule may comprise or consist of functional variants or derivatives of the chelators above, together with the radionuclide. Suitable variants/derivatives have a structure that differs to a certain limited extent and retain the ability to function as a chelator (i.e. retains sufficient activity to be used for one or more of the purposes described herein).

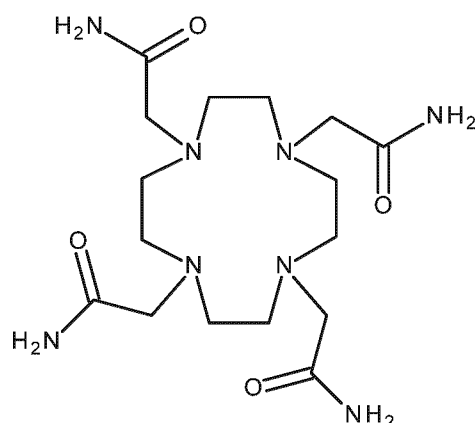
10 Functional variants/derivatives may also include a chelator as described above conjugated to one or more additional moieties or substituents, including, a small molecule, a polypeptide or a carbohydrate. This attachment may occur via one of the constituent carbons, for example in a backbone portion of the chelator. A suitable substituent can be, for example, a hydrocarbon group such as alkyl, alkenyl, aryl or alkynyl; a hydroxy group; an alcohol group; a halogen atom; a nitro group; a cyano group; a sulfonyl group; a thiol group; an amine group; an oxo group; a carboxy group; a thiocarboxy group; a carbonyl group; an amide group; an ester group; or a heterocycle including heteroaryl groups. The substituent may be, for example, one of those defined for group "R¹" below. A small molecule can be, for example, a dye (such as Alexa 647 or Alexa 488), biotin or a biotin moiety, or a phenyl or benzyl moiety. A
20 polypeptide may be, for example, an oligo peptide, e.g., an oligopeptide of two or three amino acids. Exemplary carbohydrates include dextran, linear or branched polymers or copolymers (e.g. polyalkylene, poly(ethylene-lysine), polymethacrylate, polyamino acids, poly- or oligosaccharides, dendrimers). Derivatives may also include multimers of the chelator compounds in which compounds as set out above are linked through a linker moiety. Derivatives may also include functional fragments of the above compounds, which retain the ability to chelate the metal ion.

Particular examples of derivatives include benzyl-EDTA and hydroxyethyl-thiourido-benzyl EDTA, DOTA-benzene (e.g., (S-2-(4-aminobenzyl)-1,4,7,10-tetraazacyclododecane tetraacetic acid), DOTA-biotin, and DOTA-TyrLys-DOTA.

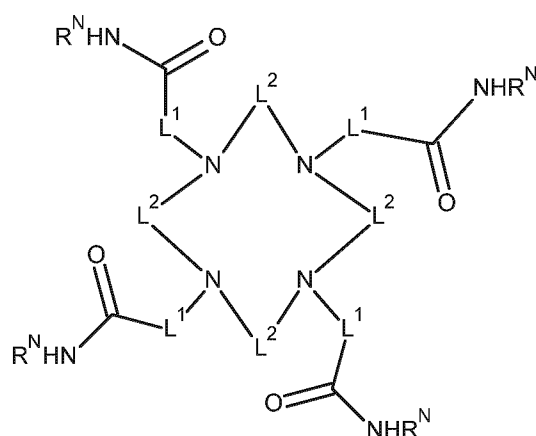
30 In some embodiments of the present invention, the functional binding site formed by association of the first and second antibody binds to a metal chelate comprising DOTAM and a metal, e.g., lead (Pb). As mentioned above, "DOTAM" has the chemical name:

1,4,7,10-Tetrakis(carbamoylmethyl)-1,4,7,10-tetraazacyclododecane,

which is a compound of the following formula:



The present invention may in certain aspects and embodiments also make use of functional variants or derivatives of DOTAM incorporating a metal ion. Suitable variants/derivatives of DOTAM have a structure that differs to a certain limited extent from the structure of DOTAM and retain the ability to function (i.e. retains sufficient activity to be used for one or more of the purposes described herein). In such aspects and embodiments, the DOTAM or functional variant/derivative of DOTAM may be one of the active variants disclosed in WO 2010/099536. Suitable functional variants/derivatives may be a compound of the following formula:



or a pharmaceutically acceptable salt thereof; wherein

R^N is H, C_{1-6} alkyl, C_{1-6} haloalkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{3-7} cycloalkyl, C_{3-7} cycloalkyl- C_{1-4} alkyl, C_{2-7} heterocycloalkyl, C_{2-7} heterocycloalkyl- C_{1-4} alkyl, phenyl, phenyl-

C₁₋₄-alkyl, C₁₋₇ heteroaryl, and C₁₋₇ heteroaryl-C₁₋₄-alkyl; wherein C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₂₋₆ alkenyl, and C₂₋₆ alkynyl are each optionally substituted by 1, 2, 3, or 4 independently selected R^w groups; and wherein said C₃₋₇ cycloalkyl, C₃₋₇ cycloalkyl-C₁₋₄-alkyl, C₂₋₇ heterocycloalkyl, C₂₋₇ heterocycloalkyl-C₁₋₄ alkyl, phenyl, phenyl-C₁₋₄-alkyl, C₁₋₇ heteroaryl, and C₁₋₇ heteroaryl-C₁₋₄-alkyl are each optionally substituted by 1, 2, 3, or 4 independently selected R^x groups;

L¹ is independently C₁₋₆ alkylene, C₁₋₆ alkenylene, or C₁₋₆ alkynylene, each of which is optionally substituted by 1, 2, or 3 groups independently selected R¹ groups;

10 L² is C₂₋₄ straight chain alkylene, which is optionally substituted by an independently selected R¹ group; and which is optionally substituted by 1, 2, 3, or 4 groups independently selected from C₁₋₄ alkyl and or C₁₋₄ haloalkyl;

R¹ is independently selected from D¹-D²-D³, halogen, cyano, nitro, hydroxyl, C₁₋₆ alkoxy, C₁₋₆ haloalkoxy, C₁₋₆ alkylthio, C₁₋₆ alkylsulfinyl, C₁₋₆ alkylsulfonyl, amino, C₁₋₆ alkylamino, di-C₁₋₆ alkylamino, C₁₋₄ alkylcarbonyl, carboxy, C₁₋₆ alkoxycarbonyl, C₁₋₆ alkylcarbonylamino, di-C₁₋₆ alkylcarbonylamino, C₁₋₆ alkoxycarbonylamino, C₁₋₆ alkoxycarbonyl-(C₁₋₆ alkyl)amino, carbamyl, C₁₋₆ alkylcarbamyl, and di-C₁₋₆ alkylcarbamyl;

20 each D¹ is independently selected from C₆₋₁₀ aryl-C₁₋₄ alkyl, C₁₋₉ heteroaryl-C₁₋₄ alkyl, C₃₋₁₀ cycloalkyl-C₁₋₄ alkyl, C₂₋₉ heterocycloalkyl-C₁₋₄ alkyl, C₁₋₈ alkylene, C₁₋₈ alkenylene, and C₁₋₈ alkynylene; wherein said C₁₋₈ alkylene, C₁₋₈ alkenylene, and C₁₋₈ alkynylene are optionally substituted by 1, 2, 3, or 4 independently selected R⁴ groups; and wherein said C₆₋₁₀ aryl-C₁₋₄ alkyl, C₁₋₉ heteroaryl-C₁₋₄ alkyl, C₃₋₁₀ cycloalkyl-C₁₋₄ alkyl, C₂₋₉ heterocycloalkyl-C₁₋₄ alkyl are each optionally substituted by 1, 2, 3, or 4 independently selected R⁵ groups;

30 each D² is independently absent or C₁₋₂₀ straight chain alkylene, wherein from 1 to 6 non-adjacent methylene groups of said C₁₋₂₀ straight chain alkylene are each optionally replaced by an independently selected -D⁴- moiety, provided that at least one methylene unit in said C₁₋₂₀ straight chain alkylene is not optionally replaced by a -D⁴- moiety; wherein said C₁₋₂₀ straight chain alkylene is optionally substituted by one or more groups independently selected from halogen, cyano, nitro, hydroxyl, C₁₋₄ alkyl, C₁₋₄ haloalkyl, C₁₋₄ alkoxy, C₁₋₄ haloalkoxy, amino, C₁₋₄ alkylamino, di-C₁₋₄ alkylamino, C₁₋₄ alkylcarbonyl, carboxy, C₁₋₄ alkoxycarbonyl, C₁₋₄ alkylcarbonylamino, di-C₁₋₄ alkylcarbonylamino,

C₁₋₄ alkoxy-carbonylamino, C₁₋₄ alkoxy-carbonyl-(C₁₋₄ alkyl)amino, carbamyl, C₁₋₄ alkylcarbamyl, and di-C₁₋₄ alkylcarbamyl;

each D³ is independently selected from H, halogen, cyano, nitro, hydroxyl, C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₃₋₁₄ cycloalkyl, C₃₋₁₄ cycloalkyl-C₁₋₄ alkyl, C₂₋₁₄ heterocycloalkyl, C₂₋₁₄ heterocycloalkyl-C₁₋₄ alkyl, C₆₋₁₄ aryl, C₆₋₁₄ aryl-C₁₋₄ alkyl, C₁₋₁₃ heteroaryl, C₁₋₁₃ heteroaryl-C₁₋₄ alkyl; wherein said C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl are each optionally substituted by 1, 2, 3, or 4 independently selected R⁶ groups; and wherein said C₃₋₁₄ cycloalkyl, C₃₋₁₄ cycloalkyl-C₁₋₄ alkyl, C₂₋₁₄ heterocycloalkyl, C₂₋₁₄ heterocycloalkyl-C₁₋₄ alkyl, C₆₋₁₄ aryl, C₆₋₁₄ aryl-C₁₋₄ alkyl, C₁₋₁₃ heteroaryl, C₁₋₁₃ heteroaryl-C₁₋₄ alkyl are each optionally substituted by 1, 2, 3 or 4 independently selected R⁷ groups;

each D⁴ is independently selected from -O-, -S-, -NR^aC(=O)-, -NR^aC(=S)-, -NR^bC(=O)NR^c-, -NR^bC(=S)NR^c-, -S(=O)-, -S(=O)₂-, -S(=O)NR^a-, -C(=O)-, -C(=S)-, -C(=O)O-, -OC(=O)NR^a-, -OC(=S)NR^a-, -NR^a-, -NR^bS(=O)NR^c-, and NR^bS(=O)₂NR^o-;

each R⁴ and R⁶ is independently selected from halogen, cyano, nitro, hydroxyl, C₁₋₄ alkoxy, C₁₋₄ haloalkoxy, C₁₋₄ alkylthio, C₁₋₄ alkylsulfinyl, C₁₋₄ alkylsulfonyl, amino, C₁₋₄ alkylamino, di-C₁₋₄ alkylamino, C₁₋₄ alkylcarbonyl, carboxy, C₁₋₄ alkoxy-carbonyl, C₁₋₄ alkylcarbonylamino, di-C₁₋₄ alkylcarbonylamino, C₁₋₄ alkoxy-carbonylamino, C₁₋₄ alkoxy-carbonyl-(C₁₋₄ alkyl)amino, carbamyl, C₁₋₄ alkylcarbamyl, and di-C₁₋₄ alkylcarbamyl;

each R⁵ is independently selected from halogen, cyano, cyanate, isothiocyanate, nitro, hydroxyl, C₁₋₄ alkyl, C₂₋₄ alkenyl, C₂₋₄ alkynyl, C₁₋₄ alkoxy, C₁₋₄ haloalkoxy, C₁₋₄ alkylthio, C₁₋₄ alkylsulfinyl, C₁₋₄ alkylsulfonyl, amino, C₁₋₄ alkylamino, di-C₁₋₄ alkylamino, C₁₋₄ alkylcarbonyl, carboxy, C₁₋₄ alkoxy-carbonyl, C₁₋₄ alkylcarbonylamino, di-C₁₋₄ alkylcarbonylamino, C₁₋₄ alkoxy-carbonylamino, C₁₋₄ alkoxy-carbonyl-(C₁₋₄ alkyl)amino, carbamyl, C₁₋₄ alkylcarbamyl, and di-C₁₋₄ alkylcarbamyl;

each R⁷ is independently selected from halogen, cyano, nitro, hydroxyl, C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₃₋₇ cycloalkyl, C₃₋₇ cycloalkyl-C₁₋₄ alkyl, C₂₋₇ heterocycloalkyl, C₂₋₇ heterocycloalkyl-C₁₋₄ alkyl, phenyl, phenyl-C₁₋₄ alkyl, C₁₋₇ heteroaryl, C₁₋₇ heteroaryl-C₁₋₄ alkyl, -OR^o, -SR^o, -S(=O)R^p, -S(=O)₂R^p, -S(=O)NR^sR^t, -C(=O)R^p, -C(=O)OR^p, -C(=O)NR^sR^t, -OC(=O)R^p, -OC(=O)NR^sR^t, -NR^sR^t, -NR^qC(=O)R^r, -NR^qC(=O)OR^r, -NR^qC(=O)NR^r, -NR^qS(=O)₂R^r, and -NR^pS(=O)₂NR^sR^t; wherein said C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl are each optionally substituted by 1, 2, 3, or 4 independently

selected R' groups; and wherein said C₃₋₇ cycloalkyl, C₃₋₇ cycloalkyl-C₁₋₄ alkyl, C₂₋₇ heterocycloalkyl, C₂₋₇ heterocycloalkyl-C₁₋₄ alkyl, phenyl, phenyl-C₁₋₄ alkyl, C₁₋₇ heteroaryl, C₁₋₇ heteroaryl-C₁₋₄ alkyl are each optionally substituted by 1, 2, 3, or 4 independently selected R'' groups;

each R^a, R^b, and R^c is independently selected from H, C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₃₋₇ cycloalkyl, C₃₋₇ cycloalkyl-C₁₋₄ alkyl, C₂₋₇ heterocycloalkyl, C₂₋₇ heterocycloalkyl-C₁₋₄ alkyl, phenyl, phenyl-C₁₋₄ alkyl, C₁₋₇ heteroaryl, C₁₋₇ heteroaryl-C₁₋₄ alkyl; wherein said C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl are each optionally substituted by 1, 2, 3, or 4 independently selected R^w groups; and wherein said C₃₋₇ cycloalkyl, C₃₋₇ cycloalkyl-C₁₋₄ alkyl, C₂₋₇ heterocycloalkyl, C₂₋₇ heterocycloalkyl-C₁₋₄ alkyl, phenyl, phenyl-C₁₋₄ alkyl, C₁₋₇ heteroaryl, C₁₋₇ heteroaryl-C₁₋₄ alkyl are each optionally substituted by 1, 2, 3, or 4 independently selected R^x groups;

each R^o, R^p, R^q, R^r, R^s and R^t is independently selected from H, C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₃₋₇ cycloalkyl, C₃₋₇ cycloalkyl-C₁₋₄ alkyl, C₂₋₇ heterocycloalkyl, C₂₋₇ heterocycloalkyl-C₁₋₄ alkyl, phenyl, phenyl-C₁₋₄ alkyl, C₁₋₇ heteroaryl, C₁₋₇ heteroaryl-C₁₋₄ alkyl; wherein said C₁₋₆ alkyl, C₁₋₆ haloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl are each optionally substituted by 1, 2, 3, or 4 independently selected R^y groups; and wherein said C₃₋₇ cycloalkyl, C₃₋₇ cycloalkyl-C₁₋₄ alkyl, C₂₋₇ heterocycloalkyl, C₂₋₇ heterocycloalkyl-C₁₋₄ alkyl, phenyl, phenyl-C₁₋₄ alkyl, C₁₋₇ heteroaryl, C₁₋₇ heteroaryl-C₁₋₄ alkyl are each optionally substituted by 1, 2, 3, or 4 independently selected R^z groups;

each R', R^w and R^y is independently selected from hydroxyl, cyano, nitro, C₁₋₄ alkoxy, C₁₋₄ haloalkoxy, amino, C₁₋₄ alkylamino, and di-C₁₋₄ alkylamino; and

each R'', R^x, and R^z is independently selected from hydroxyl, halogen, cyano, nitro, C₁₋₄ alkyl, C₁₋₄ haloalkyl, C₁₋₄ alkoxy, C₁₋₄ haloalkoxy, amino, C₁₋₄ alkylamino, and di-C₁₋₄ alkylamino;

provided that the valency of each atom in the optionally substituted moieties is not exceeded.

Suitably, the functional variants/derivatives of the above formula have an affinity for an antibody of the present invention which is comparable to or greater than that of DOTAM, and have a binding strength for Pb which is comparable to or greater than that of DOTAM

(“affinity” being as measured by the dissociation constant, as described above). For example, the dissociation constant of the functional/variant derivative with the antibody of the present invention or/Pb may be 1.1 times or less, 1.2 times or less, 1.3 times or less, 1.4 times or less, 1.5 times or less, or 2 times or less than the dissociation constant of DOTAM with the same antibody/Pb.

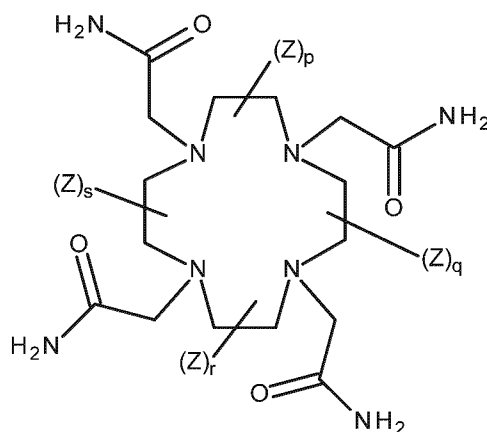
Each R^N may be H, C_{1-6} alkyl, or C_{1-6} haloalkyl; preferably H, C_{1-4} alkyl, or C_{1-4} haloalkyl. Most preferably, each R^N is H.

For DOTAM variants, it is preferred that 1, 2, 3 or most preferably each L^2 is C_2 alkylene. Advantageously, the C_2 alkylene variants of DOTAM can have particularly high affinity for Pb. The optional substituents for L^2 may be R^1 , C_{1-4} alkyl, or C_{1-4} haloalkyl. Suitably, the optional substituents for L^2 may be C_{1-4} alkyl or C_{1-4} haloalkyl.

Optionally, each L^2 may be unsubstituted C_2 alkylene $-CH_2CH_2-$.

Each L^1 is preferably C_{1-4} alkylene, more preferably C_1 alkylene such as $-CH_2-$.

The functional variant/derivative of DOTAM may be a compound of the following formula:



wherein each Z is independently R^1 as defined above; p, q, r, and s are 0, 1 or 2; and $p+q+r+s$ is 1 or greater. Preferably, p, q, r, and s are 0 or 1 and/or $p+q+r+s$ is 1. For example, the compound may have $p+q+r+s = 1$, where Z is *p*-SCN-benzyl moiety – such a compound is commercially available from Macrocylics, Inc. (Plano, Texas).

Radionuclides useful in the invention may include radioisotopes of metals, such as of lead (Pb), lutetium (Lu), or yttrium (Y).

Radionuclides particularly useful in imaging applications may be radionuclides that are gamma emitters. For instance, they may be selected from ^{203}Pb or ^{205}Bi .

Radionuclides particularly useful in therapeutic applications be radionuclides that are alpha or beta emitters. For instance, they may be selected from ^{212}Pb , ^{212}Bi , ^{213}Bi , ^{90}Y , ^{177}Lu , ^{225}Ac , ^{211}At , ^{227}Th , ^{223}Ra .

In some embodiments, it may be preferred that DOTAM (or salts or functional variants thereof) is chelated with Pb or Bi such as one of the Pb or Bi radioisotopes listed above. In other embodiments, it may be preferred that DOTA (or salts or functional variants thereof) is chelated with Lu or Y such as one of the Lu or Y radioisotopes listed above.

In some embodiments, methods and uses may comprise combined methods of therapy and imaging, which make use of a mixture of radioisotopes, e.g., a radioisotope suitable for therapy and a radioisotope suitable for imaging. For instance, these may be different radioisotopes of the same metal, chelated by the same chelator. In one embodiment, the method may comprise administering ^{203}Pb -DOTAM and ^{212}Pb -DOTAM as a mixture. In another embodiment, the method may comprise a first cycle of dosimetry using a gamma emitter such as ^{203}Pb or ^{205}Bi followed by one or more rounds of treatment using an alpha or beta emitter such as ^{212}Pb , ^{212}Bi , ^{213}Bi , ^{90}Y , ^{177}Lu , ^{225}Ac , ^{211}At , ^{227}Th , or ^{223}Ra . Such methods are described further below.

In some embodiments, the functional binding site formed by association of the first and the second antibody may bind to a Pb-DOTAM chelate.

In some embodiments, the functional binding site formed by association of the first and the second antibody may specifically bind to the radiolabelled compound. In some embodiments, it may bind to the radiolabelled compound, such as the Pb-DOTAM chelate, with a dissociation constant (K_D) to Pb-DOTAM and/or the target of $\leq 1\mu\text{M}$, $\leq 100\text{ nM}$, $\leq 10\text{ nM}$, $\leq 1\text{ nM}$, $\leq 0.1\text{ nM}$, $\leq 0.01\text{ nM}$, or $\leq 0.001\text{ nM}$ (e.g. 10^{-7}M or less, e.g. from 10^{-7} to 10^{-13} , 10^{-8} M or less, e.g. from 10^{-8} M to 10^{-13} M , e.g., from 10^{-9} M to 10^{-13} M). In some embodiments it may be preferred that it binds with a K_D value of the binding affinity of 100pM , 50pM , 20pM , 10pM , 5pM , 1pM or less, e.g., 0.9pM or less, 0.8pM or less, 0.7pM or less, 0.6pM or less or 0.5pM or less. For instance, the functional binding site may bind the metal chelate with a K_D of about 1pM - 1nM , e.g., about 1 - 10 pM , 1 - 100pM , 5 - 50 pM , 100 - 500 pM or 500pM - 1 nM .

D. Exemplary antigen binding sites for DOTA

In one particular embodiment of the invention, the first and second antibody associate to form a functional binding site for DOTA (or a functional derivative or variant thereof),

e.g., DOTA chelated with Lu or Y (e.g., ^{177}Lu or ^{90}Y). For instance, the functional binding site may bind the radiolabelled compound with a K_d of about 1pM-1nM, e.g., about 1-10 pM, 1-100pM, 5-50 pM, 100-500 pM or 500pM-1 nM.

C825 is a known scFv with high affinity for DOTA-Bn (S-2-(4-aminobenzyl)-1,4,7,10-tetraazacyclododecane tetraacetic acid) complexed with radiometals such as ^{177}Lu and ^{90}Y (see for instance Cheal et al 2018, Theranostics 2018, and WO2010099536, incorporated herein by reference). The CDR sequences and the VL and VH sequences of C825 are provided herein. In one embodiment, the heavy chain variable region forming part of the antigen binding site for the radiolabelled compound may comprise at least one, two or
10 all three CDRs selected from (a) CDR-H1 comprising the amino acid sequence of 35; (b) CDR-H2 comprising the amino acid sequence of 36; (c) CDR-H3 comprising the amino acid sequence of 37. In an alternative embodiment, CDR-H1 may have the sequence GFSLT DYGVH (SEQ ID NO.: 148). The light chain variable region forming part of the binding site for the radiolabelled compound may comprise at least one, two or all three CDRs selected from (d) CDR-L1 comprising the amino acid sequence of 38; (e) CDR-L2 comprising the amino acid sequence of 39; and (f) CDR-L3 comprising the amino acid sequence of 40.

In another embodiment, the heavy chain variable domain forming part of the functional antigen binding site for the radiolabelled compound (on the first antibody)
20 comprises the amino acid sequence of SEQ ID NO: 41, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 41. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but a binding site comprising that sequence retains the ability to bind to DOTA complexed with Lu or Y, preferably with an affinity as described herein. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:41. In certain embodiments, substitutions, insertions, or deletions occur in regions outside the CDRs (i.e., in the FRs). Optionally, the antibody comprises the VH sequence in SEQ ID NO:41, including
30 post-translational modifications of that sequence. In a particular embodiment, the VH comprises one, two or three CDRs selected from: (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:35 or the sequence GFSLT DYGVH (SEQ ID NO.: 148), (b) CDR-

H2 comprising the amino acid sequence of SEQ ID NO:36, and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:37.

Optionally, the light chain variable domain forming part of the functional antigen binding site for the radiolabelled compound (on the second antibody) comprises an amino acid sequence of SEQ ID NO: 42 or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 42. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but a binding site comprising that sequence retains the ability to bind to DOTA complexed with Lu or Y, preferably with an affinity as described herein. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO: 42. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the CDRs (i.e., in the FRs). Optionally, the antibody comprises the VL sequence in SEQ ID NO:42, including post-translational modifications of that sequence. In a particular embodiment, the VL comprises one, two or three CDRs selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:38; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:39; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:40.

Embodiments concerned with the heavy chain variable region and the light chain variable region are explicitly contemplated in combination. Thus, the functional antigen binding site may be formed from a heavy chain variable region as defined above and a light chain variable region as defined above, on the first and second antibody respectively.

In any of the above embodiments, the light and heavy chain variable regions forming the binding site for the DOTA complex may be humanized. In one embodiment, the light and heavy chain variable region comprise CDRs as in any of the above embodiments, and further comprise an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework.

In some embodiments, the heavy chain variable domain may be extended by one or more C-terminal residues such as one or more C-terminal alanine residues, or one or more residues from the N-terminus of the CH1 domain, as discussed further below.

E. Exemplary antigen binding sites for DOTAM

In another particular embodiment of the invention, the first and second antibody associate to form a functional antigen binding site for a Pb-DOTAM chelate (Pb-DOTAM).

Exemplary antigen binding sites are described in WO2019/201959, which is incorporated herein by reference in its entirety.

In certain embodiments, the functional antigen-binding site that binds to Pb-DOTAM may have one or more of the following properties:

- Binds specifically to Pb-DOTAM and to Bi-DOTAM;
- Is selective for Pb-DOTAM (and optionally Bi-DOTAM) as compared to other chelated metals, such as Cu-DOTAM;
- Binds to Pb-DOTAM with a very high affinity;
- Binds to the same epitope on Pb-DOTAM as antibodies described in

10 WO2019/201959, e.g., PRIT-0213 or PRIT-0214 and/or has the same contact residues as said antibodies.

Radioisotopes of Pb are useful in methods of diagnosis and therapy. Particular radioisotopes of lead which may be of use in the present invention include ^{212}Pb and ^{203}Pb .

Radionuclides which are α -particle emitters have the potential for more specific tumour cell killing with less damage to the surrounding tissue than β -emitters because of the combination of short path length and high linear energy transfer. ^{212}Bi is an α -particle emitter but its short half-life hampers its direct use. ^{212}Pb is the parental radionuclide of ^{212}Bi and can serve as an *in vivo* generator of ^{212}Bi , thereby effectively overcoming the short half-life of ^{212}Bi (Yong and Brechbiel, Dalton Trans. 2001 June 21; 40(23)6068-6076).

20 ^{203}Pb is useful as an imaging isotope. Thus, an antibody bound to ^{203}Pb -DOTAM may have utility in radioimmunoimaging (RII).

Generally, radiometals are used in chelated form. In certain aspects of the present invention, DOTAM is used as the chelating agent. DOTAM is a stable chelator of Pb(II) (Yong and Brechbiel, Dalton Trans. 2001 June 21; 40(23)6068-6076; Chappell et al Nuclear Medicine and Biology, Vol. 27, pp. 93-100, 2000). Thus, DOTAM is particularly useful in conjunction with isotopes of lead as discussed above, such as ^{212}Pb and ^{203}Pb .

In some embodiments, it may be preferred that the antibodies bind Pb-DOTAM with a K_d value of the binding affinity of 100pM, 50pM, 20pM, 10pM, 5pM, 1pM or less, e.g., 0.9pM or less, 0.8pM or less, 0.7pM or less, 0.6pM or less or 0.5pM or less. For instance, the
30 functional binding site may bind the radiolabelled compound with a K_d of about 1pM-1nM, e.g., about 1-10 pM, 1-100pM, 5-50 pM, 100-500 pM or 500pM-1 nM.

In certain embodiment, the antibodies additionally bind to Bi chelated by DOTAM. In some embodiments, it may be preferred that the antibodies bind Bi-DOTAM (i.e., a chelate

comprising DOTAM complexed with bismuth, also termed herein a “Bi-DOTAM chelate”) with a K_d value of the binding affinity of 1nM, 500pM, 200pM, 100pM, 50pM, 10pM or less, e.g., 9pM, 8pM, 7pM, 6pM, 5pM or less. For instance, the functional binding site may bind a metal chelate with a K_d of about 1pM-1nM, e.g., about 1-10 pM, 1-100pM, 5-50 pM, 100-500 pM or 500pM-1 nM.

In some embodiments, the antibodies may bind to Bi-DOTAM and to Pb-DOTAM with a similar affinity. For instance, it may be preferred that the ratio of affinity, e.g., the ratio of K_d values, for Bi-DOTAM/Pb-DOTAM is in the range of 0.1-10, for example 1-10.

In one embodiment, the heavy chain variable region forming part of the antigen binding site for Pb-DOTAM may comprise at least one, two or all three CDRs selected from (a) CDR-H1 comprising the amino acid sequence of GFSLSSTYSMS (SEQ ID NO:1); (b) CDR-H2 comprising the amino acid sequence of FIGSRGDTYYASWAKG (SEQ ID NO:2); (c) CDR-H3 comprising the amino acid sequence of ERDPYGGGAYPPHL (SEQ ID NO:3). The light chain variable region forming part of the binding site for Pb-DOTAM may comprise at least one, two or all three CDRs selected from (d) CDR-L1 comprising the amino acid sequence of QSSHSVYSDNDLA (SEQ ID NO:4); (e) CDR-L2 comprising the amino acid sequence of QASKLAS (SEQ ID NO:5); and (f) CDR-L3 comprising the amino acid sequence of LGGYDDSDTYG (SEQ ID NO:6).

In some embodiments, the antibodies may comprise one or more of CDR-H1, CDR-H2 and/or CDR-H3, or one or more of CDR-L1, CDR-L2 and/or CDR-L3, having substitutions as compared to the amino acid sequences of SEQ ID NOs: 1-6, respectively, e.g., 1, 2 or 3 substitutions.

In some embodiments, antibodies may share the same contact residues as the described herein: e.g., these residues may be invariant. These residues may include the following:

- a) in heavy chain CDR2: Phe50, Asp56 and/or Tyr58, and optionally also Gly52 and/or Arg 54;
 - b) in heavy chain CDR3: Glu95, Arg96, Asp97, Pro98, Tyr99, Ala100C and/or Tyr100D and optionally also Pro100E;
 - c) in light chain CDR1: Tyr28 and/or Asp32;
 - d) in light chain CDR3: Gly91, Tyr92, Asp93, Thr95c and/or Tyr96;
 - e) in light chain CDR2: optionally Gln50;
- all numbered according to Kabat.

For example, in some embodiments, CDR-H2 may comprise the amino acid sequence FIGSRGDTYYASWAKG (SEQ ID NO:2), or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 2, wherein these substitutions do not include Phe50, Asp56 and/or Tyr58, and optionally also do not include Gly52 and/or Arg 54, all numbered according to Kabat.

In some embodiments, CDR-H2 may be substituted at one or more positions as shown below. Here and in the substitution tables that follow, substitutions are based on the germline residues (underlined) or by amino acids which theoretically sterically fit and also occur in the crystallized repertoire at the site. In some embodiments, the residues as mentioned above may be fixed and other residues may be substituted according to the table below: in other
 10 embodiments, substitutions of any residue may be made according to the table below.

WolfGuy	Kabat	AA	Substitution
251	50	F	<u>Y</u> , <u>H</u>
252	51	I	
253	52	G	
254	53	S	<u>A</u> , <u>G</u> , <u>T</u> , I, N
288	54	R	<u>A</u> , <u>D</u> , <u>G</u> , <u>N</u> , <u>S</u> , T, F, Y
289	55	G	<u>D</u> , <u>S</u> , Y, T, A, N, R, V
290	56	D	
291	57	T	<u>K</u> , <u>L</u> , A, P, S
292	58	Y	<u>E</u> , W, H
293	59	Y	<u>N</u> , F, H, L, S
294	60	A	<u>G</u> , <u>N</u> , <u>S</u> , <u>T</u>
295	61	S	<u>A</u> , <u>G</u> , <u>N</u> , <u>Q</u> , <u>T</u>
296	62	W	<u>K</u> , <u>P</u> , <u>S</u> , A, T, D, N, R, Q
297	63	A	<u>E</u> , <u>L</u> , <u>V</u> , M, I
298	64	K	<u>N</u> , <u>Q</u> , R, E
299	65	G	<u>S</u> , <u>T</u> , D, N, A

Optionally, CDR-H3 may comprise the amino acid sequence ERDPYGGGAYPPHL (SEQ ID NO:3), or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 3, wherein these substitutions do not include Glu95, Arg96, Asp97, Pro98, and optionally also

do not include Ala100C, Tyr100D, and/or Pro100E and/or optionally also do not include Tyr99. For instance, in some embodiments the substitutions do not include Glu95, Arg96, Asp97, Pro98, Tyr99 Ala100C and Tyr100D.

In certain embodiments, CDR-H3 may be substituted at one or more positions as shown below. In some embodiments, the residues as mentioned above may be fixed and other residues may be substituted according to the table below: in other embodiments, substitutions of any residue may be made according to the table below.

WolfGuy	Kabat	AA	Substitution
351	95	E	
352	96	R	K, E
353	97	D	
354	98	P	
355	99	Y	F, G, S, T, D
356	100	G	
392	100A	G	
393	100B	G	
394	100C	A	S, T
395	100D	Y	F
396	100E	P	
397	100F	P	
398	101	H	A, T, V, D
399	102	L	Y, V, I, H, F

10

Optionally, CDR-L1 may comprise the amino acid sequence QSSHSVYSDNDLA (SEQ ID NO:4) or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 4, wherein these substitutions do not include Tyr28 and/or Asp32 (Kabat numbering).

In certain embodiments, CDR-L1 may be substituted at one or more positions as shown below. Again, in some embodiments, the residues as mentioned above may be fixed and other residues may be substituted according to the table below: in other embodiments, substitutions of any residue may be made according to the table below.

WolfGuy	Kabat	AA	Substitution
551	24	Q	<u>R</u> , <u>K</u>
552	25	S	<u>A</u> , <u>G</u>
554	26	S	<u>T</u>
555	27	H	<u>Q</u> , <u>S</u> , R, K
556	27A	S	Q
557	27B	V	<u>I</u> , <u>D</u> , <u>N</u>
561	28	Y	F
562	29	S	T, <u>V</u>
571	30	D	<u>R</u> , <u>S</u> , <u>N</u> , <u>G</u>
572	31	N	K
597	32	D	
598	33	L	<u>I</u> , <u>V</u> , <u>M</u>
599	34	A	<u>S</u>

Optionally, CDR-L3 may comprise the amino acid sequence LGGYDDESDTYG (SEQ ID NO:6) or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 6, wherein these substitutions do not include Gly91, Tyr92, Asp93, Thr95c and/or Tyr96 (Kabat).

In certain embodiments, CDR-L3 may be substituted at the following positions as shown below. (Since most residues are solvent exposed and without antigen contacts, many substitutions are conceivable). Again, in some embodiments, the residues as mentioned above may be fixed and other residues may be substituted according to the table below: in
10 other embodiments, substitutions of any residue may be made according to the table below.

WolfGuy	Kabat	AA	Substitution
751	89	L	A, V, Q
752	90	G	A
753	91	G	
754	92	Y	A, D, E, F, G, H, I, K, L, N, Q, R, S, T, V

755	93	D	A, E, F, G, H, I, K, L, M, N, Q, R, S, T, V, W, Y
756	94	D	A, E, F, G, H, I, K, L, M, N, Q, R, S, T, V, W, Y
794	95	E	A, D, F, G, H, I, K, L, M, N, Q, R, S, T, V, W, Y
795	95A	S	A, F, G, H, I, K, L, M, N, Q, R, T, V, W, Y
796	95B	D	A, E, F, G, H, I, L, M, N, Q, S, T, V, W, Y
797	95C	T	S
798	96	Y	F, H, R
799	97	G	A, E, I, K, L, M, N, Q, S, T, V

The antibody may further comprise CDR-H1 or CDR-L2, optionally having the sequence of SEQ ID NO: 1 or SEQ ID NO: 5 respectively, or a variant thereof having at least 1, 2 or 3 substitutions relative thereto, optionally conservative substitutions.

Thus, the heavy chain variable domain forming part of the antigen binding site for Pb-DOTAM may comprise at least:

- a) heavy chain CDR2 comprising the amino acid sequence FIGSRGDTYYASWAKG (SEQ ID NO:2), or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 2, wherein these substitutions do not include Phe50, Asp56 and/or Tyr58, and optionally also do not include Gly52 and/or Arg54;
- b) heavy chain CDR3 comprising the amino acid sequence ERDPYGGGAYPPHL (SEQ ID NO:3), or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 3, wherein these substitutions do not include Glu95, Arg96, Asp97, Pro98, and optionally also do not include Ala100C, Tyr100D, and/or Pro100E and/or optionally also do not include Tyr99.

In some embodiments, the heavy chain variable domain additionally includes a heavy chain CDR1 which is optionally:

c) a heavy chain CDR1 comprising the amino acid sequence GFSLSTYSMS (SEQ ID NO:1) or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 1.

In another embodiment, the light chain variable domain forming part of the antigen binding site for Pb-DOTAM comprises at least:

d) light chain CDR1 comprising the amino acid sequence QSSHSVYSDNDLA (SEQ ID NO:4) or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 4, wherein these substitutions do not include Tyr28 and Asp32;

10 e) light chain CDR3 comprising the amino acid sequence LGGYDDSDTYG (SEQ ID NO:6) or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 6, wherein these substitutions do not include Gly91, Tyr92, Asp93, Thr95c and Tyr96.

In some embodiments, the light chain variable domain additionally includes a light chain CDR2 which is optionally:

f) a light chain CDR2 comprising the amino acid sequence QASKLAS (SEQ ID NO: 5) or a variant thereof having at least 1, 2 or 3 substitutions in SEQ ID NO: 5, optionally not including Gln50.

20 In any embodiments of the present invention which include variants of a sequence comprising the CDRs as set out above (e.g., of a variable domain), the protein may be invariant in one or more of the CDR residues as set out above.

Optionally, the heavy chain variable domain forming part of the functional antigen binding site for Pb-DOTAM (on the first antibody) comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 7 and SEQ ID NO 9, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 7 or SEQ ID NO: 9. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but a binding site comprising that sequence retains the ability to bind to Pb-DOTAM, preferably with an affinity as described herein. The VH sequence may retain the
30 invariant residues as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO: 7 or SEQ ID NO 9. In certain embodiments, substitutions, insertions, or deletions occur in regions outside the CDRs (i.e., in the FRs). Optionally, the antibody comprises the VH sequence in SEQ ID NO:7 or SEQ ID

NO: 9, including post-translational modifications of that sequence. In a particular embodiment, the VH comprises one, two or three CDRs selected from: (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:1, (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:2, and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:3.

Optionally, the light chain variable domain forming part of the functional antigen binding site for Pb-DOTAM (on the second antibody) comprises an amino acid sequence of SEQ ID NO: 8, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 8. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but an anti-Pb-DOTAM binding site comprising that sequence retains the ability to bind to Pb-DOTAM, preferably with an affinity as described herein. The VL sequence may retain the invariant residues as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:8. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the CDRs (i.e., in the FRs). Optionally, the anti-Pb-DOTAM antibody comprises the VL sequence in SEQ ID NO:8, including post-translational modifications of that sequence. In a particular embodiment, the VL comprises one, two or three CDRs selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:4; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:5; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:6.

Embodiments concerned with the heavy chain variable region and the light chain variable region are explicitly contemplated in combination. Thus, the functional antigen binding site for Pb-DOTAM may be formed from a heavy chain variable region as defined above and a light chain variable region as defined above, on the first and second antibody respectively.

Optionally, the antigen binding site specific for the Pb-DOTAM chelate may be formed from a heavy chain variable domain comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 7 or SEQ ID NO: 9, or a variant thereof as defined above, and a light chain variable domain comprising an amino acid sequence of SEQ ID NO: 8, or a variant thereof as defined above. For example, the antigen binding site specific for the Pb-DOTAM chelate may comprise a heavy chain variable domain comprising the amino acid

sequence of SEQ ID NO: 7 or a variant thereof, and a light chain variable domain comprising the amino acid sequence of SEQ ID NO: 8 or a variant thereof, including post-translational modifications of those sequences. In another embodiment, it may comprise a heavy chain variable domain comprising the amino acid sequence of SEQ ID NO: 9 or a variant thereof and a light chain variable domain comprising the amino acid sequence of SEQ ID NO: 8 or a variant thereof, including post-translational modifications of those sequences.

In any of the above embodiments, the light and heavy chain variable regions forming the anti-Pb-DOTAM binding site may be humanized. In one embodiment, the light and heavy chain variable region comprise CDRs as in any of the above embodiments, and further
10 comprise an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework. In another embodiment, the light and/or heavy chain variable regions comprise CDRs as in any of the above embodiments, and further comprises framework regions derived from vk 1 39 and/or vh 2 26. For vk 1 39, in some embodiments there may be no back mutations. For vh 2 26, the germline Ala49 residue may be backmutated to Gly49.

F. Exemplary antigen binding sites for CEA

In another particular embodiment of the present invention, which may be combined with the embodiments discussed above, the target antigen bound by the first and/or second
20 antibody may be CEA (carcinoembryonic antigen). Antibodies that have been raised against CEA include T84.66 and humanized and chimeric versions thereof, such as T84.66-LCHA as described in WO2016/075278 A1 and/or WO2017/055389, CH1A1a, an anti-CEA antibody as described in WO2012/117002 and WO2014/131712, and CEA hMN-14 or labetuzimab (e.g., as described in US 6 676 924 and US 5 874 540). Another exemplary antibody against CEA is A5B7 (e.g., as described in M.J. Banfield et al, Proteins 1997, 29(2), 161-171), or a humanized antibody derived from murine A5B7 as described in WO 92/01059 and WO 2007/071422. See also co-pending application PCT/EP2020/067582. An example of a humanized version of A5B7 is A5H1EL1(G54A). A further exemplary antibody against CEA is MFE23 and the humanized versions thereof described in US 7 626 011 and/or co-
30 pending application PCT/EP2020/067582. A still further example of an anti-CEA antibody is 28A9. Any of these or antigen binding fragments thereof may be used to form a CEA-binding moiety in the present invention.

Optionally, the antigen-binding site which binds to CEA may bind with a K_d value of 1nM or less, 500pM or less, 200pM or less, or 100pM or less for monovalent binding.

In some embodiments, the first and/or second antibody may bind to the CH1A1a epitope, the A5B7 epitope, the MFE23 epitope, the T84.66 epitope or the 28A9 epitope of CEA.

In some embodiments, at least one of the first and second antibodies binds to a CEA epitope which is not present on soluble CEA (sCEA). Soluble CEA is a part of the CEA molecule which is cleaved by GPI phospholipase and released into the blood. An example of an epitope not found on soluble CEA is the CH1A1A epitope. Optionally, one of the first and/or second antibody binds to an epitope which is not present on soluble CEA, and the other binds to an epitope which is present on soluble CEA.

The epitope for CH1A1a and its parent murine antibody PR1A3 is described in WO2012/117002A1 and Durbin H. et al., Proc. Natl. Acad. Sci. USA, 91:4313-4317, 1994. An antibody which binds to the CH1A1a epitope binds to a conformational epitope within the B3 domain and the GPI anchor of the CEA molecule. In one aspect, the antibody binds to the same epitope as the CH1A1a antibody having the VH of SEQ ID NO: 25 and VL of SEQ ID NO 26 herein. The A5B7 epitope is described in co-pending application PCT/EP2020/067582. An antibody which binds to the A5B7 epitope binds to the A2 domain of CEA, i.e., to the domain comprising the amino acids of SEQ ID NO: 141:

20 PKPFITSNNSNPVEDEDAVALTCEPEIQNTTYLWWVNNQSLPVSRLQLSNDN
 RTLTLFSVTRNDVGP YECGIQNKLSVDHSDPVILN (SEQ ID NO: 141).

In one aspect, the antibody binds to the same epitope as the A5B7 antibody having the VH of SEQ ID NO: 49 and VL of SEQ ID NO: 50 herein.

In one aspect, the antibody binds to the same epitope as the T84.66 described in WO2016/075278. The antibody may bind to the same epitope as the antibody having the VH of SEQ ID NO: 17 and VL of SEQ ID NO:18 herein.

The MFE23 epitope is described in co-pending application PCT/EP2020/067582. An antibody which binds to the MFE23 epitope binds to the A1 domain of CEA, i.e., to the domain comprising the amino acids of SEQ ID NO: 142:

30 PKPSSNNSKPVEDKDAVAFTCEPETQDATYLWWVNNQSLPVSRLQLSNG
 NRTLTLFNVTRNDTAS YKCETQNPVSARRSDSVILN (SEQ ID NO: 142).

In one aspect, the antibody may bind to the same epitope as an antibody having the VH domain of SEQ ID NO: 127 and the VL domain of SEQ ID NO: 128 herein.

In some embodiments, the first and/or second antibody may bind to the same CEA-epitope as an antibody provided herein, e.g., P1AD8749, P1AD8592, P1AE4956, P1AE4957, P1AF0709, P1AF0298, P1AF0710 or P1AF0711.

In some embodiments, the first and the second antibody bind the same epitope of CEA as each other. Thus, for example, the first and the second antibody may both bind to the CH1A1a epitope, the A5B7 epitope, the MFE23 epitope, the T84.66 epitope or the 28A9 epitope.

In some embodiments, both the first and second antibody may have CEA binding sequences (i.e., CDRs and/or VH/VL domains) from CH1A1A; or, the first and the second antibody may both have CEA binding sequences from A5B7 or a humanized version thereof; or, the first and the second antibody may both have CEA binding sequences from T84.66 or a humanized version thereof; or the first and the second antibody may both have CEA binding sequences from MFE23 or a humanized version thereof; or the first and second antibody may both have CEA binding sequences from 28A9 or a humanized version thereof. Exemplary sequences are disclosed herein.

In other embodiments, the first and the second antibodies bind to different epitopes of CEA. Thus, for example, i) one antibody may bind the CH1A1A epitope and the other may bind the A5B7 epitope, the T84.66 epitope, the MFE23 epitope or the 28A9 epitope; ii) one antibody may bind the A5B7 epitope and the other may bind the CH1A1A epitope, T84.66 epitope, MFE23 epitope or 28A9 epitope; iii) one antibody may bind the MFE23 epitope and the other may bind the CH1A1A epitope, A5B7 epitope, T84.66 epitope or 28A9 epitope; iv) one antibody may bind the T84.66 epitope and the other may bind the CH1A1A epitope, A5B7 epitope, MFE23 epitope or 28A9 epitope; or v) one antibody may bind the 28A9 epitope and the other may bind the CH1A1a epitope, the A5B7 epitope, the MFE23 epitope, or the T84.66 epitope.

In some embodiments, i) one antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from CH1A1A and the other may have CEA binding sequences from A5B7 or a humanized version thereof, from T84.66 or a humanized version thereof, from MFE23 or a humanized version thereof, or from 28A9 or a humanized version thereof; ii) one antibody may have CEA binding sequences from A5B7 or a humanized version thereof and the other may have CEA binding sequences from CH1A1A, from T84.66 or a humanized version thereof, from MFE23 or a humanized version thereof, or from 28A9 or a humanized version thereof; iii) one antibody may have CEA binding sequences from MFE23 or a

humanized version thereof and the other may have CEA binding sequences from CH1A1A, from A5B7 or a humanized version thereof, from T84.66 or a humanized version thereof, or from 28A9 or a humanized version thereof; iv) one antibody may have CEA binding sequences from T84.66 or a humanized version thereof and the other may have CEA binding sequences from CH1A1A, from A5B7 or a humanized version thereof, from MFE23 or a humanized version thereof, or from 28A9 or a humanized version thereof; v) one antibody may have CEA-binding sequences from 28A9 or a humanized version thereof and the other may have CEA binding sequences from CH1A1A, from A5B7 or a humanized version thereof, from T84.66 or a humanized version thereof, or from MFE23 or a humanized version thereof.

10 In one particular embodiment, one antibody may bind the CH1A1A epitope and the other may bind the A5B7 epitope. The first antibody may have CEA binding sequences from the antibody CH1A1A and the second antibody may have CEA binding sequences from A5B7 (including a humanized version thereof); or, the first antibody may have CEA binding sequences from the antibody A5B7 (including a humanized version thereof) and the second antibody may have CEA binding sequences from CH1A1A.

In another particular embodiment, one antibody may bind the CH1A1A epitope and the other may bind the T84.66 epitope. The first antibody may have CEA binding sequences from the antibody CH1A1A and the second antibody may have CEA binding sequences from T84.66 (including a humanized version thereof); or, the first antibody may have CEA binding
20 sequences from the antibody T84.66 (including a humanized version thereof) and the second antibody may have CEA binding sequences from CH1A1A. In some embodiments, a first antibody may bind the T84.66 epitope and/or have an antigen binding site as described in (i) below, and the second antibody may bind the CH1A1A epitope and/or have an antigen binding site as described in (ii) below.

Exemplary CEA-binding sequences i)-v) are disclosed below. These provide examples of CEA-binding sequences from i) T84.66, ii) CH1A1A, iii) A5B7, iv) 28A9 and v) MFE23(or from humanized versions thereof).

i). In one embodiment, the antigen-binding site which binds to CEA may comprise at least one, two, three, four, five, or six CDRs selected from (a)CDR-H1 comprising the amino
30 acid sequence of SEQ ID NO:11; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:12; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:13; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:14; (e) CDR-L2 comprising the amino

acid sequence of SEQ ID NO:15; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:16.

Optionally, the antigen-binding site which binds to CEA may comprise at least one, at least two, or all three VH CDR sequences selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:11; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:12; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:13.

Optionally, the antigen-binding site which binds to CEA comprises at least one, at least two, or all three VL CDRs sequences selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:14; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:15; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:16.

Optionally, the antigen-binding site which binds to CEA comprises (a) a VH domain comprising at least one, at least two, or all three VH CDR sequences selected from (i) CDR-H1 comprising the amino acid sequence of SEQ ID NO:11, (ii) CDR-H2 comprising the amino acid sequence of SEQ ID NO:12, and (iii) CDR-H3 comprising an amino acid sequence selected from SEQ ID NO:13; and (b) a VL domain comprising at least one, at least two, or all three VL CDR sequences selected from (i) CDR-L1 comprising the amino acid sequence of SEQ ID NO:14, (ii) CDR-L2 comprising the amino acid sequence of SEQ ID NO:15, and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:16.

In another aspect, the antigen-binding site which binds to CEA comprises (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:11; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:12; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:13; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:14; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:15; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:16.

In any of the above embodiments, the multispecific antibody may be humanized. In one embodiment, the anti-CEA antigen binding site comprises CDRs as in any of the above embodiments, and further comprises an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework.

In another embodiment, the antigen-binding site which binds to CEA comprises a heavy chain variable domain (VH) sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:17. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative

substitutions), insertions, or deletions relative to the reference sequence, but the antigen binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:17. In certain embodiments, substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site which binds to CEA comprises the VH sequence in SEQ ID NO:17, including post-translational modifications of that sequence. In a particular embodiment, the VH comprises one, two or three CDRs selected from: (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:11, (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:12, and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:13.

In another embodiment, the antigen-binding site which binds to CEA comprises a light chain variable domain (VL) having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:18. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen-binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:18. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site for CEA comprises the VL sequence in SEQ ID NO:18, including post-translational modifications of that sequence. In a particular embodiment, the VL comprises one, two or three CDRs selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:14; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:15; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:16.

In another embodiment, the antigen-binding site which binds to CEA comprises a VH as in any of the embodiments provided above, and a VL as in any of the embodiments provided above. In one embodiment, the antibody comprises the VH and VL sequences in SEQ ID NO:17 and SEQ ID NO:18, respectively, including post-translational modifications of those sequences.

ii). In further particular embodiment, the antigen-binding site which binds to CEA may comprise at least one, two, three, four, five, or six CDRs selected from (a) CDR-H1

comprising the amino acid sequence of SEQ ID NO:19; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:20; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:21; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:22; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:23; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:24.

Optionally, the antigen-binding site which binds to CEA may comprise at least one, at least two, or all three VH CDR sequences selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:19; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:20; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:21.

10 Optionally, the antigen-binding site which binds to CEA comprises at least one, at least two, or all three VL CDRs sequences selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:22; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:23; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:24.

Optionally, the antigen-binding site which binds to CEA comprises (a) a VH domain comprising at least one, at least two, or all three VH CDR sequences selected from (i) CDR-H1 comprising the amino acid sequence of SEQ ID NO:19, (ii) CDR-H2 comprising the amino acid sequence of SEQ ID NO:20, and (iii) CDR-H3 comprising an amino acid sequence selected from SEQ ID NO:21; and (b) a VL domain comprising at least one, at least two, or all three VL CDR sequences selected from (i) CDR-L1 comprising the amino acid sequence of SEQ ID NO:22, (ii) CDR-L2 comprising the amino acid sequence of SEQ ID NO:23, and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:24.

20 In another aspect, the antigen-binding site which binds to CEA comprises (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:19; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:20; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:21; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:22; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:23; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:24.

30 In any of the above embodiments, the multispecific antibody may be humanized. In one embodiment, the anti-CEA antigen binding site comprises CDRs as in any of the above embodiments, and further comprises an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework.

In another embodiment, the antigen-binding site which binds to CEA comprises a heavy chain variable domain (VH) sequence having at least 90%, 91%, 92%, 93%, 94%,

95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:25. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:25. In certain embodiments, substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site which binds to CEA comprises the VH sequence in SEQ ID NO:25, including post-translational modifications of that sequence. In a particular embodiment, the VH comprises one, two or three CDRs selected from: (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:19, (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:20, and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:21.

In another embodiment, the antigen-binding site which binds to CEA comprises a light chain variable domain (VL) having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:26. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen-binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:26. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site for CEA comprises the VL sequence in SEQ ID NO:26, including post-translational modifications of that sequence. In a particular embodiment, the VL comprises one, two or three CDRs selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:22; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:23; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:24.

In another embodiment, the antigen-binding site which binds to CEA comprises a VH as in any of the embodiments provided above, and a VL as in any of the embodiments provided above. In one embodiment, the antibody comprises the VH and VL sequences in SEQ ID NO:25 and SEQ ID NO:26, respectively, including post-translational modifications of those sequences.

iii) In further particular embodiment, the antigen-binding site which binds to CEA may comprise at least one, two, three, four, five, or six CDRs selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:43; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:44; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:45; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:46; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:47; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:48. In some embodiments, CDR-H1 may have the sequence GFTFTDYMN (SEQ ID NO.: 151).

10 Optionally, the antigen-binding site which binds to CEA may comprise at least one, at least two, or all three VH CDR sequences selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:43; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:44; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:45. In some embodiments, CDR-H1 may have the sequence GFTFTDYMN (SEQ ID NO.: 151).

 Optionally, the antigen-binding site which binds to CEA comprises at least one, at least two, or all three VL CDRs sequences selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:46; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:47; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:48.

20 Optionally, the antigen-binding site which binds to CEA comprises (a) a VH domain comprising at least one, at least two, or all three VH CDR sequences selected from (i) CDR-H1 comprising the amino acid sequence of SEQ ID NO:43, (ii) CDR-H2 comprising the amino acid sequence of SEQ ID NO:44, and (iii) CDR-H3 comprising an amino acid sequence selected from SEQ ID NO:45; and (b) a VL domain comprising at least one, at least two, or all three VL CDR sequences selected from (i) CDR-L1 comprising the amino acid sequence of SEQ ID NO:46, (ii) CDR-L2 comprising the amino acid sequence of SEQ ID NO:47, and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:48. In some embodiments, CDR-H1 may have the sequence GFTFTDYMN (SEQ ID NO.: 151).

30 In another aspect, the antigen-binding site which binds to CEA comprises (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:43; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:44; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:45; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:46; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:47; and (f) CDR-L3 comprising

the amino acid sequence of SEQ ID NO:48. In some embodiments, CDR-H1 may have the sequence GFTFTDYMN (SEQ ID NO.: 151).

In any of the above embodiments, the multispecific antibody may be humanized. In one embodiment, the anti-CEA antigen binding site comprises CDRs as in any of the above embodiments, and further comprises an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework.

In another embodiment, the antigen-binding site which binds to CEA comprises a heavy chain variable domain (VH) sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:49. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:49. In certain embodiments, substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site which binds to CEA comprises the VH sequence in SEQ ID NO:49, including post-translational modifications of that sequence. In a particular embodiment, the VH comprises one, two or three CDRs selected from: (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:43 or the sequence GFTFTDYMN (SEQ ID NO.: 151), (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:44, and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:45.

In another embodiment, the antigen-binding site which binds to CEA comprises a light chain variable domain (VL) having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:50. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen-binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:50. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site for CEA comprises the VL sequence in SEQ ID NO:50, including post-

translational modifications of that sequence. In a particular embodiment, the VL comprises one, two or three CDRs selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:46; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:47; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:48.

In another embodiment, the antigen-binding site which binds to CEA comprises a VH as in any of the embodiments provided above, and a VL as in any of the embodiments provided above. In one embodiment, the antibody comprises the VH and VL sequences in SEQ ID NO:49 and SEQ ID NO:50, respectively, including post-translational modifications of those sequences.

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iv) In a still further particular embodiment, the antigen-binding site which binds to CEA may comprise at least one, two, three, four, five, or six CDRs selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:59; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:60; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:61; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:62; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:63; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:64.

Optionally, the antigen-binding site which binds to CEA may comprise at least one, at least two, or all three VH CDR sequences selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:59; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:60; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:61.

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Optionally, the antigen-binding site which binds to CEA comprises at least one, at least two, or all three VL CDRs sequences selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:62; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:63; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:64.

Optionally, the antigen-binding site which binds to CEA comprises (a) a VH domain comprising at least one, at least two, or all three VH CDR sequences selected from (i) CDR-H1 comprising the amino acid sequence of SEQ ID NO:59, (ii) CDR-H2 comprising the amino acid sequence of SEQ ID NO:60, and (iii) CDR-H3 comprising an amino acid sequence selected from SEQ ID NO:61; and (b) a VL domain comprising at least one, at least two, or all three VL CDR sequences selected from (i) CDR-L1 comprising the amino acid sequence of SEQ ID NO:62, (ii) CDR-L2 comprising the amino acid sequence of SEQ ID NO:63, and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:64.

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In another aspect, the antigen-binding site which binds to CEA comprises (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:59; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:60; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:61; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:62; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:63; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:64.

In any of the above embodiments, the multispecific antibody may be humanized. In one embodiment, the anti-CEA antigen binding site comprises CDRs as in any of the above embodiments, and further comprises an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework.

In another embodiment, the antigen-binding site which binds to CEA comprises a heavy chain variable domain (VH) sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:65. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:65. In certain embodiments, substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site which binds to CEA comprises the VH sequence in SEQ ID NO:65, including post-translational modifications of that sequence. In a particular embodiment, the VH comprises one, two or three CDRs selected from: (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:59, (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:60, and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:61.

In another embodiment, the antigen-binding site which binds to CEA comprises a light chain variable domain (VL) having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:66. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen-binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted,

inserted and/or deleted in SEQ ID NO:66. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site for CEA comprises the VL sequence in SEQ ID NO:66, including post-translational modifications of that sequence. In a particular embodiment, the VL comprises one, two or three CDRs selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:62; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:63; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:64.

In another embodiment, the antigen-binding site which binds to CEA comprises a VH as in any of the embodiments provided above, and a VL as in any of the embodiments provided above. In one embodiment, the antibody comprises the VH and VL sequences in SEQ ID NO:65 and SEQ ID NO:66, respectively, including post-translational modifications of those sequences.

v). In a still further particular embodiment, the antigen-binding site which binds to CEA may comprise at least one, two, three, four, five, or six CDRs selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:116; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:117 or 118; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:119; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:120, 121 or 122; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:123, 124 or 125; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:126.

Optionally, the antigen-binding site which binds to CEA may comprise:

VH CDR sequences (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:116; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:117 or 118; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:119; and/or

VL CDRs sequences (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:120, 121 or 122; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:123, 124 or 125; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:126.

In one embodiment, the antigen binding site for CEA comprises a heavy chain variable region (VH) comprise the amino acid sequence of SEQ ID NO: 127, or (more preferably) selected from SEQ ID NO: 129, 130, 131, 132, 133 or 134, and a light chain variable region (VL) comprising the amino acid sequence of SEQ ID NO: 128 or (more preferably) selected from SEQ ID NO: 135, 136, 137, 138, 139 or 140.

In any of the above embodiments, the multispecific antibody may be humanized. In one embodiment, the anti-CEA antigen binding site comprises CDRs as in any of the above embodiments, and further comprises an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework.

In a particular aspect, the antigen binding domain capable of binding to CEA comprises:

- (a) a VH domain comprising an amino acid sequence of SEQ ID NO:129 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or
- (b) a VH domain comprising an amino acid sequence of SEQ ID NO:133 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or
- (c) a VH domain comprising an amino acid sequence of SEQ ID NO:130 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or
- (d) a VH domain comprising an amino acid sequence of SEQ ID NO:134 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
- (e) a VH domain comprising an amino acid sequence of SEQ ID NO:133 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
- (f) a VH domain comprising an amino acid sequence of SEQ ID NO:131 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
- (g) a VH domain comprising an amino acid sequence of SEQ ID NO:129 and a VL domain comprising an amino acid sequence of SEQ ID NO:138.

In another embodiment, the antigen-binding site which binds to CEA comprises a heavy chain variable domain (VH) sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence as mentioned in a) to g) above. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted. In certain embodiments, substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs).

In another embodiment, the antigen-binding site which binds to CEA comprises a light chain variable domain (VL) having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence as mentioned in a) to

g) above. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen-binding site comprising that sequence retains the ability to bind to CEA, preferably with the affinity set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs).

In another embodiment, the antigen-binding site which binds to CEA comprises a VH as in any of the embodiments provided above, and a VL as in any of the embodiments provided above.

G. Exemplary antigen binding sites for GPRC5D or FAP

In another particular embodiment of the present invention, which may be combined with the embodiments discussed above (e.g., the binding sites for DOTA or DOTAM), the target antigen bound by the first and second antibody may be GPRC5D or FAP.

Optionally, the antigen-binding site which binds to GPRC5D or FAP may bind with a K_d value of 1nM or less, 500pM or less, 200pM or less, or 100pM or less for monovalent binding.

Exemplary GPRC5D-binding sequences are described below.

In one embodiment, the antigen-binding site which binds to GPRC5D may comprise at least one, two, three, four, five, or six CDRs selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:67; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:68; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:69; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:70; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:71; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:72.

Optionally, the antigen-binding site which binds to GPRC5D may comprise at least one, at least two, or all three VH CDR sequences selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:67; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:68; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:69.

Optionally, the antigen-binding site which binds to GPRC5D comprises at least one, at least two, or all three VL CDRs sequences selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:70; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:71; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:72.

Optionally, the antigen-binding site which binds to GPRC5D comprises (a) a VH domain comprising at least one, at least two, or all three VH CDR sequences selected from (i) CDR-H1 comprising the amino acid sequence of SEQ ID NO:67, (ii) CDR-H2 comprising the amino acid sequence of SEQ ID NO:68, and (iii) CDR-H3 comprising an amino acid sequence selected from SEQ ID NO:69; and (b) a VL domain comprising at least one, at least two, or all three VL CDR sequences selected from (i) CDR-L1 comprising the amino acid sequence of SEQ ID NO:70, (ii) CDR-L2 comprising the amino acid sequence of SEQ ID NO:71, and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:72.

10 In another aspect, the antigen-binding site which binds to GPRC5D comprises (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:67; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:68; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:69; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:70; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:71; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:72.

In any of the above embodiments, the multispecific antibody may be humanized. In one embodiment, the anti- GPRC5D antigen binding site comprises CDRs as in any of the above embodiments, and further comprises an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework.

20 In another embodiment, the antigen-binding site which binds to GPRC5D comprises a heavy chain variable domain (VH) sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:73. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen binding site comprising that sequence retains the ability to bind to GPRC5D, preferably with the affinity as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:73. In certain embodiments, substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs).
30 Optionally, the antigen-binding site which binds to GPRC5D comprises the VH sequence in SEQ ID NO:73, including post-translational modifications of that sequence. In a particular embodiment, the VH comprises one, two or three CDRs selected from: (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:67, (b) CDR-H2 comprising the amino

acid sequence of SEQ ID NO:68, and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:69.

In another embodiment, the antigen-binding site which binds to GPRC5D comprises a light chain variable domain (VL) having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:74. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen-binding site comprising that sequence retains the ability to bind to GPRC5D, preferably with the affinity set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:74. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site for GPRC5D comprises the VL sequence in SEQ ID NO:74, including post-translational modifications of that sequence. In a particular embodiment, the VL comprises one, two or three CDRs selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:70; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:71; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:72.

In another embodiment, the antigen-binding site which binds to GPRC5D comprises a VH as in any of the embodiments provided above, and a VL as in any of the embodiments provided above. In one embodiment, the antibody comprises the VH and VL sequences in SEQ ID NO:73 and SEQ ID NO:74, respectively, including post-translational modifications of those sequences.

Exemplary FAP-binding sequences are described below.

In one embodiment, the antigen-binding site which binds to FAP may comprise at least one, two, three, four, five, or six CDRs selected from (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:75; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:76; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:77; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:78; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:79; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:80.

Optionally, the antigen-binding site which binds to FAP may comprise at least one, at least two, or all three VH CDR sequences selected from (a) CDR-H1 comprising the amino

acid sequence of SEQ ID NO:75; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:76; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:77.

Optionally, the antigen-binding site which binds to FAP comprises at least one, at least two, or all three VL CDRs sequences selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:78; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:79; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:80.

Optionally, the antigen-binding site which binds to FAP comprises (a) a VH domain comprising at least one, at least two, or all three VH CDR sequences selected from (i) CDR-H1 comprising the amino acid sequence of SEQ ID NO:75, (ii) CDR-H2 comprising the amino acid sequence of SEQ ID NO:76, and (iii) CDR-H3 comprising an amino acid sequence selected from SEQ ID NO:77; and (b) a VL domain comprising at least one, at least two, or all three VL CDR sequences selected from (i) CDR-L1 comprising the amino acid sequence of SEQ ID NO:78, (ii) CDR-L2 comprising the amino acid sequence of SEQ ID NO:79, and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:80.

In another aspect, the antigen-binding site which binds to FAP comprises (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:75; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:76; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:77; (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:78; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:79; and (f) CDR-L3 comprising an amino acid sequence SEQ ID NO:80.

In any of the above embodiments, the multispecific antibody may be humanized. In one embodiment, the anti- FAP antigen binding site comprises CDRs as in any of the above embodiments, and further comprises an acceptor human framework, e.g. a human immunoglobulin framework or a human consensus framework.

In another embodiment, the antigen-binding site which binds to FAP comprises a heavy chain variable domain (VH) sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:81. In certain embodiments, a VH sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen binding site comprising that sequence retains the ability to bind to FAP, preferably with the affinity as set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:81. In certain embodiments, substitutions,

insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site which binds to FAP comprises the VH sequence in SEQ ID NO:81, including post-translational modifications of that sequence. In a particular embodiment, the VH comprises one, two or three CDRs selected from: (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:75, (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:76, and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:77.

In another embodiment, the antigen-binding site which binds to FAP comprises a light chain variable domain (VL) having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% sequence identity to the amino acid sequence of SEQ ID NO:82. In certain embodiments, a VL sequence having at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identity contains substitutions (e.g., conservative substitutions), insertions, or deletions relative to the reference sequence, but the antigen-binding site comprising that sequence retains the ability to bind to FAP, preferably with the affinity set out above. In certain embodiments, a total of 1 to 10 amino acids have been substituted, inserted and/or deleted in SEQ ID NO:82. In certain embodiments, the substitutions, insertions, or deletions occur in regions outside the HVRs (i.e., in the FRs). Optionally, the antigen-binding site for FAP comprises the VL sequence in SEQ ID NO:82, including post-translational modifications of that sequence. In a particular embodiment, the VL comprises one, two or three CDRs selected from (a) CDR-L1 comprising the amino acid sequence of SEQ ID NO:78; (b) CDR-L2 comprising the amino acid sequence of SEQ ID NO:79; and (c) CDR-L3 comprising the amino acid sequence of SEQ ID NO:80.

In another embodiment, the antigen-binding site which binds to FAP comprises a VH as in any of the embodiments provided above, and a VL as in any of the embodiments provided above. In one embodiment, the antibody comprises the VH and VL sequences in SEQ ID NO:81 and SEQ ID NO:82, respectively, including post-translational modifications of those sequences.

H. Exemplary antibodies

Aspects and embodiments concerning target antigen binding (e.g., CEA, GPRC5D or FAP and aspects and embodiments concerning effector moiety binding (e.g. DOTA, DOTAM) are expressly contemplated in combination. Any of the exemplary target antigen binding sites describes above may be used in combination with any of the effector moiety binding sites described above.

In some specific embodiments, aspects and embodiments concerning target antigen binding (e.g., CEA, GPRC5D or FAP) and aspects and embodiments concerning DOTAM binding are combined. In some embodiments it may be preferred that the target antigen is CEA.

In some embodiments, the first antibody may comprise:

a) a Fab fragment, wherein the Fab fragment binds to an antigen expressed on the surface of a target cell selected from CEA, GPRC5D or FAP, optionally CEA;

10 b) a polypeptide comprising or consisting of an antibody heavy chain variable domain (VH) of an antigen binding site for Pb-DOTAM, wherein the heavy chain variable domain comprises heavy chain CDRs of SEQ ID NOs 1-3, and/or wherein the heavy chain variable domain has at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 7; and

 c) an Fc domain comprising two subunits,
wherein the polypeptide of (b) is fused by its N-terminus to the C-terminus of one of the chains of the Fab fragment of (a) and by its C-terminus to the N-terminus of one of the subunits of the Fc domain of (c);
and wherein the first antibody does not comprise a VL domain of an antigen binding site for Pb-DOTAM.

The second antibody may comprise:

20 d) a Fab fragment, wherein the Fab fragment binds to an antigen expressed on the surface of a target cell selected from CEA, GPRC5D or FAP, optionally CEA;

 e) a polypeptide comprising or consisting of an antibody light chain variable domain (VL) of an antigen binding site for Pb-DOTAM wherein the light chain variable domain comprises CDRs of SEQ ID NO: 4-6 and/or wherein the light chain variable domain has at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 8; and

 f) an Fc domain comprising two subunits,
wherein the polypeptide of (e) is fused by its N-terminus to the C-terminus of one of the chains of the Fab fragment of (d) and by its C-terminus to the N-terminus of one of the subunits of the Fc domain of (f);

30 and wherein the second antibody does not comprise a VH domain of an antigen binding site for Pb-DOTAM.

Said VH domain of the first antibody and said VL domain of the second antibody are together capable of forming a functional antigen binding site for Pb-DOTAM.

In one particular embodiment, the first and/or second antibodies (e.g., each of the first and second antibodies) may comprise an additional Fab fragment binding to CEA, GPRC5D or FAP, optionally CEA. It may be preferred that each of the target antigen-binding Fabs in the first antibody bind to the same target antigen as each other (e.g., in some embodiments CEA), and that each of the target antigen-binding Fabs in the second antibody bind to the same target antigen as each other (e.g., in some embodiments CEA), which may further be the same as that bound by the first antibody. It may further be preferred that each of the target antigen-binding Fabs in the first antibody bind to the same epitope of target antigen, e.g., CEA (i.e., that the first antibody is monospecific in respect of the target antigen) and that each of the target antigen-binding Fabs in the second antibody bind to the same epitope of target antigen, e.g., CEA (i.e., that the second antibody is monospecific in respect of the target antigen). The epitope may the bound by the two antibodies may be the same or different.

In some embodiments, the first antibody may comprise the following peptides:

- i) a first heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VH domain of an antigen binding site for Pb-DOTAM wherein the heavy chain variable domain comprises heavy chain CDRs of SEQ ID NOs 1-3, and/or wherein the heavy chain variable domain has at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 7; an optional linker; and an Fc subunit (e.g., CH2-CH3);
- ii) a Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (i) to form a binding site for a target antigen selected from CEA, GPRC5D, or FAP, optionally CEA;
- iii) a second heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; and an Fc subunit (e.g., CH2-CH3); and
- iv) a further Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (iii) to form a binding site for a target antigen selected from CEA, GPRC5D or FAP, optionally CEA.

Optionally the Fab heavy chain in (i) has the same sequence as the Fab heavy chain in (iii) and the Fab light chains of (ii) and (iv) have the same sequence as each other.

The second antibody may comprise the following peptides:

- v) a first heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VL domain of an antigen binding site for Pb-DOTAM wherein the light chain variable domain comprises CDRs of SEQ ID NO: 4-6 and/or wherein the light chain variable domain has at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 8; an optional linker; and an Fc subunit (e.g., CH2-CH3);
- vi) a Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (v) to form a binding site for a target antigen selected from CEA, GPRC5D or FAP, optionally CEA;
- vii) a second heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; and an Fc subunit (e.g., CH2-CH3); and
- viii) a further Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (vii) to form a binding site for a target antigen selected from CEA, GPRC5D or FAP, optionally CEA.

Optionally the Fab heavy chain in (v) has the same sequence as the Fab heavy chain in (vii) and the Fab light chains of (vi) and (viii) have the same sequence as each other. Optionally these are also the same as for the first antibody.

In a particular embodiment, the first antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody CH1A1A.

For example, the two Fab light chain polypeptides in (ii) and (iv) may comprise the CDRs of SEQ ID Nos 22-24 and/or may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26. In some embodiments they may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 34. In some embodiments, it may be preferred that the two light chains in (ii) and (iv) are identical to each other.

The two Fab heavy chains in (i) and (iii) may comprise the CDRs of SEQ ID NOs: 19-21 and/or the two Fab heavy chains in (i) and (iii) may comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25. In some

embodiments, it may be preferred that the two Fab heavy chains in (i) and (iii) are identical to each other.

In another particular embodiment, the first antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody A5B7 (including a humanized version thereof).

For example, the two Fab light chain polypeptides in (ii) and (iv) may comprise the CDRs of SEQ ID Nos 46-48 and/or may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 50. In some embodiments they may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to
10 SEQ ID NO: 54. In some embodiments, it may be preferred that the two Fab light chain polypeptides in (ii) and (iv) are identical to each other.

In some embodiments, the two Fab heavy chains in (i) and (iii) may comprise the CDRs of SEQ ID NOs: 43-45 and/or the two Fab heavy chains in (i) and (iii) may comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 49. In some embodiments, it may be preferred that the two Fab heavy chains in (i) and (iii) are identical to each other.

In another particular embodiment, the first antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody T84.66 (including a humanized
20 version thereof).

For example, the two Fab light chain polypeptides in (ii) and (iv) may comprise the CDRs of SEQ ID Nos 14-16 and/or may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 18. In some embodiments they may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to
SEQ ID NO: 89.

In some embodiments, the two Fab heavy chains in (i) and (iii) may comprise the CDRs of SEQ ID NOs: 11-13 and/or the two Fab heavy chains in (i) and (iii) may comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 17. In some embodiments, it may be preferred that the two Fab heavy chains in (i) and (iii) are identical to each other.

30 In another particular embodiment, the first antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody 28A9 (including a humanized version thereof).

For example, the two Fab light chain polypeptides in (ii) and (iv) may comprise the CDRs of SEQ ID Nos 62-64 and/or may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 66. In some embodiments they may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 96.

In some embodiments, the two Fab heavy chains in (i) and (iii) may comprise the CDRs of SEQ ID NOs: 59-61 and/or the two Fab heavy chains in (i) and (iii) may comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 65. In some embodiments, it may be preferred that the two Fab heavy chains in (i) and (iii) are identical to each other.

In some embodiments, the second antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody CH1A1A.

For example, the two Fab light chains in (vi) and (viii) may comprise the CDRs of SEQ ID Nos 22-24 and/or may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26. In some embodiments they may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 34.

In some embodiments, the two Fab heavy chains in (v) and (vii) comprise the CDRs of SEQ ID NOs: 19-21 and/or the two Fab heavy chains in (v) and (vii) comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25. In some embodiments, it may be preferred that the two Fab heavy chains in (v) and (vii) are identical to each other.

In another particular embodiment, the second antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from A5B7 (including a humanized version thereof).

For example, the two Fab light chain polypeptides in (vi) and (viii) may comprise the CDRs of SEQ ID Nos 46-48 and/or may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 50. In some embodiments they may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 58.

In some embodiments, the two Fab heavy chains in (v) and (vii) comprise the CDRs of SEQ ID NOs: 43-45 and/or the two Fab heavy chains in (v) and (vii) comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO

49. In some embodiments, it may be preferred that the two Fab heavy chains in (v) and (vii) are identical to each other.

In another particular embodiment, the second antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody T84.66 (including a humanized version thereof).

For example, the two light chain polypeptides in (vi) and (viii) may comprise the CDRs of SEQ ID Nos 14-16 and/or may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 18. In some embodiments they may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to
10 SEQ ID NO: 89.

In some embodiments, the two Fab heavy chains in (v) and (vii) may comprise the CDRs of SEQ ID NOs: 11-13 and/or the two Fab heavy chains in (v) and (vii) comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 17. In some embodiments, it may be preferred that the two Fab heavy chains in (v) and (vii) are identical to each other.

In another particular embodiment, the second antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody 28A9 (including a humanized version thereof).

For example, the two Fab light chains in (vi) and (vii) may comprise the CDRs of
20 SEQ ID Nos 62-64 and/or may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 66. In some embodiments they may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 96.

In some embodiments, the two Fab heavy chains in (v) and (vii) may comprise the CDRs of SEQ ID NOs: 59-61 and/or two Fab heavy chains in (v) and (vii) comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 65. In some embodiments, it may be preferred that the two Fab heavy chains in (v) and (vii) are identical to each other.

In some embodiments, the first and the second antibody bind the same epitope of CEA. Thus, for example, the first and the second antibody may both have CEA binding
30 sequences from the antibody CH1A1A; or, the first and the second antibody may both have CEA binding sequences from A5B7 (including a humanized version thereof); or, the first and the second antibody may both have CEA binding sequences from T84.66 (including a humanized version thereof); or, the first and the second antibody may both have CEA binding

sequences from 28A9 (including a humanized version thereof); or, the first and the second antibody may both have CEA binding sequences from MFE23 (including a humanized version thereof). In some embodiments, it may be preferred that the two light chain polypeptides in (vi) and (viii) have the same sequence as the light chains in (ii) and (iv) of the first antibody, e.g., that all said light chains have the same sequence.

In some embodiments, both the two Fab light chain polypeptides of the first antibody, in (ii) and (iv), may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26; both the two Fab heavy chains of the first antibody in (i) and (iii) may comprise a heavy chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25; both the two Fab light chain polypeptides of the second antibody, in (vi) and (viii), may comprise light chain variable domains having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26; and both the two Fab heavy chains of the second antibody, in (v) and (vii), may comprise a heavy chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25.

In one particular embodiment, the first antibody comprises:

- a first heavy chain polypeptide of SEQ ID NO: 146;
- a second heavy chain polypeptide of SEQ ID NO: 147;
- a first and second light chain polypeptide of SEQ ID NO: 143; and/or

the second antibody comprises

- a first heavy chain polypeptide of SEQ ID NO: 145;
- a second heavy chain polypeptide of SEQ ID NO: 147;
- a first and second light chain polypeptide of SEQ ID NO: 143.

In other embodiments, the first and the second antibodies bind to different epitopes of CEA, as discussed above. Thus, for instance, the first antibody may have CEA binding sequences from the antibody CH1A1A and the second antibody may have CEA binding sequences from A5B7; or, the first antibody may have CEA binding sequences from the antibody A5B7 and the second antibody may have CEA binding sequences from CH1A1A.

In other embodiments, the antibodies are one-armed antibodies. For example, in some embodiments, the first antibody comprises the following polypeptides:

- i) a polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VH domain of an antigen binding site for Pb-DOTAM wherein

the heavy chain variable domain comprises heavy chain CDRs of SEQ ID NOs 1-3, and/or wherein the heavy chain variable domain has at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 7; an optional linker; and an Fc subunit (e.g., CH2-CH3);

ii) a Fab light chain polypeptide (e.g., VL-CL); and

iii) an Fc subunit polypeptide (e.g., CH2-CH3);

wherein the Fab heavy chain of (i) and the Fab light chain of (ii) form a Fab fragment capable of binding to a target antigen selected from CEA, GPRC5D or FAP, optionally CEA.

The second antibody may comprise the following polypeptides:

iv) a polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VL domain of an antigen binding site for Pb-DOTAM wherein the light chain variable domain comprises CDRs of SEQ ID NO: 4-6 and/or wherein the light chain variable domain has at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 8; an optional linker; and an Fc subunit (e.g., CH2-CH3);

v) a Fab light chain polypeptide (e.g., VL-CL), and

vi) an Fc subunit polypeptide (e.g., CH2-CH3);

wherein the Fab heavy chain of (iv) and the Fab light chain of (v) form a Fab fragment capable of binding to a target antigen selected from CEA, GPRC5D or FAP, optionally CEA.

In some embodiments of these one-armed antibodies, the Fab heavy chain of (i) and of (iv) may have the same sequence as each other; and the Fab light chain polypeptide of (ii) and (v) may have the same sequence as each other.

In a particular embodiment, the first antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody CH1A1A.

For example, the Fab light chain polypeptide in (ii) may comprise the CDRs of SEQ ID Nos 22-24 and/or may comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26. In some embodiments it may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 34.

The Fab heavy chain in (i) may comprise the CDRs of SEQ ID NOs: 19-21 and/or the Fab heavy chain in (i) may comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25.

In another particular embodiment, the first antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody A5B7 (including a humanized version thereof).

For example, the Fab light chain polypeptide in (ii) may comprise the CDRs of SEQ ID Nos 46-48 and/or may comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 50. In some embodiments it may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 54.

In some embodiments, the Fab heavy chain in (i) may comprise the CDRs of SEQ ID NOs: 43-45 and/or the Fab heavy chain in (i) may comprise a variable domain having at least
10 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 49.

In another particular embodiment, the first antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody T84.66 (including a humanized version thereof).

For example, the Fab light chain polypeptide in (ii) may comprise the CDRs of SEQ ID Nos 14-16 and/or may comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 18. In some embodiments it may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 89.

In some embodiments, the Fab heavy chain in (i) may comprise the CDRs of SEQ ID NOs: 11-13 and/or the Fab heavy chain in (i) may comprise a variable domain having at least
20 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 17.

In another particular embodiment, the first antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody 28A9 (including a humanized version thereof).

For example, the Fab light chain polypeptide in (ii) may comprise the CDRs of SEQ ID Nos 62-64 and/or may comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 66. In some embodiments it may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 96.

In some embodiments, the Fab heavy chain in (i) may comprise the CDRs of SEQ ID NOs: 59-61 and/or the Fab heavy chain in (i) may comprise a variable domain having at least
30 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 65.

In some embodiments, the second antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody CH1A1A.

For example, the Fab light chain in (v) may comprise the CDRs of SEQ ID Nos 22-24 and/or may comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26. In some embodiments it may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 34.

In some embodiments, the Fab heavy chain in (iv) comprises the CDRs of SEQ ID NOs: 19-21 and/or the Fab heavy chain in (iv) comprises a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25.

10 In another particular embodiment, the second antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from A5B7 (including a humanized version thereof).

For example, the Fab light chain polypeptide in (v) may comprise the CDRs of SEQ ID Nos 46-48 and/or may comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 50. In some embodiments it may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 58.

In some embodiments, the Fab heavy chain in (iv) comprises the CDRs of SEQ ID NOs: 43-45 and/or the Fab heavy chain in (iv) comprises a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 49.

20 In another particular embodiment, the second antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody T84.66 (including a humanized version thereof).

For example, the Fab light chain polypeptide in (v) may comprise the CDRs of SEQ ID Nos 14-16 and/or may comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 18. In some embodiments it may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 89.

In some embodiments, the Fab heavy chain in (iv) may comprise the CDRs of SEQ ID NOs: 11-13 and/or the Fab heavy chain in (iv) may comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 17.

30 In another particular embodiment, the second antibody may have CEA binding sequences (i.e., CDRs or VH/VL domains) from the antibody 28A9 (including a humanized version thereof).

For example, the Fab light chain in (v) may comprise the CDRs of SEQ ID Nos 62-64 and/or may comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96,

97, 98, 99 or 100% identity to SEQ ID NO 66. In some embodiments it may have at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO: 96.

In some embodiments, the Fab heavy chain in (iv) may comprise the CDRs of SEQ ID NOs: 59-61 and/or the Fab heavy chain in (iv) may comprise a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 65.

10 In some embodiments, the first and the second antibody bind the same epitope of CEA. Thus, for example, the first and the second antibody may both have CEA binding sequences from the antibody CH1A1A; or, the first and the second antibody may both have CEA binding sequences from A5B7 (including a humanized version thereof); or, the first and the second antibody may both have CEA binding sequences from T84.66 (including a humanized version thereof); or, the first and the second antibody may both have CEA binding sequences from 28A9 (including a humanized version thereof); or, the first and the second antibody may both have CEA binding sequences from MFE23 (including a humanized version thereof). In some embodiments, it may be preferred that the light chain polypeptides in (ii) has the same sequence as the light chains in (v).

In a particular embodiment, the Fab light chains in (ii) and (v) both comprise a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26; and the Fab heavy chain in (i) and (iv) comprises a variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25.

20 In one particular embodiment,
the first antibody comprises:

- a first heavy chain of SEQ ID NO: 146;
- a second heavy chain of SEQ ID NO: 144;
- a light chain of SEQ ID NO: 143; and/or

the second antibody comprises:

- a first heavy chain of SEQ ID NO: 145;
- a second heavy chain of SEQ ID NO: 144;
- a light chain of SEQ ID NO: 143.

30 In other embodiments, the first and the second antibodies bind to different epitopes of CEA, as discussed above. Thus, for instance, the first antibody may have CEA binding sequences from the antibody CH1A1A and the second antibody may have CEA binding

sequences from A5B7; or, the first antibody may have CEA binding sequences from the antibody A5B7 and the second antibody may have CEA binding sequences from CH1A1A.

I. Antibody Variants

In certain embodiments, amino acid sequence variants of the antibodies provided herein are contemplated. For example, it may be desirable to improve the binding affinity and/or other biological properties of the antibody. Amino acid sequence variants of an antibody may be prepared by introducing appropriate modifications into the nucleotide sequence encoding the antibody, or by peptide synthesis. Such modifications include, for example, deletions from, and/or insertions into and/or substitutions of residues within the amino acid sequences of the antibody. Any combination of deletion, insertion, and substitution can be made to arrive at the final construct, provided that the final construct possesses the desired characteristics, e.g., antigen-binding.

Substitution, Insertion, and Deletion Variants

In certain embodiments, antibody variants having one or more amino acid substitutions are provided. Sites of interest for substitutional mutagenesis include the HVRs (CDRs) and FRs. Conservative substitutions are shown in Table 1 under the heading of "preferred substitutions." More substantial changes are provided in Table 1 under the heading of "exemplary substitutions," and as further described below in reference to amino acid side chain classes. Amino acid substitutions may be introduced into an antibody of interest and the products screened for a desired activity, e.g., retained/improved antigen binding, decreased immunogenicity, or reduced or eliminated ADCC or CDC.

TABLE 1

Original Residue	Exemplary Substitutions	Preferred Substitutions
Ala (A)	Val; Leu; Ile	Val
Arg (R)	Lys; Gln; Asn	Lys
Asn (N)	Gln; His; Asp, Lys; Arg	Gln
Asp (D)	Glu; Asn	Glu
Cys (C)	Ser; Ala	Ser
Gln (Q)	Asn; Glu	Asn

Original Residue	Exemplary Substitutions	Preferred Substitutions
Glu (E)	Asp; Gln	Asp
Gly (G)	Ala	Ala
His (H)	Asn; Gln; Lys; Arg	Arg
Ile (I)	Leu; Val; Met; Ala; Phe; Norleucine	Leu
Leu (L)	Norleucine; Ile; Val; Met; Ala; Phe	Ile
Lys (K)	Arg; Gln; Asn	Arg
Met (M)	Leu; Phe; Ile	Leu
Phe (F)	Trp; Leu; Val; Ile; Ala; Tyr	Tyr
Pro (P)	Ala	Ala
Ser (S)	Thr	Thr
Thr (T)	Val; Ser	Ser
Trp (W)	Tyr; Phe	Tyr
Tyr (Y)	Trp; Phe; Thr; Ser	Phe
Val (V)	Ile; Leu; Met; Phe; Ala; Norleucine	Leu

Amino acids may be grouped according to common side-chain properties:

- (1) hydrophobic: Norleucine, Met, Ala, Val, Leu, Ile;
- (2) neutral hydrophilic: Cys, Ser, Thr, Asn, Gln;
- (3) acidic: Asp, Glu;
- (4) basic: His, Lys, Arg;
- (5) residues that influence chain orientation: Gly, Pro;
- (6) aromatic: Trp, Tyr, Phe.

10 Non-conservative substitutions will entail exchanging a member of one of these classes for another class.

One type of substitutional variant involves substituting one or more hypervariable region residues of a parent antibody (e.g., a humanized or human antibody). Generally, the resulting variant(s) selected for further study will have modifications (e.g., improvements) in certain biological properties (e.g., increased affinity, reduced immunogenicity) relative to the parent antibody and/or will have substantially retained certain biological properties of the

parent antibody. An exemplary substitutional variant is an affinity matured antibody, which may be conveniently generated, e.g., using phage display-based affinity maturation techniques such as those described herein. Briefly, one or more CDR residues are mutated and the variant antibodies displayed on phage and screened for a particular biological activity (e.g., binding affinity).

Alterations (e.g., substitutions) may be made in CDRs, e.g., to improve antibody affinity. Such alterations may be made in CDR “hotspots”, i.e., residues encoded by codons that undergo mutation at high frequency during the somatic maturation process (see, e.g., Chowdhury, *Methods Mol. Biol.* 207:179-196 (2008)), and/or residues that contact antigen, with the resulting variant VH or VL being tested for binding affinity. Affinity maturation by constructing and reselecting from secondary libraries has been described, e.g., in Hoogenboom et al. in *Methods in Molecular Biology* 178:1-37 (O’Brien et al., ed., Human Press, Totowa, NJ, (2001).) In some aspects of affinity maturation, diversity is introduced into the variable genes chosen for maturation by any of a variety of methods (e.g., error-prone PCR, chain shuffling, or oligonucleotide-directed mutagenesis). A secondary library is then created. The library is then screened to identify any antibody variants with the desired affinity. Another method to introduce diversity involves CDR-directed approaches, in which several CDR residues (e.g., 4-6 residues at a time) are randomized. CDR residues involved in antigen binding may be specifically identified, e.g., using alanine scanning mutagenesis or modelling. CDR-H3 and CDR-L3 in particular are often targeted.

In certain aspects, substitutions, insertions, or deletions may occur within one or more CDRs so long as such alterations do not substantially reduce the ability of the antibody to bind antigen. For example, conservative alterations (e.g., conservative substitutions as provided herein) that do not substantially reduce binding affinity may be made in the CDRs. Such alterations may, for example, be outside of antigen contacting residues in the CDRs. In certain variant VH and VL sequences provided above, each CDR either is unaltered, or contains no more than one, two or three amino acid substitutions.

A useful method for identification of residues or regions of an antibody that may be targeted for mutagenesis is called “alanine scanning mutagenesis” as described by Cunningham and Wells (1989) *Science*, 244:1081-1085. In this method, a residue or group of target residues (e.g., charged residues such as arg, asp, his, lys, and glu) are identified and replaced by a neutral or negatively charged amino acid (e.g., alanine or polyalanine) to determine whether the interaction of the antibody with antigen is affected. Further

substitutions may be introduced at the amino acid locations demonstrating functional sensitivity to the initial substitutions. Alternatively, or additionally, a crystal structure of an antigen-antibody complex may be used to identify contact points between the antibody and antigen. Such contact residues and neighbouring residues may be targeted or eliminated as candidates for substitution. Variants may be screened to determine whether they contain the desired properties.

Amino acid sequence insertions include amino- and/or carboxyl-terminal fusions ranging in length from one residue to polypeptides containing a hundred or more residues, as well as intrasequence insertions of single or multiple amino acid residues. Examples of
10 terminal insertions include an antibody with an N-terminal methionyl residue. Other insertional variants of the antibody molecule include the fusion to the N- or C-terminus of the antibody to an enzyme (e.g., for ADEPT (antibody directed enzyme prodrug therapy)) or a polypeptide which increases the serum half-life of the antibody.

Glycosylation variants

In certain aspects, an antibody provided herein is altered to increase or decrease the extent to which the antibody is glycosylated. Addition or deletion of glycosylation sites to an antibody may be conveniently accomplished by altering the amino acid sequence such that one or more glycosylation sites is created or removed.

Where the antibody comprises an Fc region, the oligosaccharide attached thereto may
20 be altered. Native antibodies produced by mammalian cells typically comprise a branched, biantennary oligosaccharide that is generally attached by an N-linkage to Asn297 of the CH2 domain of the Fc region. See, e.g., Wright et al. *TIBTECH* 15:26-32 (1997). The oligosaccharide may include various carbohydrates, e.g., mannose, N-acetyl glucosamine (GlcNAc), galactose, and sialic acid, as well as a fucose attached to a GlcNAc in the “stem” of the biantennary oligosaccharide structure. In some aspects, modifications of the oligosaccharide in an antibody of the invention may be made in order to create antibody variants with certain improved properties.

In one aspect, antibody variants are provided having a non-fucosylated oligosaccharide, i.e. an oligosaccharide structure that lacks fucose attached (directly or indirectly) to an Fc
30 region. Such non-fucosylated oligosaccharide (also referred to as “afucosylated” oligosaccharide) particularly is an N-linked oligosaccharide which lacks a fucose residue attached to the first GlcNAc in the stem of the biantennary oligosaccharide structure. In one

aspect, antibody variants are provided having an increased proportion of non-fucosylated oligosaccharides in the Fc region as compared to a native or parent antibody. For example, the proportion of non-fucosylated oligosaccharides may be at least about 20%, at least about 40%, at least about 60%, at least about 80%, or even about 100% (i.e. no fucosylated oligosaccharides are present). The percentage of non-fucosylated oligosaccharides is the (average) amount of oligosaccharides lacking fucose residues, relative to the sum of all oligosaccharides attached to Asn 297 (e. g. complex, hybrid and high mannose structures) as measured by MALDI-TOF mass spectrometry, as described in WO 2006/082515, for example. Asn297 refers to the asparagine residue located at about position 297 in the Fc region (EU numbering of Fc region residues); however, Asn297 may also be located about $\pm$ 3 amino acids upstream or downstream of position 297, i.e., between positions 294 and 300, due to minor sequence variations in antibodies. Such antibodies having an increased proportion of non-fucosylated oligosaccharides in the Fc region may have improved Fc γ RIIIa receptor binding and/or improved effector function, in particular improved ADCC function. See, e.g., US 2003/0157108; US 2004/0093621.

Examples of cell lines capable of producing antibodies with reduced fucosylation include Lec13 CHO cells deficient in protein fucosylation (Ripka et al. *Arch. Biochem. Biophys.* 249:533-545 (1986); US 2003/0157108; and WO 2004/056312, especially at Example 11), and knockout cell lines, such as alpha-1,6-fucosyltransferase gene, *FUT8*, knockout CHO cells (see, e.g., Yamane-Ohnuki et al. *Biotech. Bioeng.* 87:614-622 (2004); Kanda, Y. et al., *Biotechnol. Bioeng.*, 94(4):680-688 (2006); and WO 2003/085107), or cells with reduced or abolished activity of a GDP-fucose synthesis or transporter protein (see, e.g., US2004259150, US2005031613, US2004132140, US2004110282).

In a further aspect, antibody variants are provided with bisected oligosaccharides, e.g., in which a biantennary oligosaccharide attached to the Fc region of the antibody is bisected by GlcNAc. Such antibody variants may have reduced fucosylation and/or improved ADCC function as described above. Examples of such antibody variants are described, e.g., in Umana et al., *Nat Biotechnol* 17, 176-180 (1999); Ferrara et al., *Biotechn Bioeng* 93, 851-861 (2006); WO 99/54342; WO 2004/065540, WO 2003/011878.

Antibody variants with at least one galactose residue in the oligosaccharide attached to the Fc region are also provided. Such antibody variants may have improved CDC function. Such antibody variants are described, e.g., in WO 1997/30087; WO 1998/58964; and WO 1999/22764.

It may be preferred that the antibody is modified to reduce the extent of glycosylation. In some embodiments the antibody may be aglycosylated or deglycosylated. The antibody may include a substitution at N297, e.g., N297D/A.

Fc region variants

In certain embodiments, one or more amino acid modifications may be introduced into the Fc region of an antibody provided herein, thereby generating an Fc region variant. The Fc region variant may comprise a human Fc region sequence (e.g., a human IgG1, IgG2, IgG3 or IgG4 Fc region) comprising an amino acid modification (e.g. a substitution) at one or
10 more amino acid positions.

In certain embodiments, the invention contemplates an antibody variant with reduced Fc effector function, e.g., reduced or eliminated CDC, ADCC and/or FcγR binding. In certain aspects, the invention contemplates an antibody variant that possesses some but not all Fc effector functions, which make it a desirable candidate for applications in which the half life of the antibody in vivo is important yet certain Fc effector functions (such as complement-dependent cytotoxicity (CDC) and antibody-dependent cell-mediated cytotoxicity (ADCC)) are unnecessary or deleterious.

In vitro and/or *in vivo* cytotoxicity assays can be conducted to confirm the reduction/depletion of CDC and/or ADCC activities. For example, Fc receptor (FcR) binding
20 assays can be conducted to ensure that the antibody lacks FcγR binding (hence likely lacking ADCC activity), but retains FcRn binding ability. The primary cells for mediating ADCC, NK cells, express FcγRIII only, whereas monocytes express FcγRI, FcγRII and FcγRIII. FcR expression on hematopoietic cells is summarized in Table 3 on page 464 of Ravetch and Kinet, *Annu. Rev. Immunol.* 9:457-492 (1991). Non-limiting examples of *in vitro* assays to assess ADCC activity of a molecule of interest is described in U.S. Patent No. 5,500,362 (see, e.g., Hellstrom, I. et al. *Proc. Nat'l Acad. Sci. USA* 83:7059-7063 (1986)) and Hellstrom, I et al., *Proc. Nat'l Acad. Sci. USA* 82:1499-1502 (1985); 5,821,337 (see Bruggemann, M. et al., *J. Exp. Med.* 166:1351-1361 (1987)). Alternatively, non-radioactive assays methods may be employed (see, for example, ACT1™ non-radioactive cytotoxicity assay for flow cytometry
30 (CellTechnology, Inc. Mountain View, CA; and CytoTox 96® non-radioactive cytotoxicity assay (Promega, Madison, WI). Useful effector cells for such assays include peripheral blood mononuclear cells (PBMC) and Natural Killer (NK) cells. Alternatively, or additionally,

ADCC activity of the molecule of interest may be assessed *in vivo*, e.g., in an animal model such as that disclosed in Clynes et al. *Proc. Nat'l Acad. Sci. USA* 95:652-656 (1998). C1q binding assays may also be carried out to confirm that the antibody is unable to bind C1q and hence lacks CDC activity. See, e.g., C1q and C3c binding ELISA in WO 2006/029879 and WO 2005/100402. To assess complement activation, a CDC assay may be performed (see, for example, Gazzano-Santoro *et al.*, *J. Immunol. Methods* 202:163 (1996); Cragg, M.S. et al., *Blood* 101:1045-1052 (2003); and Cragg, M.S. and M.J. Glennie, *Blood* 103:2738-2743 (2004)). FcRn binding and *in vivo* clearance/half life determinations can also be performed using methods known in the art (see, e.g., Petkova, S.B. et al., *Int'l. Immunol.* 18(12):1759-1769 (2006); WO 2013/120929 A1).

Antibodies with reduced Fc effector function include those with substitution of one or more of Fc region residues 238, 265, 269, 270, 297, 327 and 329 (U.S. Patent No. 6,737,056), e.g., P329G. Such Fc mutants include Fc mutants with substitutions at two or more of amino acid positions 265, 269, 270, 297 and 327, including the so-called "DANA" Fc mutant with substitution of residues 265 and 297 to alanine (US Patent No. 7,332,581).

In certain aspects, an antibody variant comprises an Fc region with one or more amino acid substitutions which diminish FcγR binding, e.g., substitutions at positions 234 and 235 of the Fc region (EU numbering of residues). In one aspect, the substitutions are L234A and L235A (LALA). In certain aspects, the antibody variant further comprises D265A and/or P329G in an Fc region derived from a human IgG1 Fc region. In one aspect, the substitutions are L234A, L235A and P329G (LALA-PG) in an Fc region derived from a human IgG1 Fc region. (See, e.g., WO 2012/130831). In another aspect, the substitutions are L234A, L235A and D265A (LALA-DA) in an Fc region derived from a human IgG1 Fc region. Alternative substitutions include L234F and/or L235E, optionally in combination with D265A and/or P329G and/or P331S.

In other embodiments, it may be possible to use a IgG subtype with reduced Fc effector function such as IgG4 or IgG2.

Certain antibody variants with improved or diminished binding to FcRs are described. (See, e.g., U.S. Patent No. 6,737,056; WO 2004/056312, and Shields et al., *J. Biol. Chem.* 276(9): 6591-6604 (2001).)

In some embodiments, alterations are made in the Fc region that result in altered (*i.e.*, either improved or diminished, preferably diminished) C1q binding and/or Complement

Dependent Cytotoxicity (CDC), e.g., as described in US Patent No. 6,194,551, WO 99/51642, and Idusogie et al. *J. Immunol.* 164: 4178-4184 (2000).

In certain aspects, an antibody variant comprises an Fc region with one or more amino acid substitutions, which reduce FcRn binding, e.g., substitutions at positions 253, and/or 310, and/or 435 of the Fc-region (EU numbering of residues). In certain aspects, the antibody variant comprises an Fc region with the amino acid substitutions at positions 253, 310 and 435. In one aspect, the substitutions are I253A, H310A and H435A in an Fc region derived from a human IgG1 Fc-region. See, e.g., Grevys, A., et al., *J. Immunol.* 194 (2015) 5497-5508.

10 In certain aspects, an antibody variant comprises an Fc region with one or more amino acid substitutions, which reduce FcRn binding, e.g., substitutions at positions 310, and/or 433, and/or 436 of the Fc region (EU numbering of residues). In certain aspects, the antibody variant comprises an Fc region with the amino acid substitutions at positions 310, 433 and 436. In one aspect, the substitutions are H310A, H433A and Y436A in an Fc region derived from a human IgG1 Fc-region. (See, e.g., WO 2014/177460 A1). For instance, in some embodiments, normal FcRn binding may be used.

See also Duncan & Winter, *Nature* 322:738-40 (1988); U.S. Patent No. 5,648,260; U.S. Patent No. 5,624,821; and WO 94/29351 concerning other examples of Fc region variants.

20 The C-terminus of a heavy chain of the full-length antibody as reported herein can be a complete C-terminus ending with the amino acid residues PGK. The C-terminus of the heavy chain can be a shortened C-terminus in which one or two of the C terminal amino acid residues have been removed. The C-terminus of the heavy chain may be a shortened C-terminus ending PG. In one aspect of all aspects as reported herein, an antibody comprising a heavy chain including a C-terminal CH3 domain, as specified herein, comprises a C-terminal glycine residue (G446, EU index numbering of amino acid positions). This is still explicitly encompassed with the term “full length antibody” or “full length heavy chain” as used herein.

Antibody Derivatives

30 In certain aspects, an antibody provided herein may be further modified to contain additional nonproteinaceous moieties that are known in the art and readily available. The moieties suitable for derivatization of the antibody include but are not limited to water

soluble polymers. Non-limiting examples of water soluble polymers include, but are not limited to, polyethylene glycol (PEG), copolymers of ethylene glycol/propylene glycol, carboxymethylcellulose, dextran, polyvinyl alcohol, polyvinyl pyrrolidone, poly-1, 3-dioxolane, poly-1,3,6-trioxane, ethylene/maleic anhydride copolymer, polyaminoacids (either homopolymers or random copolymers), and dextran or poly(n-vinyl pyrrolidone)polyethylene glycol, propylene glycol homopolymers, polypropylene oxide/ethylene oxide copolymers, polyoxyethylated polyols (e.g., glycerol), polyvinyl alcohol, and mixtures thereof. Polyethylene glycol propionaldehyde may have advantages in manufacturing due to its stability in water. The polymer may be of any molecular weight, and may be branched or unbranched. The number of polymers attached to the antibody may vary, and if more than one polymer are attached, they can be the same or different molecules. In general, the number and/or type of polymers used for derivatization can be determined based on considerations including, but not limited to, the particular properties or functions of the antibody to be improved, whether the antibody derivative will be used in a therapy under defined conditions, etc.

J. Recombinant Methods and Compositions

Antibodies may be produced using recombinant methods and compositions, e.g., as described in U.S. Patent No. 4,816,567. In one embodiment, an isolated nucleic acid or a set of isolated nucleic acids encoding a set of antibodies described herein is provided.

For instance, a set of nucleic acids may comprise the following nucleic acids encoding the first antibody:

- i) a nucleic acid encoding a first heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VH domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3); ;
- ii) a nucleic acid encoding a Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (i) to form a binding site for a target antigen;
- iii) a nucleic acid encoding a second heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; and an Fc subunit (e.g., CH2-CH3)
- iv) a nucleic acid encoding a further Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (iii) to form a binding site for a target antigen.

A set of nucleic acids according to the invention may additionally or alternatively comprise the following nucleic acids encoding the second antibody:

- v) a nucleic acid encoding a first heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VL domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);
- vi) a nucleic acid encoding a Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (v) to form a binding site for a target antigen ;
- 10 vii) a nucleic acid encoding a second heavy chain polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; and an Fc subunit (e.g., CH2-CH3);
- viii) a nucleic acid encoding a further Fab light chain polypeptide (e.g., VL-CL) which pairs with the Fab heavy chain of (vii) to form a binding site for a target antigen.

In some embodiments, certain of these nucleic acids may be the same as each other. For instance, the nucleic acid in (ii) may be the same as in (iv), and/or the nucleic acid in (vi) may be the same as in (viii) such that the overall set comprises fewer than 8 distinct nucleic acid sequences.

In another embodiment a set of nucleic acids may comprise the following nucleic acids encoding the first antibody:

- i) a nucleic acid encoding a polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VH domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);
 - ii) a nucleic acid encoding a Fab light chain polypeptide (e.g., VL-CL); and
 - iii) a nucleic acid encoding an Fc subunit polypeptide (e.g., CH2-CH3),
- wherein the Fab heavy chain of (i) and the Fab light chain of (ii) form a Fab fragment
- 30 capable of binding to a target antigen.

A set of nucleic acids according to the invention may additionally or alternatively comprise the following nucleic acids encoding the second antibody:

iv) a nucleic acid encoding a polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VL domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);

v) a nucleic acid encoding a Fab light chain polypeptide (e.g., VL-CL), and

vi) a nucleic acid encoding an Fc subunit polypeptide (e.g., CH2-CH3);

wherein the Fab heavy chain of (iv) and the Fab light chain of (v) form a Fab fragment capable of binding to a target antigen.

Again, in some embodiments, certain of these nucleic acids may be the same as each other. For instance, the nucleic acid in (ii) may be the same as in (v), such that the overall set
10 comprises only 5 distinct nucleic acid sequences.

The nucleic acids can be comprised in one or more nucleic acid molecules or expression vectors.

Thus, in a further embodiment, one or more vectors (e.g., expression vectors) comprising such nucleic acid(s) are provided. In one embodiment, each respective heavy and light chain is expressed from an individual plasmid.

In a further embodiment, a host cell or a set of host cells comprising such nucleic acid(s) or vector(s) is provided. In one embodiment, a first host cell is provided expressing the first antibody, and a second host cell is provided expressing the second antibody.

In one such embodiment, a first host cell comprises one or more vectors which
20 collectively encode the nucleic acids of the first antibody. A second host cell comprises (e.g., has been transformed with) one or more vectors which collectively encode the nucleic acids of the second antibody.

In one embodiment, the host cell is eukaryotic, e.g. a Chinese Hamster Ovary (CHO) cell or lymphoid cell (e.g., Y0, NS0, Sp20 cell). In one embodiment, a method of making an antibody according to the invention is provided, wherein the method comprises culturing a host cell comprising nucleic acids encoding the antibody, as provided above, under conditions suitable for expression of the antibody, and optionally recovering the antibody from the host cell (or host cell culture medium).

For recombinant production of an antibody, nucleic acid encoding an antibody, e.g.,
30 as described above, is isolated and inserted into one or more vectors for further cloning and/or expression in a host cell. Such nucleic acid may be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of the antibody).

Suitable host cells for cloning or expression of antibody-encoding vectors include prokaryotic or eukaryotic cells described herein. For example, antibodies may be produced in bacteria, in particular when glycosylation and Fc effector function are not needed. For expression of antibody fragments and polypeptides in bacteria, see, e.g., US 5,648,237, US 5,789,199, and US 5,840,523. (See also Charlton, K.A., In: *Methods in Molecular Biology*, Vol. 248, Lo, B.K.C. (ed.), Humana Press, Totowa, NJ (2003), pp. 245-254, describing expression of antibody fragments in *E. coli*.) After expression, the antibody may be isolated from the bacterial cell paste in a soluble fraction and can be further purified.

10 In addition to prokaryotes, eukaryotic microbes such as filamentous fungi or yeast are suitable cloning or expression hosts for antibody-encoding vectors, including fungi and yeast strains whose glycosylation pathways have been “humanized”, resulting in the production of an antibody with a partially or fully human glycosylation pattern. See Gerngross, T.U., *Nat. Biotech.* 22 (2004) 1409-1414; and Li, H. et al., *Nat. Biotech.* 24 (2006) 210-215.

Suitable host cells for the expression of (glycosylated) antibody are also derived from multicellular organisms (invertebrates and vertebrates). Examples of invertebrate cells include plant and insect cells. Numerous baculoviral strains have been identified which may be used in conjunction with insect cells, particularly for transfection of *Spodoptera frugiperda* cells.

20 Plant cell cultures can also be utilized as hosts. See, e.g., US 5,959,177, US 6,040,498, US 6,420,548, US 7,125,978, and US 6,417,429 (describing PLANTIBODIESTM technology for producing antibodies in transgenic plants).

Vertebrate cells may also be used as hosts. For example, mammalian cell lines that are adapted to grow in suspension may be useful. Other examples of useful mammalian host cell lines are monkey kidney CV1 line transformed by SV40 (COS-7); human embryonic kidney line (293 or 293T cells as described, e.g., in Graham, F.L. et al., *J. Gen Virol.* 36 (1977) 59-74); baby hamster kidney cells (BHK); mouse sertoli cells (TM4 cells as described, e.g., in Mather, J.P., *Biol. Reprod.* 23 (1980) 243-252); monkey kidney cells (CV1); African green monkey kidney cells (VERO-76); human cervical carcinoma cells (HELA); canine kidney cells (MDCK; buffalo rat liver cells (BRL 3A); human lung cells (W138); human liver cells (Hep G2); mouse mammary tumor (MMT 060562); TRI cells (as described, e.g., in Mather, J.P. et al., *Annals N.Y. Acad. Sci.* 383 (1982) 44-68); MRC 5 cells; and FS4 cells. Other useful mammalian host cell lines include Chinese hamster ovary (CHO) cells, including DHFR- CHO cells (Urlaub, G. et al., *Proc. Natl. Acad. Sci. USA* 77 (1980) 4216-4220); and

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myeloma cell lines such as Y0, NS0 and Sp2/0. For a review of certain mammalian host cell lines suitable for antibody production, see, e.g., Yazaki, P. and Wu, A.M., *Methods in Molecular Biology*, Vol. 248, Lo, B.K.C. (ed.), Humana Press, Totowa, NJ (2004), pp. 255-268.

In one aspect, the host cell is eukaryotic, e.g., a Chinese Hamster Ovary (CHO) cell or lymphoid cell (e.g., Y0, NS0, Sp20 cell).

K. Assays

Antibodies provided herein may be identified, screened for, or characterized for their physical/chemical properties and/or biological activities by various assays known in the art.

10 In one aspect, an antibody of the invention is tested for its antigen binding activity, e.g., by known methods such as ELISA, Western blot, etc.

Antibody affinity

In certain embodiments, an antibody provided herein has a dissociation constant (K_D) for the target antigen of $\leq 1\mu\text{M}$, $\leq 100\text{ nM}$, $\leq 10\text{ nM}$, $\leq 1\text{ nM}$, $\leq 0.1\text{ nM}$, $\leq 0.01\text{ nM}$, or $\leq 0.001\text{ nM}$ (e.g., 10^{-8} M or less, e.g., from 10^{-8} M to 10^{-13} M , e.g., from 10^{-9} M to 10^{-13} M), or as otherwise stated herein.

In certain embodiments, an antigen binding site for the effector moiety, e.g., radiolabelled compound has a dissociation constant (K_D) for the effector moiety/radiolabelled compound of $\leq 1\mu\text{M}$, $\leq 100\text{ nM}$, $\leq 10\text{ nM}$, $\leq 1\text{ nM}$, $\leq 0.1\text{ nM}$, $\leq 0.01\text{ nM}$, or $\leq 0.001\text{ nM}$ (e.g., 10^{-8} M or less, e.g., from 10^{-8} M to 10^{-13} M , e.g., from 10^{-9} M to 10^{-13} M). In some
20 embodiments, the K_D is 1 nM or less, 500pM or less, 200pM or less, 100pM or less, 50pM or less, 20pM or less, 10pM or less, 5pM or less or 1pM or less, or as otherwise stated herein. For instance, the functional binding site may bind the radiolabelled compound/metal chelate with a K_D of about 1pM - 1nM , e.g., about 1 - 10 pM , 1 - 100pM , 5 - 50 pM , 100 - 500 pM or 500pM - 1 nM .

In one embodiment, K_D is measured by a radiolabelled antigen binding assay (RIA). In one embodiment, an RIA is performed with the Fab version of an antibody of interest and its antigen. For example, solution binding affinity of Fabs for antigen is measured by equilibrating Fab with a minimal concentration of (^{125}I)-labelled antigen in the presence of a
30 titration series of unlabelled antigen, then capturing bound antigen with an anti-Fab antibody-coated plate (see, e.g., Chen et al., *J. Mol. Biol.* 293:865-881(1999)). To establish conditions for the assay, MICROTITER[®] multi-well plates (Thermo Scientific) are coated overnight

with 5 µg/ml of a capturing anti-Fab antibody (Cappel Labs) in 50 mM sodium carbonate (pH 9.6), and subsequently blocked with 2% (w/v) bovine serum albumin in PBS for two to five hours at room temperature (approximately 23°C). In a non-adsorbent plate (Nunc #269620), 100 pM or 26 pM [¹²⁵I]-antigen are mixed with serial dilutions of a Fab of interest (e.g., consistent with assessment of the anti-VEGF antibody, Fab-12, in Presta et al., *Cancer Res.* 57:4593-4599 (1997)). The Fab of interest is then incubated overnight; however, the incubation may continue for a longer period (e.g., about 65 hours) to ensure that equilibrium is reached. Thereafter, the mixtures are transferred to the capture plate for incubation at room temperature (e.g., for one hour). The solution is then removed and the plate washed eight times with 0.1% polysorbate 20 (TWEEN-20[®]) in PBS. When the plates have dried, 150 µl/well of scintillant (MICROSCINT-20[™]; Packard) is added, and the plates are counted on a TOPCOUNT[™] gamma counter (Packard) for ten minutes. Concentrations of each Fab that give less than or equal to 20% of maximal binding are chosen for use in competitive binding assays.

According to another embodiment, K_D is measured using a BIACORE[®] surface plasmon resonance assay. For example, an assay using a BIACORE[®]-2000 or a BIACORE[®]-3000 (BIAcore, Inc., Piscataway, NJ) is performed at 25°C with immobilized antigen CM5 chips at ~10 response units (RU). In one embodiment, carboxymethylated dextran biosensor chips (CM5, BIACORE, Inc.) are activated with *N*-ethyl-*N*'-(3-dimethylaminopropyl)-carbodiimide hydrochloride (EDC) and *N*-hydroxysuccinimide (NHS) according to the supplier's instructions. Antigen is diluted with 10 mM sodium acetate, pH 4.8, to 5 µg/ml (~0.2 µM) before injection at a flow rate of 5 µl/minute to achieve approximately 10 response units (RU) of coupled protein. Following the injection of antigen, 1 M ethanolamine is injected to block unreacted groups. For kinetics measurements, two-fold serial dilutions of Fab (0.78 nM to 500 nM) are injected in PBS with 0.05% polysorbate 20 (TWEEN-20[™]) surfactant (PBST) at 25°C at a flow rate of approximately 25 µl/min. Association rates (k_{on}) and dissociation rates (k_{off}) are calculated using a simple one-to-one Langmuir binding model (BIACORE[®] Evaluation Software version 3.2) by simultaneously fitting the association and dissociation sensorgrams. The equilibrium dissociation constant (K_D) is calculated as the ratio k_{off}/k_{on} . See, e.g., Chen et al., *J. Mol. Biol.* 293:865-881 (1999). If the on-rate exceeds $10^6 \text{ M}^{-1} \text{ s}^{-1}$ by the surface plasmon resonance assay above, then the on-rate can be determined by using a fluorescent quenching technique that measures the increase or decrease in fluorescence emission intensity (excitation = 295 nm; emission = 340 nm, 16

nm band-pass) at 25°C of a 20 nM anti-antigen antibody (Fab form) in PBS, pH 7.2, in the presence of increasing concentrations of antigen as measured in a spectrometer, such as a stop-flow equipped spectrophotometer (Aviv Instruments) or a 8000-series SLM-AMINCO™ spectrophotometer (ThermoSpectronic) with a stirred cuvette.

In another embodiment, K_D is measured using a SET (solution equilibration titration) assay. According to this assay, test antibodies are typically applied in a constant concentration and mixed with serial dilutions of the test antigen. After incubation to establish an equilibrium, the portion of free antibodies is captured on an antigen coated surface and detected with labelled/tagged anti-species antibody, generally using electrochemiluminescence (e.g., as described in Haenel *et al* Analytical Biochemistry 339 (2005) 182-184).

For example, in one embodiment 384-well streptavidin plates (Nunc, Microcoat #11974998001) are incubated overnight at 4°C with 25 µl/well of an antigen-Biotin-Isomer Mix in PBS-buffer at a concentration of 20 ng/ml. For equilibration of antibody samples with free antigen: 0.01 nM - 1 nM of antibody is titrated with the relevant antigen in 1:3, 1:2 or 1:1.7 dilution steps starting at a concentration of 2500 nM, 500 nM or 100 nM of antigen. The samples are incubated at 4°C overnight in sealed REMP Storage polypropylene microplates (Brooks). After overnight incubation, streptavidin plates are washed 3x with 90 µl PBST per well. 15 µl of each sample from the equilibration plate is transferred to the assay plate and incubated for 15 min at RT, followed by 3x 90 µl washing steps with PBST buffer. Detection is carried out by adding 25 µl of a goat anti-human IgG antibody-POD conjugate (Jackson, 109-036-088, 1:4000 in OSEP), followed by 6x 90 µl washing steps with PBST buffer. 25 µl of TMB substrate (Roche Diagnostics GmbH, Cat. No.: 11835033001) are added to each well. Measurement takes place at 370/492 nm on a Safire2 reader (Tecan).

In another embodiment, K_D is measured using a KinExA (kinetic exclusion) assay. According to this assay, the antigen is typically titrated into a constant concentration of antibody binding sites, the samples are allowed to equilibrate, and then drawn quickly through a flow cell where free antibody binding sites are captured on antigen-coated beads, while the antigen-saturated antibody complex is washed away. The bead-captured antibody is then detected with a labelled anti-species antibody, e.g., fluorescently labelled (Bee *et al* PloS One, 2012; 7(4): e36261). For example, in one embodiment, KinExA experiments are performed at room temperature (RT) using PBS pH 7.4 as running buffer. Samples are prepared in running buffer supplemented with 1 mg/ml BSA ("sample buffer"). A flow rate of 0.25 ml/min is used. A constant amount of antibody with 5 pM binding site concentration

is titrated with antigen by twofold serial dilution starting at 100 pM (concentration range 0.049 pM – 100 pM). One sample of antibody without antigen serves as 100% signal (i.e. without inhibition). Antigen–antibody complexes are incubated at RT for at least 24 h to allow equilibrium to be reached. Equilibrated mixtures are then drawn through a column of antigen-coupled beads in the KinExA system at a volume of 5 ml permitting unbound antibody to be captured by the beads without perturbing the equilibrium state of the solution. Captured antibody is detected using 250 ng/ml Dylight 650©-conjugated anti-human Fc-fragment specific secondary antibody in sample buffer. Each sample is measured in duplicates for all equilibrium experiments. The KD is obtained from non-linear regression analysis of the data using a one-site homogeneous binding model contained within the KinExA software (Version 4.0.11) using the “standard analysis” method.

L. Therapeutic Methods and Compositions

The set of antibodies as described herein may be used in therapeutic methods. In one aspect a set of antibodies as described herein is provided for use as a medicament. In certain aspects, a set of antibodies for use in a method of treatment is provided.

In some aspects, a set of antibodies as described herein can be used as immunotherapeutic agents, for example in the treatment of proliferative disease, e.g., cancers. The treatment may induce lysis of a target cell, particularly a tumour cell.

As discussed above, in some aspects, sets of antibodies according to the present invention are suitable for any treatment in which it is desired to deliver a radionuclide to a target cell in a subject. For example, there is provided a set of antibodies as described herein for use in a method of pre-targeted radioimmunotherapy, e.g., for cancer treatment.

In certain aspects, the invention provides the set of antibodies for use in a method of immunotherapy or pre-targeted radioimmunotherapy in an individual comprising administering to the individual an effective amount of the set of antibodies. An “individual” according to any of the above aspects is preferably a human.

As noted above, the treatment may be of any condition that is treatable by cytotoxic activity targeted to diseased cells of the patient. The treatment is preferably of a tumour or cancer. However, the applicability of the invention is not limited to tumours and cancers. For example, the treatment may also be of viral infection, or infection by another pathogenic organism, e.g., a prokaryote. Optionally, targeting may also be to T-cells for treatment of T-cell driven autoimmune disease or T-cell blood cancers. Thus, conditions to be treated may include viral infections such as HIV, rabies, EBV and Kaposi’s sarcoma-associated

herpesvirus, and autoimmune diseases such as multiple sclerosis and graft-versus-host disease drugs.

The term “cancer” as used herein include both solid and hematologic cancers, such as lymphomas, lymphocytic leukemias, lung cancer, non small cell lung (NSCL) cancer, bronchioloalviolar cell lung cancer, bone cancer, pancreatic cancer including pancreatic ductal adenocarcinoma (PDAC), skin cancer, cancer of the head or neck, cutaneous or intraocular melanoma, uterine cancer, ovarian cancer, cancer of the anal region, stomach cancer, gastric cancer, colorectal cancer, which may be colon cancer and/or rectal cancer, breast cancer, uterine cancer, carcinoma of the fallopian tubes, carcinoma of the
10 endometrium, carcinoma of the cervix, carcinoma of the vagina, carcinoma of the vulva, Hodgkin's Disease, cancer of the esophagus, cancer of the small intestine, cancer of the endocrine system, cancer of the thyroid gland, cancer of the parathyroid gland, cancer of the adrenal gland, sarcoma of soft tissue, cancer of the urethra, cancer of the penis, prostate cancer, cancer of the bladder, cancer of the kidney or ureter, renal cell carcinoma, carcinoma of the renal pelvis, mesothelioma, hepatocellular cancer, biliary cancer, neoplasms of the central nervous system (CNS), spinal axis tumours, brain stem glioma, glioblastoma multiforme, astrocytomas, schwannomas, ependymomas, medulloblastomas, meningiomas, squamous cell carcinomas, pituitary adenoma and Ewings sarcoma, including refractory
20 versions of any of the above cancers, checkpoint-inhibitor experienced versions of any of the above cancers, or a combination of one or more of the above cancers.

Methods of treatment may comprise administering the first and second antibody simultaneously or sequentially.

In some embodiments, the antibodies described herein may be administered as part of a combination therapy. For example, they may be administered in combination with one or more chemotherapeutic agents: the chemotherapeutic agent and the antibody may be administered simultaneously or sequentially, in either order. Additionally or alternatively, they may be administered in combination with one or more immunotherapeutic: the immunotherapeutic agent and the antibody may be administered simultaneously or sequentially, in either order.

30 Antibodies of the invention (and any additional therapeutic agent, e.g., the radiolabelled compound) can be administered by any suitable means, including parenteral, intrapulmonary, and intranasal, and, if desired for local treatment, intralesional administration. Parenteral infusions include intramuscular, intravenous, intraarterial,

intraperitoneal, or subcutaneous administration. Dosing can be by any suitable route, e.g., by injections, such as intravenous or subcutaneous injections.

In one example, a method of targeting a radioisotope to a cell, tissue or organ for therapy may comprise:

- i) administering to the subject a first and a second antibody as described herein (simultaneously or sequentially, in either order), wherein the antibodies bind to the target antigen and localise to the surface of a cell expressing the target antigen; and wherein association of the first and second antibody forms a functional binding site for the radiolabelled compound;
- and
- ii) subsequently administering a radiolabelled compound, wherein the radiolabelled compound binds to functional binding site for the radiolabelled compound.

The radiolabelled compound is labelled with a radioisotope which is cytotoxic to cells. Suitable radioisotopes include alpha and beta emitters as discussed above.

In methods of pre-targeted radioimmunotherapy which make use of a bispecific antibody (i.e., not a “split” antibody according to the present invention) it is common practice to administer a clearing agent or a blocking agent, between administration of the antibody and administration of the radiolabelled compound. Clearing agents bind to the antibodies and enhance their rate of clearance from the body. They include anti-idiotypic antibodies.

Blocking agents are typically agents which bind to the antigen binding site for the radiolabelled compound, but which are not themselves radiolabelled. For example, where the radiolabelled compound comprises a chelator loaded with a radioisotope of a certain chemical element (e.g., a metal), the blocking agent may comprise the same chelator loaded with a non-radioactive isotope of the same element (e.g., metal), or may comprise a non-loaded chelator or a chelator loaded with a different non-radioactive moiety (e.g., a non-radioactive isotope of a different element), provided that it can still be bound by the antigen-binding site. In some cases, the blocking agent may additionally comprise a moiety which increases the size and/or hydrodynamic radius of the molecule. These hinder the ability of the molecule to access the tumour, without interfering with the ability of the molecule to bind to the antibody in the circulation. Exemplary moieties include hydrophilic polymers. The moiety may be a polymer or co-polymer e.g., of dextran, dextrin, PEG, polysialic acids (PSAs), hyaluronic acid, hydroxyethyl-starch (HES) or poly(2-ethyl 2-oxazoline) (PEOZ). In other embodiments

the moiety may be a non-structured peptide or protein such as XTEN polypeptides (unstructured hydrophilic protein polymers), homo-amino acid polymer (HAP), proline-alanine-serine polymer (PAS), elastin-like peptide (ELP), or gelatin-like protein (GLK). Further exemplary moieties include proteins such as albumin e.g., bovine serum albumin, or IgG. Suitable molecular weights for the moieties/polymers may be in the range e.g., of at least 50 kDa, for example between 50 kDa to 2000 kDa. For example, the molecular weight may be 200-800kDa, optionally greater than 300, 350, 400 or 450 kDa, and optionally less than 700, 650, 600 or 550kDa, optionally about 500kDa.

10 According to certain aspects of the present invention, there is no step of administering a clearing agent or a blocking agent to the subject. In certain aspects, there is no step of administering any agent which binds to the first or the second antibody, between the administration of the antibodies and the administration of the radiolabelled compound. In certain aspects, there is no step of administering any agent between the administration of the antibodies and the radiolabelled compound, except optionally a compound selected from a chemotherapeutic agent, immunotherapeutic and a radiosensitizer. In some embodiments, no agent is administered between the administration of the antibodies and the administration of the radiolabelled compound. In some embodiments there may be no injection or infusion of any other agent to the subject, between the administration of the antibody and the administration of the radiolabelled compound.

20 In some embodiments, the method may be a two-step method of pre-targeted radioimmunotherapy consisting or consisting essentially of the steps of i) administering the set of antibodies (wherein the first and second antibody may be administered simultaneously or sequentially in either order) and ii) subsequently administering the radiolabelled compound. The treatment may involve multiple cycles of such therapy, i.e., multiple cycles of these two steps. An exemplary treatment cycle duration is 28 days, in which the set of antibodies is administered on day 1 of the cycle, and the radiolabelled compound is optionally administered on day 1,2,3,4,5,6,7, or 8 of the cycle, e.g., on day 7. The number of therapeutic cycles may vary. In one embodiment, there may be 4, 5, or 6 treatment cycles.

30 The present inventors have surprisingly determined that using antibodies according to the invention, it is possible to obtain therapeutically effective uptake of the radiolabelled compound into the tumour, while avoiding excessive accumulation of radioactivity in normal tissues. Indeed, in the examples the level of accumulation of radioactivity in non-target

tissues was found to be lower than in a three-step PRIT method, using bispecific antibodies and a clearing step, while also making use of a simpler procedure.

In some embodiments, the radiolabelled compound may be administered to the subject once the first and second antibody have been given a suitable period of time to localise to the target cells. For instance, in some embodiments, the radiolabelled compound may be administered to the subject immediately after the first and second antibodies or at least 4 hours, 8 hours, 1 day, or 2 days, after the first and second antibodies. Optionally, it may be administered no more than 3 days, 5 days, or 7 days after the first and second antibodies. In one particular embodiment, the radiolabelled compound may be administered to the subject 2 to 7 days after the first and second antibodies.

In some embodiments, the antibodies described herein may additionally or alternatively be administered in combination with radiosensitizers. The radiosensitizer and the antibody may be administered simultaneously or sequentially, in either order.

In some embodiments, one or more dosimetry cycles may be used prior to one or more treatment cycles as described above. A dosimetry cycle may comprise the steps of i) administering the set of antibodies (wherein the first and second antibody may be administered simultaneously or sequentially in either order) and ii) subsequently administering a compound suitable for imaging radiolabelled with a gamma-emitter (wherein said radiolabelled compound binds to functional binding site for the radiolabelled compound). The compound may be the same as the compound used in the subsequent treatment cycles, except that it is labelled with a gamma emitter rather than an alpha or beta emitter. For example, in one embodiment, the radiolabelled compound used in the dosimetry cycle may be ^{203}Pb -DOTAM and the radiolabelled compound used in the treatment cycle may be ^{212}Pb -DOTAM. The patient may be subject to imaging to determine the uptake of the compound into the tumour and/or to estimate the absorbed dose of the compound. This information may be used to estimate the expected radiation exposure in subsequent treatment steps and to adjust the dose of the radiolabelled compound used in the treatment steps to a safe level.

M. Pharmaceutical Formulations

The first and second antibody described herein may be formulated in a single pharmaceutical composition or in separate pharmaceutical compositions. Thus, in a further aspect, the present invention provides a pharmaceutical composition comprising the first and second antibodies of the invention, or a first pharmaceutical formulation comprising the first

antibody of the invention and a second pharmaceutical composition comprising the second antibody of the invention, e.g., for use in any of the therapeutic or diagnostic methods described herein. In one embodiment, the pharmaceutical composition further comprises a pharmaceutically acceptable carrier. In another embodiment, the pharmaceutical composition further comprises at least one additional therapeutic agent, e.g., as described below.

Pharmaceutical formulations of antibodies as described herein may be prepared by mixing such antibody having the desired degree of purity with one or more optional pharmaceutically acceptable carriers (*Remington's Pharmaceutical Sciences* 16th edition, Osol, A. Ed. (1980)), in the form of lyophilized formulations or aqueous solutions.

10 Pharmaceutically acceptable carriers are generally nontoxic to recipients at the dosages and concentrations employed, and include, but are not limited to: buffers such as histidine, phosphate, citrate, acetate, and other organic acids; antioxidants including ascorbic acid and methionine; preservatives (such as octadecyldimethylbenzyl ammonium chloride; hexamethonium chloride; benzalkonium chloride; benzethonium chloride; phenol, butyl or benzyl alcohol; alkyl parabens such as methyl or propyl paraben; catechol; resorcinol; cyclohexanol; 3-pentanol; and m-cresol); low molecular weight (less than about 10 residues) polypeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids such as glycine, glutamine, asparagine, histidine, arginine, or lysine; monosaccharides, disaccharides, and other carbohydrates
20 including glucose, mannose, or dextrans; chelating agents such as EDTA; sugars such as sucrose, mannitol, trehalose or sorbitol; salt-forming counter-ions such as sodium; metal complexes (e.g., Zn-protein complexes); and/or non-ionic surfactants such as polyethylene glycol (PEG). Exemplary pharmaceutically acceptable carriers herein further include interstitial drug dispersion agents such as soluble neutral-active hyaluronidase glycoproteins (sHASEGP), for example, human soluble PH-20 hyaluronidase glycoproteins, such as rHuPH20 (HYLENEX®, Halozyme, Inc.). Certain exemplary sHASEGPs and methods of use, including rHuPH20, are described in US Patent Publication Nos. 2005/0260186 and 2006/0104968. In one aspect, a sHASEGP is combined with one or more additional glycosaminoglycanases such as chondroitinases.

30 Exemplary lyophilized antibody compositions are described in US Patent No. 6,267,958. Aqueous antibody compositions include those described in US Patent No. 6,171,586 and WO 2006/044908, the latter compositions including a histidine-acetate buffer.

The formulation herein may also contain more than one active ingredients as necessary for the particular indication being treated, preferably those with complementary activities that do not adversely affect each other. For example, it may be desirable to further provide chemotherapeutic agents, immunotherapeutic agents and/or radiosensitizers as discussed above. Such active ingredients are suitably present in combination in amounts that are effective for the purpose intended.

Active ingredients may be entrapped in microcapsules prepared, for example, by coacervation techniques or by interfacial polymerization, for example, hydroxymethylcellulose or gelatin-microcapsules and poly-(methylmethacrylate) microcapsules, respectively, in colloidal drug delivery systems (for example, liposomes, albumin microspheres, microemulsions, nano-particles and nanocapsules) or in macroemulsions. Such techniques are disclosed in *Remington's Pharmaceutical Sciences* 16th edition, Osol, A. Ed. (1980).

Sustained-release preparations may be prepared. Suitable examples of sustained-release preparations include semipermeable matrices of solid hydrophobic polymers containing the antibody, which matrices are in the form of shaped articles, e.g. films, or microcapsules.

The formulations to be used for *in vivo* administration are generally sterile. Sterility may be readily accomplished, e.g., by filtration through sterile filtration membranes.

N. Methods and Compositions for Diagnosis and Detection

Where the effector agent is a radiolabelled moiety, the set of antibodies as described herein may also be used in methods of diagnosis or imaging, preferably methods of or comprising pre-targeted radioimmunoimaging. Accordingly, the present invention provides methods of diagnosis and imaging. It further provides use of the set of antibodies in a method of imaging as described herein, and a set of antibodies as described herein (i.e., the first and second antibody as described herein) for use in a method of diagnosis carried out on a subject, e.g., on the human or animal body.

The imaging methods are suitable for imaging the presence and/or distribution of the target antigen in the body. For example, the method may be a method of imaging cells expressing an antigen associated with a disease, such as any of the disease conditions discussed above. Optionally the method is for imaging tumours or cancer. The method may

be for the purpose of diagnosing a subject suspected of having a proliferative disorder such as cancer, or an infectious disease.

In some embodiments it may be preferred that the subject is a human.

A method of targeting a radioisotope to a tissue or organ for imaging or diagnosis may comprise:

- i) administering to the subject a first and a second antibody as described herein (simultaneously or sequentially, in either order), wherein the antibodies bind to a target antigen and localise to the surface of a cell expressing the target antigen, wherein association of the first and second antibody forms a functional binding site for the radiolabelled compound;
- and
- ii) subsequently administering a radiolabelled compound, wherein the radiolabelled compound binds to the functional binding site for the radiolabelled compound.

Optionally, the method may further comprise:

- iii) imaging the tissue or organ where the radiolabelled compound has localised, or is expected to be localised.

Optionally, the method may further comprise one or more steps of forming a diagnosis, delivering a diagnosis to the subject, and/or determining and/or administering a suitable treatment on the basis of the diagnosis.

In another embodiment, a method of the invention may comprise imaging a tissue or organ of a subject, wherein the subject has been previously administered with:

- i) a first and a second antibody as described herein (simultaneously or sequentially, in either order), wherein the antibodies bind to a target antigen and localise to the surface of a cell expressing the target antigen, and wherein association of the first and second antibody forms a functional binding site for the radiolabelled compound; and
- ii) a radiolabelled compound, wherein the radiolabelled compound binds to the antigen binding site for said radiolabelled compound formed by association of the first and second antibody.

In imaging and/or diagnostic methods as described herein, the radiolabelled compound is labelled with a radioisotope which is suitable for imaging. Suitable radioisotopes include gamma emitters as discussed above.

In conventional methods of pre-targeted radioimaging, it is common practice to administer a clearing or blocking agent between the administration of the antibody and the administration of the radiolabelled compound, e.g., a clearing or blocking agent as described above.

In certain embodiments of the present invention, there is no step of administering a clearing agent or a blocking agent. In certain aspects, there is no step of administering any agent which binds to the first or the second antibody, between the administration of the antibodies and the administration of the radiolabelled compound. In certain aspects, there is no step of administering any agent between the administration of the antibodies and the radiolabelled compound, except optionally a compound selected from a chemotherapeutic agent, immunotherapeutic agent and a radiosensitizer. In some embodiments, no agent is administered between the administration of the antibodies and the administration of the radiolabelled compound. In some embodiments there may be no injection or infusion of any other agent to the subject, between the administration of the antibody and the administration of the radiolabelled compound.

In some embodiments, the radiolabelled compound may be administered to the subject once the first and second antibody have been given a suitable period of time to localise to the target cells. For instance, in some embodiments, the radiolabelled compound may be administered to the subject immediately after the first and second antibodies or at least 4 hours, 8 hours, 1 day, or 2 days after the first and second antibodies. Optionally, it may be administered no more than 3 days, 5 days, or 7 days after the first and second antibodies. In one particular embodiment, the radiolabelled compound may be administered to the subject 2 to 7 days after the first and second antibodies.

In some embodiments, the imaging method may be a method of pre-targeted radioimaging consisting or consisting essentially of the steps of i) administering the set of antibodies (wherein the first and second antibody may be administered simultaneously or sequentially in either order) ii) subsequently administering the radiolabelled compound and iii) imaging the tissue or organ of interest. A diagnostic method may consist or consist essentially of said steps followed by steps of forming a diagnosis, which may then be delivered to the patient and may be used as the basis for selected and/or administering a treatment regimen.

The target antigen may be any target antigen as discussed herein. In some embodiments, the target antigen may be a tumour-specific antigen as discussed above, and

the imaging may be a method of imaging a tumour or tumours. The individual may be known to or suspected of having a tumour.

For example, the method may be a method of imaging tumours in an individual having or suspected of having lung cancer, non small cell lung (NSCL) cancer, bronchioloalviolar cell lung cancer, bone cancer, pancreatic cancer including PDAC, skin cancer, cancer of the head or neck, cutaneous or intraocular melanoma, uterine cancer, ovarian cancer, colorectal cancer which may be rectal cancer and/or colon cancer, cancer of the anal region, stomach cancer, gastric cancer, breast cancer, uterine cancer, carcinoma of the fallopian tubes, carcinoma of the endometrium, carcinoma of the cervix, carcinoma of the
10 vagina, carcinoma of the vulva, Hodgkin's Disease, cancer of the esophagus, cancer of the small intestine, cancer of the endocrine system, cancer of the thyroid gland, cancer of the parathyroid gland, cancer of the adrenal gland, sarcoma of soft tissue, cancer of the urethra, cancer of the penis, prostate cancer, cancer of the bladder, cancer of the kidney or ureter, renal cell carcinoma, carcinoma of the renal pelvis, mesothelioma, hepatocellular cancer, biliary cancer, neoplasms of the central nervous system (CNS), spinal axis tumours, brain stem glioma, glioblastoma multiforme, astrocytomas, schwannomas, ependymomas, medulloblastomas, meningiomas, squamous cell carcinomas, pituitary adenoma and Ewings sarcoma, including refractory versions of any of the above cancers, or check-point inhibitor experienced versions of any of these cancers, or a combination of one or more of the above
20 cancers.

III. SEQUENCES

SEQ ID NO	Description	Sequence
1	heavy chain CDR1, <Pb-Dotam>	GFSLSTYSMS
2	heavy chain CDR2 <Pb-Dotam>	FIGSRGDTYYASWAKG
3	heavy chain CDR3 <Pb-Dotam>	ERDPYGGGAYPPHL
4	light chain CDR1, <Pb-Dotam>	QSSHVSYSNDLA
5	light chain CDR2 <Pb-Dotam>	QASKLAS
6	light chain CDR3 <Pb-Dotam>	LGGYDDESDTYG
7	heavy chain variable domain	VTLKESGPVLVKPTETLTLTCTVSGFSLSTYSMSWIRQPPGKALEWLGFIGSRGDTYYASWAKGRLTISKDTSKSQ VVLTMTNMDPVDATATYYCARERDPYGGGAYPPHLWGRGTLVTVSS

	<Pb-Dotam> PRIT-0213	
8	light chain variable domain <Pb-Dotam> PRIT-0213	IQMTQSPSSLASVGDRLVTTTCQSSHVSYSNDLAWYQQKPGKAPKLLIYQASKLASGVPSRFSGSGTGDTFTLI SSLQPEDFATYYCLGGYDDESDTYGFGGGTKVEIK
9	heavy chain variable domain <Pb-Dotam> PRIT-0214	VQLQQWGAGLLKPSETLSLTCAVYGFSLSTYSMSWIRQQPGKGLEWIGFIGSRGDTYYASWAKGRVTISR DTSKNQVSLKLSVTAADTAVYYCARERDPYGGGAYPPHLWGRGTLVTVSS
10	light chain variable domain <Pb-Dotam> PRIT-0214	IQMTQSPSSLASVGDRLVTTTCQSSHVSYSNDLAWYQQKPGKAPKLLIYQASKLASGVPSRFSGSGSGT DFTLTISLQPEDFATYYCLGGYDDESDTYGFGGGTKVEIK
11	heavy chain CDR1 <CEA> T84.66	GFNIKDTYMH
12	heavy chain CDR2 <CEA> T84.66	RIDPANGNSKYVPKFQG
13	heavy chain CDR3 <CEA> T84.66	FGYYVSDYAMAY

14	light chain CDR1 <CEA> T84.66	RAGESVDIFGVGFLH
15	light chain CDR2 <CEA> T84.66	RASNRAT
16	light chain CDR3 <CEA> T84.66	QQTNEDPYT
17	heavy chain variable domain <CEA> T84.66	QVQLVQSGAEVKKPGSSVKVCKASGFNIKDTYMHWVRQAPGQGLEWMGRIDPANGNSKYVPKFQGRVTITA DTSTSTAYMELSSLRSEDTAVYYCAPFGYYVSDYAMAYWGQGTLVTVSS
18	light chain variable domain <CEA> T84.66	EIVLTQSPATLSLSPGERATLSCRAGESVDIFGVGFLHWYQQKPGQAPRLIYRASNRATGIPARFSGSGGTDFTL TISSLEPEDFAVYYCQQTNEDPYTFGQGTKLEIK
19	heavy chain CDR1 <CEA> CH1A1A	GYTFTEFGMN
20	heavy chain CDR2 <CEA> CH1A1A	WINTKTGEATYVEEFKG
21	heavy chain CDR3 <CEA> CH1A1A	WDFAYYVEAMDY
22	light chain CDR1	KASAAVGTYYA

	<CEA> CH1A1A	
23	light chain CDR2 <CEA> CH1A1A	SASYRKR
24	light chain CDR3 <CEA> CH1A1A	HQYYTYPLFT
25	heavy chain variable domain <CEA> CH1A1A	QVQLVQSGAEVKKPGASVKVCKKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTKTEATYVEEFKGRVTFRTT DTSTSTAYMELRSLRSDDTAVYYCARWDFAYYVEAMDYWGQGTITVTVSS
26	light chain variable domain <CEA> CH1A1A	DIQMTQSPSSLSASVGDRTVTITCKASAAVGTYYVAWYQQKPGKAPKLLIYSASYRKRGVPSRFSGSGGTDFTLTIS SLQPEDFATYYCHQYYTYPLFTFGQGTKLEIK
27	Heavy chain <CEA> of PIAD8749 without linker and <DOTAM- VH>→ Same Plasmid as SeqID32, lacking linker and <DOTAM>	QVQLVQSGAEVKKPGASVKVCKKASGYTFTEFGMNWVRQAPGQGLEWMGW INTKTKTEATYVEEFKGRVTFRTTDTSTSTAYMELRSLRSDDTAVYYCARWD FAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLV KDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQ TYICNVNHKPSNTKVDKKVEPKSCDKTHTCTPPCPAPEAAGGSPVFLFPPK PKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTKSKAKGQPREP QVYTLPPCRDELTKNQVSLWCLVKGFYPSDIAVEWESNGQPENNYKTTTPP VLDSDGSFFFLYSKLTVDKSRWQQGNVVFSCVMHEALHNHYTQKSLSLSPG

28	PIAD8749 Heavy chain hole <CEA> CH1A1A	QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTKTEATYVEEFKGRVTFIT DTSTSTAYMELSLRSDDDTAIVYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTSKAKGQPREPQVCTLPPSRDELTKNQVS LSCAVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLVSKLTVDKSRWQQGNVFSCVMHEALHNHYTQ KSLSLSPG
29	Heavy chain <CEA> of PIAD8592 without linker and <DOTAM- VL>→ Same Plasmid as SeqID33, lacking linker and <DOTAM>	QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPGQGLEWMGW INTKTKTEATYVEEFKGRVTFITDTSTSTAYMELSLRSDDDTAIVYCARWD FAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLV KDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQ TYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPEAAGGPSVFLFPPK PKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTSKAKGQPREP QVCTLPPSRDELTKNQVSLSCAVKGFYPSDIAVEWESNGQPENNYKTTTP VLDSDGSFFLVSKLTVDKSRWQQGNVFSCVMHEALHNHYTQKSLSLSPG
30	PIAD8592 Heavy chain Knob <CEA>CH1A1A	QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTKTEATYVEEFKGRVTFIT DTSTSTAYMELSLRSDDDTAIVYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTSKAKGQPREPQVCTLPPSRDELTKNQVS LWCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLVSKLTVDKSRWQQGNVFSCVMHEALHNHYT

		QKSLSPG	
31	Linker	GGGGSGGGSGGGSGGGGS	
32	P1AD8749 heavy chain knob <CEA>CH1A1A <Dotam-VH>	QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPQGQGLEWMGWINTKTGEATYVEEFKGRVTFTTDTSTSTAYMELRSLRSDDDTAVYYCARWDFAYYVEAMDYWGGQTTVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSVVTVPPSSSLGTQTYICNVNHHKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPKPDKDLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIISKAKGQPREPQVYTLPPCRDELTKNQVS LWCLVKGFYPSDIAVEWESNGQPENNYKTTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT QKSLSLSPG <u>GGGGSGGGSGGGSGGGGS</u> VTLKESGPVLVKPTETLTCTVSGFSLSYSTYSMSWIRQPPGKALE WLGFIGSRGDTYYASWAKRLTISKDTSKSQVVLTMTNMDPVDATYYCAREDPYGGGAYPPHLWGRTLV TVSS	
33	P1AD8592 heavy chain hole <CEA>CH1A1A <Dotam-VL>	QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPQGQGLEWMGWINTKTGEATYVEEFKGRVTFTTDTSTSTAYMELRSLRSDDDTAVYYCARWDFAYYVEAMDYWGGQTTVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSVVTVPPSSSLGTQTYICNVNHHKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPKPDKDLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIISKAKGQPREPQVCTLPPSRDELTKNQVS LSCAVKGFYPSDIAVEWESNGQPENNYKTTTPPVLDSDGSFFLVSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQ KLSLSPG <u>GGGGSGGGSGGGSGGGSIQM</u> TQSPSSLASVGDRVTITCQSSHVSYSNDNLAWYQQKPGKAP KLLIYQASKLASGVPSRFSGSGGTDFLTITSLQPEDFATYYCLGGYDDESDTYGFGGGTKVEIK	
34	P1AD8749 and P1AD8592 light chain <CEA> CH1A1A	DIQMTQSPSSLSASVGDRVITITCKASAAVGTYYVAWYQQKPGKAPKLLIYS ASYRKRGVPSPRSFGSGSGTDFTLTITSLQPEDFATYYCHQYITYTYPLFTFG QGTKLEIKRTVAAPSVEIFPPPSDEQLKSGTASVVCLLNNFYPREAKVQWK VDNALQSGNSQESVTEQDSKDSYSTYLSSTLTLSKADYEKHKVYACEVTHQ	

		GLSSPVTKSFNRGEC	
35	Heavy chain CDR 1, <C825>	DYGVH	
36	Heavy chain CDR 2, <C825>	VIWSGGGTAYNTALIS	
37	Heavy chain CDR 3, <C825>	RGSYPYNYFDA	
38	Light chain CDR 1, <C825>	GSSTGAVTASNYAN	
39	Light chain CDR 2, <C825>	GHNNRPP	
40	Light chain CDR 3, <C825>	ALWYSDHWV	
41	Heavy chain variable domain <C825>	HVKLQESGPGLVQPSQSLTCTVSGFSLTDYGVHVVRRQSPGKGLEWLGVIWSSGGGTAYNTALISRLNIYRDNS KNQVFLEMNSLQAEDTAMYYCARRGSYPYNYFDAWGQGTTVTVSS	
42	Light chain variable domain, <C825>	QAVVIQESALTTTPGETVTLTCGSSTGAVTASNYANWVQEKPDHLFTGLIGGHNNRPPGVPARFSGSLIGDKAAL TIAGTQTEDEAIYFCALWYSDHWVIGGGTKLTVL	
43	heavy chain CDR1 <CEA> A5B7	DYYMN	

44	heavy chain CDR2 <CEA> A5B7	FIGNKANAYTTTEYSASVKG
45	heavy chain CDR3 <CEA> A5B7	DRGLRFYFDY
46	light chain CDR1 <CEA> A5B7	RASSSVTYIH
47	light chain CDR2 <CEA> A5B7	ATSNLAS
48	light chain CDR3 <CEA> A5B7	QHWSSKPPT
49	heavy chain variable domain <CEA> A5B7	EVQLLESGGGLVQPGGSLRLSCAASGFTFTDYYMNWVVRQAPGKGLEWLGFIGNKANAYTTTEYSASVKG RFTISR DKSKNTLYLQMNSLRAEDTATYYCTRDRLRFYFDYWGGGTTTVTVSS
50	light chain variable domain <CEA> A5B7	EIVLTQSPATLSLSPGERATLSCRASSSVTYIHVYQQKPGQAPRSWIYATSNLASGIPARFSGSGGTDFLTITSSLE PEDFAVYYCQHWSSKPPFTFGQGTKLEIK

[illegible]

[illegible]

	and DOTAM-VL> → Same Plasmid as SeqID 56, lacking linker and <DOTAM>	EQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTKSKAKGQPREPQVCTLPSPSRDELTKNQVSLSCAVKGFYPSDIAVEWESNGQPENNYKTTTPPVLDSDGSFFLVSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSP
58	P1AE4957 light chain <CEA> A5B7	EIVLTQSPATLSLSPGERATLSCRASSSVTYIHVYQQKPGQAPRSWIYATSNLASGIPARFSGSGGTDFLTISSEPEDFAVYYCQHWSSKPTFGGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC
59	heavy chain CDR1 <CEA> 28A9	GGTFSYYAIS
60	heavy chain CDR2 <CEA> 28A9	GILPAFGAANYAQKFQG
61	heavy chain CDR3 <CEA> 28A9	LPPLPGAGLDY
62	light chain CDR1 <CEA> 28A9	RASQSISSWLA

63	light chain CDR2 <CEA> 28A9	DASSLES
64	light chain CDR3 <CEA> 28A9	QQNTQYPMT
65	heavy chain variable domain <CEA> 28A9	QVQLVQSGAEVKKPGSSVKVCKASGGTFSYYAISWVRQAPGQGLEWMGGILPAFGAANYAQKFQGRVTITAD KSTSTAYMELSSLRSEDTAVYYCARLPPLPGAGLDYWGGGTTVTVSS
66	light chain variable domain <CEA> 28A9	DIQMTQSPSTLSASVGDRVTITCRASQSISSWLAWYQQKPGKAPKLLIYDASSLES <u>GVPSRFS</u> SGSGSGTE FTLTISLQPD ~~DF~~ ATYYCQQNTQYPM <u>TF</u> GQGGTKVEIK
67	heavy chain CDR1 <GPRC5D>	GFTFSKYAMA
68	heavy chain CDR2 <GPRC5D>	SISTGGVNTYYADSVKG
69	heavy chain CDR3 <GPRC5D>	HTGDYFDY

70	light chain CDR1 <GPRC5D>	RASQSVSISGINLMN
71	light chain CDR2 <GPRC5D>	HASILAS
72	light chain CDR3 <GPRC5D>	QQTRESPLT
73	Heavy chain variable domain <GPRC5D>	EVQLLESGGGLVQPGGSLRLSCAASGFTFSKYAMAWVRQAPGKGLEWVASISTGGVNTYYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCAHTGTGDFDYWGQGTMTVTVSS
74	Light chain variable domain <GPRC5D>	EIVLTQSPGTLSPGERATLSCRASQSVSISGINLMNWYQQKPGQPKLLIYHASILASGIPDRFSGSGSGTDFTLT ISRLEPEDFAVYYCQQTRESPLTFGQGTRLEIK
75	heavy chain CDR1 <FAP> 4B9	GFTFSSYAMS
76	heavy chain CDR2 <FAP> 4B9	AIIGSGASTYYADSVKGG

77	heavy chain CDR3 <FAP> 4B9	GWFGGFNY
78	light chain CDR1 <FAP> 4B9	RASQSVTSSYLA
79	light chain CDR2 <FAP> 4B9	VGSRRAT
80	light chain CDR3 <FAP> 4B9	QQGIMLPPT
81	Heavy chain variable domain <FAP> 4B9	EVQLLESGGGLVQPGGSLRLSCAASGFTSSYAMSWVRQAPGKGLWVSAIIGSGASTYYADSVKGRFTISRDN KNTLYLQMNSLRAEDTAVYYCAKGWFGGFNYWGQGTLLVTVSS
82	Light chain variable domain <FAP> 4B9	EIVLTQSPGTLSPGERATLSCRASQSVTSSYLAWYQQKPGQAPRLLINVGSRRAATGIPDRFSGSGSGTDFTLTISR LEPEDFAVYYCQQGIMLPPTFGQGTKVEIK
83	P1AF0709 HCknob <CEA> T84.66	QVQLVQSGAEVKKPGSSVKVCKASGFNIKDTYIHWVRQAPGQGLEWMGRIDPANGNSKYVPKFQGRVTITA DTSTSTAYMELSSLRSEDTAVYYCAPFGYYVSDYAMAYWGQGTLLVTVSSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHNKPSNTKVDKKVEP

	(D1AE4688)	KSCDKTHTCPPCPAEEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKP REEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTISKAKGQPREPQVYTLPPCRDELTKNQVSL WCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNNHYYTQ KSLSLSPG
84	P1AF0709 HChole <CEA> T84.66 Dotam- VL (D1AA4920)	QVQLVQSGAEVKKPGSSVKVCKASGFNIKDTYMHWVRQAPGQGLEWMGRIDPANGNSKYVPKFQGRVTITA DTSTSTAYMELSSLRSEDTAVYYCAPFGYYVSDYAMAYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVDKKVEP KSCDKTHTCPPCPAEEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKP REEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTISKAKGQPREPQVCTLPSPRDELTKNQVSL SCAVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNNHYYTQ KSLSLPGGGGGGGGGGGGGGGGGGGSIQMTQSPSSLSASVGDRTITCQSSSHSVYSDNDLAWYQQKPGKAPK LLIYQASKLASGVPSRFSGSGGTDFITLTISSLQPEDFATYYCLGGYDDESDTYGFGGGTKVEIK
85	P1AF0709 HChole <CEA> T84.66 without linker and DOTAM	QVQLVQSGAEVKKPGSSVKVCKASGFNIKDTYMHWVRQAPGQGLEWMGRIDPANGNSKYVPKFQGRVTITA DTSTSTAYMELSSLRSEDTAVYYCAPFGYYVSDYAMAYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVDKKVEP KSCDKTHTCPPCPAEEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKP REEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTISKAKGQPREPQVCTLPSPRDELTKNQVSL SCAVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNNHYYTQ KSLSLSPG
86	P1AF0298 HChole <CEA> T84.66 (D1AE4687)	QVQLVQSGAEVKKPGSSVKVCKASGFNIKDTYMHWVRQAPGQGLEWMGRIDPANGNSKYVPKFQGRVTITA DTSTSTAYMELSSLRSEDTAVYYCAPFGYYVSDYAMAYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVDKKVEP KSCDKTHTCPPCPAEEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKP

		REEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTISKAKGQPREPQVCTLPSPRDELTKNQVSL SCAVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLVSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQ KSLSLSPG
87	PIAF0298 HCknob <CEA> T84.66 Dotam- VH-AST (D1AE3668)	QVQLVQSGAEVKKPGSSVKVCKASGFNIKDTYMHWVRQAPGQGLEWMGRIDPANGNSKYVPKFQGRVTITA DTSTSTAYMELSSLRSEDTAVYYCAPFGYYVSDYAMAYWGQGTLLVTVSSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVDKKVEP KSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKP REEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTISKAKGQPREPQVYTLPPCRDELTKNQVSL WCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQ KSLSLSPGGGGGGGGGGGGGGVTLKESGPVLVKPTETLLTCTVSGFSLSTYSMSWIRQPPGKALEWL GFIGSRGDTYYASWAKGRLTISKDTKSQVLTMTNMDPVDATYYCARERDPYGGGAYPPHLWGRGTLVTVS SAST
88	PIAF0298 HCknob <CEA> T84.66 without linker and DOTAM	QVQLVQSGAEVKKPGSSVKVCKASGFNIKDTYMHWVRQAPGQGLEWMGRIDPANGNSKYVPKFQGRVTITA DTSTSTAYMELSSLRSEDTAVYYCAPFGYYVSDYAMAYWGQGTLLVTVSSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVDKKVEP KSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKP REEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTISKAKGQPREPQVYTLPPCRDELTKNQVSL WCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQ KSLSLSPG
89	PIAF0709 and PIAF0298 light chain (D1AA4120)	EIVLTQSPATLSLSPGERATLSCRAGESVDIFGVGFLHWYQQKPGQAPRLLIYRASNRATGIPARFSGSGSGTDFTL TISSLEPEDFAVYYCQQTNEPDYTFGQGTKEIKRTVAAPSVFIFPPSDEQLKSGTASVVCCLLNNFYPREAKVQWK VDNALQSGNSQESVTEQDSKDSITYSLSSITLTSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

90	P1AF0710 HC'nob <CEA> 28A9 (D1AE4690)	QVQLVQSGAEVKKPGSSVKVCKASGGTFSYYAISWVRQAPGQGLEWMGGILPAFGAANYAQKFQGRVTITAD KSTSTAYMELSSLRSEDTAVYYCARLPPLPGAGLDYWGGGTTVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCL VKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVVDKKVEPKSC DKTHTCPCPAPEAAAGGPSVFLFPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVYTLPPCRDELTKNQVSLWCL VKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQGNVVFSCVMHEALHNHYTQKSLS LSPG
91	P1AF0710 HC'hole <CEA> 28A9 Dotam- VL (D1AC3172)	QVQLVQSGAEVKKPGSSVKVCKASGGTFSYYAISWVRQAPGQGLEWMGGILPAFGAANYAQKFQGRVTITAD KSTSTAYMELSSLRSEDTAVYYCARLPPLPGAGLDYWGGGTTVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCL VKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVVDKKVEPKSC DKTHTCPCPAPEAAAGGPSVFLFPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVCTLPPSRDELTKNQVSLSCA VKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQGNVVFSCVMHEALHNHYTQKSLS LSPGGGGGGGGGGGGGGGGGGSIQMTQSPSSLSASVGDRTITTCQSSHSVYSDNDLAWYQQKPGKAPKLLIY QASKLASGVPSRFSGSGGTDFTLTISLQPEDFATYYCLGGYDDESDTYGFGGGTKVEIK
92	P1AF0710 HC'hole <CEA> 28A9 without linker or DOTAM	QVQLVQSGAEVKKPGSSVKVCKASGGTFSYYAISWVRQAPGQGLEWMGGILPAFGAANYAQKFQGRVTITAD KSTSTAYMELSSLRSEDTAVYYCARLPPLPGAGLDYWGGGTTVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCL VKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVVDKKVEPKSC DKTHTCPCPAPEAAAGGPSVFLFPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVCTLPPSRDELTKNQVSLSCA VKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQGNVVFSCVMHEALHNHYTQKSLS LSPG

93	P1AF0711 HChole <CEA> 28A9 (D1AE4689)	QVQLVQSGAEVKKPGSSVKVCKASGGTFSYYAISWVRQAPGQGLEWMGGILPAFGAANYAQKFQGRVTITAD KSTSTAYMELSSLRSEDTAVYYCARLPPLPGAGLDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCL VKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVDKKVEPKSC DKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVCTLPPSRDELTKNQVSLSCA VKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQKSLS LSPG
94	P1AF0711 HCknob <CEA> 28A9 Dotam- VH-AST (D1AE3671)	QVQLVQSGAEVKKPGSSVKVCKASGGTFSYYAISWVRQAPGQGLEWMGGILPAFGAANYAQKFQGRVTITAD KSTSTAYMELSSLRSEDTAVYYCARLPPLPGAGLDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCL VKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVDKKVEPKSC DKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVYTLPPCRDELTKNQVSLWCL VKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQKSLS LSPGGGGGGGGGGGGGGSVTLKESGPVLVKPTETLTLCTVSGFSLSTYSMSWIRPPGKALEWLFIG SRGDTYYASWAKGRLTISKDTSKSQVVLTMNMDPVDATYYCARERDPYGGGAYPPHLWGRGTLVTVSSAST
95	P1AF0711 HCknob <CEA> 28A9 without linker and DOTAM	QVQLVQSGAEVKKPGSSVKVCKASGGTFSYYAISWVRQAPGQGLEWMGGILPAFGAANYAQKFQGRVTITAD KSTSTAYMELSSLRSEDTAVYYCARLPPLPGAGLDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCL VKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTKVDKKVEPKSC DKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVYTLPPCRDELTKNQVSLWCL VKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQKSLS LSPG
96	P1AF0710 and	DIQMTQSPSTLSASVGDRTVITCRASQSISSWLAWYQQKPGKAPKLLIYDASSLESGVPSRFSGSGSGTEFTLTISL

	P1AF0711 light chain (D1AA2299)	QPDDEATYYCQONTQYPMFTGGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDN ALQSGNSQESVTEQDSKDSITYSLSSITLSKADYEKHKVYACEVTHQGLSPVTKSFNRGEC
97	P1AF0712 HCnob <CEA> CH1A1A (D1AC4023)	QVQLVQSGAEVKKPGASVKVCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTKTEATYVEEFKGRVTFTT DTSTSTAYMELSLRSDDDTAVYYCARWDFAYYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHNKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVYTLPPCRDELTKNQVS LWCLVKGYFSPDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNHYT QKSLSLSPG
98	P1AF0712 HChole <CEA> CH1A1A DOTA-VL (D1AE4684)	QVQLVQSGAEVKKPGASVKVCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTKTEATYVEEFKGRVTFTT DTSTSTAYMELSLRSDDDTAVYYCARWDFAYYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHNKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVCTLPPSRDELTKNQVS LSCAVKGYFSPDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQ KSLSLSPGGGGGGGGGGGGGGGSAVVIQESALITPPGETVTLTCSGSTGAVTASNYYANWVQEKPDHLF TGLIGGHNNRPPGVPARFSGSLIGDKAALTIAGTQTEDEAIYFCALWYSDHWVIGGGTKLTVL
99	P1AF0712 HChole <CEA> without linker or DOTA	QVQLVQSGAEVKKPGASVKVCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTKTEATYVEEFKGRVTFTT DTSTSTAYMELSLRSDDDTAVYYCARWDFAYYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHNKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVCTLPPSRDELTKNQVS

		LSCAVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLVSKLTVDKSRWQQGNVFSCVMHEALHNHYTQ KSLSLSPG
100	P1AF0713 HCHole <CEA> CH1A1A (D1AC4022)	QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTGEATYVVEEFKGRVTFTT DTSTSTAYMELSLRSDDDTAVYYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTTISKAKGQPREPQVCTLPPSRDELTKNQVS LSCAVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLVSKLTVDKSRWQQGNVFSCVMHEALHNHYTQ KSLSLSPG
101	P1AF0713 HCknob <CEA> CH1A1A DOTA-VH-AST (D1AE3670)	QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTGEATYVVEEFKGRVTFTT DTSTSTAYMELSLRSDDDTAVYYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTTISKAKGQPREPQVYTLPPCRDELTKNQVS LWCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLVSKLTVDKSRWQQGNVFSCVMHEALHNHYT QKLSLSPGGGGGGGGGGGGGGGGGGGSHVKLQESGPGLVQPSQSLSLTCTVSGFSLTDYGVHWVVRQSPGKGL EWLGVIWSSGGGTAYNTALISRLNIYRDNSKNQVFLEMNSLQAEDTAMYYCARRGSYPYNYFDAGWGQGTITVTVS SAST
102	P1AF0713 HCknob <CEA> CH1A1A without linker	QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTGEATYVVEEFKGRVTFTT DTSTSTAYMELSLRSDDDTAVYYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVE PKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTTISKAKGQPREPQVYTLPPCRDELTKNQVS

	and DOTA	LWCLVKGFYPSDIAVEWESNGQPENNYKTTTPPVLSDSDGSFFLYSKLTVDKSRWQQQGNVFCSCVMHEALHNHYT QKSLSPG
103	P1AF0712 and P1AF0713 light chain (D1AA3384)	DIQMTQSPSSLSASVGDRVITITCKASAAVGTVAWYQQKPGKAPKLLIYSASVYRKRGVPSRFSGSGTDFTLTIS SLQPEDFATYYCHQYYTYPLFTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVCLLNNFYPREAKVQWKV DNALQSGNSQESVTEQDSKDSYSTLSSTLTLISKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC
104	P1AF8284 and P1AF8285 HCknob <GPRC5D> (D1AF6517)	EVQLLESGGGLVQPGGSLRLSCAASGFTFSKYAMAWVRQAPGKGLEWVASISTGGVNTYYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCAHTHTGDYFDYWGGQTMVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLVK DYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHHKPSNTKVDKKVEPKSCDK THTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVYTLPPCRDELTKNQVSLWCLVK GFYPSDIAVEWESNGQPENNYKTTTPPVLSDSDGSFFLYSKLTVDKSRWQQQGNVFCSCVMHEALHNHYTQKSLSL PGK
105	P1AF8284 HChole Dotam- VL (D1AG3592)	DKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVCTLPPSRDELTKNQVSLSCA VKGFYPSDIAVEWESNGQPENNYKTTTPPVLSDSDGSFFLYSKLTVDKSRWQQQGNVFCSCVMHEALHNHYTQKSL LSPGKGGGGGGGGGGGGSSIQMTQSPSSLSASVGDRVITITCQSSHVSYSNDLAWYQQKPGKAPKLLI YQASKLASGVPSRFSGSGTDFTLTISSLQPEDFATYYCLGGYDDESDTYGFGGGTKVEIK
106	P1AF8285 Hhole Dotam- VHA (D1AG3591)	DKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTIKAKGQPREPQVCTLPPSRDELTKNQVSLSCA VKGFYPSDIAVEWESNGQPENNYKTTTPPVLSDSDGSFFLYSKLTVDKSRWQQQGNVFCSCVMHEALHNHYTQKSL LSPGKGGGGGGGGGGGGSVTLKESGPVLVKPTETLTLCTVSGFSLSTYSMSWIRQPPGKALEWLFI GSRGDTYYASWAKGRLTISKDTSKSQVVLTMNMDPVDATATYTCARERDPYGGGAYPPHLWGRGTLVTVSSA

[illegible]

	P1AF8287 light chain (D1AB9974)	ALQSGNSQESVTEQDSKDYSLSSLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC
112	P1AF7782 and P1AF7784 HCknob <CEA> CH1A1A (D1AD3419)	QVQLVQSGAEVKKPGASVKVCKASGYTFTEFGMNWVRQAPGQGQGLEWMGWINTKTGEATYVVEEFKGRVTFTTDTSTSTAYMELRSLRSDDTAVYYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHKPSNTKVDKKVE PKSCDKTHTCTPPCPAPEAAAGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTK PREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTKAKGQPREPQVYTLPPCRDELTKNQVS LWCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCSCVMHEALHNHYT QKSLSLSPGK
113	P1AF7782 HChole Dotam-VL (D1AG2237)	SIQMTQSPSSLSASVGDRVTITCQSSHVYSDNDLAWYQQKPGKAPKLLIYQASKLASGVPSRFSGSGSGTDFTLT ISSLPEDFATYYCLGGYDDESDTYGFGGGTKVEIKGGGGSGGGSGGGGDKTHTCTPPCPAPEAAGGP SVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQD WNLNGKEYKCKVSNKALGAPIEKTKAKGQPREPQVCTLPSPRDELTKNQVSLSCAVKGFYPSDIAVEWESNGQP ENNYKTTPPVLDSDGSFFLVSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQKSLSLSPGK
114	P1AF7784 HChole Dotam-VH (D1AG2236)	GVTLKESGPVLVKPTETLTLTCTVSGFSLSTYSMSWIRPPGKALEWLGFIGSRGDTYYASWAKGRLTISKDTSKS QVVLTMTNMDPVDTATYYCARERDPYGGGAYPPHLWGRGTLVTVSSGGGGSGGGSGGGGDKTHT CPPCPAPEAAAGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNST YRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTKAKGQPREPQVCTLPSPRDELTKNQVSLSCAVKGFY PSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLVSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQKSLSLSPGK
115	P1AF7782 and P1AF7784 light chain	DIQMTQSPSSLSASVGDRVTITCKASAAVGTYYVAWYQQKPGKAPKLLIYSASYRKRGVPSRFSGSGSGTDFLTITIS SLQPEDFATYYCHQYYTYPLFTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKV DNALQSGNSQESVTEQDSKDYSLSSLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC

	chain (D1AD3421)	
116	heavy chain CDR1 <CEA> MFE23	DSYMH
117	heavy chain CDR2 <CEA> MFE23	WIDPENGDT EYAPKFQG
118	heavy chain CDR2 <CEA> MFE23-H26	WIDPENGGTNYAQKFQG
119	heavy chain CDR3 <CEA> MFE23	GTP TGPYYFDY
120	light chain CDR1 <CEA> MFE23	SASSSVSYMH
121	light chain CDR1 <CEA> MFE23-L24, L25	RASSSVSYMH

122	light chain CDR1 <CEA> MFE23-L26	RASQSISSYM
123	light chain CDR2 <CEA> MFE23	STSNLAS
124	Light chain CDR2 <CEA> MFE23-L26	YTSNLAS
125	Light chain CDR2 <CEA> MFE23-L29	STSSLQS
126	light chain CDR3 <CEA> MFE23	QQRSSYPLT
127	Heavy chain variable domain <CEA> MFE23	QVKLQQSGAELVRSGTSVKLSCTASGFNIKDSYMHWLRQGPEQGLEWIGWIDPENGDTEYAPKFGQKATFTTDT SSNTAYLQLSSLTSEDTA VYYCNEGTPGPYYFDYWGQGTTVTVSS
128	Light chain variable domain <CEA> MFE23	ENVLTQSPAIMASPGKEKVTITCSASSSVSYMHWFFQKPGTSPKLVWYSTSNLASGVPARFSGSGSGTSYSLTISRM EAEDAATYYCQQRSSYPLTFGAGTKLELK

129	MFE-H24	QVQLVQSGAEVKKPGASVKVSCKASGFNIKDSYMHWVRQAPGQGLEWMGWIDPENGDTTEYAPKFQGRVTMT TDTSTAYMELSRSDDTAVYYCNEGTPPTGPIYFDYWGQGTLLVTVSS
130	MFE-H25	QVQLVQSGAEVKKPGASVKVSCKASGYTFKDSYMHWVRQAPGQGLEWMGWIDPENGDTTEYAPKFQGRVTMT TDTSTAYMELSRSDDTAVYYCNEGTPPTGPIYFDYWGQGTLLVTVSS
131	MFE-H26	QVQLVQSGAEVKKPGASVKVSCKASGFNIKDSYMHWVRQAPGQGLEWMGWIDPENGDTTEYAPKFQGRVTMT TDTSTAYMELSRSDDTAVYYCNEGTPPTGPIYFDYWGQGTLLVTVSS
132	MFE-H27	QVQLVQSGAEVKKPGASVKVSCKASGFNIKDSYMHWVRQAPGQGLEWMGWIDPENGDTTEYAPKFQGRVTMT TDTSTAYMELSRSDDTAVYYCARGTPPTGPIYFDYWGQGTLLVTVSS
133	MFE-H28	QVQLVQSGAEVKKPGASVKVSCKASGFNIKDSYMHWVRQAPGQGLEWMGWIDPENGDTTEYAPKFQGRVTMT RDTSTAYMELSRSDDTAVYYCNEGTPPTGPIYFDYWGQGTLLVTVSS
134	MFE-H29	QVQLVQSGAEVKKPGSSVKVSCKASGFNIKDSYMHWVRQAPGQGLEWMGWIDPENGDTTEYAPKFQGRVTITT DESTSTAYMELSSLRSEDTAVYYCNEGTPPTGPIYFDYWGQGTLLVTVSS
135	MFE-L24	DIQMTQSPSSLSASVGDRVTITCRASSSVSYMHWYQQKPGKAPKLLIYSTSNLASGVPSRFSGSGGTDFTLTISSL QPEDFATYYCQQRSSYPPLTFGGGKLEIK
136	MFE-L25	EIQMTQSPSSLSASVGDRVTITCRASSSVSYMHWYQQKPGKAPKLLIYSTSNLASGVPSRFSGSGGTDFTLTISSL QPEDFATYYCQQRSSYPPLTFGGGKLEIK
137	MFE-L26	EIQMTQSPSSLSASVGDRVTITCRASQSISSYMHWYQQKPGKAPKLLIYSTSNLASGVPSRFSGSGGTDFTLTISSL QPEDFATYYCQQRSSYPPLTFGGGKLEIK
138	MFE-L27	EIQMTQSPSSLSASVGDRVTITCRASSSVPYMHWYQQKPGKAPKLLIYSTSNLASGVPSRFSGSGGTDFTLTISSV QPEDFATYYCQQRSSYPPLTFGGGKLEIK
139	MFE-L28	EIQMTQSPSSLSASVGDRVTITCRASSSVPYMHWYQQKPGKAPKLLIYSTSNLASGVPSRFSGSGGTDFTLTISSV QPEDFATYYCQQRSSYPPLTFGGGKLEIK
140	MFE-L29	EIQMTQSPSSLSASVGDRVTITCRASSSVPYMHWYQQKPGKAPKLLIYSTSNLASGVPSRFSGSGGTDFTLTISSV

		QPEDFATYYCQQRSSYP LTFGGG TKLEIK
141	A2 domain of CEA	PKPFITSNNSNPVEDEDAVALTCEPEIQNTTYLWWVNNQSLPVPRLQLSNDNR TLTLLSVTRNDVGP YECGIQNKLSVDHSDPVILN
142	A1 domain of CEA	PKPSISSNNKPVEDKDAVAFTCEPETQDATYLVWVNNQSLPVPRLQLSNGNR TLT LFNVTRNDTAS YKCETQNPVSARRSDSVILN
143	D1AD3421 LC <CEA> CH1A1A	DIQMTQSPSSLSASVGDRTVTITCKASAAVGTVAWYQQKPGKAPKLLIYSASYRKRGVPSRFSGSGTDFLTIS SLQPEDFATYYCHQYYTYPLFTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVCLLNFFYPREAKVQWKV DNALQSGNSQESVTEQDSKDSYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC
144	D1AC3981 HCHole	DKTHTCPPCPAEEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREE QYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEK TISKAKGQPREPQVCTLPPSRDELTKNQVSLSCA VKGFYPSDIAVEWESNGQPENNYKTTTPVLDSGDSFFLVSKLTVDKSRWQQGNVVFSCVMHEALHNHYTQKSLS LSPGK
145	D1AF7527 HCnob <CEA> CH1A1A Dotam-VL	QVQLVQSGAEVKKPGASVKVCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTGEATYVEEFKGRVTFTT DTSTSTAYMELRSLRSDDTAVYYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHNKPSNTKVDKKVE PKSCGGGGSGGGSGGGSGGSIQMTQSPSSLSASVGDRVTITCQSSHVSVDNDLAWYQQKPGKAPKLLI YQASKLASGVPSRFSGSGGTDFLTITSLQPEDFATYYCLGGYDDESDTYGFGGGTKVEIKGGGGSGGGSGGG GSGGGGDKTHTCCPPCPAEEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNA KTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEK TISKAKGQPREPQVYTLPPCRDELTKN QVSLWCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSGDSFFLVSKLTVDKSRWQQGNVVFSCVMHEALHN HYTQKSLSLSPGK
146	D1AF7526	QVQLVQSGAEVKKPGASVKVCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTKTGEATYVEEFKGRVTFTT

	HCknob <CEA> CH1A1A Dotam-VH	DTSTSTAYMELRSLRSDDDTAVYYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHKPSNTKVDKKVE PKSCGGGGGGGGGGGGGGGVTLKESGVPVLVKPTETLTLTCTVSGFSLSTYSMSWIRQPPGKALEWLGFI GSRGDTYYASWAKGRLTISKDTSKSQVVLTMTNMDPVDATATYCARERDPYGGGAYPPHLWGRGTLVTVSSG GGGGGGGGGGGGGGGGDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDVSHEDPEVKFN WYVVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPIEKTSKAKGQPREPQVY TLPPCRDELTKNQVSLWCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDPKSRWQQGNVVF SCSVMHEALHNHYTQKSLSLSPGK
147	D1AC6742 HCHole <CEA> CH1A1A	QVQLVQSGAEVKKPGASVKVSCKASGYTFTTEFGMNWVVRQAPGQGLEWMGWINTKGTGEATYVEEFK GRVTFRTDSTSTAYMELRSLRSDDDTAVYYCARWDFAYYVEAMDYWGQGTITVTVSSASTKGPSVFPL APSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPEAAGGPSVFLFPPKPKDITLMISRTPEVTCVVVDV SHEDPEVKFNWYVVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPI EKTISKAKGQPREPQVCTLPSPRDELTKNQVSLSCAVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSD GSFFLVSKLTVDPKSRWQQGNVVFSCSVMHEALHNHYTQKSLSLSPGK

IV. EXAMPLES

The following are examples of methods and compositions of the invention. It is understood that various other embodiments may be practiced, given the general description provided above.

GLOSSARY OF ABBREVIATIONS

	ADA	Anti-drug antibody
	AST	Alanine, serine, threonine
10	BsAb	Bispecific antibody
	CA	Clearing agent
	CEA	Carcinoembryonic antigen
	DOTAM	1,4,7,10-tetrakis(carbamoylmethyl)-1,4,7,10-tetraazacyclododecane
	ID	Injected dose
15	ELISA	Enzyme-linked immunosorbent assay
	FAP	Fibroblast activation protein
	GPRC5D	G-protein coupled receptor family C group 5 member D
	IV	Intravenous
	MW	Molecular weight
20	PBS	Phosphate-buffered saline
	p.i.	Post injection
	PK	Pharmacokinetic
	PRIT	Pretargeted radioimmunotherapy
	RIT	Radioimmunotherapy
25	RT	Room temperature
	SC	Subcutaneous
	SCID	Severe combined immunodeficiency
	SD	Standard deviation
	SOPF	Specific and opportunistic pathogen-free
30	SPLIT	SeParated v-domains LInkage Technology
	TA	Target antigen
	TGI	Tumor growth inhibition
	TR	Tumor regression

Example 1: Generation of CEA-Split-DOTAM VH/VL antibodies

Methods of PRIT (Pretargeted radioimmunotherapy) using bispecific antibodies having a binding site for the target antigen and a binding site for the radiolabelled compound commonly use a clearing agent (CA) between the administrations of antibody and radioligand, to ensure effective targeting and high tumour-to-normal tissue absorbed dose ratios (see Figure 3). In an example of one such method, injected BsAb is allowed sufficient time for penetrating into the tumours, generally 4–10 days, after which circulating BsAb is neutralized using a Pb-DOTAM-dextran-500 CA. The CA blocks ^{212}Pb -DOTAM binding to nontargeted BsAb without penetrating into the tumour, which would block the pretargeted sites. This pretargeting regimen allows efficient tumour accumulation of the subsequently administered radiolabelled chelate, ^{212}Pb -DOTAM.

However, in methods involving a clearing agent, the use of a CA introduces a further step to the method which is inefficient. Moreover, it can be important to choose the timing and dosing of the CA administration with care, which is a complicating factor.

To address the problems associated with use of a clearing agent, the present inventors have proposed a strategy of splitting the DOTAM VL and VH domains, such that they are found on separate antibodies.

The generation of exemplary split DOTAM VH/VL antibodies is discussed further below

Generation of plasmids for the recombinant expression of antibody heavy or light chains

Desired proteins were expressed by transient transfection of human embryonic kidney cells (HEK 293). For the expression of a desired gene/protein (e.g. full length antibody heavy chain, full length antibody light chain, or a full length antibody heavy chain containing an additional domain (e.g. an immunoglobulin heavy or light chain variable domain at its C-terminus) a transcription unit comprising the following functional elements was used:

- the immediate early enhancer and promoter from the human cytomegalovirus (P-CMV) including intron A,
- a human heavy chain immunoglobulin 5'-untranslated region (5'UTR),
- a murine immunoglobulin heavy chain signal sequence (SS),
- a gene/protein to be expressed, and

- the bovine growth hormone polyadenylation sequence (BGH pA).

In addition to the expression unit/cassette including the desired gene to be expressed the basic/standard mammalian expression plasmid contained

- an origin of replication from the vector pUC18 which allows replication of this plasmid in *E. coli*, and
- a beta-lactamase gene which confers ampicillin resistance in *E. coli*.

a) Expression plasmid for antibody heavy chains

Antibody heavy chain encoding genes including C-terminal fusion genes comprising a complete and functional antibody heavy chain, followed by an additional antibody V-heavy or V-light domain was assembled by fusing a DNA fragment coding for the respective sequence elements (V-heavy or V-light) separated each by a G4Sx4 linker to the C-terminus of the CH3 domain of a human IgG molecule (VH-CH1-hinge-CH2-CH3-linker-VH or VH-CH1-hinge-CH2-CH3-linker-VL). Recombinant antibody molecules bearing one VH and one VL domain at the C-termini of the two CH3 domains, respectively, were expressed using the knob-into-hole technology.

The expression plasmids for the transient expression of an antibody heavy chain with a C-terminal VH or VL domain in HEK293 cells comprised besides the antibody heavy chain fragment with C-terminal VH or VL domain expression cassette, an origin of replication from the vector pUC18, which allows replication of this plasmid in *E. coli*, and a beta-lactamase gene which confers ampicillin resistance in *E. coli*. The transcription unit of the antibody heavy chain fragment with C-terminal VH or VL domain fusion gene comprises the following functional elements:

- the immediate early enhancer and promoter from the human cytomegalovirus (P-CMV) including intron A,
- a human heavy chain immunoglobulin 5'-untranslated region (5'UTR),
- a murine immunoglobulin heavy chain signal sequence,
- an antibody heavy chain (VH-CH1-hinge-CH2-CH3-linker-VH or VH-CH1-hinge-CH2-CH3-linker-VL) encoding nucleic acid, and
- the bovine growth hormone polyadenylation sequence (BGH pA).

b) Expression plasmid for antibody light chains

Antibody light chain encoding genes comprising a complete and functional antibody light chain was assembled by fusing a DNA fragment coding for the respective sequence elements.

The expression plasmid for the transient expression of an antibody light chain comprised besides the antibody light chain fragment an origin of replication from the vector pUC18, which allows replication of this plasmid in *E. coli*, and a beta-lactamase gene which confers ampicillin resistance in *E. coli*. The transcription unit of the antibody light chain fragment comprises the following functional elements:

- the immediate early enhancer and promoter from the human cytomegalovirus (P-CMV) including intron A,
- a human heavy chain immunoglobulin 5'-untranslated region (5'UTR),
- a murine immunoglobulin heavy chain signal sequence,
- an antibody light chain (VL-CL) encoding nucleic acid, and
- the bovine growth hormone polyadenylation sequence (BGH pA).

Transient expression of the antibody molecules

The antibody molecules were generated in transiently transfected HEK293 cells (human embryonic kidney cell line 293-derived) cultivated in F17 Medium (Invitrogen Corp.). For transfection "293-Free" Transfection Reagent (Novagen) was used. The respective antibody heavy- and light chain molecules as described above were expressed from individual expression plasmids. Transfections were performed as specified in the manufacturer's instructions. Immunoglobulin-containing cell culture supernatants were harvested three to seven (3-7) days after transfection. Supernatants were stored at reduced temperature (e.g. -80°C) until purification.

General information regarding the recombinant expression of human immunoglobulins in e.g. HEK293 cells is given in: Meissner, P. et al., *Biotechnol. Bioeng.* 75 (2001) 197-203.

The PRIT Hemibodies (split antibodies) were purified by a MabSelect Sure (Affinity Chromatography) and followed by Superdex 200 (Size Exclusion Chromatography).

Sequences of exemplary antibodies/hemibodies are summarised below.

Antibody name	First heavy chain	Second heavy chain	Light chain
P1AD8592	SEQ ID NO: 30	SEQ ID NO: 33	SEQ ID NO: 34
P1AD8749	SEQ ID NO: 28	SEQ ID NO: 32	SEQ ID NO: 34

P1AE4956	SEQ ID NO: 51	SEQ ID NO: 52	SEQ ID NO: 54
P1AE4957	SEQ ID NO 55	SEQ ID NO 56	SEQ ID NO: 58

For the PRIT Split Antibody with DOTAM-VL -P1AD8592 5mg with a concentration of 1.372mg/mL and a purity >96% based on analytical SEC and CE-SDS were produced. For the PRIT Split Antibody with DOTAM-VH - P1AD8749 14mg with a concentration of 2.03mg/mL and a purity >91% based on analytical SEC and CE-SDS were produced.

Antibodies P1AE4956 and P1AE4957 were also generated. For the PRIT Split Antibody with DOTAM-VL -P1AE4957, 19 mg with a concentration of 2.6mg/mL and a purity >81.6% based on analytical SEC and CE-SDS were produced. For the PRIT Split Antibody with DOTAM-VH - P1AE4956, 6.9mg with a concentration of 1.5mg/mL and a purity >90% based on analytical SEC and CE-SDS were produced. ESI-MS was used too confirm the identity of the PRIT hemibodies.

Example 2: FACS Analysis of Split Antibody Functionality

To assess the functionality of the split antibodies or hemibodies, MKN-45 cells were detached from the culture vessel using accutase at 37°C for 10 minutes. Subsequently, the cells were washed twice in PBS, and seeded into 96 well v-bottom plates to a final density of 4×10^6 cells/well.

The hemibodies P1AD8749 and P1AD8592 and a human ISO control were mixed 1:1 added to the cells in concentrations as indicated in Fig 5. Subsequently, the cells were incubated for 1 h on ice and washed twice in PBS. The cell pellet was resuspended and 40µl/well of detection reagent was added, either <human IgG(H+L)>FITC, (10µg/ml) or Pb_Dotam_FITC 1:100 => (10µg/ml) in PBS / 5% FCS. After 60 min incubation on ice, the cells were washed twice in PBS and resuspended in 200µl PBS / 5% FCS for measurement of FITC fluorescence using a FACS canto.

To assess the binding capability of the hemibodies to CEA on MKN-45 cells, they were detected using of antibodies using human IgG specific secondary antibodies (Figure 5). As expected, no significant binding of the human ISO control is observed on these cells.

When adjusted to the same IgG concentration, both hemibodies as well as the combination of both shows a dose dependent binding to MKN-45 cells, with a pronounced hook effect at very high concentrations as expected. This experiment demonstrates that the CEA binding is functional in the hemibodies.

To assess the binding capability of the hemibodies to DOTAM, they were bound to the cells either in the presence of a human ISO control or their respective split antibody partner in a 1:1 ratio. After their binding to MKN-45 cells, the cells were washed to remove unbound antibody. Subsequently, Pb-DOTAM-FITC (fluorescently labelled Pb-DOTAM) was added to detect DOTAM binding competent cell bound antibodies (Figure 6). As expected, no significant FITC is observed on these cells when one of the split antibody partners is combined with the of the human ISO control. Only a combination of both hemibodies in a 1:1 ratio shows a dose dependent FITC signal. This experiment shows that the DOTAM binding site becomes functional when both hemibodies come together on one cell.

EXAMPLE 3: IN VIVO STUDIES

Example 3a: Materials and Methods - General

All experimental protocols were reviewed and approved by the local authorities (Comité Régional d’Ethique de l’Expérimentation Animale du Limousin [CREEAL], Laboratoire Départemental d’Analyses et de Recherches de la Haute-Vienne). Female severe combined immunodeficiency (SCID) mice (Charles River) were maintained under specific and opportunistic pathogen free (SOPF) conditions with daily cycles of light and darkness (12 h/12 h), in line with ethical guidelines. No manipulations were performed during the first 5 days after arrival, to allow the animals to acclimatize to the new environment. Animals were controlled daily for clinical symptoms and detection of adverse events.

Solid xenografts were established by subcutaneous (SC) injection of CEA-expressing tumor cells in cell culture media mixed 1:1 with Corning® Matrigel® basement membrane matrix (growth factor reduced; cat No. 354230). Tumor volumes were estimated through manual caliper 3 times per week, calculated according to the formula: $volume = 0.5 \times length \times width^2$. Additional tumor measurements were made as needed depending on the tumor growth rate.

Mice were euthanized before the scheduled endpoint if they showed signs of unamenable distress or pain due to tumor burden, side effects of the injections, or other

causes. Indications of pain, distress, or discomfort include, but are not limited to, acute body weight (BW) loss, scruffy fur, diarrhea, hunched posture, and lethargy. The BW of treated animals was measured 3 times per week, with additional measurements as needed depending on the health status. Wet food was provided to all mice starting the day after the radioactive injection, for 7 days or until all individuals had recovered sufficiently from any acute BW loss. Mice whose BW loss exceeded 20% of their initial BW or whose tumor volume reached 3000 mm³ were euthanized immediately. Other factors taken into account for euthanasia for ethical reasons were tumor status (e.g. necrotic areas, blood/liquid leaking out, signs of automutilation) and general appearance of the animal (e.g. fur, posture, movement).

To minimize re-ingestion of radioactive urine/feces, all efficacy study mice were placed in cages with gridded floors for 4 hours after ²¹²Pb-DOTAM administration, before being transferred to new cages with standard bedding. All cages were then changed at 24 hours post injection (p.i.). This procedure was not performed for mice sacrificed for biodistribution purposes within 24 hours after the radioactive injection.

Blood was collected at the time of euthanasia from the venous sinus using retro-orbital bleeding on anesthetized mice, before termination through cervical dislocation followed by additional tissue harvest for radioactive measurements and/or histological analysis, as mandated by the protocols. Unexpected or abnormal conditions were documented. Tissues collected for formalin fixation were immediately put in 10% neutral buffered formalin (4°C) and then transferred to phosphate-buffered saline (PBS; 4°C) after 5 days. Organs and tissues collected for biodistribution purposes were weighed and measured for radioactivity using a 2470 WIZARD² automatic gamma counter (PerkinElmer), and the percent injected dose per gram of tissue (% ID/g) subsequently calculated, including corrections for decay and background.

Statistical analysis was performed using GraphPad Prism 7 (GraphPad Software, Inc.) and JMP 12 (SAS Institute Inc.). Curve analysis of tumor growth inhibition (TGI) was performed based on mean tumor volumes using the formula:

$$TGI = 100 - \frac{V_{\text{treatment}d} - V_{\text{treatment}0}}{V_{\text{ref},d} - V_{\text{ref},0}} \times 100$$

where *d* indicates study day and *0* the baseline value. Vehicle was selected as the reference group. Tumor regression (TR) was calculated according to:

$$TR = \frac{V_{\text{treatment}0} - V_{\text{treatment}d}}{V_{\text{treatment}0}}$$

where positive values indicated tumor regression, and values below -1 growth beyond the double baseline value.

Test compounds

The compounds utilized in the described studies are presented in the tables below, respectively for bispecific antibodies, clearing agents, and radiolabeled chelates.

CEA-DOTAM (RO7198427, PRIT-0213) is a fully humanized BsAb targeting the T84.66 epitope of CEA (see also WO2019/201959). PRIT-0213 is composed of

- i) a first heavy chain as shown below;
- ii) a second heavy chain as shown below; and
- iii) two antibody light chains as shown below.

Description	SEQUENCE
light chain of PRIT-00213	1 eivltqspat lslspgerat lscragesvd ifgvglhwy qqkpgqaprl 51 liyrasnrat giparfsgsg sgtdfilitis slepedfavy ycqqtndepy 101 tfgqgtklei krtvaapsvf ifppsdeqlk sgtasvvcil nnfypreakv 151 qwkvdnalqs gnsqesvteq dskdstysls stltlskady ekhkvyacev 201 thqglsspvt ksfnrgec (SEQ ID NO: 89)
heavy chain 1 PRIT-0213	1 qvqlvqsgae vkkpgssvkv sckasgfnik dymhwvrqa pgqglewmgr 51 idpangnsky vpkfqgrvti tadtststay melsslrse tavyycapfg 101 yyvsdyamay wgqgtlvtvs sastkgpsvf plapssksts ggtaalgclv 151 kdyfpepvtv swnsgaltsg vhtfpavlqs sglyslssv tvpssslgtq 201 tyicnvnhkp sntkvdkkve pkscdkthtc ppcpapeaag gpsvflfppk 251 pkdtlmisrt pevtcvvvdv shedpevkfn

	wyvdgvevhn aktkpreeqy 301 nstyrvvsvl tvlhqdwlng keykckvsnk algapiekti skakgqprep 351 qvytlppcrd eltknqvslw clvkgfypsd iavewesngq pennykttp 401 vldsdsffl yskltvdkr wqqgnvfscs vmhealhnhy tqkslslspg 451 gggsgggsgs gggsgggsgs vtlkesgpvl vkptetltlt ctvsgfslst 501 ysmwirqpp gkalewlgfi gsgdtyyas wakgrltisk dtsksqvlt 551 mtnmdpvdta tyycarerdp ygggaypphl wgrgtlvtvs s (SEQ ID NO: 152)
heavy chain 2 of PRIT-0213	1 qvqlvqsgae vkkpgsskv sckasgnik dtymhwvrqa pgqglewmgr 51 idpangnsky vpkfqgrvti tadtststay melsslr sed tavyycapfg 101 yyvsdyamay wgqgtlvtvs sastkgpsvf plapssksts ggtaalgclv 151 kdyfpepvtv swnsgaltsg vhtfpavlqs sglyslssv tvpssslgtq 201 tyicvnhkp sntkvdkkve pkscdkthtc ppcpapeaag gpsvflfppk 251 pkdtlmisrt pevtevvvdv shedpevkfn wyvdgvevhn aktkpreeqy 301 nstyrvvsvl tvlhqdwlng keykckvsnk algapiekti skakgqprep 351 qvctlpssrd eltknqvsls cavkgfypsd iavewesngq pennykttp 401 vldsdsffl vskltvdkr wqqgnvfscs vmhealhnhy tqkslslspg 451 gggsgggsgs gggsgggsgs iqmtqspssl sasvgdrvti tcqsshsvs

	501 dndlawyqqk pgkapklliy qasklasgvp srfsgsgsgt dftltisslq 551 pedfatyycl ggyddesdy gfgggtkvei k (SEQ ID NO: 84)
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DIG-DOTAM (RO7204012) is a non-CEA-binding BsAb used as a negative control.

P1AD8749, P1AD8592, P1AE4956, and P1AE4957 are CEA-split-DOTAM-VH/VL antibodies targeting the CH1A1A or A5B7 epitopes of CEA. Their sequences are described
5 above. All antibody constructs were stored at -80°C until the day of injection when they were thawed and diluted in standard vehicle buffer (20 mM Histidine, 140 mM NaCl; pH 6.0) or 0.9% NaCl to their final respective concentrations for intravenous (IV) or intraperitoneal (IP) administration.

10 The Pb-DOTAM-dextran-500 CA (RO7201869) was stored at -20°C until the day of injection when it was thawed and diluted in PBS for IV or IP administration.

The DOTAM chelate for radiolabeling was provided by Macrocyclics and maintained at -20°C before radiolabeling, performed by Orano Med (Razès, France). ^{212}Pb -DOTAM (RO7205834) was generated by elution with DOTAM from a thorium generator, and subsequently quenched with Ca after labeling. The ^{212}Pb -DOTAM solution was diluted with
15 0.9% NaCl to obtain the desired ^{212}Pb activity concentration for IV injection. Mice in vehicle control groups received multiple injections of vehicle buffer instead of BsAb, CA, and ^{212}Pb -DOTAM.

Bispecific antibodies

Compound	Target	Protocols
CEA-DOTAM (RO7198427, PRIT-0213)	T84.66	144, 158, 160
DIG-DOTAM (RO7204012)	Digoxigenin	160
CEA-split-DOTAM-VH P1AD8749	CH1A1A	144, 158
CEA-split-DOTAM-VH-AST P1AF0171	CH1A1A	175, 185, 189
CEA-split-DOTAM-VL P1AD8592	CH1A1A	144, 158, 175, 185, 189
CEA-split-DOTAM-VH	A5B7	158

P1AE4956		
CEA-split-DOTAM-VL P1AE4957	A5B7	158
CEA-split-DOTAM- VH-AST P1AF0298	T84.66	185, 189
CEA-split-DOTAM-VL P1AF0709	T84.66	185, 189

Clearing agents

Compound	Protocols
Ca-DOTAM-dextran-500 (RO7201869)	144, 158, 160

Radiolabeled chelates

Compound	Quenching	Protocols
²¹² Pb-DOTAM (RO7205834)	Ca	144, 158, 160, 175, 185, 189
²¹² Pb-DOTAM-CEA- DOTAM	Ca	160

5

Tumor models

The tumor cell line used and the injected amount for inoculation in mice is described in the table below. BxPC3 is a human primary pancreatic adenocarcinoma cell line, naturally expressing CEA. Cells were cultured in RPMI 1640 Medium, GlutaMAX™ Supplement, HEPES (Gibco, ref. No. 72400-021) enriched with 10% fetal bovine serum (GE Healthcare Hyclone SH30088.03). Solid xenografts were established in each SCID mouse on study day 0 by subcutaneous injection of cells in RPMI media mixed 1:1 with Corning® Matrigel® basement membrane matrix (growth factor reduced; cat No. 354230), into the right flank.

10

Tumor cell lines

Cell line	Cells per mouse	Injected volume	Protocols	Supplier
BxPC3	5×10 ⁶	100 µL	144, 158, 160, 175, 185, 189	ECACC*

15 *European Collection of Authenticated Cell Cultures (Salisbury, UK)

EXAMPLE 3b: Protocol 144

The aim of protocol 144 was to provide PK and in vivo distribution data of pretargeted ^{212}Pb -DOTAM in SCID mice carrying SC BxPC3 tumors after 2-step PRIT using CEA-split-DOTAM-VH/VL BsAbs.

- 5 Two-step PRIT was performed by injection of the CEA-split-DOTAM-VH and CEA-split-DOTAM-VL (P1AD8749 and P1AD8592), separately or together, followed 7 days later by ^{212}Pb -DOTAM. Mice were sacrificed 6 hours after the radioactive injection, and blood and organs harvested for radioactive measurement. The 2-step scheme was compared with 3-step PRIT using the standard CEA-DOTAM bispecific antibody, followed 7 days later by Ca-
- 10 DOTAM-dextran-500 CA, and ^{212}Pb -DOTAM 24 hours after the CA.

PK data of CEA-split-DOTAM-VH/VL clearance was collected by repeated blood sampling from 1 hour to 7 days after the antibody injection, and subsequently analyzed by an ELISA.

- 15 The study outline is shown in Figure 7. Figure 7a shows the outline of the 2-step PRIT regimen, including blood sampling for CEA-split-DOTAM-VH/VL PK, in SCID mice carrying SC BxPC3 tumors. Figure 7b shows the outline of the 3-step PRIT regimen, performed in SCID mice carrying SC BxPC3 tumors (h = hours, d = days).

Study design

The time course and design of protocol 144 is shown in the tables below.

20 Time course of protocol 144

Study day	Date	Experimental procedure
0	2018-05-02	Preparation of BxPC3 cells and filling of syringes
0	2018-05-02	SC injection of BxPC3 cells
14	2018-05-16	IV injection of CEA-DOTAM BsAb (group D)
15	2018-05-17	IV injection of CEA-split-DOTAM-VH/VL BsAbs (groups Aa, Ab, Ba, Bb, Ca, Cb)
15	2018-05-17	Retro-orbital bleeding (1 and 4 h p.i.; groups Aa, Ba, Ca, and Ab, Bb, Cb, respectively)
16	2018-05-18	Retro-orbital bleeding (24 h p.i.; groups Aa, Ba, Ca)
18	2018-05-20	Retro-orbital bleeding (72 h p.i.; groups Ab, Bb, Cb)
21	2018-05-23	IV injection of CA (group D)
21	2018-05-23	Elution of ^{212}Pb -DOTAM and filling of syringes
22	2018-05-24	IV injection of ^{212}Pb -DOTAM (groups Aa, Ba, Ca, D)
22	2018-05-24	Retro-orbital bleeding (168 h p.i.) and euthanasia (groups Ab, Bb, Cb)

22	2018-05-24	Euthanasia and tissue harvest, incl. retro-orbital bleeding (6 h p.i.) + gamma counting (groups Aa, Ba, Ca, D)
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Study groups in protocol 144

Group	P1AD8749 (VH) CH1A1A (μ g)	P1AD8592 (VL) CH1A1A (μ g)	CEA-DOTAM BsAb (μ g)	PK (h p.i.)	CA (μ g)	^{212}Pb (μ Ci)	BD (h p.i.)	n (mice)
Aa	100	0	0	1, 24, 168	0	10	6	4
Ab	100	0	0	4, 72, 168	0	0	—	4
Ba	0	100	0	1, 24, 168	0	10	6	4
Bb	0	100	0	4, 72, 168	0	0	—	4
Ca	100	100	0	1, 24, 168	0	10	6	4
Cb	100	100	0	4, 72, 168	0	0	—	4
D	0	0	100	—	25	10	6	4

Solid xenografts were established in each SCID mouse on study day 0 by SC injection of 5×10^6 cells (passage 26) in RPMI/Matrigel into the right flank. Fourteen days after tumor cell injection, mice were sorted into experimental groups with an average tumor volume of 116 mm^3 . The ^{212}Pb -DOTAM was injected on day 22 after inoculation; the average tumor volume was 140 mm^3 on day 21.

Blood from mice in groups Aa, Ba, and Ca was collected through retro-orbital bleeding under anesthesia 1 h (right eye), 24 h (left eye), and 168 h (right eye, at termination) after CEA-split-DOTAM-VH/VL injection. Similarly, samples were taken from mice in groups Ab, Bb, and Cb 4 h (right eye), 72 h (left eye), and 168 h (right eye, at termination) after CEA-split-DOTAM-VH/VL injection.

Mice in groups Aa, Ba, Ca, and D were sacrificed and necropsied 6 hours after injection of ^{212}Pb -DOTAM, and the following organs and tissues harvested for measurement of radioactive content: blood, skin, bladder, stomach, small intestine, colon, spleen, pancreas, kidneys, liver, lung, heart, femoral bone, muscle, brain, tail, ears, and tumor.

Results

The average ^{212}Pb accumulation and clearance in all collected tissues 6 hours after injection is displayed in Figure 8. Pretargeting with either CEA-split-DOTAM-VH or CEA-split-DOTAM-VL alone resulted in no accumulation of radioactivity in tumors. Combined, the two complimentary antibodies resulted in a tumor uptake after 2-step PRIT of $65 \pm 12\%$

ID/g, to be compared with $87 \pm 15\%$ ID/g for the standard 3-step PRIT regimen. Two-way analysis of variance (ANOVA) with Tukey's multiple comparisons test showed that the difference in tumor uptake between the two PRIT treatments was significant, as was the difference in bladder ($1 \pm 2\%$ ID/g and $38 \pm 17\%$ ID/g for 2- and 3-step PRIT, respectively);
 5 no other differences in tissue accumulation were statistically significant using this test ($p = 0.05$).

The clearance of IV injected CEA-split-DOTAM-VH/VL constructs as analyzed by an enzyme-linked immunosorbent assay (ELISA) is shown in Figure 9.

Adverse events and toxicity

10 There were no adverse events or toxicity associated with this study.

Conclusion

The results of the study demonstrated proof-of-concept of CA-independent 2-step pretargeting using complimentary CEA-split-DOTAM-VH/VL antibodies. High and specific tumor uptake of ^{212}Pb -DOTAM was achieved using 2-step PRIT and standard 3-step PRIT,
 15 with very little accumulation of radioactivity in normal tissues using the complimentary CEA-split-DOTAM-VH/VL antibodies.

Example 3c: Protocol 158

The aim of protocol 158 was to assess the association of ^{212}Pb -DOTAM to subcutaneous BxPC3 tumors in mice pretargeted by bi-paratopic (CH1A1A and A5B7) pairs
 20 of CEA-split-DOTAM-VH/VL antibodies for clearing agent-independent 2-step CEA-PRIT. The tumor uptake was compared with that of standard 3-step CEA-PRIT.

Mice carrying subcutaneous BxPC3 tumors were injected with either

- CEA-split-DOTAM-VH/VL antibodies followed 7 days later by the radiolabeled ^{212}Pb -DOTAM (2-step PRIT), or
- 25 • CEA-DOTAM BsAb followed 7 days later by the CA, and finally the radiolabeled ^{212}Pb -DOTAM 24 hours later (3-step PRIT).

The in vivo distribution of ^{212}Pb -DOTAM was assessed 6 hours after the radioactive injection. The study outline is shown in Figure 10.

Study design

The time course and design of protocol 158 is shown in the tables below.

Time course of protocol 158

Study day	Date	Experimental procedure
0	2018-11-26	Preparation of BxPC3 cells and filling of syringes
0	2018-11-26	SC injection of BxPC3 cells
15	2018-12-11	IV injection of CEA-DOTAM BsAb (group C)
16	2018-12-12	IV injection of CEA-split-DOTAM-VH/VL BsAbs (groups A, B)
22	2018-12-18	IV injection of CA (group C)
22	2018-12-18	Elution of ^{212}Pb -DOTAM and filling of syringes
23	2018-12-19	IV injection of ^{212}Pb -DOTAM (all)
23	2018-12-19	Euthanasia and tissue harvest, incl. retro-orbital bleeding (6 h p.i.) + gamma counting (all)

5 Study groups in protocol 158

Group	P1AD8749 (VH) CH1A1A (μg)	P1AD8592 (VL) CH1A1A (μg)	P1AE4956 (VH) A5B7 (μg)	P1AE4957 (VL) A5B7 (μg)	CEA- DOTAM BsAb (μg)	CA (μg)	^{212}Pb (μCi)	n (mice)
A	154*	0	0	100	0	0	10	4
B	0	100	167**	0	0	0	10	4
C	0	0	0	0	100	25	10	4

*P1AD8749 dose adjusted to 154 μg to compensate for a 35% hole/hole impurity;

**P1AD8592 dose adjusted to 167 μg to compensate for a 40% hole/hole impurity.

10 Solid xenografts were established in each SCID mouse on study day 0 by SC injection of 5×10^6 cells (passage 27) in RPMI/Matrigel into the right flank. Fourteen days after tumor cell injection, mice were sorted into experimental groups with an average tumor volume of 177 mm^3 . The ^{212}Pb -DOTAM was injected on day 20 after inoculation; the average tumor volume was 243 mm^3 on day 21.

15 Mice in all groups were sacrificed and necropsied 6 hours after injection of ^{212}Pb -DOTAM, and the following organs and tissues harvested for measurement of radioactive content: blood, skin, bladder, stomach, small intestine, colon, spleen, pancreas, kidneys, liver, lung, heart, femoral bone, muscle, brain, tail, and tumor.

Results

The average ^{212}Pb distribution in all collected tissues 6 hours after injection is shown in Figure 11. Two-way ANOVA with Tukey's multiple comparisons test showed that there was no significant difference in normal tissue uptake of ^{212}Pb between the three treatments, except for bladder, where both bi-paratopic CEA-split-DOTAM-VH/VL pairs yielded lower accumulation than the standard 3-step PRIT. The kidney uptake was 3–4% ID/g for all three treatments. Either bi-paratopic combination resulted in tumor accumulation of approximately 56% ID/g, compared with 67% ID/g for 3-step PRIT; the difference between 2- and 3-step PRIT was statistically significant ($p < 0.0001$).

Adverse events and toxicity

There were no adverse events or toxicity associated with this study.

Conclusion

This study assessed the association of ^{212}Pb -DOTAM to SC BxPC3 tumors in mice pretargeted by bi-paratopic pairs of CEA-split-DOTAM-VH/VL antibodies for CA-independent 2-step CEA-PRIT, compared with standard 3-step PRIT. The distribution of ^{212}Pb 6 hours after injection was comparable for 2- and 3-step PRIT, with high accumulation in tumor and very little radioactivity in healthy tissues. This demonstrated proof of concept of bi-paratopic pretargeting of CEA-expressing tumors for 2-step CEA-PRIT using CEA-split-DOTAM-VH/VL antibodies.

Example 3d: Protocol 160

The aim of protocol 160 was to compare the therapeutic efficacy after 3 cycles of CA-independent 2-step CEA-PRIT using complimentary CEA-split-DOTAM-VH/VL antibodies, with that of standard 3-step CEA-PRIT in mice bearing SC BxPC3 tumors. A comparison was also made with 1-step CEA-RIT, using BsAbs that were pre-incubated with ^{212}Pb -DOTAM before injection.

Mice carrying SC BxPC3 tumors were injected with either

- CEA-DOTAM BsAb followed 7 days later by the CA, and finally the radiolabeled ^{212}Pb -DOTAM 24 hours later (3-step PRIT),

- CEA-split-DOTAM-VH/VL antibodies followed 7 days later by the radiolabeled ^{212}Pb -DOTAM (2-step PRIT), or
- ^{212}Pb -DOTAM-CEA-DOTAM BsAb (pre-incubated; 1-step RIT).

The therapy was administered in 3 repeated cycles of 20 μCi of ^{212}Pb -DOTAM, also including comparison with a non-CEA binding control antibody (DIG-DOTAM), and no treatment (vehicle). Dedicated mice were sacrificed for biodistribution purposes to confirm ^{212}Pb -DOTAM targeting and clearance at each treatment cycle. The treatment efficacy was assessed in terms of TGI and TR, and the mice were carefully monitored for the duration of the study to assess the tolerability of the treatment. The study outline is shown in Figure 12.

The time course and design of protocol 160 are shown in the tables below.

Time course of protocol 160

Study day	Date	Experimental procedure
0	2019-01-29	Preparation of BxPC3 cells and filling of syringes
0	2019-01-29	SC injection of BxPC3 cells
15	2019-02-13	IP injection of BsAb or histidine buffer (groups A, B, C, F, G, H, I)
16	2019-02-14	IP injection of CEA-split-DOTAM-VH/VL or histidine buffer (groups D, J, K, L)
22	2019-02-20	IP injection of CA or PBS (groups A, B, C, F, G, H, I)
23	2019-02-21	Elution of ^{212}Pb -DOTAM and filling of syringes
23	2019-02-21	IV injection of ^{212}Pb -DOTAM-CEA-DOTAM or histidine buffer (groups E, M)
23	2019-02-21	IV injection of ^{212}Pb -DOTAM or 0.9% NaCl (groups B, C, D, F, G, H, I, J, K, L)
24	2019-02-22	Euthanasia and tissue harvest (24 h p.i.) + gamma counting (groups F, G, J, M)
29	2019-02-27	IP injection of PRIT BsAb or PBS (groups A, B, C, H, I)
30	2019-02-28	IP injection of CEA-split-DOTAM-VH/VL or histidine buffer (groups D, K, L)
36	2019-03-06	IP injection of CA or PBS (groups A, B, C, H, I)
37	2019-03-07	Elution of ^{212}Pb -DOTAM and filling of syringes
37	2019-03-07	IV injection of ^{212}Pb -DOTAM or 0.9% NaCl (groups B, C, D, H, I, K, L)
38	2019-03-08	Euthanasia and tissue harvest (24 h p.i.) + gamma counting (groups H, K)
43	2019-03-13	IP injection of PRIT BsAb or PBS (groups A, B, C, I)
44	2019-03-14	IP injection of CEA-split-DOTAM-VH/VL or histidine buffer (groups D, L)

50	2019-03-20	IP injection of CA or PBS (groups A, B, C, I)
51	2019-03-21	Elution of ^{212}Pb -DOTAM and filling of syringes
51	2019-03-21	IV injection of ^{212}Pb -DOTAM or 0.9% NaCl (groups B, C, D, I, L)
52	2019-03-22	Euthanasia and tissue harvest (24 h p.i.) + gamma counting (groups I, L)

Study groups in protocol 160

Group	BsAb	BsAb per cycle (μg)	CA per cycle (μg)	^{212}Pb -DOTAM per cycle (μCi)	Cycles (#)	n (mice)
A	—	0	0	0	3	10
B	DIG-DOTAM	100	25	20	3	10
C	CEA-DOTAM	100	25	20	3	10
D	CEA-split-DOTAM	154* + 100**	0	20	3	10
E	^{212}Pb -DOTAM-CEA-DOTAM	100	0	20	1***	10
F	DIG-DOTAM	100	25	20	1	3
G	CEA-DOTAM	100	25	20	1	3
H	CEA-DOTAM	100	25	20	2	3
I	CEA-DOTAM	100	25	20	3	3
J	CEA-split-DOTAM	154* + 100**	0	20	1	3
K	CEA-split-DOTAM	154* + 100**	0	20	2	3
L	CEA-split-DOTAM	154* + 100**	0	20	3	3
M	^{212}Pb -DOTAM-CEA-DOTAM	100	0	20	1	3

*P1AD8749: dose adjusted to 154 μg to compensate for a 35% hole/hole impurity in the stock solution; **P1AD8592; ***Adjusted from 3 cycles to 1 cycle due to acute radiation-induced toxicity at the first treatment cycle.

Solid xenografts were established in SCID mice on study day 0 by SC injection of 5×10^6 cells (passage 24) in RPMI/Matrigel into the right flank. Fifteen days after tumor cell injection, mice were sorted into experimental groups with an average tumor volume of 122 mm³. The ^{212}Pb -DOTAM was injected on day 23 after inoculation; the average tumor volume was 155 mm³ on day 22.

The CEA-DOTAM and DIG-DOTAM antibodies were diluted in vehicle buffer to a final concentration of 100 μg per 200 μL for IP administration according to the table above

(Study groups in protocol 160). The CEA-split-DOTAM-VH/VL antibodies were mixed together into one single injection solution for IP administration, containing 100 µg of each construct per 200 µL. For P1AD8749, the dosing was adjusted to 154 µg to compensate for a 35% hole/hole impurity in the stock solution (the side of the molecule that does not carry the VH/VL). The Ca-DOTAM-dextran-500 CA was administered IP (25 µg per 200 µL of PBS) 7 days after the BsAb injection, followed 24 hours later by ^{212}Pb -DOTAM (RO7205834) according to the experimental schedule in Figure 12. PRIT-treated mice (2- and 3-step) were injected IV with 100 µL of the Ca-quenched ^{212}Pb -DOTAM solution (20 µCi in 100 µL 0.9% NaCl).

Mice treated with 1-step RIT received only one injection: pre-bound ^{212}Pb -DOTAM-CEA-DOTAM (20 µCi/20 µg BsAb in 100 µL 0.9% NaCl for IV injection). The direct-labeled antibody was prepared by incubating the ^{212}Pb -DOTAM with the CEA-DOTAM BsAb for 10 minutes at 37°C.

The following organs and tissues were harvested from mice in groups A–E at the time of euthanasia: serum, liver, spleen, kidneys, pancreas, and tumor. Before euthanasia, the live mouse was anesthetized for retro-orbital blood collection. The collected blood samples were centrifuged at 10 000 rcf during 5 minutes and the resulting serum fractions isolated, frozen, and stored at –20°C. The excised tissues were immediately put in 10% neutral buffered formalin (4°C) and then transferred to 1X PBS (4°C) after 24 hours. The formalin-fixed samples were shipped to Roche Pharma Research and Early Development, Roche Innovation Center Basel, for further processing and analysis.

Mice in groups F, G, J, and M were sacrificed and necropsied 24 hours after their first and only injection of ^{212}Pb -DOTAM or ^{212}Pb -DOTAM-BsAb; groups H and K were sacrificed and necropsied 24 hours after their second ^{212}Pb -DOTAM injection; groups I and L were sacrificed and necropsied 24 hours after their third ^{212}Pb -DOTAM injection. Blood was collected at the time of euthanasia from the venous sinus using retro-orbital bleeding on anesthetized mice, before termination through cervical dislocation. The following organs and tissues were also harvested for biodistribution purposes: bladder, spleen, kidneys, liver, lung, muscle, tail, skin, and tumor.

Results

The average ^{212}Pb accumulation and clearance in all collected tissues 24 hours after injection is shown for each therapy and treatment cycle in Figure 13. The negative control resulted in no uptake (0.4% ID/g) in tumor. Two-way analysis of variance (ANOVA) with

Tukey's multiple comparisons test showed that the distributions were not significantly different at any cycle for the 2-step and 3-step PRIT; however, the differences were at all cycles statistically significant compared with the negative control and the 1-step RIT ($p < 0.05$). The tumor uptake was 25–45% ID/g for 3-step PRIT and 25–30% ID/g for 2-step PRIT, without any statistically significant difference between either treatment or cycle. For 1-step RIT, the tumor uptake was 99% at the one and only treatment cycle. The uptake in normal tissues was very low for both PRIT regimens, but significantly higher in all organs and tissues after 1-step RIT, due to the much longer circulating time of the pre-incubated antibody compared with the small, radiolabeled DOTAM chelate.

The average tumor development and the individual tumor growth curves are shown in Figure 14 and Figure 15, respectively. Tumors in the non-treated vehicle group and the DIG-DOTAM group grew steadily, albeit with slightly lower doubling rate in the latter after the third treatment. In contrast, tumors in the PRIT and RIT groups decreased in size after the first treatment cycle, and maintained tumor control until approximately 10 weeks after inoculation, when the tumors started to increase in size. The 2-step and 3-step PRIT treatments resulted in near identical tumor control. No tumors regressed completely.

On study day 83, the last day on which all treatment groups could be analyzed based on means, the TGI was 91.7% and 88.4% for PRIT using CEA-DOTAM (3-step) and CEA-split-DOTAM-VH/VL (2-step), respectively, compared with the vehicle control. The corresponding number for 1-step RIT was 72.6%, whereas the TGI was –59.7% for the non-specific DIG-DOTAM control. On the same day, the TR based on means was –1.9 for 3-step CEA-DOTAM PRIT, –2.9 for 2-step CEA-split-DOTAM-VH/VL PRIT, –4.7 for 1-step RIT, –28.8 for DIG-DOTAM PRIT, and –39.3 for the vehicle control.

Due to the adverse events described below, survival analysis was not considered statistically relevant.

Adverse events and toxicity

The BW development in all therapy groups is shown in Figure 16. The multiple cycles of 2- and 3-step PRIT with 20 μCi of ^{212}Pb -DOTAM were well tolerated, but acute BW loss occurred in mice receiving 1-step RIT, with 8/10 mice in group E euthanized after the first RIT cycle (6–11 days after ^{212}Pb irradiation) due to a drop in BW of 20% or more. The remaining 2 RIT mice were not given any further ^{212}Pb -DOTAM-CEA-DOTAM injections but were continuously followed up for tumor growth assessment.

In addition, a number of mice were sacrificed for ethical reasons due to declining tumor status, i.e. tumors opening up or leaking. In the DIG-DOTAM group, 9/10 mice were euthanized before reaching a tumor volume of 3000 mm³ for this reason; for the non-treated vehicle control, the corresponding number was 5/10. The problem was less pronounced in the PRIT and RIT groups, with 1/10, 2/10, and 2/10 mice euthanized for this reason in the 3-step PRIT, 2-step PRIT, and 1-step RIT groups, respectively. This is reflected in the individual tumor growth curves in Figure 15.

Finally, 1 mouse in group C was euthanized due to a degrading wound under the anus. All adverse events are listed in the table below.

Adverse events in protocol 160

Group	Mice sacrificed (n per group)	Study day	Reason for termination
A: Vehicle	5(10)	53, 71, 71, 73, 75	Declining tumor status
B : DIG-DOTAM	9(10)	55, 55, 55, 55, 61, 73, 73, 73, 74	Declining tumor status
C : CEA -DOTAM	1(10)	83	Wound under the anus
C : CEA -DOTAM	1(10)	85	Declining tumor status
D : CEA-split-DOTAM	2(10)	83, 83	Declining tumor status
E : ²¹² Pb-DOTAM-CEA-DOTAM	8(10)	29, 29, 30, 31, 31, 32, 32, 34	BW loss ≥ 20%
E : ²¹² Pb-DOTAM-CEA-DOTAM	2(10)	83, 83	Declining tumor status

²¹²Pb irradiation was performed on study day 23 (cycle 1), 37 (cycle 2), and 51 (cycle 3).

Conclusion

No difference was seen between CEA-PRIT using the 3-step scheme (CEA-DOTAM BsAb, CA, and ²¹²Pb-DOTAM) and the 2-step scheme (CEA-split-DOTAM-VH/VL antibodies and ²¹²Pb-DOTAM); the TGI was significant and near identical for the two treatments, and 3 cycles of 20 µCi could be safely administered in both cases. Contrastingly, 20 µCi of ²¹²Pb-DOTAM pre-bound to CEA-DOTAM before injection (1-step RIT) was not tolerated by a large majority of the treated mice.

The study thus demonstrated tolerability and therapeutic efficacy of CA-independent 2-step PRIT using the developed CEA-split-DOTAM-VH/VL constructs.

EXAMPLE 4: Protocol 175

The aim of protocol 175 was to assess the impact of increased injected pretargeting antibody amount on the subsequent ^{212}Pb accumulation in tumor and healthy tissues. Two different doses of CEA-split-DOTAM-VH/VL antibodies were compared: the standard amount (100 μg) and 2.5 times higher dose (250 μg). Moreover, a modification was made to the CEA-split-DOTAM-VH construct to extend its VH to avoid anti-drug antibody (ADA) formation (this was used together with a previously tested CEA-split-DOTAM-VL construct). The VH was extended to comprise the first three amino acids from the antibody CH1 domain: alanine, serine, and threonine (AST), and the construct hereafter referred to as CEA-split-DOTAM-VH-AST.

Antibody P1AD8592 has already been described above, in example 1. P1AF0171 is the same as P1AD8749 except that the fusion HC is extended by the residues AST – thus, antibody P1AD0171 consists of the light chain D1AA3384 as described above (SEQ ID NO: 34), the first heavy chain D1AC4022 as described above (SEQ ID NO: 28), and a second heavy chain D1AE3669 as shown below:

D1AE3669 (HCknob <CEA> CH1A1A Dotam-VH-AST)

QVQLVQSGAEVKKPGASVKVSCKASGYTFTEFGMNWVRQAPGQGLEWMGWINTK
TGEATYVEEFKGRVTFTTDTSTSTAYMELRSLRSDDTAVYYCARWDFAYYVEAMD
YWGQGTTVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGA
LTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSC
DKTHTCPPCPAPEAAGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNW
YVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALGAPI
EKTISKAKGQPREPQVYTLPPCRDELTKNQVSLWCLVKGFYPSDIAVEWESNGQPEN
NYKTTTPVLDSGDSFFLYSKLTVDKSRWQQGNVFSQSVSMHEALHNHYTQKSLSLSP
GGGGGSGGGGSGGGGSGGGGSVTLKESGPVLVKPTETLTCTVSGFSLSTYSMSWI
RQPPGKALEWLGFISRGDTYYASWAKGRLTISKDTSKSQVVLMTNMDPVDATY
YCARERDPYGGGAYPPHLWGRGTLVTVSSAST (SEQ ID NO: 153)

Mice carrying SC BxPC3 tumors were injected with either

- 1× the standard dose of CEA-split-DOTAM-VH/VL BsAb followed 7 days later by the radiolabeled ^{212}Pb -DOTAM, or

- 2.5× the standard dose of CEA-split-DOTAM-VH/VL BsAb followed 7 days later by the radiolabeled ^{212}Pb -DOTAM.

The in vivo distribution of ^{212}Pb -DOTAM was assessed 24 hours after the radioactive injection. The study outline is shown in figure 17.

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Study design

The time course and design of protocol 175 are shown below.

Time course of protocol 175

Study day	Experimental procedure
0	Preparation of BxPC3 cells and filling of syringes
0	SC injection of BxPC3 cells
22	IP* injection of CEA-split-DOTAM-VH/VL BsAbs
29	Elution of ^{212}Pb -DOTAM and filling of syringes
29	IV injection of ^{212}Pb -DOTAM
30	Euthanasia and tissue harvest (24 h p.i.) + gamma counting

10 *IP injection required due to low compound concentration (200 μL per construct = 400 μL in total)

Study groups in protocol 175

Group	P1AF0171 (VH-AST) (μg)	P1AD85 92 (VL) (μg)	^{212}Pb (μCi)	BD (h p.i.)	n (mice)
A	143*	100	10	24	4
B	357*	250	10	24	4

*P1AF0171 dose adjusted to 143 and 357 μg to compensate for a ~30% hole/hole impurity.

15 Solid xenografts were established in each SCID mouse on study day 0 by SC injection of 5×10^6 cells (passage 24) in RPMI/Matrigel into the right flank. Twenty-one days after tumor cell injection, mice were sorted into experimental groups with an average tumor volume of 310 mm^3 . The ^{212}Pb -DOTAM was injected on day 29 after inoculation; the average tumor volume was 462 mm^3 on day 30.

20 All mice were sacrificed and necropsied 24 hours after injection of ^{212}Pb -DOTAM, and the following organs and tissues harvested for measurement of radioactive content: blood, skin, spleen, pancreas, kidneys, liver, muscle, tail, and tumor.

Results

The average ^{212}Pb distribution in all collected tissues 24 hours after injection is shown in Figure 18. There was no significant difference in tumor or normal tissue uptake of ^{212}Pb between the two dose levels. The tumor accumulation was 30–31% ID/g for both treatment groups, with a kidney uptake of <2% ID/g at this time point. One mouse had ~1 %ID/g in the tail due to ^{212}Pb -DOTAM injection issues, but no other collected healthy tissues showed any appreciable ^{212}Pb accumulation.

Adverse events and toxicity

There were no adverse events or toxicity associated with this study.

Conclusion

Increasing the dose of the pretargeting CEA-split-DOTAM-VH/VL antibodies by 2.5-fold did not improve the tumor accumulation of subsequently administered ^{212}Pb -DOTAM in this in vivo model. However, it also did not increase the accumulation of radioactivity in normal tissues, highlighting the strong specificity achieved using this 2-step pretargeting regimen.

Finally, the results verified the function of the extended-VH CEA-split-DOTAM-VH-AST construct.

EXAMPLE 5: Protocol 185

The aim of protocol 185 was to assess a CEA-split-DOTAM-VH/VL targeting the T84.66 epitope. Sequences of P1AF0709 and P1AF0298 are provided herein. P1AF0709 has a first heavy chain of D1AE4688 (SEQ ID NO: 83) and a second heavy chain of D1AA4920 (SEQ ID NO: 84). P1AF0298 has a first heavy chain of D1AE4687 (SEQ ID NO: 86) and a second heavy chain of D1AE3668 (SEQ ID NO: 87). Both have the light chain of D1AA4120 (SEQ ID NO: 89).

Mice carrying SC BxPC3 tumors were injected with the standard dose of CEA-split-DOTAM-VH/VL BsAb (100 µg per antibody) followed 6 days later by the radiolabeled ^{212}Pb -DOTAM. The in vivo distribution of ^{212}Pb -DOTAM was assessed 6 hours after the radioactive injection. The study outline is shown in figure 19.

Study design

The time course and design of protocol 185 is shown below.

Time course of protocol 185

Study day	Date	Experimental procedure
0	2020-03-04	Preparation of BxPC3 cells and filling of syringes
0	2020-03-04	SC injection of BxPC3 cells
22	2020-03-26	IV injection of CEA-split-DOTAM-VH/VL BsAbs
27	2020-03-31	Elution of ^{212}Pb -DOTAM and filling of syringes
28	2020-04-01	IV injection of ^{212}Pb -DOTAM
28	2020-04-01	Euthanasia and tissue harvest (6 h p.i.) + gamma counting

Study groups in protocol 185

Group	P1AF0298 T84.66 (VH-AST) (μg)	P1AF0709 T84.66 (VL) (μg)	P1AF0171 CH1A1A (VH-AST) (μg)	P1AD8592 CH1A1A (VL) (μg)	^{212}Pb (μCi)	BD (h p.i.)	n (mice)
A	100	100	0	0	10	6	5
B	0	0	143*	100	10	6	5

*P1AF0171 dose adjusted to 143 μg to compensate for a ~30% hole/hole impurity.

5

Solid xenografts were established in each SCID mouse on study day 0 by SC injection of 5×10^6 cells (passage 27) in RPMI/Matrigel into the right flank. Twenty-two days after tumor cell injection, mice were sorted into experimental groups with an average tumor volume of 224 mm^3 . The ^{212}Pb -DOTAM was injected on day 28 after inoculation, at which point the average tumor volume had reached 385 mm^3 .

10

All mice were sacrificed and necropsied 6 hours after injection of ^{212}Pb -DOTAM, and the following organs and tissues harvested for measurement of radioactive content: blood, skin, spleen, pancreas, kidneys, liver, muscle, tail, and tumor. Collected tumors were split in two pieces: one was measured for radioactive content, and the other put in a cryomold containing Tissue-Tek® optimum cutting temperature (OCT) embedding medium, and put on dry ice for rapid freezing. Frozen samples in OCT were maintained at -80°C before cryosectioning,

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immunofluorescence staining, and analysis using a Zeiss Axio Scope.A1 modular microscope.

Results

The average ^{212}Pb distribution in all collected tissues 6 hours after injection is shown in Figure 20. The tumor accumulation was 40% ID/g (CH1A1A) or 44% ID/g (T84.66). The only other appreciable accumulation of radioactivity was found in kidneys: 3–5% ID/g at 6 h p.i. for the two groups.

Examples of the intratumoral distribution of CEA-split-DOTAM-VH/VL pairs targeting either T84.66 (group A) or CH1A1A (group B) are shown in Figure 21. Panels A and C show that the CEA expression is high and homogeneous in BxPC3 tumors, and panels B and D demonstrate that the antibody distribution 7 days after injection is distributed similarly. However, the samples from group A displayed a stronger signal overall, compared with tumor samples from group B, providing evidence that T84.66 is a stronger binder than CH1A1A.

Adverse events and toxicity

There were no adverse events or toxicity associated with this study.

Conclusion

The results verified the function of CEA-split-DOTAM-VH/VL constructs targeting the T84.66 epitope of CEA. The resulting accumulation of ^{212}Pb in pretargeted CEA-expressing tumors was high and specific, and CEA-split-DOTAM-VH/VL pairs targeting either the CH1A1A or T84.66 epitope were homogeneously distributed inside the CEA-expressing tumors.

EXAMPLE 6: protocol 189

The aim of protocol 189 was to assess bi-paratopic CEA-split-DOTAM-VH/VL antibody pairs targeting T84.66 VH-AST/CH1A1A VL and T84.66 VL/CH1A1 VH-AST, compared with the positive control pair targeting CH1A1A VH-AST/VL. This bi-paratopic combination precludes formation of the full Pb-DOTAM binder on soluble CEA that only expresses one of the two epitopes (e.g. T84.66), thereby mitigating potential adverse effects thereof, such as increased circulating radioactivity and associated radiation-induced toxicity, and decreased efficacy from competition with off-tumor targets.

Mice carrying SC BxPC3 tumors were injected with the standard dose of CEA-split-DOTAM-VH/VL BsAb (100 µg per antibody) followed 7 days later by the radiolabeled ^{212}Pb -DOTAM. The in vivo distribution of ^{212}Pb -DOTAM was assessed 6 hours after the radioactive injection. The study outline is shown in Figure 22.

5 Study design

The time course and design of protocol 189 is shown below.

Time course of protocol 189

Study day	Experimental procedure
0	Preparation of BxPC3 cells and filling of syringes
0	SC injection of BxPC3 cells
15	IV injection of CEA-split-DOTAM-VH/VL BsAbs
21	Elution of ^{212}Pb -DOTAM and filling of syringes
22	IV injection of ^{212}Pb -DOTAM
22	Euthanasia and tissue harvest (6 h p.i.) + gamma counting

Study groups in protocol 189

Group	P1AF0298 T84.66 (VH-AST) (µg)	P1AF0709 T84.66 (VL) (µg)	P1AF0171 CH1A1A (VH-AST) (µg)	P1AD8592 CH1A1A (VL) (µg)	^{212}Pb (µCi)	BD (h p.i.)	n (mice)
A	100	0	0	100	10	6	5
B	0	100	143*	0	10	6	5
C	0	0	143*	100	10	6	3

10 *P1AF0171 dose adjusted to 143 µg to compensate for a ~30% hole/hole impurity.

15 Solid xenografts were established in each SCID mouse on study day 0 by SC injection of 5×10^6 cells (passage 31) in RPMI/Matrigel into the right flank. Fourteen days after tumor cell injection, mice were sorted into experimental groups with an average tumor volume of 343 mm³. The ^{212}Pb -DOTAM was injected on day 22 after inoculation; the average tumor volume had reached 557 mm³ on day 21.

All mice were sacrificed and necropsied 6 hours after injection of ^{212}Pb -DOTAM, and the following organs and tissues harvested for measurement of radioactive content: blood, skin, spleen, pancreas, kidneys, liver, muscle, tail, and tumor.

20 Results

The average ^{212}Pb distribution in all collected tissues 6 hours after injection is shown in Figure 23. The tumor accumulation of the bi-paratopic variations was 71% ID/g and 46% ID/g for T84.66 VH-AST + CH1A1A VL and T84.66 VL + CH1A1A VH-AST, respectively. The positive CH1A1A control resulted in 37% ID/g. Two-way ANOVA with Tukey's multiple comparisons test showed that all three groups were significantly different from each other in terms of tumor uptake ($p < 0.0001$ for T84.66 VH-AST + CH1A1A VL versus the two other groups; $p = 0.0020$ for T84.66 VL + CH1A1A VH-AST versus CH1A1A only). No other organs showed statistically significant differences between groups, although a slightly higher retention in blood was indicated for the T84.66 VH-AST + CH1A1A VL combination compared with the two other groups: 2% ID/g compared with $<1\%$ ID/g. The kidney uptake was similarly slightly higher, although not statistically significantly so: 4.5% ID/g for T84.66 VH-AST + CH1A1A compared with 3% ID/g for the other two.

Adverse events and toxicity

There were no adverse events or toxicity associated with this study. However, the BxPC3 tumor growth was significantly faster, and with greater variability, in this study compared with the standard growth rate. On necropsy, it was concluded that the big tumors (a majority) were filled with liquid, which was emptied when tumors were cut in half before radioactive measurement; this liquid likely caused the accelerated growth rate, but did not affect the %IA/g to any great extent as the tumors were weighed and measured after being opened.

Conclusion

The results verified the function of bi-paratopic targeting of the T84.66 and CH1A1A epitopes of CEA using the tested CEA-split-DOTAM-VH/VL constructs and demonstrated surprisingly high efficacy for this combination as compared to the positive CH1A1A control. The resulting accumulation of ^{212}Pb in pretargeted CEA-expressing tumors was high and specific, with indications of a particular advantage for the T84.66 VH-AST + CH1A1A VL pair.

EXAMPLE 7

These examples investigate recruitment of Pb-DOTA to cells by split antibodies as described herein.

P1AF0712 has a first heavy chain of SEQ ID NO:97, a second heavy chain of SEQ ID NO: 98 and a light chain of SEQ ID NO: 103. P1AF0713 has a first heavy chain of SEQ ID NO: 100, a second heavy chain of SEQ ID NO: 101 and a light chain of SEQ ID NO: 103.

5

MKN-45 cells were detached from the culture bottle using Trypsin and were counted using a Casy cell counter. After pelleting at 4°C, 300g the cells were resuspended in FACS Buffer (2.5% FCS in PBS), adjusted to 2.0E+06 cells /mL dispensed to 96-well PP V-bottom-Platte (25 µL/well = 5.0E+04Zellen/well).

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FACS staining using DOTA-FITC

The CEA specific SPLIT antibodies (P1AF0712 or P1AF0713 respectively) were adjusted to 40 µg/mL in FACS buffer, resulting in a final concentration of 10 µg/mL. Both antibodies were added to the cells either combined or separated and followed by buffer and incubated at 4°C for 1 h. Subsequently, Pb-DOTA labeled with FITC was added to the cells in equimolar ratio to the antibodies and incubated for 1 h at 4°C. The cells were then washed twice in FACS buffer and resuspended in 70 µl/well FACS buffer for measurement using a FACS Canto (BD, Pharmingen). It was shown (Fig. 24) that neither of the SPLIT halves was giving rise to a fluorescence signal, indicating a lack of Pb-DOTA binding capability. Only a combination of both SPLIT halves was able to recruit Pb-DOTAM-FITC to the target cells (Fig 24).

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FACS staining using <huIgG(H+L)A488>

The CEA specific SPLIT antibodies (P1AF0712 or P1AF0713 respectively) were adjusted to 40 µg/mL in FACS buffer, resulting in a final concentration of 10 µg/mL. Both antibodies were added to the cells either separated followed by buffer or combined and incubated at 4°C for 1 h. The cells were then washed twice in FACS buffer. After washing, the cells were resuspended in 50 µL FACS-buffer containing secondary antibody (<huIgG(H+L)>-Alexa488, c=10 µg/mL) and incubated 1h at 4°C. The cells were then washed twice in FACS buffer and resuspended in 70 µl/well FACS buffer for measurement using a FACS Canto (BD, Pharmingen). EC50 for both SPLIT antibodies was comparable, indicating CEA

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specific cell binding of both SPLIT antibodies. Due to the higher amount of antibody in the mixture, a lower EC50 was obtained under these circumstances, as shown in the table below.

EC50 Determination of SPLIT antibodies

		EC50 μg/ml absolute
<hu>A488	P1AF0712+ PBS	2.7
	P1AF0713+ PBS	2.3
	P1AF0712+ P1AF0713	0.9
	hu ISO + PBS	
DOTA-FITC	P1AF0712+ PBS	na
	P1AF0713+ PBS	na
	P1AF0712+ P1AF0713	2.4
	hu ISO + PBS	

5

EC50 was determined for the SPLIT antibodies using either secondary antibody based detection (<hu>488, top panel) or Pb-DOTA-FITC (DOTA-FITC, bottom panel)

EXAMPLE 8: Biacore binding experiments

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This example tests binding of TA-split-DOTAM-VH and TA-split-DOTAM-VL individually to DOTAM, as compared to the reference antibody CEA-DOTAM (RO7198427, PRIT-0213). It further tests binding of DOTAM to the TA-split-DOTAM-VH/VL pairs, as compared to the reference antibody.

15

The correspondence between the coding used in these examples and the protein numbers used elsewhere in this application is shown below. Sequences are also provided. The reference antibody is coded as “PRIT_RS” in this example.

Target binder	SPR Code (Prodrug	SPR Code	DOTAM	Protein	LC (SEQ	HC (SEQ	Fusion HC
---------------	----------------------	-------------	-------	---------	------------	------------	--------------

	A+B)				ID NO)	ID NO)	(SEQ ID NO)
<CEA> CH1A1A	P1_AB	P1_A	VL	P1AD8592	34	30	33
		P1_B	VH	P1AD8749	34	28	32
<CEA> CH1A1A	P2_AB	P2_A	VL	P1AD8592	34	30	33
		P2_B	VH	P1AF0171	34	28	147
<CEA> T84.66	P3_AB	P3_A	VL	P1AF0709	89	83	84
		P3_B	VH	P1AF0298	89	86	89
<CEA> 28A9	P4_AB	P4_A	VL	P1AF0710	96	90	91
		P4_B	VH	P1AF0711	96	93	94
<CEA> A5H1EL1(G54A)	P5_AB	P5_A	VL	P1AE4957	58	55	56
		P5_B	VH	P1AE4956	54	51	52
<CEA> CH1A1A	P6_AB	P6_A	VL	P1AF0712	103	97	98
		P6_B	VH	P1AF0713	103	100	101
<GPRC5D>	P7_AB	P7_A	VL	P1AF8284	107	104	105
		P7_B	VH	P1AF8285	107	104	106

For these experiments, the PRIT SPLIT antibodies were purified by a first step of MabSelect Sure (Affinity Chromatography) and a second step of ion exchange chromatography (e.g. POROS XS), and then polished by Superdex 200 (Size Exclusion Chromatography).

5

The experiments were performed with Biacore T200 at 25°C measuring temperature. All Biacore T200 experiments were carried out in HBS-P+ (GE Healthcare, Br-1008-27) pH 7.4 running buffer. Two experiments were performed for each test antibody/antibody pair, using different DOTAM fractions.

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1. In a first experiment, the binding of individual TA-split-DOTAM-VH and TA-split-DOTAM-VL antibodies to biotinylated DOTAM captured on a chip was assessed, relative to the reference antibody.

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DOTAM (120 nM solution in HBS-P+) was captured in high density on CAP Chip Surface (10µl/min, 60Sec). Then the 600 nM solutions in HBS-P+ of Prodrug_A or Prodrug_B were

injected over the DOTAM surface (10µl/min, 90 sec). The dissociation was monitored for 240sec at a flow rate of 10 µl/min. The relative maximum response determination was evaluated using T200 evaluation software.

5 The results are shown in figure 26. None of the individual antibodies showed binding to the captured DOTAM.

2. In a second experiment, individual TA-split-DOTAM-VH and TA-split-DOTAM-VL antibodies were first captured in a chip using an immobilized anti-hFab, and then binding of a
10 DOTAM-monoStreptavidin complex (DOTAM +monoStreptavidin coupling 600nM, 1:1 mol, 1h at RT) was assessed.

The 600 nM solution in HBS-P+ of Prodrug_A or Prodrug B was injected over the anti hFab (GE Healthcare, BR-1008-27) CM5 Chip surface (10µl/min, 120 sec). After the high density
15 capturing of Prodrug A or B solution the DOTAM-monoStreptavidin complex was injected (20µl/min, 90 sec). The dissociation was monitored for 180 sec at a flow rate of 20 µl/min. For new cycle the surface was regenerated by using of Glycin 2.1 and 75 sec regeneration time with 10µl/min. The relative maximum response determination was evaluated using T200
20 evaluation software.

The results are shown in figure 27. Low percentage max. responses (as marked with * in the figure) are believed to be “traces” or unspecific interactions with DOTAM-SA, and reflect a need to optimize the assay.

25 3. In a third experiment, binding of the TA-split-DOTAM-VH/VL pairs to DOTAM is assessed, as compared to the reference antibody. Antibodies were first captured in a chip using an immobilized anti-hFab, and then binding of a DOTAM-monoStreptavidin complex (DOTAM +monoStreptavidin coupling 600nM, 1:1 mol, 1h at RT) was assessed.

30 The 300 nM solution in HBS-P+ of Prodrug_A and Prodrug B was injected over the anti hFab (GE Healthcare, BR-1008-27) CM5 Chip surface (10µl/min, 120 sec). After the high density capturing of Prodrug A and B solution the DOTAM-monoStreptavidin complex was injected (20µl/min, 90 sec). The dissociation was monitored for 180 sec at a flow rate of 20 µl/min. For new cycle the surface was regenerated by using of Glycin 2.1 and 75 sec

regeneration time with 10µl/min. The relative maximum response determination was evaluated using T200 evaluation software.

The results are shown in figure 28. All TA-split-DOTAM-VH/VL pairs showed a significant amount of binding for DOTAM, except for the P6_AB (P1AF0712/P1AF0713) pair, which are DOTA binders.

Similar results showing a significant amount of DOTAM binding for the TA-split-DOTAM-VH/VL pair but not for the individual members of the pair have also been obtained for the FAP-binders P1AF8286 and P1AF8287. P1AF8286 is composed of a first heavy chain of SEQ ID NO: 108, a second heavy chain of SEQ ID NO: 109 and a light chain of SEQ ID NO: 111, and P1AF8287 is composed of a first heavy chain of SEQ ID NO: 108, a second heavy chain of SEQ ID NO: 110 and a light chain of SEQ ID NO: 111. However, this assay still needs to be optimised.

EXAMPLE 9

MATERIALS AND METHODS

General

All experimental protocols were reviewed and approved by the local authorities (Comité Régional d’Ethique de l’Expérimentation Animale du Limousin [CREEAL], Laboratoire Départemental d’Analyses et de Recherches de la Haute-Vienne). Female severe combined immunodeficiency (SCID) mice (Charles River) were maintained under specific and opportunistic pathogen free (SOPF) conditions with daily cycles of light and darkness (12 h/12 h), in line with ethical guidelines. No manipulations were performed during the first 5 days after arrival, to allow the animals to acclimatize to the new environment. Animals were controlled daily for clinical symptoms and detection of adverse events.

Solid xenografts were established by subcutaneous (SC) injection of CEA-expressing tumor cells in cell culture media mixed 1:1 with Corning® Matrigel® basement membrane matrix (growth factor reduced; cat No. 354230). Tumor volumes were estimated through manual caliper 3 times per week, calculated according to the formula: $volume = 0.5 \times length \times width^2$. Additional tumor measurements were made as needed depending on the tumor growth rate.

Mice were euthanized before the scheduled endpoint if they showed signs of unamenable distress or pain due to tumor burden, side effects of the injections, or other causes. Indications of pain, distress, or discomfort include, but are not limited to, acute body weight (BW) loss, scruffy fur, diarrhea, hunched posture, and lethargy. General criteria for immediate

- 5 euthanasia are BW loss exceeding 20% of the initial BW or a tumor volume reaching 3000 mm³. The BW of treated animals was measured 3 times per week, with additional measurements as needed depending on the health status. Other factors taken into account for euthanasia for ethical reasons were tumor status (e.g. necrotic areas, blood/liquid leaking out, signs of automutilation) and general appearance of the animal (e.g. fur, posture, movement).
- 10 Blood was collected at the time of euthanasia from the venous sinus using retro-orbital bleeding on anesthetized mice, before termination through cervical dislocation followed by additional tissue harvest for radioactive measurements and histological analysis. Unexpected or abnormal conditions were documented. Organs and tissues collected for biodistribution purposes were weighed and measured for radioactivity using a 2470 WIZARD² automatic
- 15 gamma counter (PerkinElmer), and the percent injected dose per gram of tissue (% ID/g) subsequently calculated, including corrections for decay and background.

Statistical analysis was performed using GraphPad Prism 7 (GraphPad Software, Inc.).

Test compounds

- P1AF6241, P1AF6239, P1AF7887, and P1AF7889 are N-terminal CEA-split-DOTAM-
 20 VH/VL antibodies targeting the CH1A1A epitope of CEA. P1AF6241 and P1AF6239 are monovalent for CEA, whereas P1AF7887 and P1AF7889 are CEA bivalent. Their sequences are as below.

Target binder	Effector	Protein	LC	HC	FusionHC
<CEA> CH1A1A	Dotam VL	P1AF6241	D1AD3421 (SEQ ID NO: 143)	D1AC3981 (SEQ ID NO: 144)	D1AF7527 (SEQ ID NO: 145)
<CEA> CH1A1A	Dotam VH	P1AF6239	D1AD3421 (SEQ ID NO: (SEQ ID NO:	D1AC3981 (SEQ ID NO: (SEQ ID NO:	D1AF7526 (SEQ ID NO:

			143)	144)	146)
<CEA> CH1A1A	Dotam VL	P1AF7887	D1AD3421 (SEQ ID NO: 143)	D1AC6742 (SEQ ID NO: 147)	D1AF7527 (SEQ ID NO: 145)
<CEA> CH1A1A	Dotam VH	P1AF7889	D1AD3421 (SEQ ID NO: 143)	D1AC6742 (SEQ ID NO: 147)	D1AF7526 (SEQ ID NO: 146)

All antibody constructs were stored at -80°C until the day of injection when they were thawed and diluted in standard vehicle buffer (20 mM Histidine, 140 mM NaCl; pH 6.0) or 0.9% NaCl to their final respective concentrations for intravenous (IV) administration.

- 5 The DOTAM chelate for radiolabeling was provided by Macrocyclics and maintained at -20°C before radiolabeling, performed by Orano Med (Razès, France). ^{212}Pb -DOTAM (RO7205834) was generated by elution with DOTAM from a thorium generator, and subsequently quenched with Ca after labeling. The ^{212}Pb -DOTAM solution was diluted with 0.9% NaCl to obtain the desired ^{212}Pb activity concentration for IV injection.

10 Tumor models

BxPC3 is a human primary pancreatic adenocarcinoma cell line, naturally expressing CEA. Cells were supplied by ECACC (European Collection of Authenticated Cell Cultures (Salisbury, UK)) and cultured in RPMI 1640 Medium, GlutaMAX™ Supplement, HEPES (Gibco, ref. No. 72400-021) enriched with 10% fetal bovine serum (GE Healthcare Hyclone SH30088.03). Solid xenografts were established in each SCID mouse on study day 0 by subcutaneous injection of cells in RPMI media mixed 1:1 with Corning® Matrigel® basement membrane matrix (growth factor reduced; cat No. 354230), into the right flank. 5×10^6 BxPC3 cells were injected per mouse in an injected volume of 100 μL .

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Studies

Protocol 193

The aim of protocol 193 was to provide in vivo distribution data of pretargeted ^{212}Pb -DOTAM in SCID mice carrying SC BxPC3 tumors after 2-step PRIT using N-terminal
5 SPLIT constructs.

Two-step PRIT was performed by co-injection of the CEA-split-DOTAM-VH and CEA-split-DOTAM-VL, followed 7 days later by ^{212}Pb -DOTAM. Mice were sacrificed 6 hours after the radioactive injection, and blood and organs harvested for radioactive measurement.

The study outline is shown in Figure 30.

Study design

The time course and design of protocol 193 is shown in the tables below.

Time course of protocol 193

Study day	Date	Experimental procedure
0	2020-10-22	Preparation of BxPC3 cells and filling of syringes
0	2020-10-22	SC injection of BxPC3 cells
15	2020-11-06	IV injection of SPLIT constructs
22	2020-11-13	Elution of ^{212}Pb -DOTAM and filling of syringes
22	2020-11-13	IV injection of ^{212}Pb -DOTAM
22	2020-11-13	Euthanasia and tissue harvest (6 h p.i.) + gamma counting

Study groups in protocol 193

Group	P1AF6241 (VL) (μg)	P1AF6239 (VH) (μg)	P1AF7887 (VL) (μg)	P1AF7889 (VH) (μg)	^{212}Pb (μCi)	n (mice)
A	100	100	0	0	10	5
B	0	0	100	100	10	5

Solid xenografts were established in each SCID mouse on study day 0 by SC injection of 5×10^6 cells (passage 28) in RPMI/Matrigel into the right flank. Fifteen days after tumor cell injection, mice were sorted into experimental groups with a mean tumor volume of 272 mm^3 . The ^{212}Pb -DOTAM was injected on day 22 after inoculation, by which time the mean tumor
20 volume was 400 mm^3 .

All mice were sacrificed and necropsied 6 hours after injection of ^{212}Pb -DOTAM, and the following organs and tissues harvested for measurement of radioactive content: blood, skin, ovaries, stomach, small intestine, colon, spleen, pancreas, kidneys, liver, lung, heart, femoral bone, muscle, brain, tail, and tumor.

5 **Results**

The mean ^{212}Pb accumulation and clearance in all collected tissues 6 hours after injection is displayed in Figure 31. Pretargeting with CEA monovalent N-terminal SPLITs (P1AF6241 + P1AF6239) resulted in a mean tumor uptake of $24.3 \pm 5.4\%$ ID/g. The CEA bivalent N-terminal SPLITs (P1AF7887 + P1AF7889) resulted in a mean tumor uptake of $7.1 \pm 2.2\%$ ID/g. Two-way analysis of variance (ANOVA) with Sidak's multiple comparisons test showed that the difference in tumor ^{212}Pb accumulation was statistically significant for the treatments ($p < 0.0001$). No other differences in tissue accumulation were statistically significant using this test ($p = 0.05$), and no unexpected uptake was seen for either SPLIT pair. The uptake in kidneys was 2.7–3.8% ID/g for the two treatment groups.

15 **Adverse events and toxicity**

There were no adverse events or toxicity associated with this study.

Conclusion

The results of the study demonstrated proof-of-concept of CA-independent 2-step pretargeting using N-terminal SPLIT constructs. High and specific tumor uptake of ^{212}Pb -DOTAM was achieved, with very little accumulation of radioactivity in normal tissues.

Although the foregoing invention has been described in some detail by way of illustration and example for purposes of clarity of understanding, the descriptions and examples should not be construed as limiting the scope of the invention. The disclosures of all patent and scientific literature cited herein are expressly incorporated in their entirety by reference.

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WHAT IS CLAIMED IS:

1. A set of antibodies comprising:

i) a first antibody comprising:

a) an antigen binding moiety, wherein the antigen binding moiety binds to a target antigen;

b) a polypeptide comprising or consisting of an antibody heavy chain variable domain (VH) of an antigen binding site for an effector moiety; and

c) an Fc domain comprising two subunits,

10 wherein the polypeptide of (b) is fused by its N-terminus to the C-terminus of the antigen binding moiety of (a) and by its C-terminus to the N-terminus of one of the subunits of the Fc domain of (c);

and wherein the first antibody does not comprise a VL domain of an antigen binding site for the effector moiety; and

ii) a second antibody comprising:

d) an antigen binding moiety, wherein the antigen binding moiety binds to a target antigen;

e) a polypeptide comprising or consisting of an antibody light chain variable domain (VL) of an antigen binding site for the effector moiety; and

20 f) an Fc domain comprising two subunits,

wherein the polypeptide of (e) is fused by its N-terminus to the C-terminus of the antigen binding moiety of (d) and by its C-terminus to the N-terminus of one of the subunits of the Fc domain of (f);

and wherein the second antibody does not comprise a VH domain of an antigen binding site for the effector moiety;

wherein said VH domain of the first antibody and said VL domain of the second antibody are together capable of forming a functional antigen binding site for the effector moiety.

2. The set of antibodies of claim 1, wherein the antigen binding moiety of (a) and the
30 antigen binding moiety of (d) is a Fab.

3. The set of antibodies of claim 1 or claim 2, wherein the first and second antibodies are one-armed antibodies.

4. The set of antibodies of claim 3, wherein the first antibody and second antibody are monospecific and monovalent for a target antigen.

5. The set of antibodies according to any one of the preceding claims, wherein:
the first antibody comprises the following polypeptides:

i) a polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VH domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);

10 ii) a Fab light chain polypeptide (e.g., VL-CL); and

iii) an Fc subunit polypeptide (e.g., CH2-CH3);

wherein the Fab heavy chain of (i) and the Fab light chain of (ii) form a Fab fragment capable of binding to a target antigen; and

the second antibody comprises the following polypeptides:

iv) a polypeptide comprising from N-terminus to C-terminus: a Fab heavy chain (e.g., VH-CH1); an optional linker; a VL domain of an antigen binding site for an effector moiety; an optional linker; and an Fc subunit (e.g., CH2-CH3);

v) a Fab light chain polypeptide (e.g., VL-CL), and

vi) an Fc subunit polypeptide (e.g., CH2-CH3);

20 wherein the Fab heavy chain of (iv) and the Fab light chain of (v) form a Fab fragment capable of binding to a target antigen.

6. The set of antibodies of claim 1 or claim 2, wherein the first antibody comprises a further antigen binding moiety binding to a target antigen.

7. The set of antibodies of claim 6, wherein the further antibody moiety of the first antibody binds to the same target antigen as the antigen binding moiety of (a).

8. The set of antibodies of any one of claims 1, 2, 6 or 7, wherein the second antibody
30 comprises a further antigen binding moiety binding to a target antigen.

9. The set of antibodies of claim 8, wherein the further antigen binding moiety of the second antibody binds to the same target antigen as the antigen binding moiety of (d).

10. The set of antibodies according to any one of claims 6 to 9, wherein:

the first antibody comprises:

a) a first Fab fragment, wherein the Fab fragment binds to an antigen expressed on the surface of a target cell;

b) a polypeptide comprising or consisting of an antibody heavy chain variable domain (VH) of an antigen binding site for an effector moiety; and

c) an Fc domain comprising a first and a second subunit,

wherein the polypeptide of (b) is fused by its N-terminus to the C-terminus of one of the chains of the Fab fragment of (a) and by its C-terminus to the N-terminus of the first subunit of the Fc domain of (c);

and further comprising a second Fab fragment which binds to an antigen expressed on the surface of a target cell, wherein the second Fab is fused by the C-terminus of one of its chains to the N-terminus of the second subunit of the Fc domain of (c);

and the second antibody comprises:

d) a first Fab fragment, wherein the Fab fragment binds to an antigen expressed on the surface of a target cell;

e) a polypeptide comprising or consisting of an antibody light chain variable domain (VL) of an antigen binding site for the effector moiety; and

f) an Fc domain comprising a first and a second subunit,

wherein the polypeptide of (e) is fused by its N-terminus to the C-terminus of one of the chains of the Fab fragment of (d) and by its C-terminus to the N-terminus of the first subunit of the Fc domain of (f);

and further comprising a second Fab fragment which binds to an antigen expressed on the surface of a target cell, wherein the second Fab is fused by the C-terminus of one of its chains to the N-terminus of the second subunit of the Fc domain of (f);

wherein the second antibody does not comprise a VH domain of an antigen binding site for the effector moiety; and

wherein said VH domain of the first antibody and said VL domain of the second antibody are together capable of forming a functional antigen binding site for the effector moiety.

11. The set of antibodies of any one of the preceding claims, wherein the Fc domain is an IgG Fc domain.

12. The set of antibodies of any one of the preceding claims, wherein the Fc region is engineered to reduce or eliminate Fc effector function.
13. The set of antibodies according to any one of the preceding claims, wherein
in the first antibody the polypeptide of (b) is fused by its N-terminus via a linker to the C-terminus of one of the chains of the Fab fragment of (a) and by its C-terminus via a linker to the N-terminus of one of the subunits of the Fc domain of (c); and
in the second antibody the polypeptide of (e) is fused by its N-terminus via a linker to the C-terminus of one of the chains of the Fab fragment of (d) and by its C-terminus via a
10 linker to the N-terminus of one of the subunits of the Fc domain of (f).
14. The set of antibodies of claim 13, wherein the linker is a flexible linker of 10 to 70, 10 to 60, 10 to 50 amino acids or of 15 to 30 amino acids.
15. The set of antibodies of claim 13, wherein the linker is a flexible linker comprising 15 to 30 consecutive residues selected from Gly and Ser, optionally, 15-25 consecutive residues selected from Gly and Ser, optionally 16, 17, 18, 19, 20, 21, 22, 23 or 24 consecutive residues selected from Gly and Ser.
- 20 16. The set of antibodies of claim 15, wherein the linker is or comprises
GGGGSGGGSGGGSGGSGG (SEQ ID NO: 148) or GGGGSGGGSGGGSGGSGGS
(SEQ ID NO: 149) or GGGGSGGGSGGGSGGSGGG (SEQ ID NO: 150), optionally
GGGGSGGGSGGGSGGSGG (SEQ ID NO: 148) or GGGGSGGGSGGGSGGSGGG
(SEQ ID NO: 150).
17. The set of antibodies according to any of the preceding claims, wherein the effector moiety is selected from a drug, a cytotoxin, an imaging agent or a radiolabelled compound.
18. The set of antibodies according to any one of the preceding claims wherein the
30 effector moiety is a radiolabelled compound.
19. The set of antibodies according to claim 18, wherein the radiolabelled compound comprises radiolabelled DOTA or a salt or functional variant thereof.

20. The set of antibodies of claims 19, wherein the radiolabelled compound is DOTA or a salt or functional variant thereof radiolabelled with a Lu or Y radioisotope.
21. The set of antibodies according to claim 19 or 20, wherein the VH domain of the antigen binding site for the radiolabelled compound comprises (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO: 35; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO: 36; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO: 37.
22. The set of antibodies according to any one of claims 19 to 21, wherein the VH domain
10 of the antigen binding site for the radiolabelled compound comprises the amino acid sequence of SEQ ID NO: 41, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 41.
23. The set of antibodies of any one of claims 19 to 22, wherein the VL domain of the antigen binding site for the radiolabelled compound comprises (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO: 38; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO: 39; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO: 40.
24. The set of antibodies of any one of claims 19 to 23, wherein the VL domain of the
20 antigen binding site for the radiolabelled compound comprises an amino acid sequence of SEQ ID NO: 42 or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 42.
25. The set of antibodies according to claim 18, wherein the radiolabelled compound comprises Pb-DOTAM.
26. The set of antibodies according to claim 25, wherein the functional binding site for Pb-DOTAM binds with a K_D value of the binding affinity of 100pM, 50pM, 20pM, 10pM, 5pM, 1pM or less, e.g. 0.9pM or less, 0.8pM or less, 0.7pM or less, 0.6pM or less or 0.5pM
30 or less.
27. The set of antibodies according to claim 25 or claim 26, wherein the functional binding site for Pb-DOTAM binds to Pb-DOTAM and to Bi-DOTAM.

28. The set of antibodies of any one of claims 25 to 27, wherein the VH domain of the antigen binding site for the radiolabelled compound comprises:

a) heavy chain CDR2 comprising the amino acid sequence FIGSRGDTYYASWAKG (SEQ ID NO:2), or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 2, wherein these substitutions do not include Phe50, Asp56 and/or Tyr58, and optionally also do not include Gly52 and/or Arg54;

b) heavy chain CDR3 comprising the amino acid sequence ERDPYGGGAYPPHL (SEQ ID NO:3), or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 3, wherein these substitutions do not include Glu95, Arg96, Asp97, Pro98, and optionally also do not include Ala100C, Tyr100D, and/or Pro100E and/or optionally also do not include Tyr99;

and a heavy chain CDR1 which is optionally:

c) a heavy chain CDR1 comprising the amino acid sequence GFSLSTYSMS (SEQ ID NO:1) or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 1;

29. The set of antibodies of claim 28, wherein the VH domain of the antigen binding site for the radiolabelled compound comprises (a) CDR-H1 comprising the amino acid sequence of GFSLSTYSMS (SEQ ID NO:1); (b) CDR-H2 comprising the amino acid sequence of FIGSRGDTYYASWAKG (SEQ ID NO:2) and (c) CDR-H3 comprising the amino acid sequence of ERDPYGGGAYPPHL (SEQ ID NO:3).

30. The set of antibodies of any one of claims 25 to 29, wherein the VH domain of the antigen binding site for the radiolabelled compound comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 7 and SEQ ID NO 9, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 7 or SEQ ID NO: 9.

31. The set of antibodies of any one of claims 25 to 30, wherein the VL domain of the antigen binding site for the radiolabelled compound comprises

d) light chain CDR1 comprising the amino acid sequence QSSHSVYSDNDLA (SEQ ID NO:4) or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 4, wherein these substitutions do not include Tyr28 and Asp32;

- e) light chain CDR3 comprising the amino acid sequence LGGYDDESDTYG (SEQ ID NO:6) or a variant thereof having up to 1, 2, or 3 substitutions in SEQ ID NO: 6, wherein these substitutions do not include Gly91, Tyr92, Asp93, Thr95c and Tyr96, and a light chain CDR2 which is optionally
- f) light chain CDR 2 comprising the amino acid sequence QASKLAS (SEQ ID NO: 5) or a variant thereof having at least 1, 2 or 3 substitutions in SEQ ID NO: 5, optionally not including Gln50.

32. The set of antibodies of claim 31, wherein the VL domain of the antigen binding site for the radiolabelled compound comprises (d) CDR-L1 comprising the amino acid sequence of QSSHSVYSDNDLA (SEQ ID NO:4); (e) CDR-L2 comprising the amino acid sequence of QASKLAS (SEQ ID NO:5); and (f) CDR-L3 comprising the amino acid sequence of LGGYDDESDTYG (SEQ ID NO:6).

33. The set of antibodies of any one of claims 25 to 32, wherein the VL domain of the antigen binding site for the radiolabelled compound comprises an amino acid sequence of SEQ ID NO: 8, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 8.

34. The set of antibodies according to any one of the preceding claims, wherein the first and second antibody bind to the same target antigen.

35. The set of antibodies according to claim 34, wherein the first and second antibody bind to the same epitope of the same target antigen.

36. The set of antibodies according to claim 34, wherein the first and second antibody bind to different epitopes of the same target antigen.

37. The set of antibodies of any one of the preceding claims, wherein the target antigen is a tumour-associated antigen.

38. The set of antibodies of any one of the preceding claims, wherein the target antigen is selected from the group consisting of carcinoembryonic antigen (CEA), CD20, HER2, EGP-1

(epithelial glycoprotein-1, also known as trophoblast-2), colon-specific antigen-p (CSAp), a pancreatic mucin MUC1, GPRC5D and FAP.

39. The set of antibodies of any one of the preceding claims, wherein the target antigen is selected from the group consisting of CEA, GPRC5D and FAP.

40. The set of antibodies of any one of claims 1 to 39, wherein the target antigen is CEA

41. The set of antibodies according to claim 40, wherein the first antibody comprises an antigen-binding site for CEA comprising:

a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:19; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:20; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:21; and

a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:22; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:23; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:24.

42. The set of antibodies according to claim 40 or claim 41, wherein the first antibody comprises an antigen-binding site for CEA comprising a VH sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 25, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 25.

43. The set of antibodies according to any one of claims 40-42, wherein the first antibody comprises an antigen-binding site for CEA comprising a VL sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 26, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO: 26.

44. The set of antibodies according to claim 40, wherein the first antibody comprises an antigen-binding site which binds to CEA, comprising a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:43; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:44; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:45; and a light chain variable region comprising (d)

CDR-L1 comprising the amino acid sequence of SEQ ID NO:46; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:47; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:48.

45. The set of antibodies according to claim 40 or 44, wherein the first antibody comprises an antigen-binding site for CEA comprising a VH sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 49, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:49.

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46. The set of antibodies according to any one of claims 40, 44 or 45, wherein the first antibody comprises an antigen-binding site for CEA comprising a VL sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 50, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:50.

47. The set of antibodies according to claim 40, wherein the first antibody comprises an antigen-binding site which binds to CEA, comprising

20 a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:11; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:12; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:13; and

a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:14; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:15; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:16.

48. The set of antibodies according to claim 40 or 47, wherein the first antibody comprises an antigen-binding site for CEA comprising a VH sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 17, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:17.

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49. The set of antibodies according to claim 40, 47 or 48, wherein the first antibody comprises an antigen-binding site for CEA comprising a VL sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 18, or a variant thereof

comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:18.

50. The set of antibodies according to claim 40, wherein the first antibody comprises an antigen-binding site which binds to CEA, comprising

a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:59; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:60; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:61; and a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:62; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:63; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:64.

51. The set of antibodies according to claim 40 or claim 50, wherein the first antibody comprises an antigen-binding site for CEA comprising a VH sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 65, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:65.

52. The set of antibodies according to claim 40, 50 or 51, wherein the first antibody comprises an antigen-binding site for CEA comprising a VL sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 66, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:66.

53. The set of antibodies according to claim 40, wherein the first antibody comprises an antigen-binding site which binds to CEA, comprising a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:116; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:117 or 118; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:119; and a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:120, 121 or 122; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:123, 124 or 125; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:126.

54. The set of antibodies according to claim 40 or 53, wherein the first antibody comprises an antigen-binding site which binds to CEA, comprising a heavy chain variable region (VH) comprising an amino acid sequence selected from SEQ ID NO: 129, 130, 131, 132, 133 or 134, or a sequence with 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity thereto; and a light chain variable region (VL) comprising an amino acid sequence selected from SEQ ID NO: 135, 136, 137, 138, 139 or 140, or a sequence with 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity thereto.

55. The set of antibodies according to claim 40, 53 or 54, wherein the first antibody comprises an antigen-binding site which binds to CEA, comprising:
- (a) a VH domain comprising an amino acid sequence of SEQ ID NO:129 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or
 - (b) a VH domain comprising an amino acid sequence of SEQ ID NO:133 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or
 - (c) a VH domain comprising an amino acid sequence of SEQ ID NO:130 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or
 - (d) a VH domain comprising an amino acid sequence of SEQ ID NO:134 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
 - (e) a VH domain comprising an amino acid sequence of SEQ ID NO:133 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
 - (f) a VH domain comprising an amino acid sequence of SEQ ID NO:131 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
 - (g) a VH domain comprising an amino acid sequence of SEQ ID NO:129 and a VL domain comprising an amino acid sequence of SEQ ID NO:138.

56. The set of antibodies according to any one of claims 40 to 55, wherein the second antibody comprises an antigen-binding site which binds to CEA, comprising

a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:19; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:20; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:21; and

a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:22; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:23; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:24.

57. The set of antibodies according to any one of claims 40-56, wherein the second antibody comprises an antigen-binding site for CEA comprising a VH sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 25, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:25.

58. The set of antibodies according to any one of claims 40-57, wherein the second antibody comprises an antigen-binding site for CEA comprising a VL sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 26, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:26.

59. The set of antibodies according to any one of claim 40 to 55, wherein the second antibody comprises an antigen-binding site which binds to CEA, comprising a heavy chain variable region comprising (a)CDR-H1 comprising the amino acid sequence of SEQ ID NO:43; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:44; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:45; and a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:46; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:47; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:48.

60. The set of antibodies according to any one of claims 40 to 55 or 59, wherein the second antibody comprises an antigen-binding site for CEA comprising a VH sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 49, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:49.

61. The set of antibodies according to any one of claims 40 to 55, 59 or 60, wherein the second antibody comprises an antigen-binding site for CEA comprising a VL sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 50, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:50.

62. The set of antibodies according to any one of claims 40 to 55, wherein the second antibody comprises an antigen-binding site which binds to CEA, comprising a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:11; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:12; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:13; and a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:14; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:15; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:16.

10 63. The set of antibodies according to any one of claims 40 to 55 or 62, wherein the second antibody comprises an antigen-binding site for CEA comprising a VH sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 17, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:17.

64. The set of antibodies according to any one of claims 40 to 55, 62 or 63, wherein the second antibody comprises an antigen-binding site for CEA comprising a VL sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 18, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96,
20 97, 98, or 99% identity to SEQ ID NO:18.

65. The set of antibodies according to any one of claim 40 to 55, wherein the second antibody comprises an antigen-binding site which binds to CEA, comprising a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:59; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:60; and (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:61; and a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:62; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:63; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:64.

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66. The set of antibodies according to any one of claims 40 to 55 or 65, wherein the second antibody comprises an antigen-binding site for CEA comprising a VH sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 65, or

a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:65.

67. The set of antibodies according to claim 40 to 55, 65 or 66, wherein the second antibody comprises an antigen-binding site for CEA comprising a VL sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 66, or a variant thereof comprising an amino acid sequence having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity to SEQ ID NO:66.

10 68. The set of antibodies according to claim 40 to 55, wherein the second antibody comprises an antigen-binding site which binds to CEA, comprising a heavy chain variable region comprising (a) CDR-H1 comprising the amino acid sequence of SEQ ID NO:116; (b) CDR-H2 comprising the amino acid sequence of SEQ ID NO:117 or 118; (c) CDR-H3 comprising the amino acid sequence of SEQ ID NO:119; and a light chain variable region comprising (d) CDR-L1 comprising the amino acid sequence of SEQ ID NO:120, 121 or 122; (e) CDR-L2 comprising the amino acid sequence of SEQ ID NO:123, 124 or 125; and (f) CDR-L3 comprising the amino acid sequence of SEQ ID NO:126.

20 69. The set of antibodies according to claim 40 to 55 or 68, wherein the second antibody comprises an antigen-binding site which binds to CEA, comprising a heavy chain variable region (VH) comprising an amino acid sequence selected from SEQ ID NO: 129, 130, 131, 132, 133 or 134, or a sequence with 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity thereto; and a light chain variable region (VL) comprising an amino acid sequence selected from SEQ ID NO: 135, 136, 137, 138, 139 or 140, or a sequence with 90, 91, 92, 93, 94, 95, 96, 97, 98, or 99% identity thereto.

70. The set of antibodies according to claim 40 to 55, 68 or 69, wherein the second antibody comprises an antigen-binding site which binds to CEA, comprising:
30 (a) a VH domain comprising an amino acid sequence of SEQ ID NO:129 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or
(b) a VH domain comprising an amino acid sequence of SEQ ID NO:133 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or
(c) a VH domain comprising an amino acid sequence of SEQ ID NO:130 and a VL domain comprising an amino acid sequence of SEQ ID NO:139, or

- (d) a VH domain comprising an amino acid sequence of SEQ ID NO:134 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
- (e) a VH domain comprising an amino acid sequence of SEQ ID NO:133 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
- (f) a VH domain comprising an amino acid sequence of SEQ ID NO:131 and a VL domain comprising an amino acid sequence of SEQ ID NO:138, or
- (g) a VH domain comprising an amino acid sequence of SEQ ID NO:129 and a VL domain comprising an amino acid sequence of SEQ ID NO:138.

10 71. The set of antibodies according to claim 40, wherein the first antibody is an antibody according to any one of claims 47 to 49 and the second antibody is an antibody according to any one of claims 56 to 58.

72. The set of antibodies according to claim 71, wherein the first and second antibody comprise an antigen-binding site which binds to CEA comprising a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26 and a heavy chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25.

20 73. The set of antibodies according to claim 71 or 72, wherein the first antibody and second antibody are monospecific and monovalent for the target antigen.

74. The set of antibodies according to claim 73, wherein the first and second antibody each comprise two antigen binding sites which bind to CEA comprising a light chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 26 and a heavy chain variable domain having at least 90, 91, 92, 93, 94, 95, 96, 97, 98, 99 or 100% identity to SEQ ID NO 25.

30 75. The set of antibodies according to claim 1 or 72, wherein the first antibody comprises: a first heavy chain of SEQ ID NO: 146; a second heavy chain of SEQ ID NO: 144; and a light chain of SEQ ID NO: 143; and/or the second antibody comprises a first heavy chain of SEQ ID NO: 145; a second heavy chain of SEQ ID NO: 144; and a light chain of SEQ ID NO: 143.

76. The set of antibodies according to claim 1 or claim 74, wherein:
the first antibody comprises: a first heavy chain polypeptide of SEQ ID NO: 146; a second heavy chain polypeptide of SEQ ID NO: 147; and a first and second light chain polypeptide of SEQ ID NO: 143; and/or
the second antibody comprises: a first heavy chain polypeptide of SEQ ID NO: 145; a second heavy chain polypeptide of SEQ ID NO: 147; and a first and second light chain polypeptide of SEQ ID NO: 143.

10 77. A set of nucleic acids expressing the set of antibodies of any one of the preceding claims.

78. An expression vector or set of expression vectors comprising the set of nucleic acids according to claim 77.

79. A host cell or set of host cells comprising the expression vector or set of expression vectors of claim 78.

80. A method of pretargeting radioimmunotherapy, comprising
i) administering to a subject the set of antibodies according to any one of claims 1 to
20 76, wherein the first and second antibodies are administered simultaneously or sequentially, in either order, wherein the antibodies bind to the target antigen and localise to the surface of a cell expressing the target antigen; and wherein association of the first and second antibody forms a functional binding site for the radiolabelled compound;
and
ii) subsequently administering a radiolabelled compound, wherein the radiolabelled compound binds to functional binding site for the radiolabelled compound.

81. The method of claim 80, wherein the method does not comprise a step of
administering a clearing or blocking agent.

30

82. The method of claim 80 or 81, wherein the subject is human.

83. The method of any one of claims 80 to 82, wherein the target antigen is a cancer- or tumour-associated antigen and the method is a method of radioimmunotherapy of a tumour or cancer.

84. The set of antibodies according to any one of claims 1 to 76, for use in a method of pretargeting radioimmunotherapy according to any one of claims 80 to 83.

85. A method of targeting a radioisotope to a tissue or organ for radioimaging comprising:

10 i) administering to a subject a set of antibodies according to any one of claims 1 to 76, wherein the first and second antibodies are administered simultaneously or sequentially, in either order, wherein the antibodies bind to a target antigen and localise to the surface of a cell expressing the target antigen, wherein association of the first and second antibody forms a functional binding site for the radiolabelled compound;

and

ii) subsequently administering a radiolabelled compound, wherein the radiolabelled compound binds to the functional binding site for the radiolabelled compound.

20 86. The method of claim 85, wherein the method does not comprise a step of administering a clearing or blocking agent.

87. The method of claim 85 or 86, wherein the method further comprises a step of imaging.

88. The method of claim 87, wherein the target antigen is a cancer or tumour-associated antigen and the method is a method of imaging tumours or cancer.

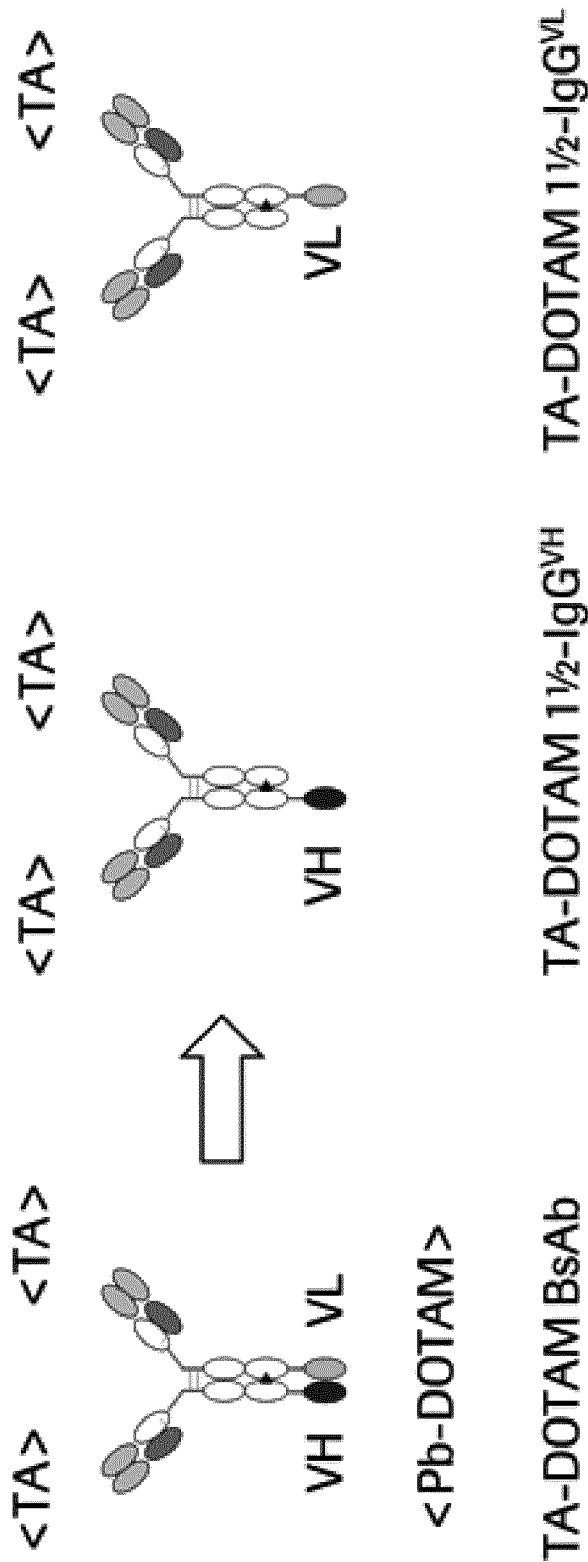


FIG. 1

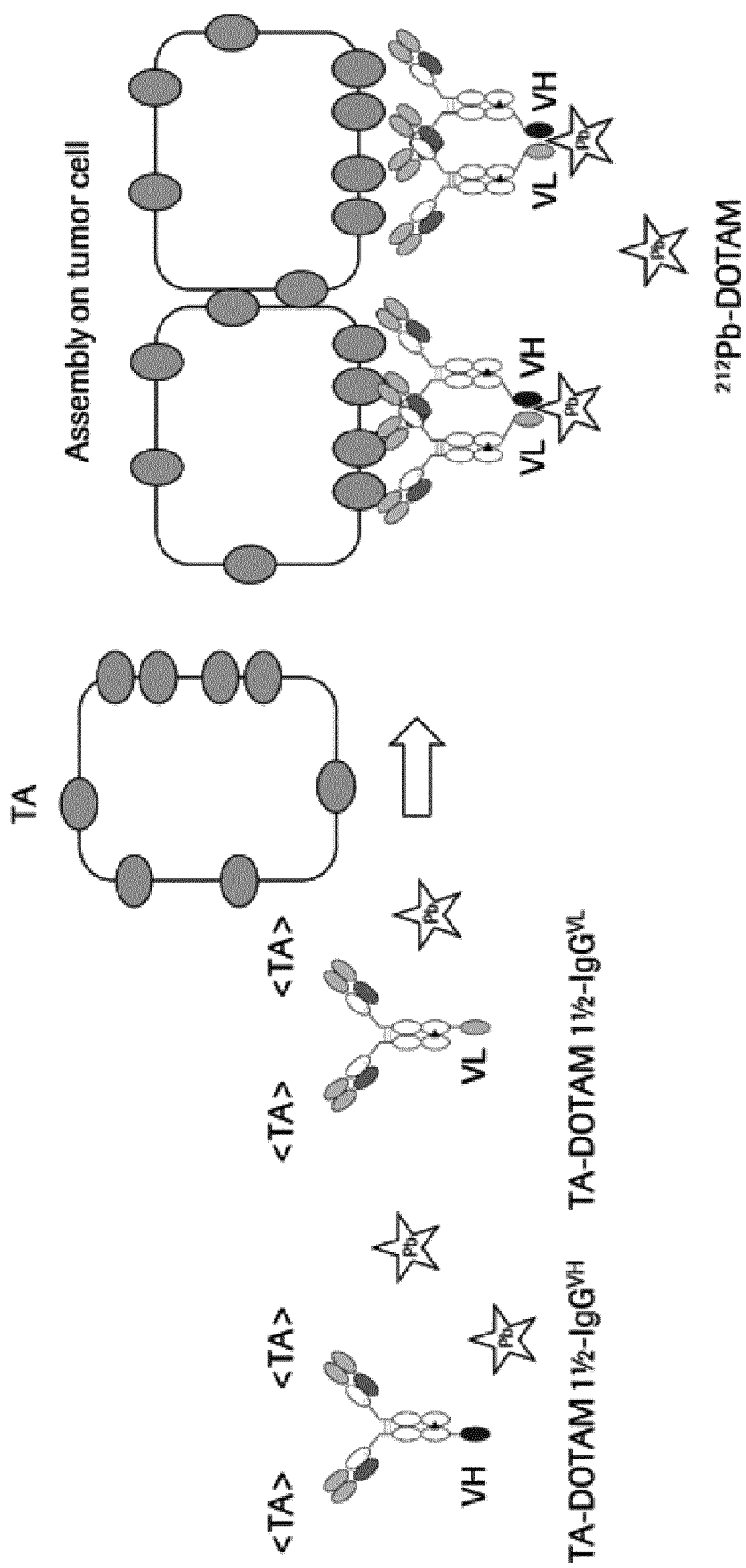


FIG. 2

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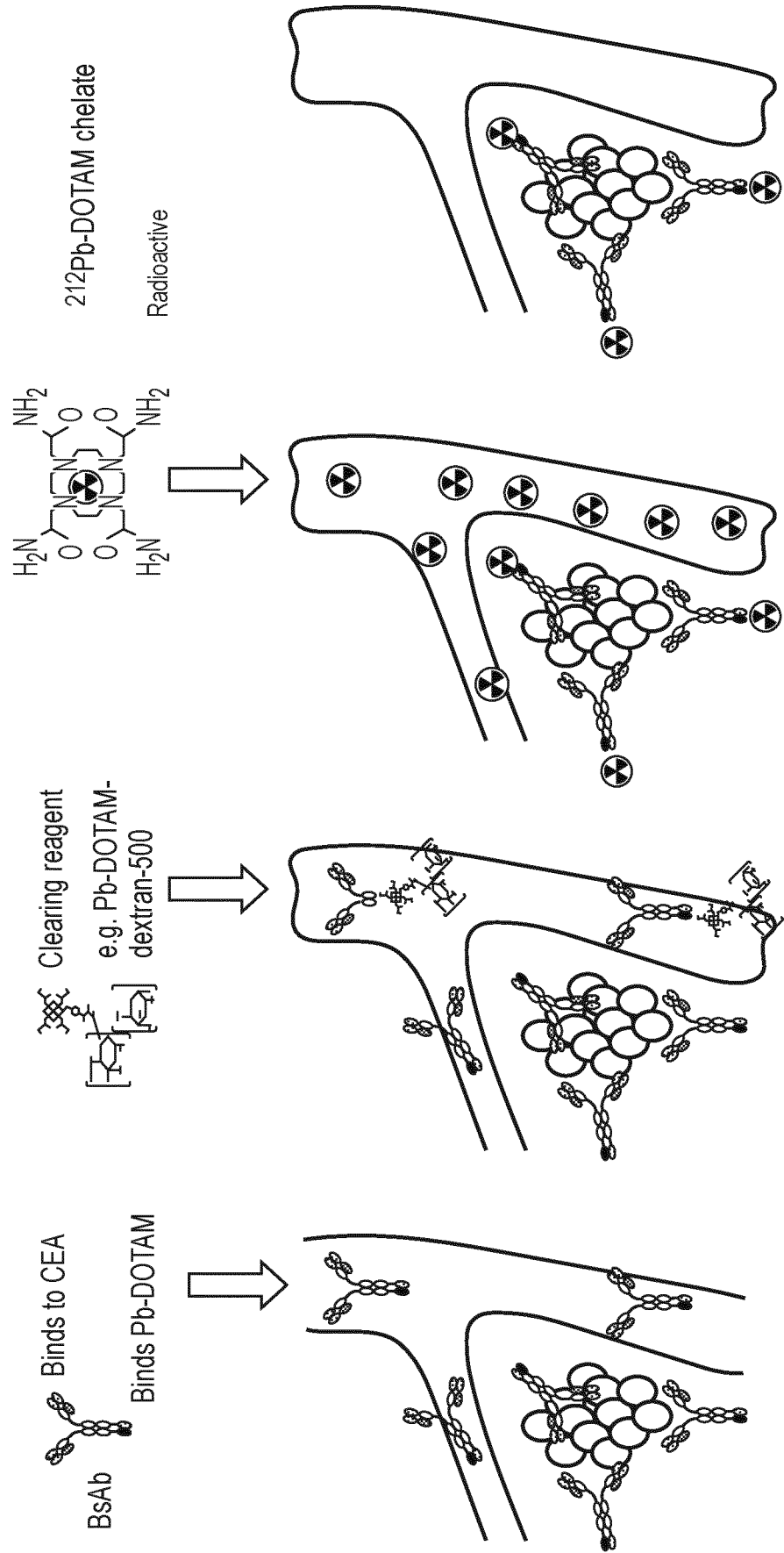


FIG. 3

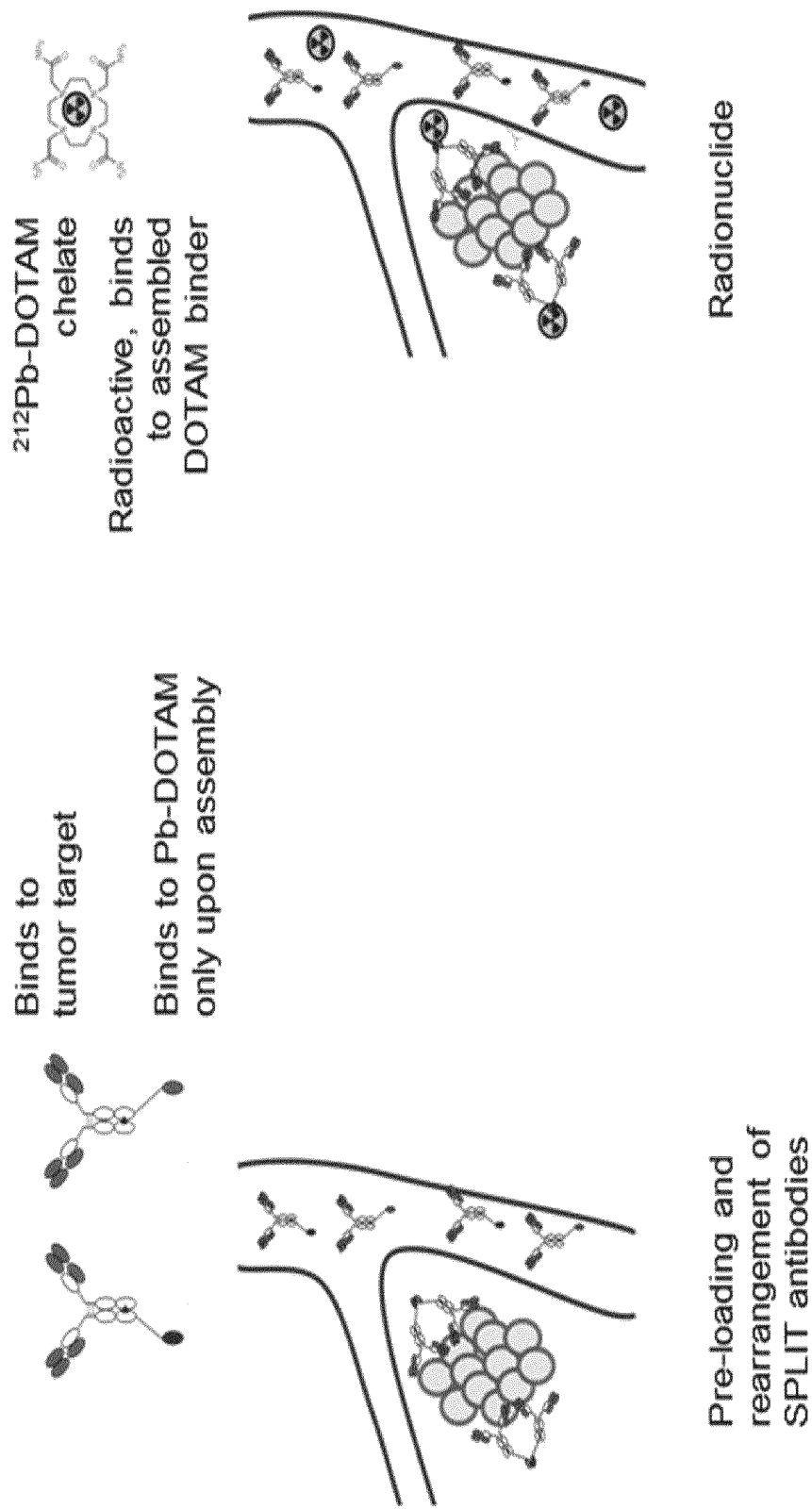


FIG. 4

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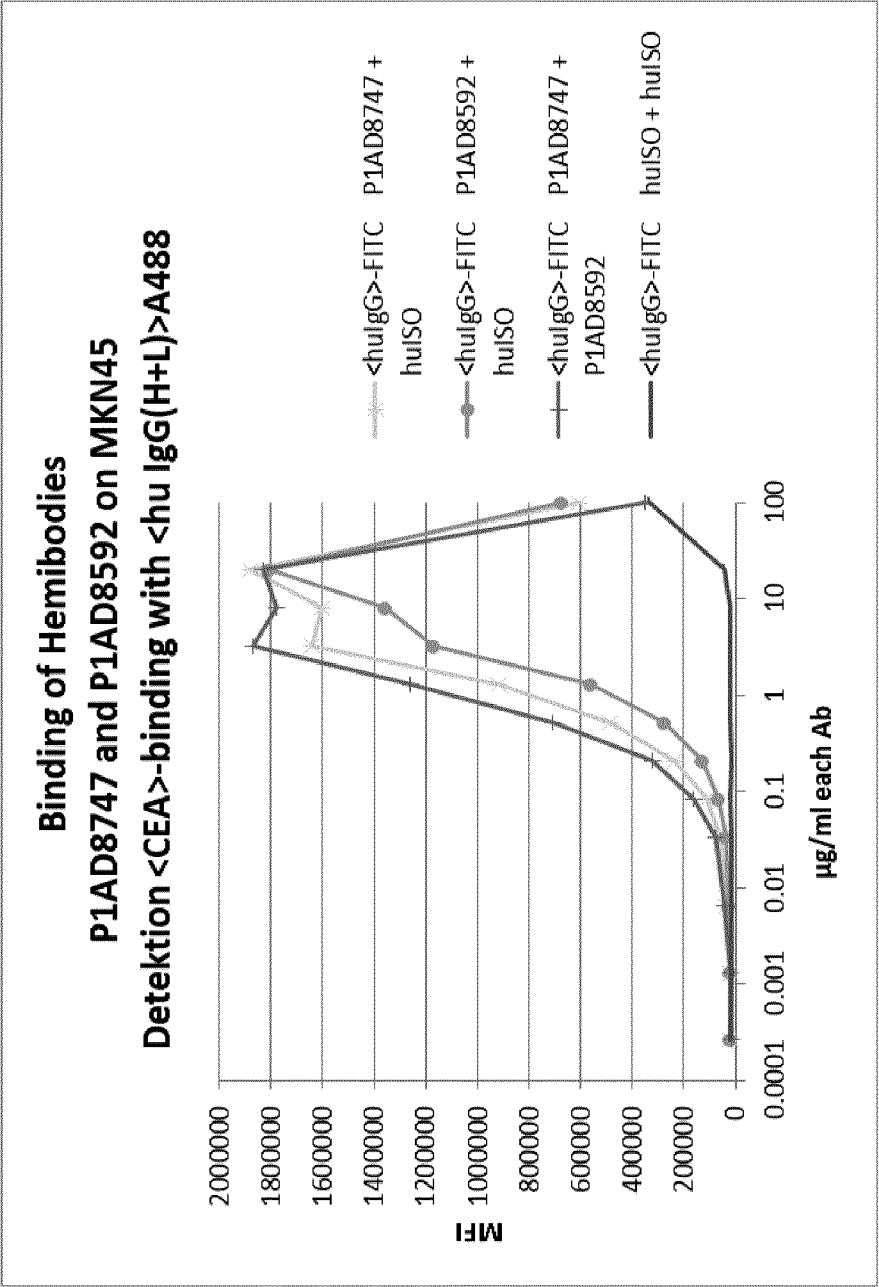


FIG. 5

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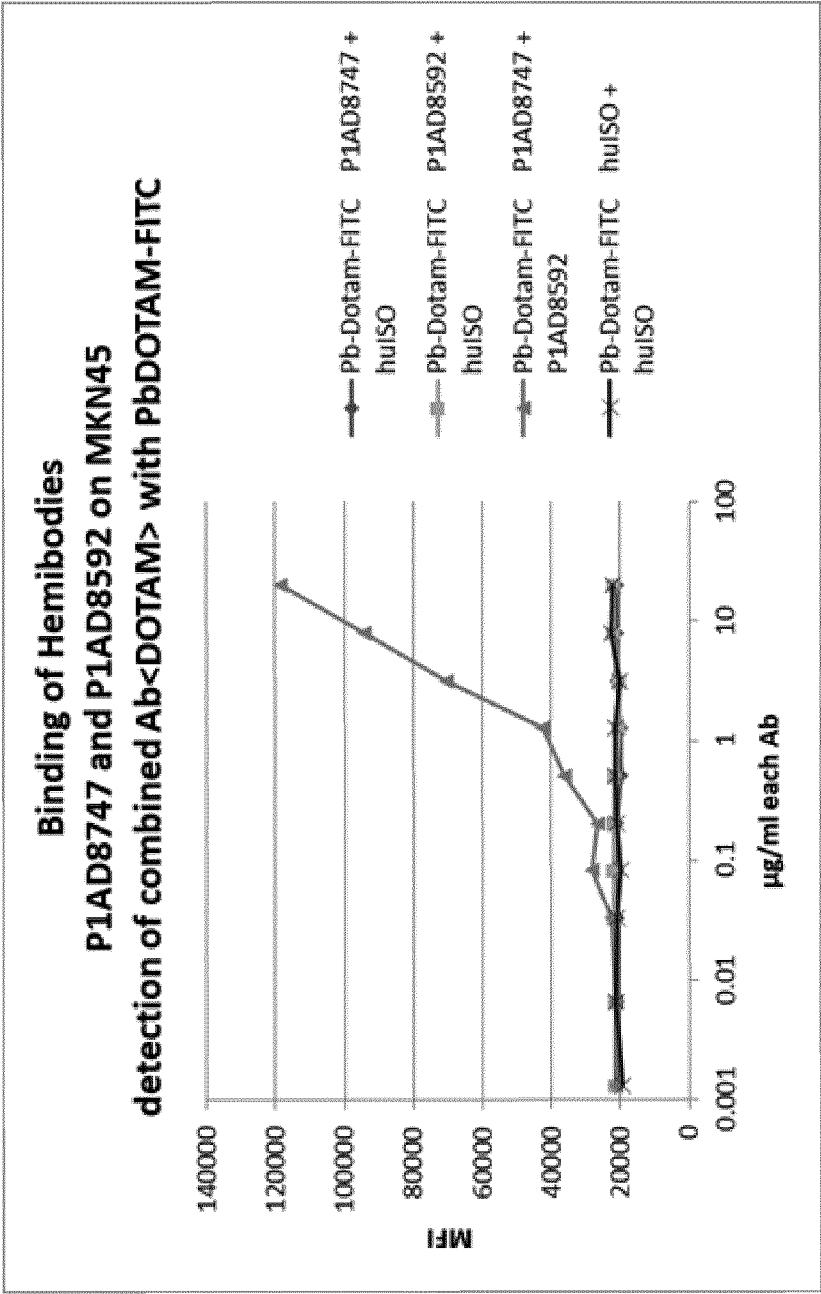


FIG. 6

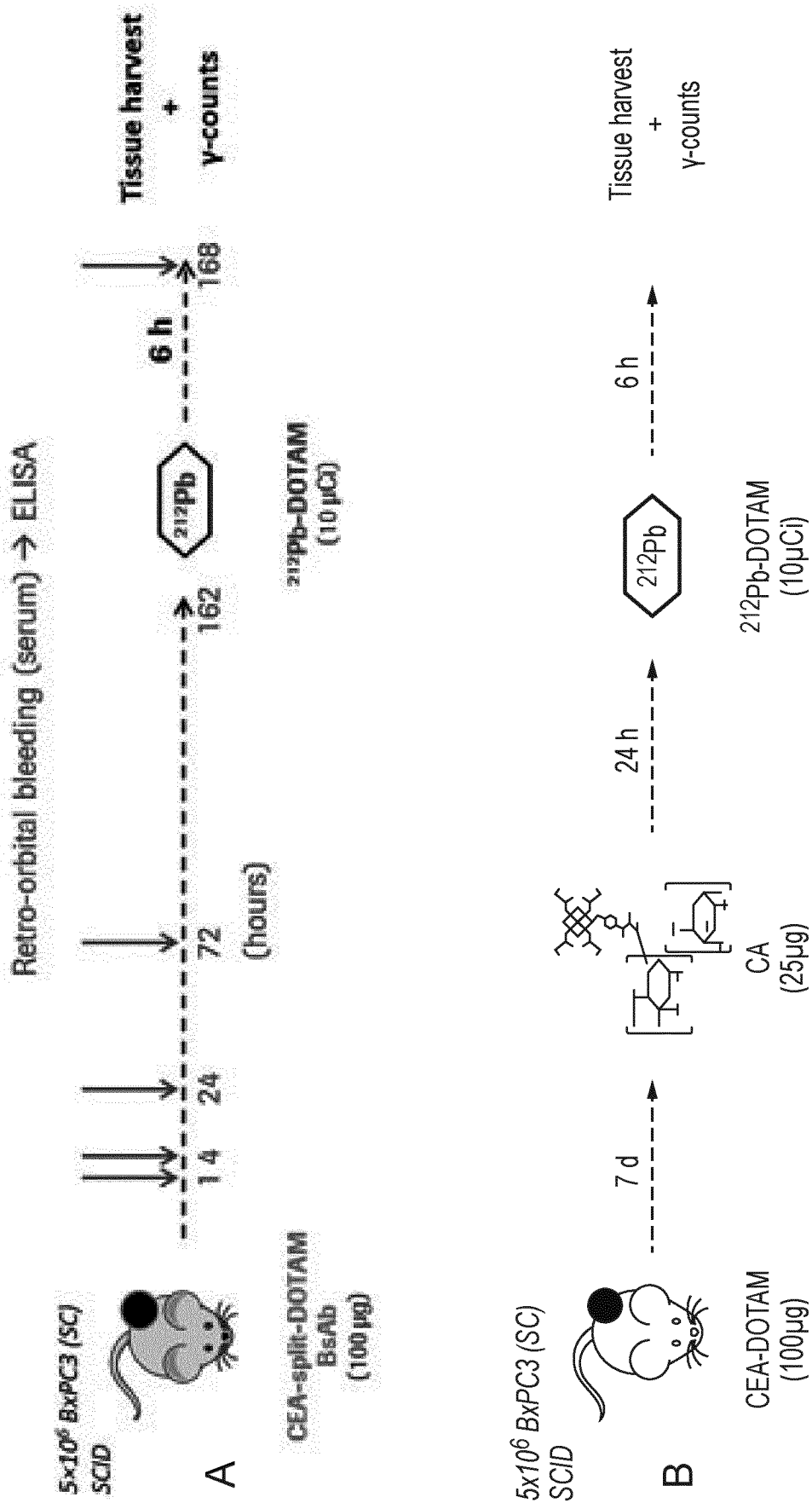


FIG. 7

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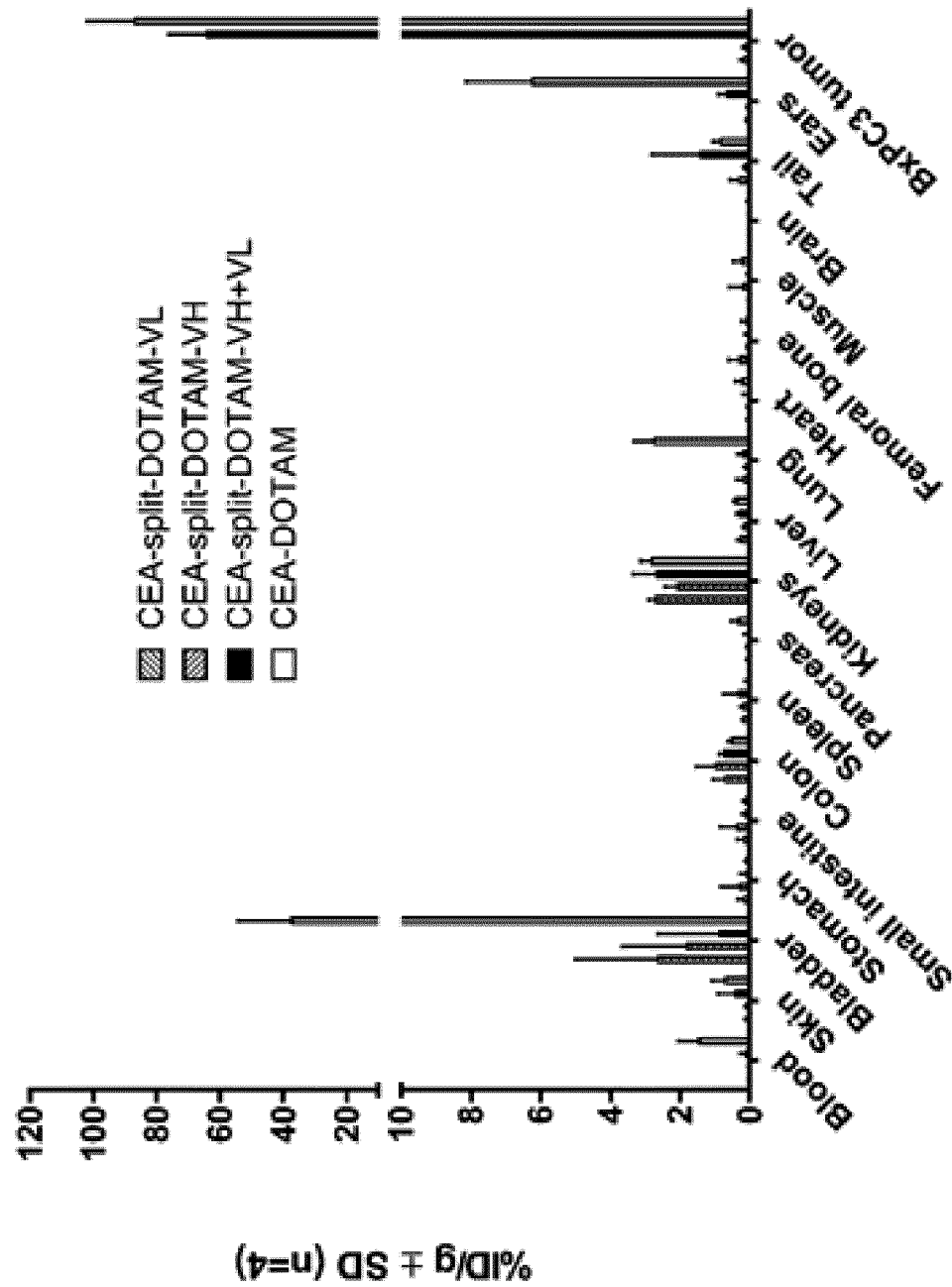


FIG. 8

Split Antibody Pharmacokinetics

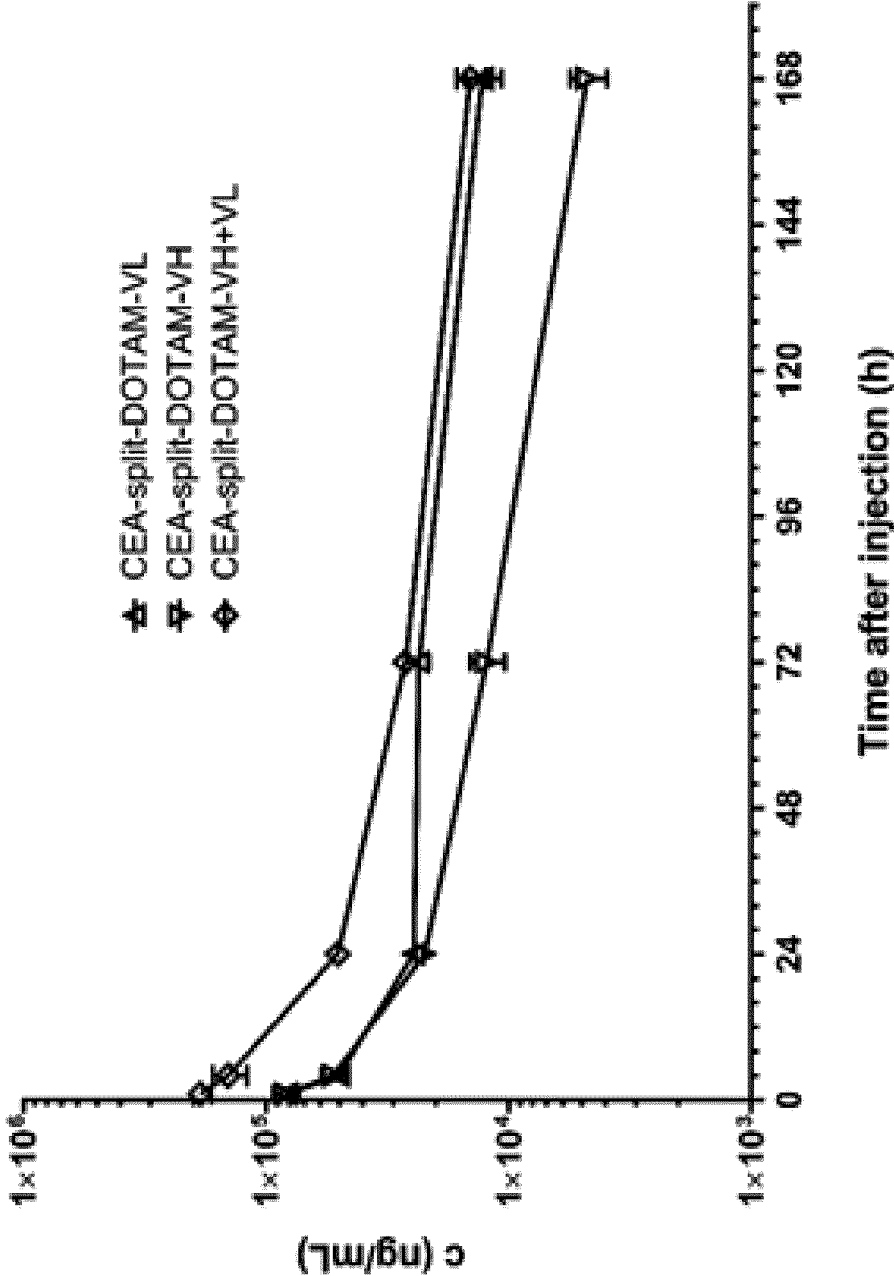


FIG. 9

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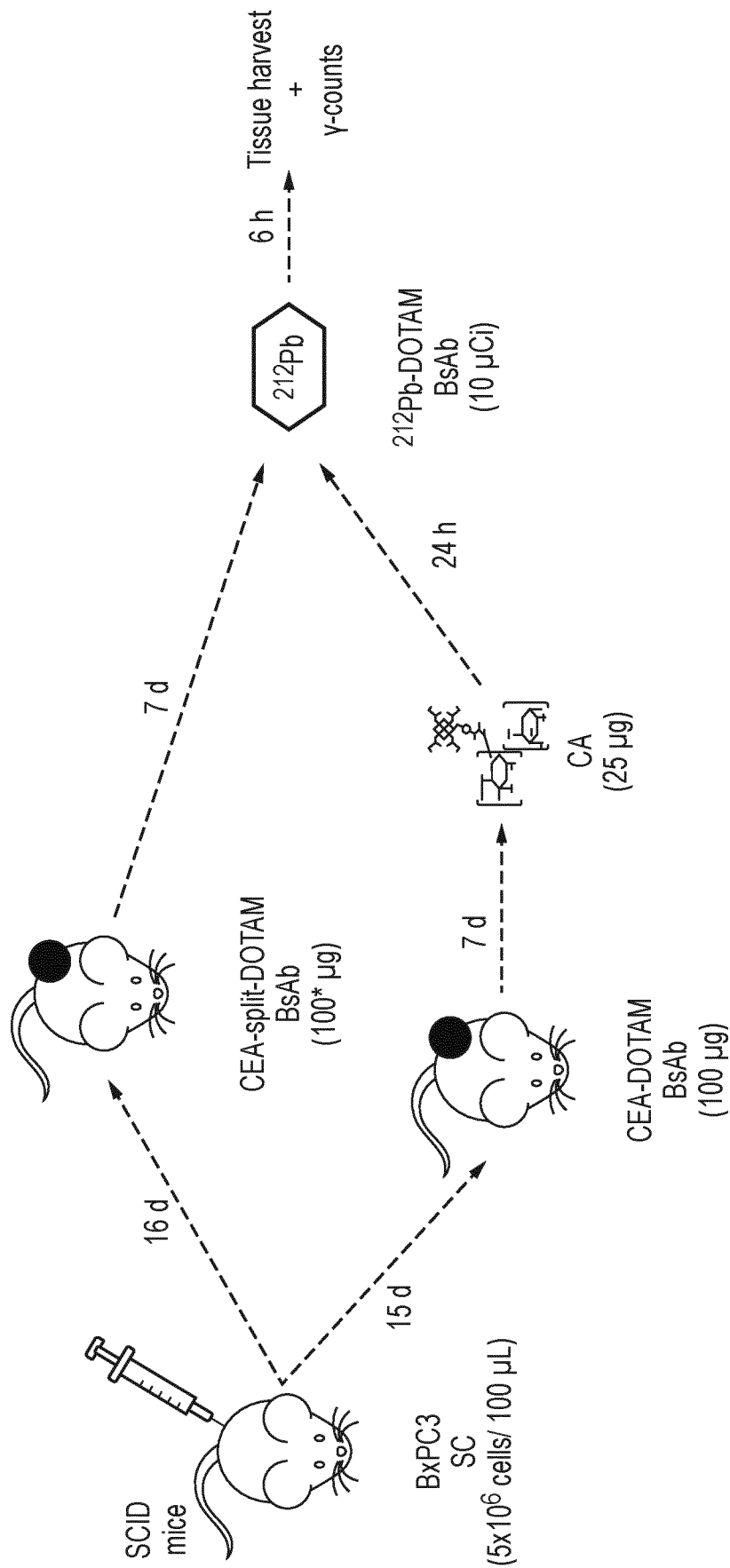


FIG. 10

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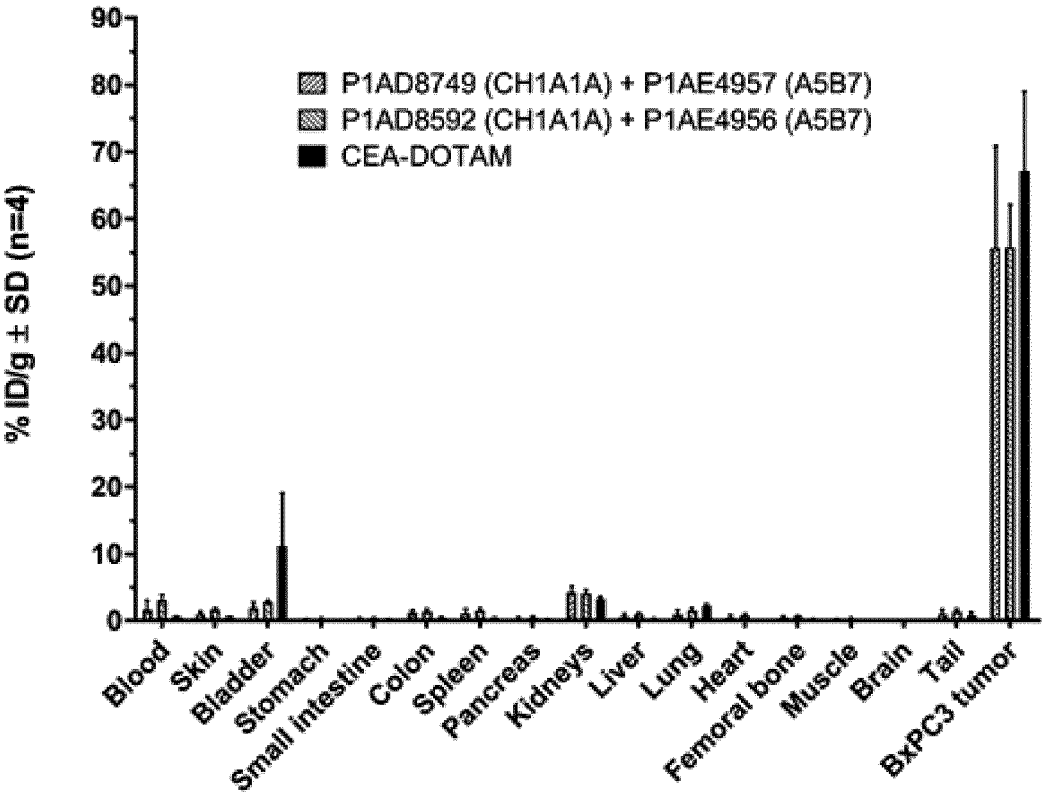


FIG. 11

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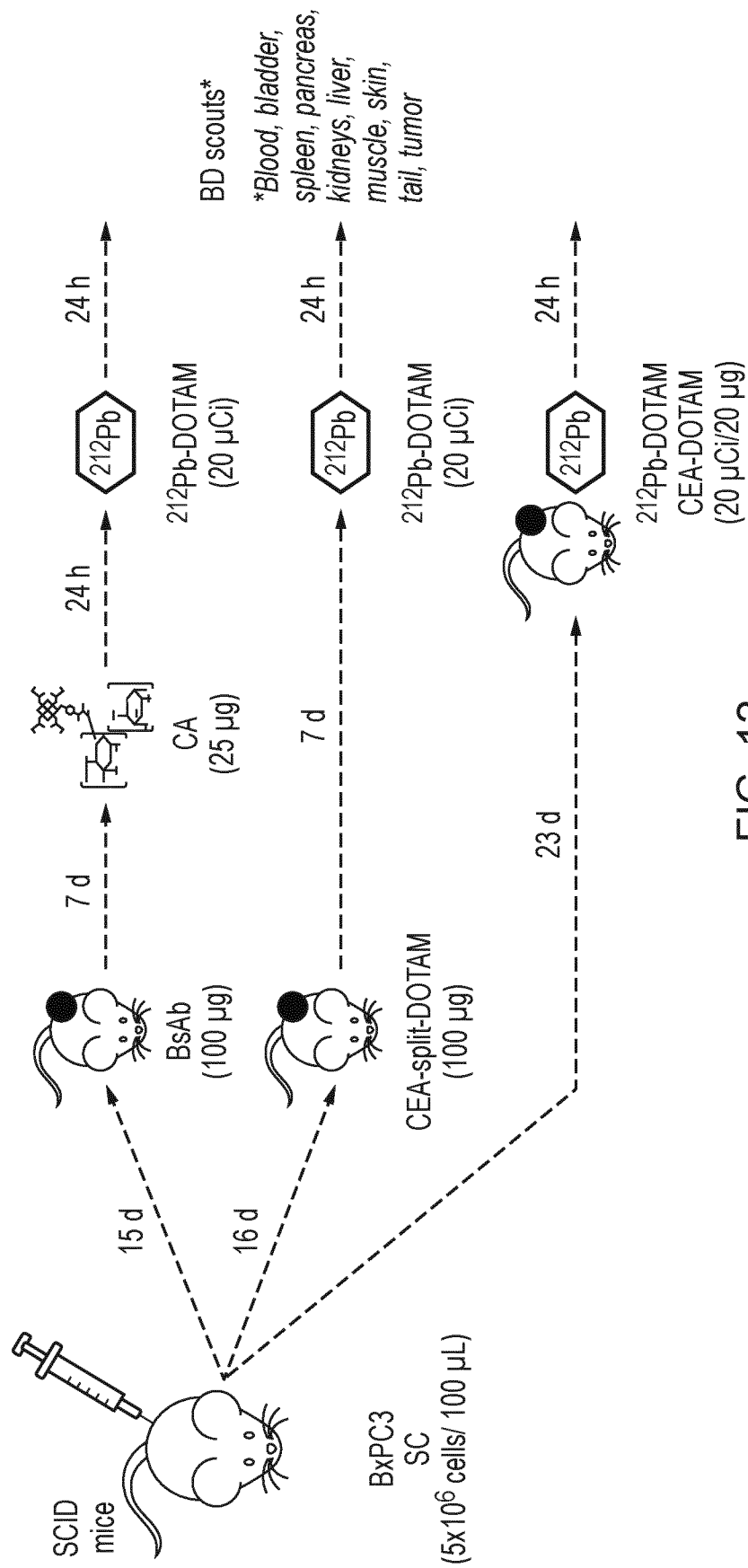


FIG. 12

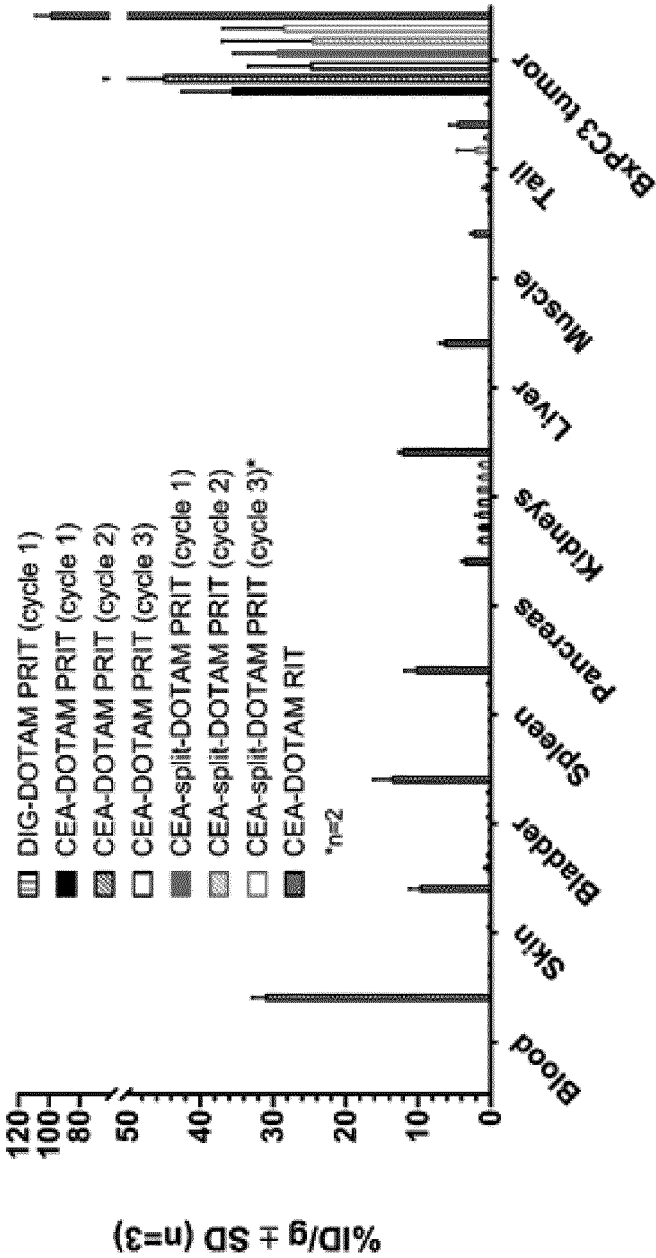


FIG. 13

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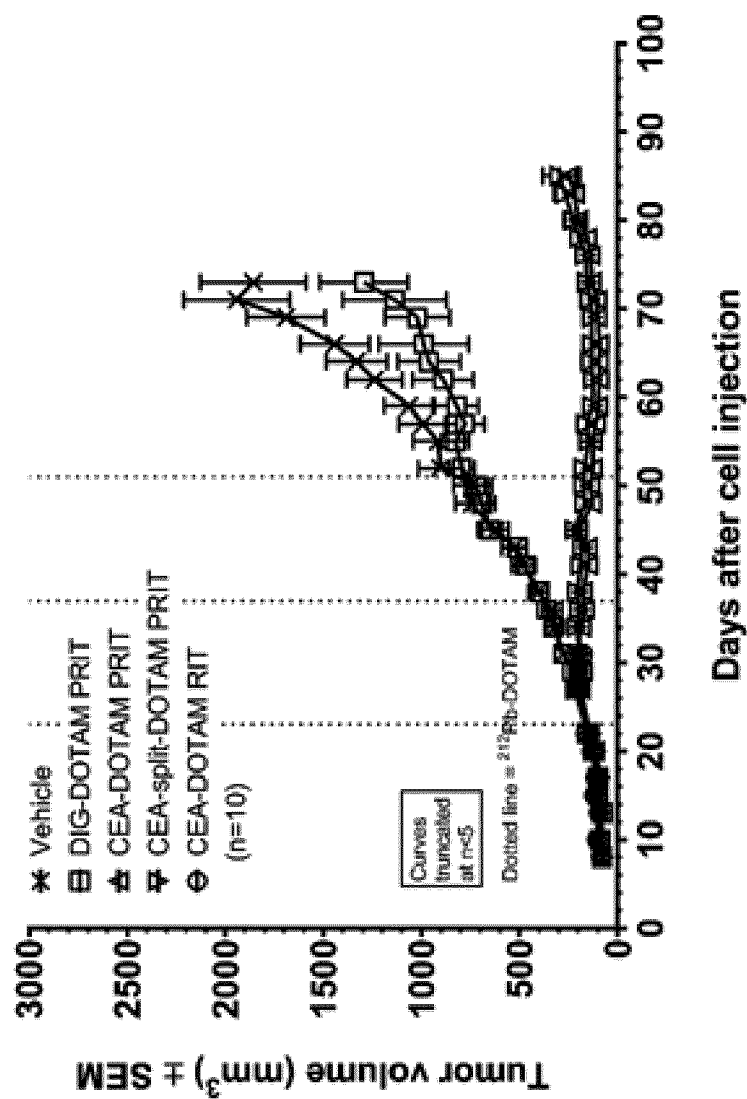


FIG. 14

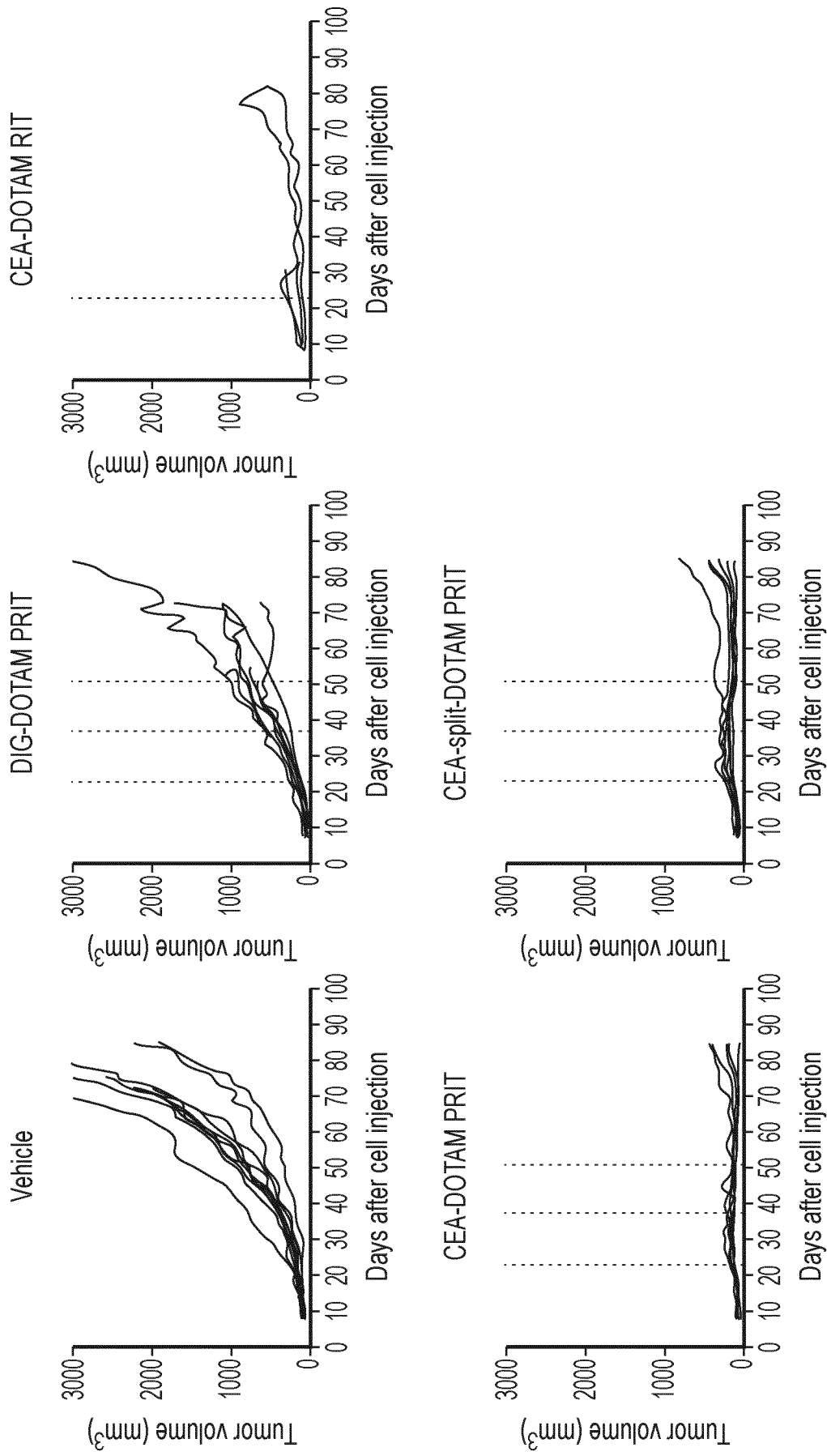


FIG. 15

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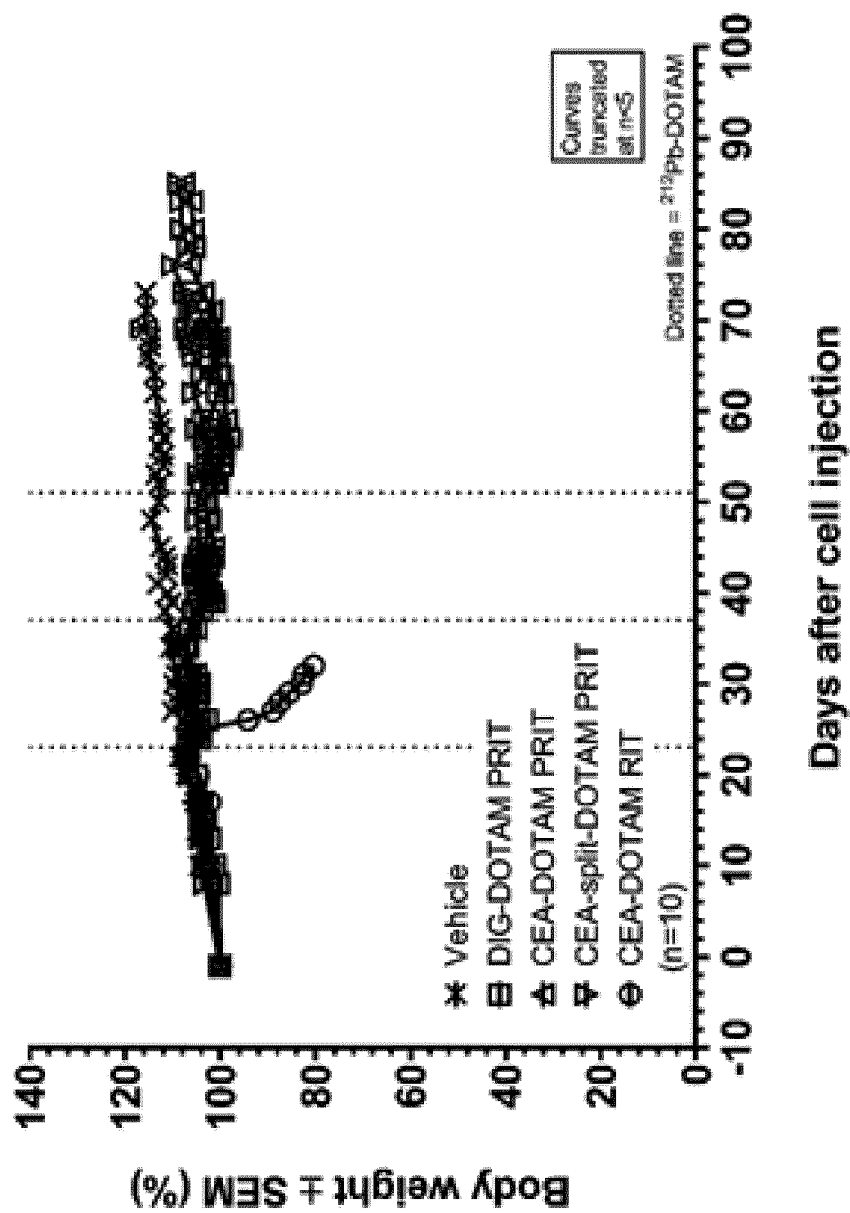


FIG. 16

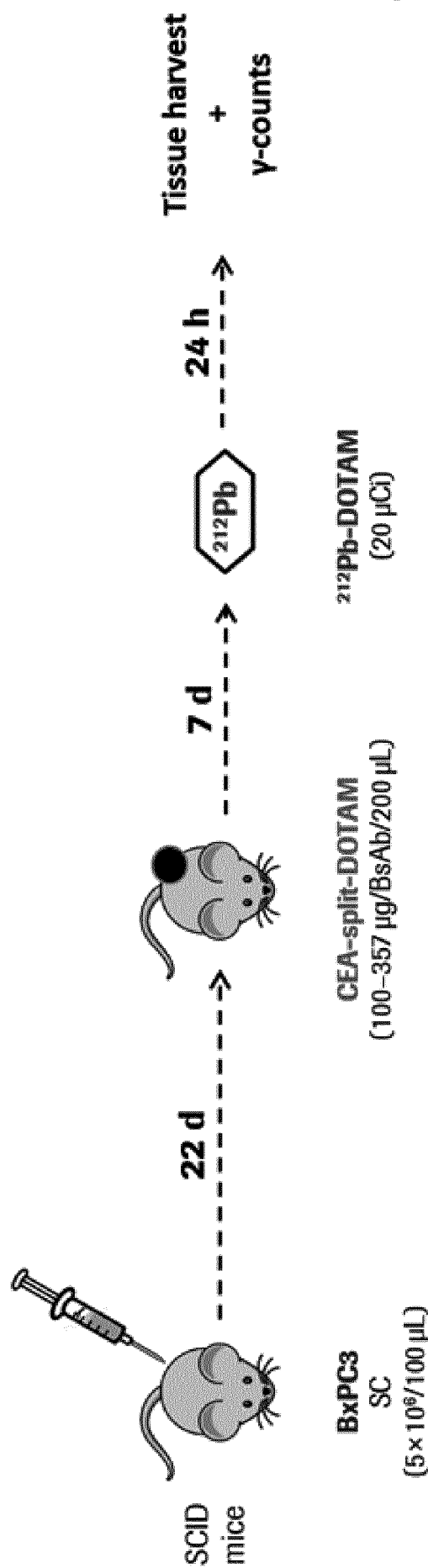


FIG. 17

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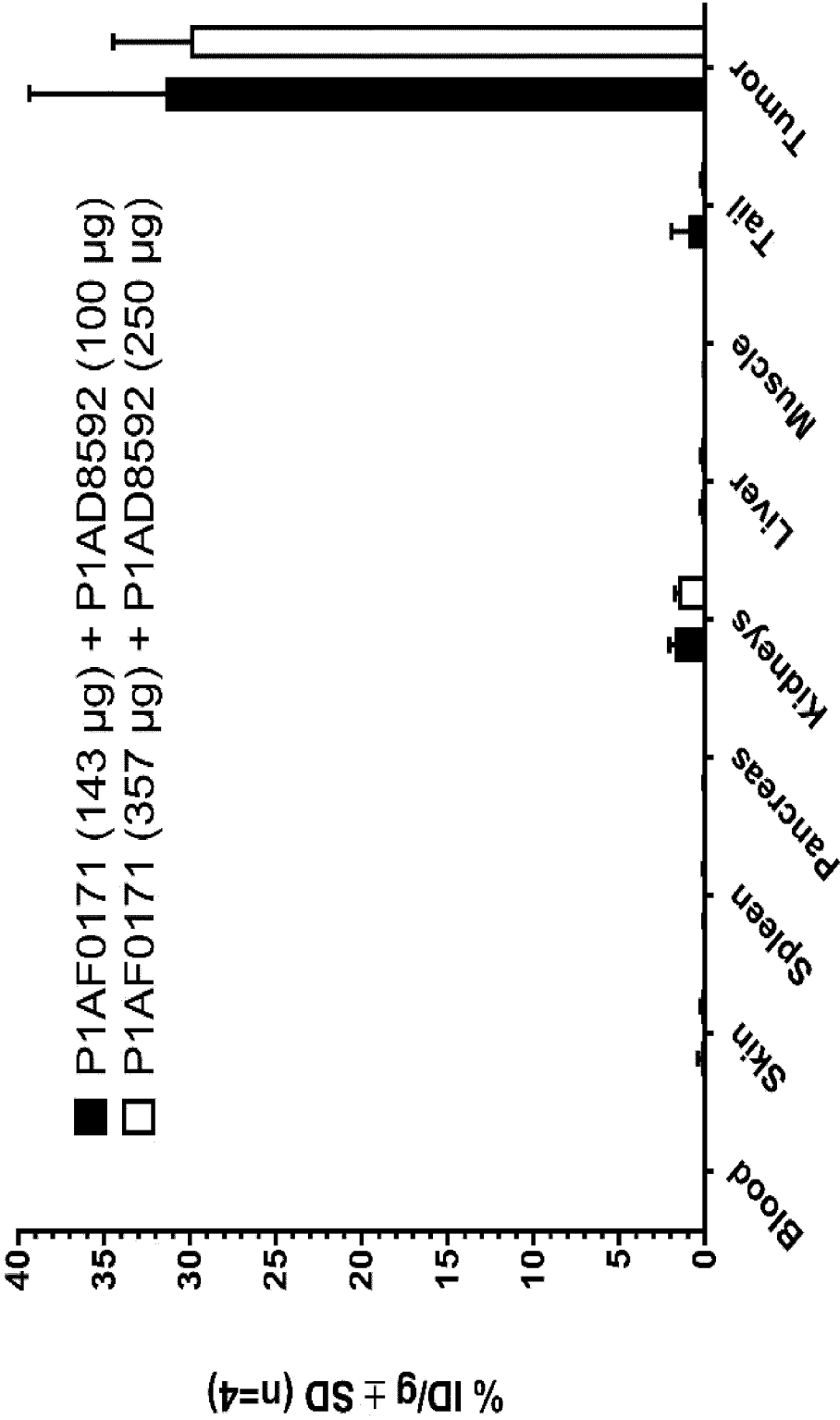


FIG. 18

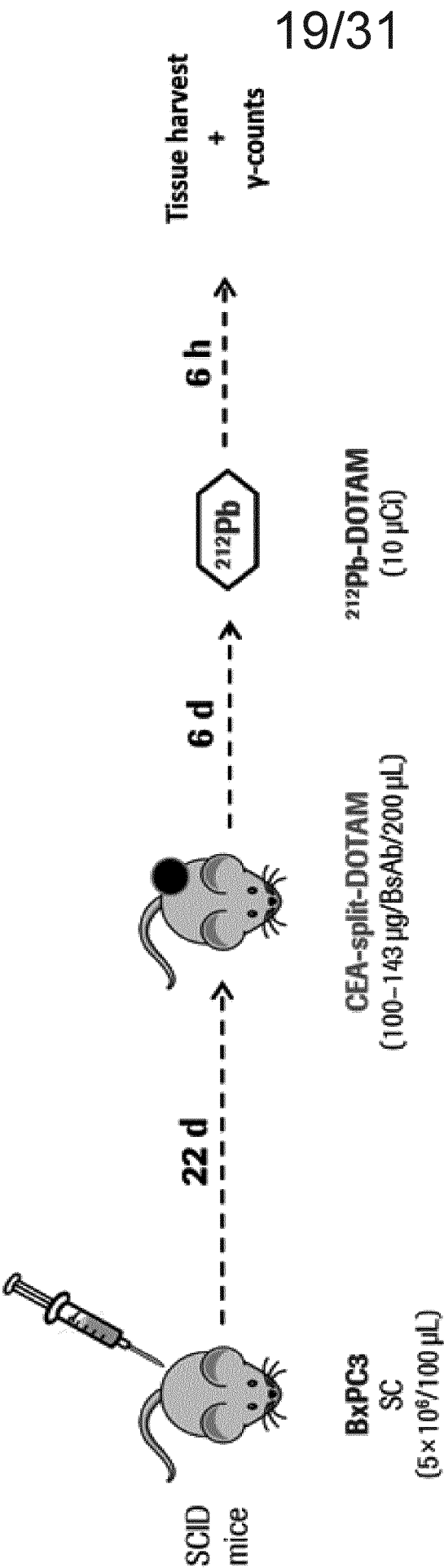


FIG. 19

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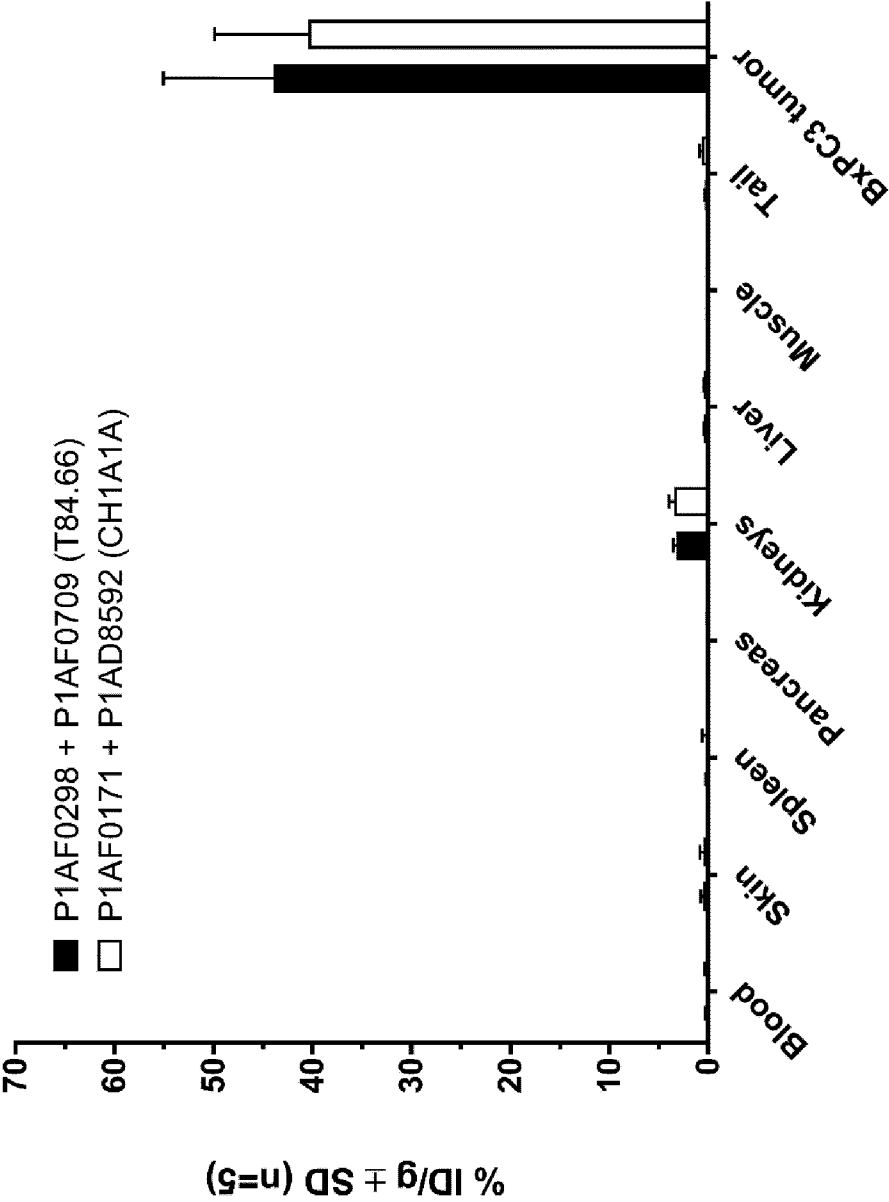


FIG. 20

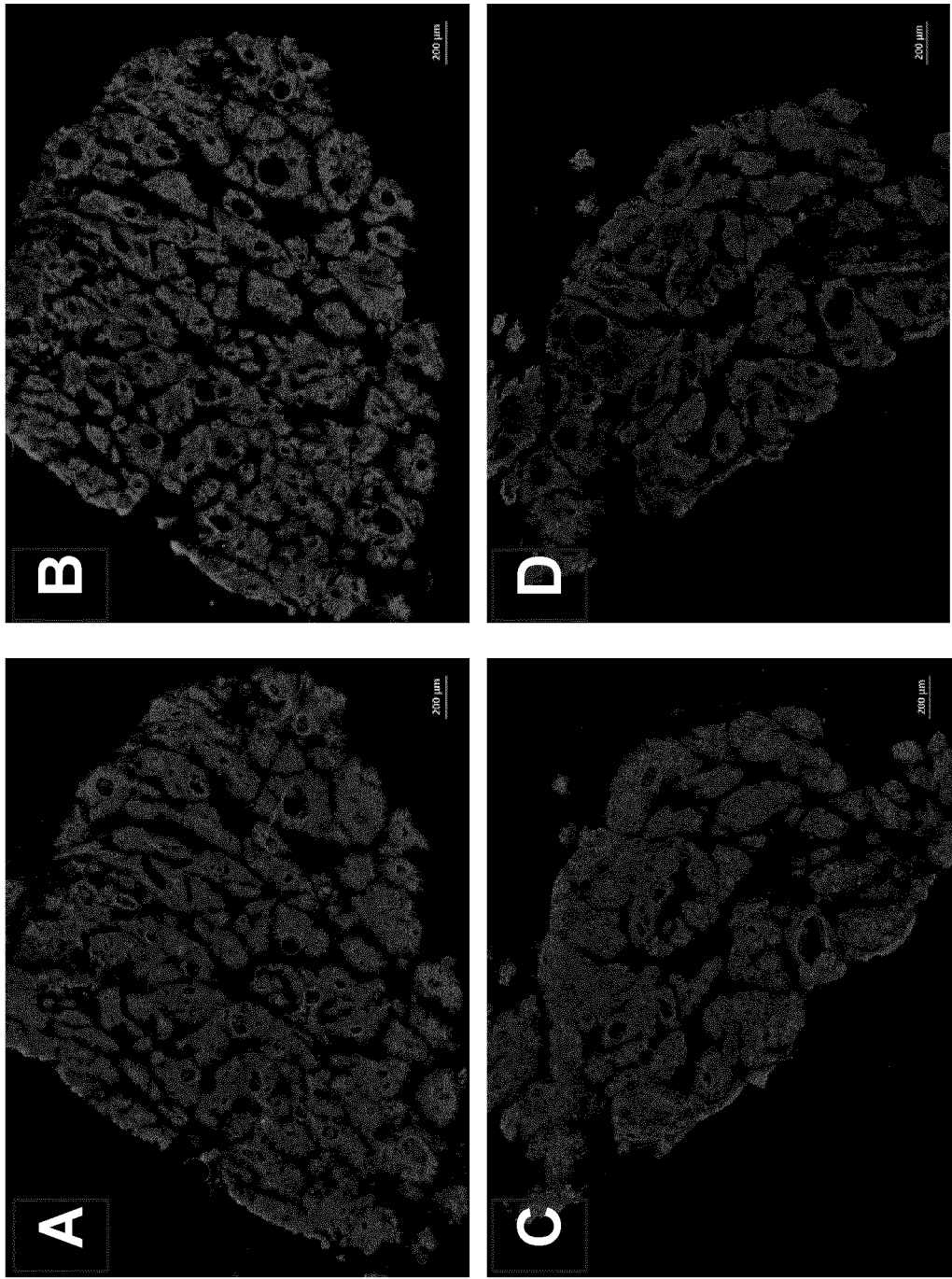


FIG. 21

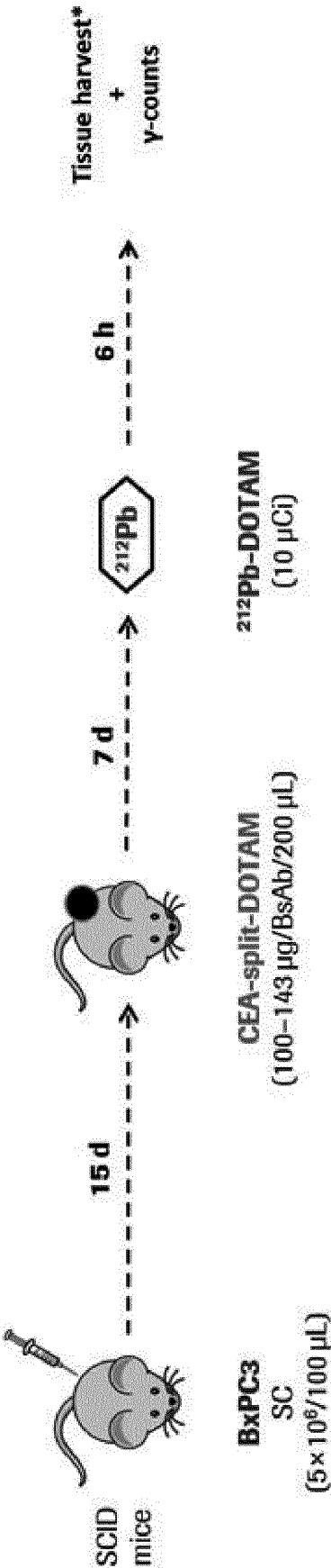


FIG. 22

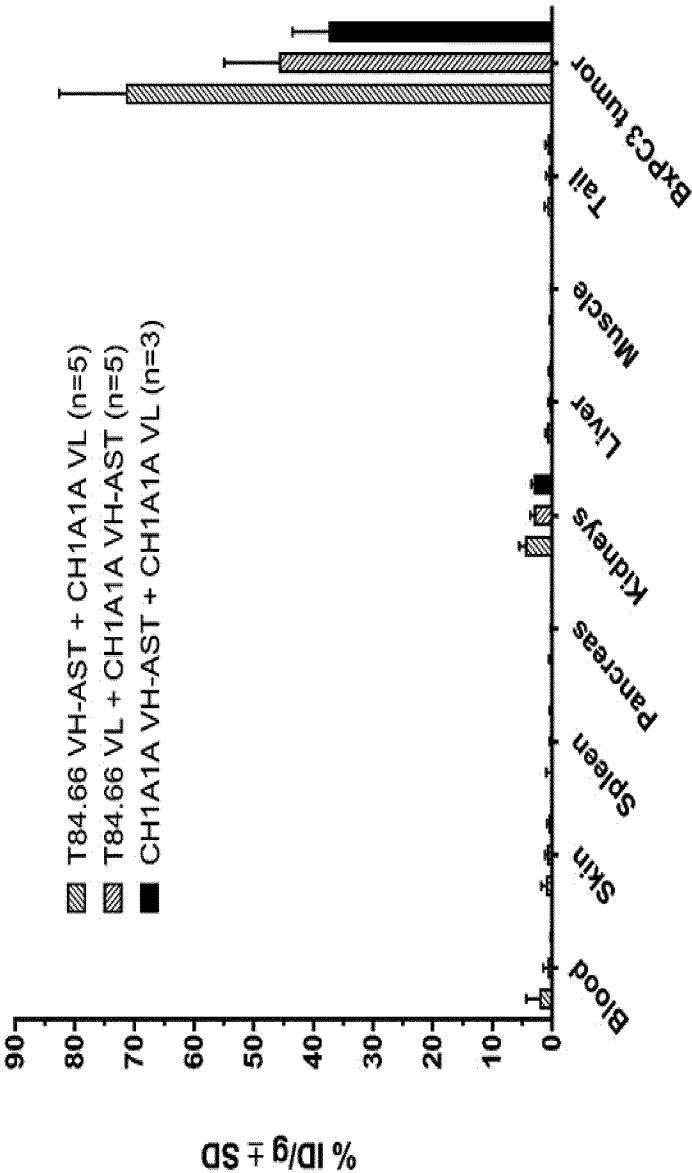


FIG. 23

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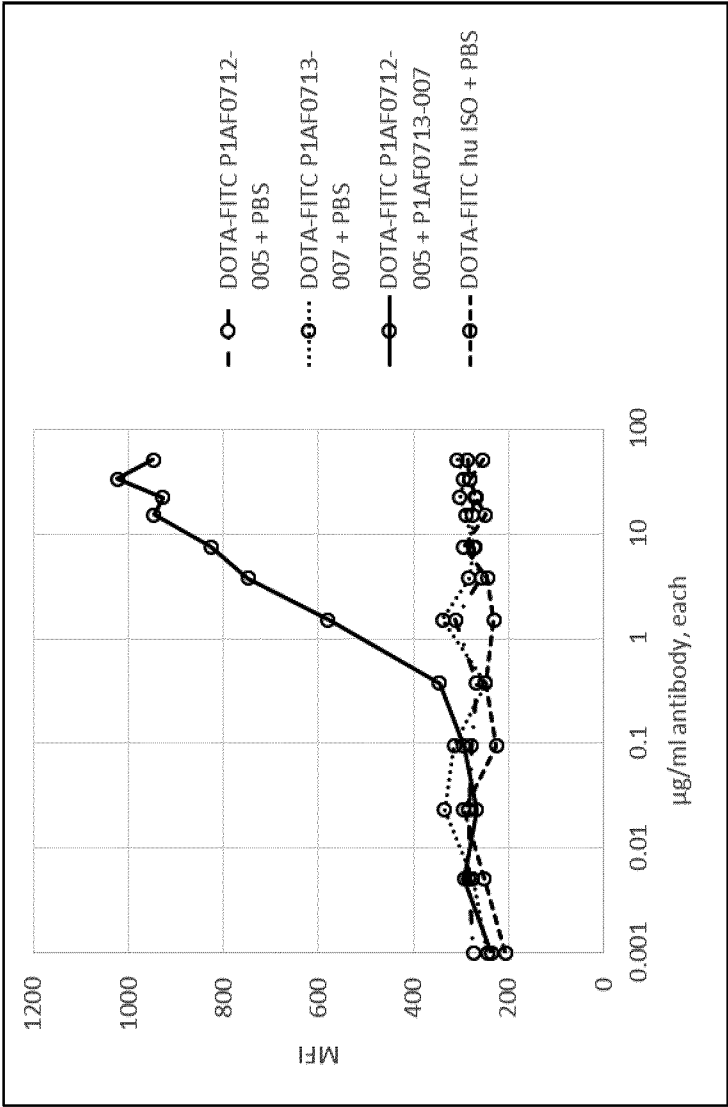


FIG. 24

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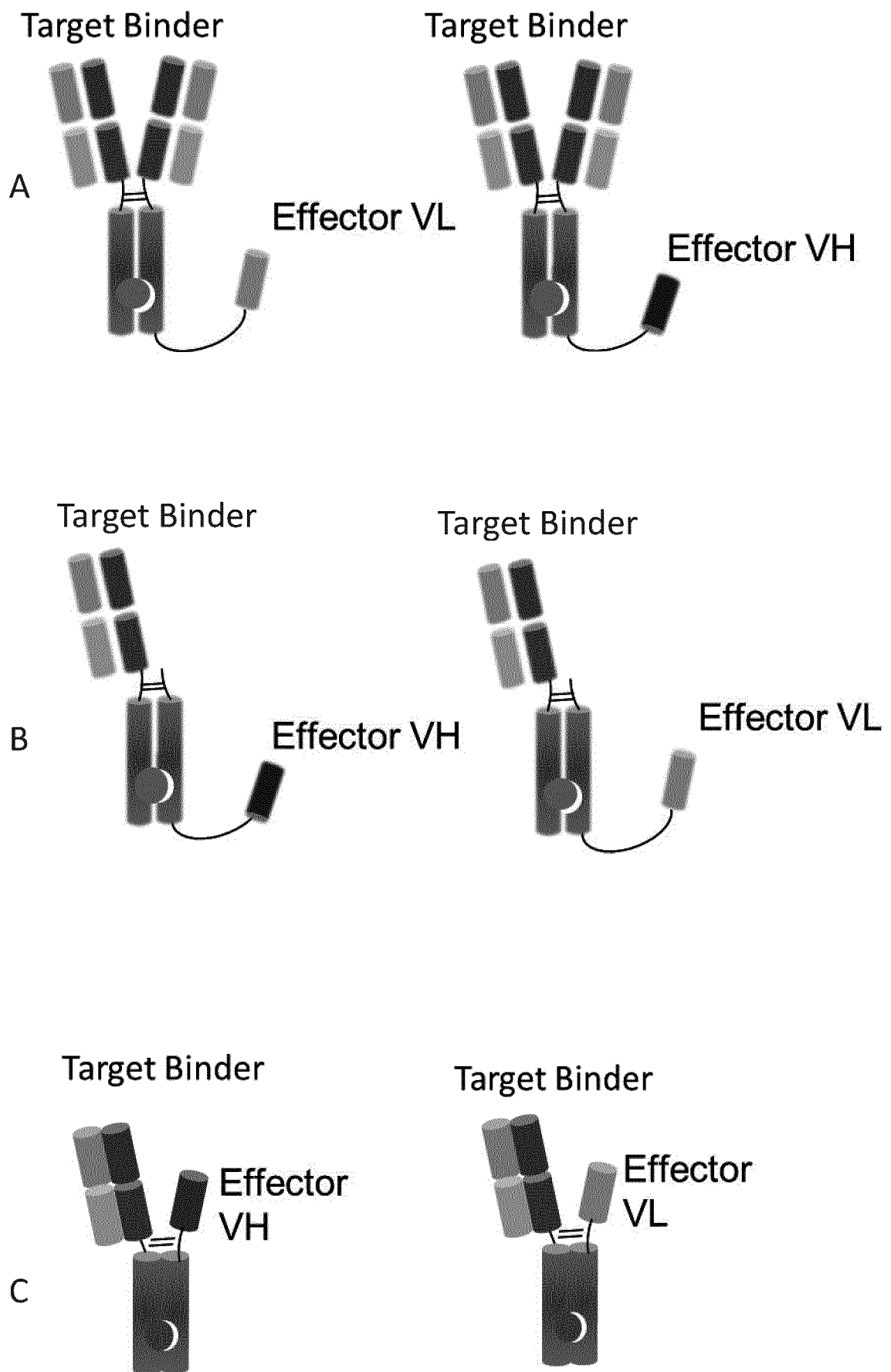


FIG. 25

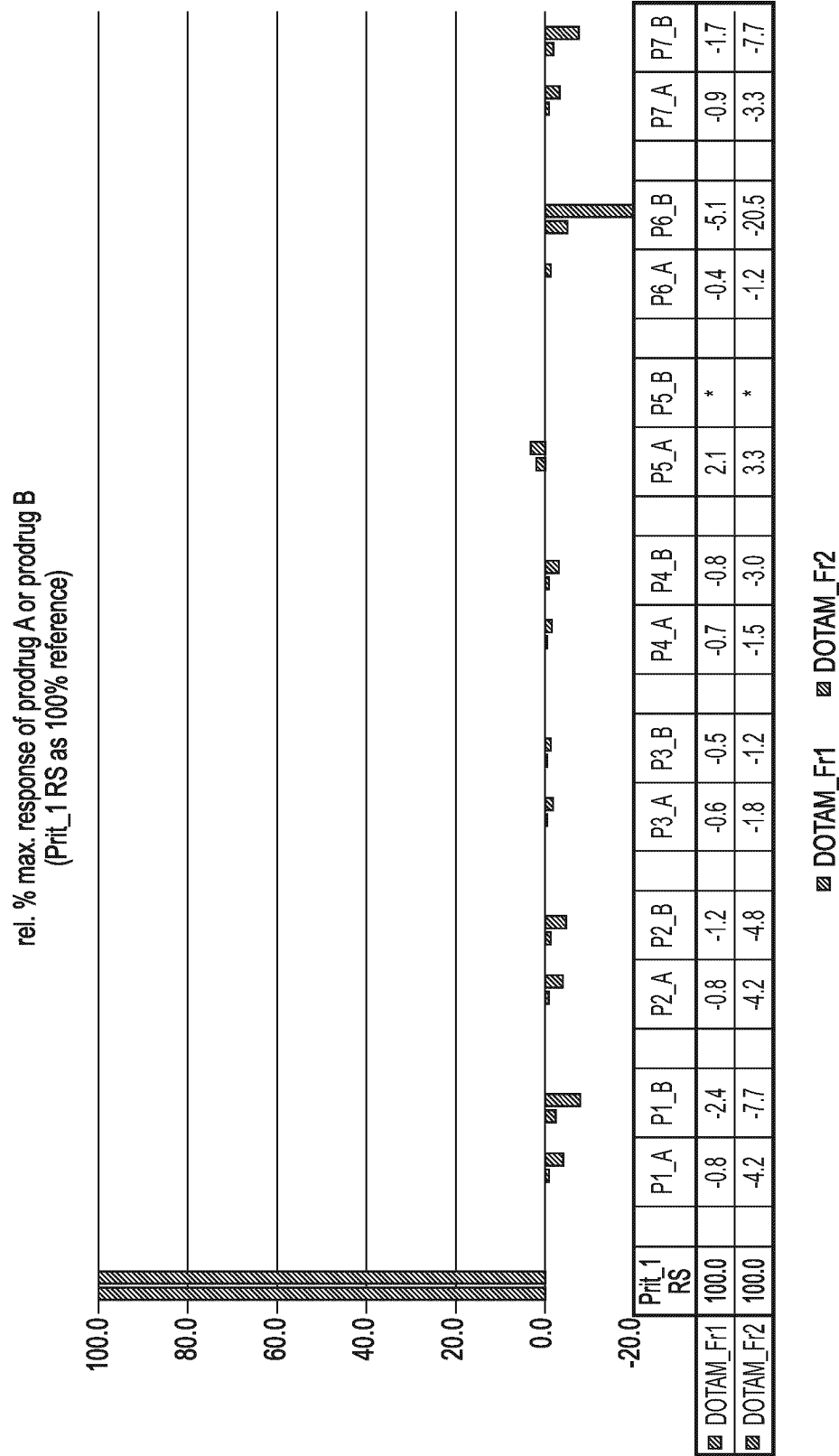


FIG. 26

rel. % max. response of DOTAM (DOTAM coupled with mono streptavidin)
Prodrugs A + B captured by <hFab> on surface in very high density
(Prit_1 RS as 100% reference)

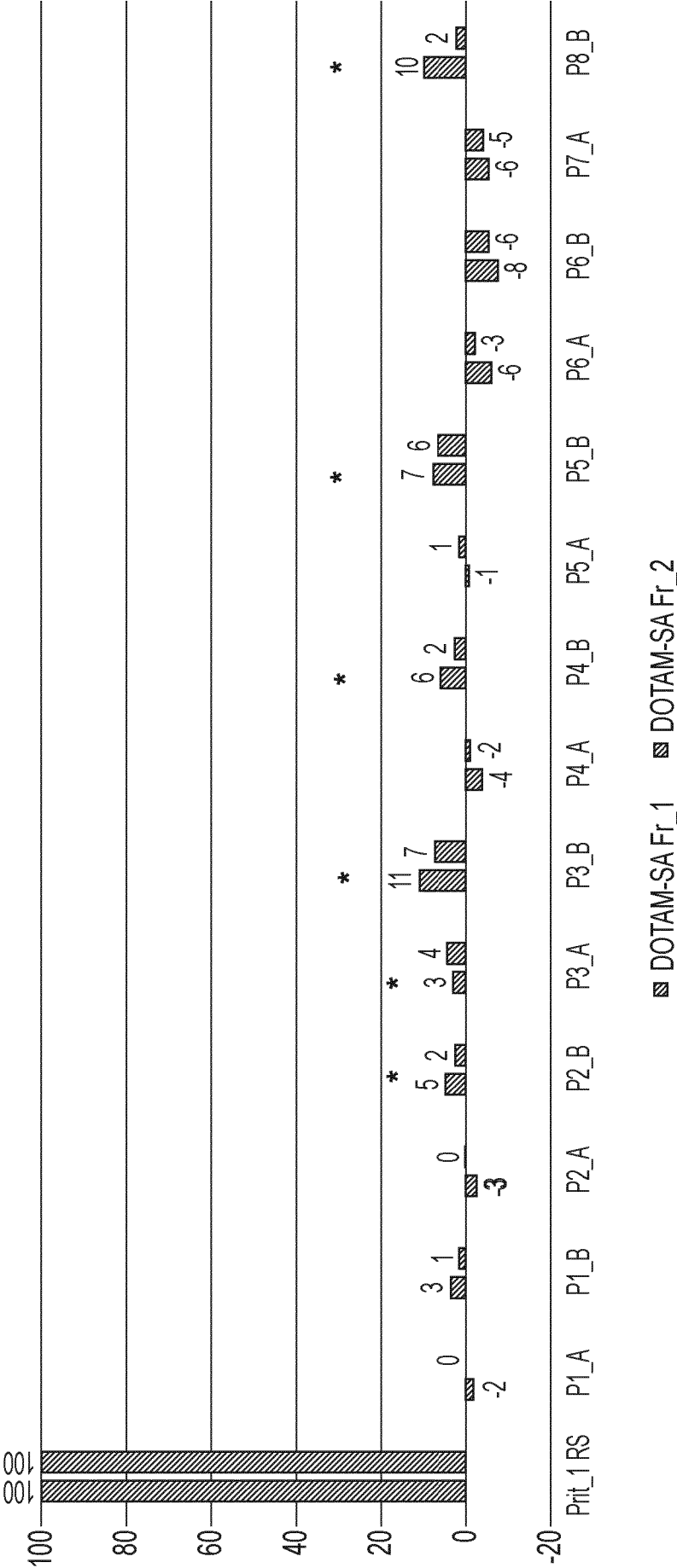


FIG. 27

rel. % max. response of DOTAM (DOTAM coupled with mono streptavidin)
Solution of Prodrugs A + B captured on <hFab> surface in very high density
(Prit_1 RS as 100% reference)

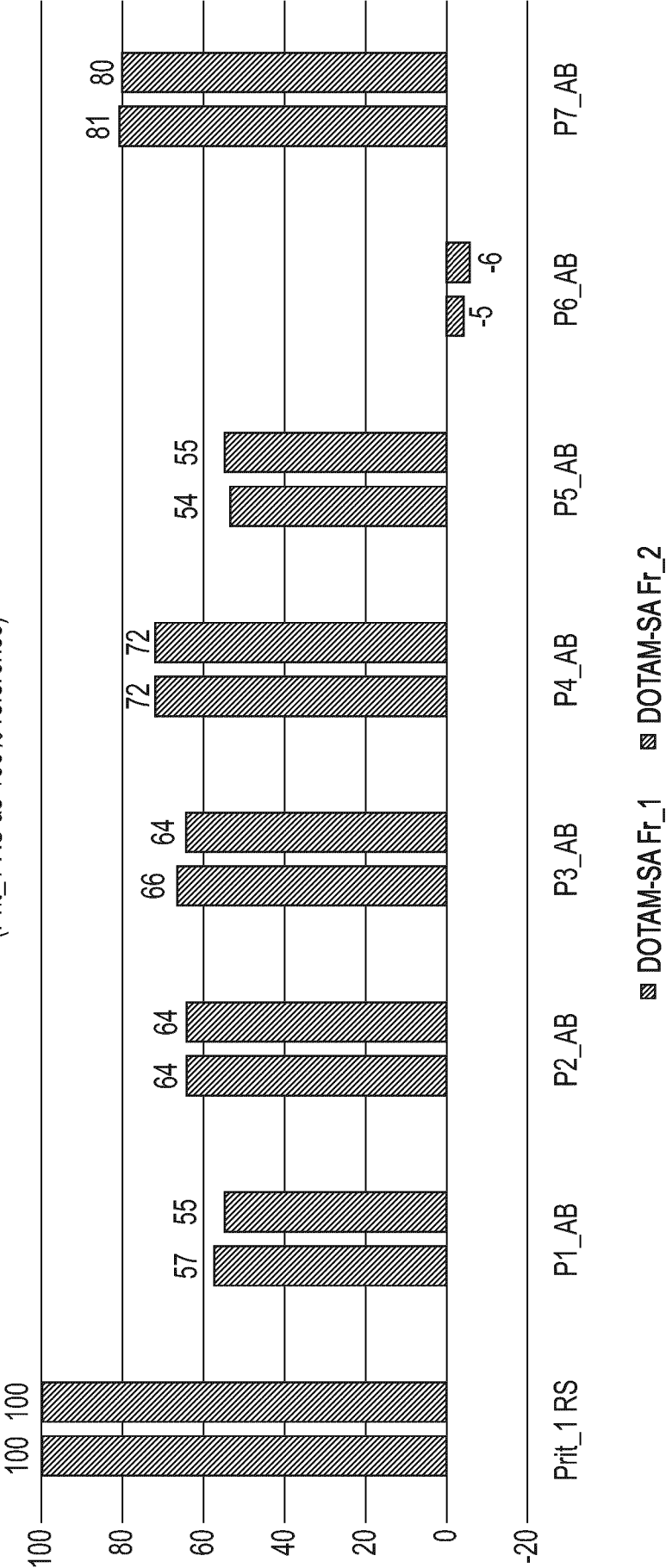


FIG. 28

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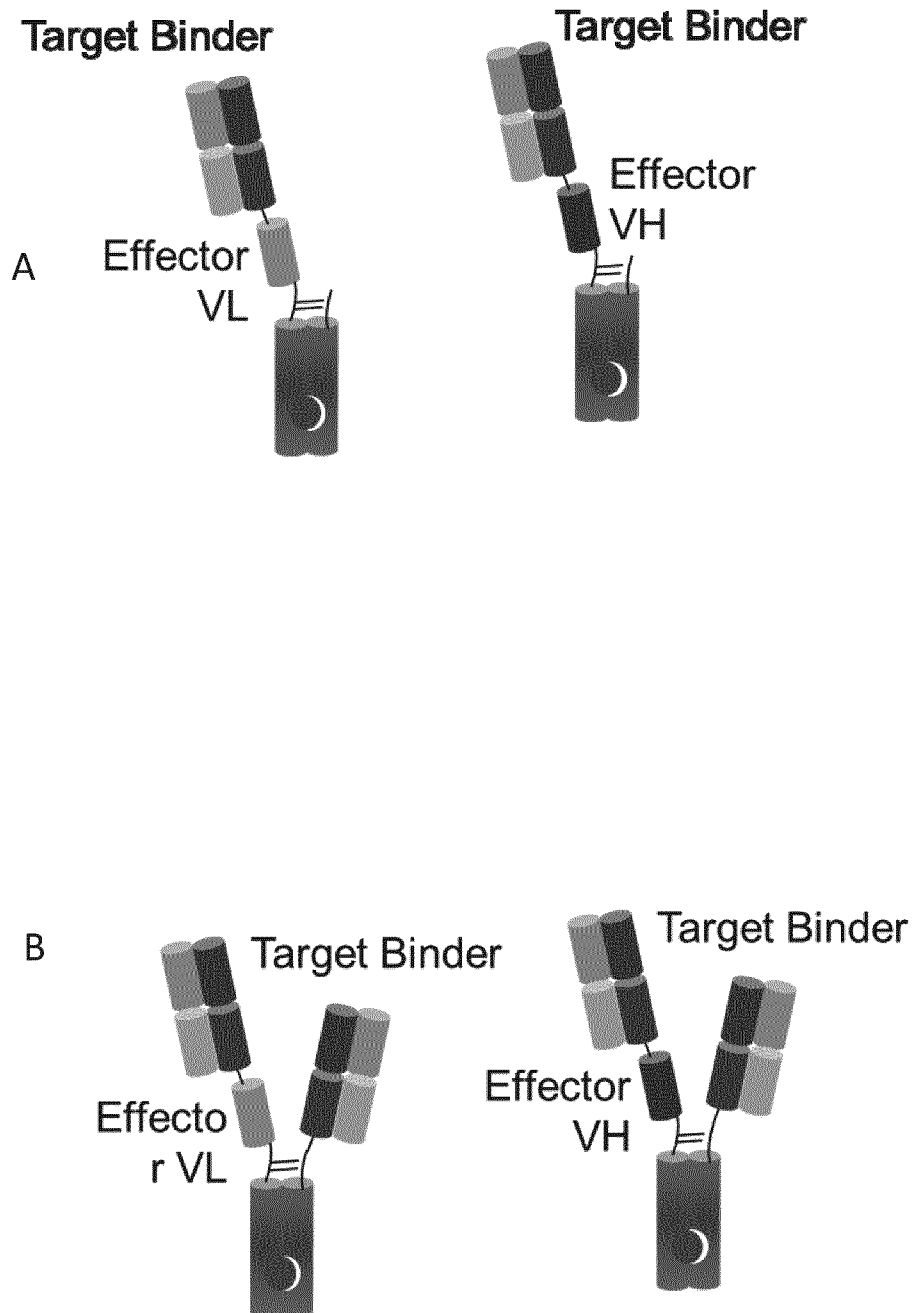


FIG. 29

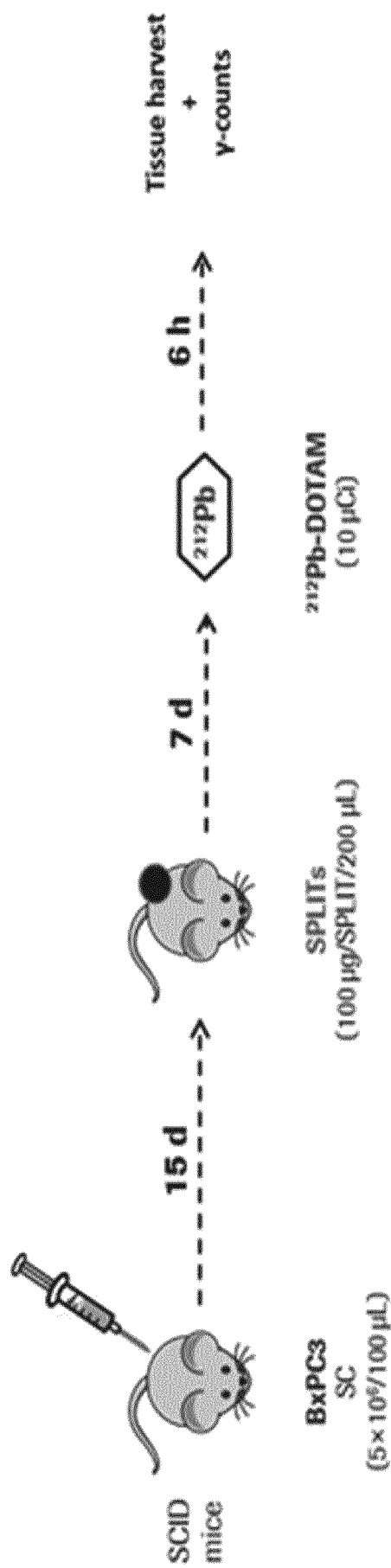


FIG. 30

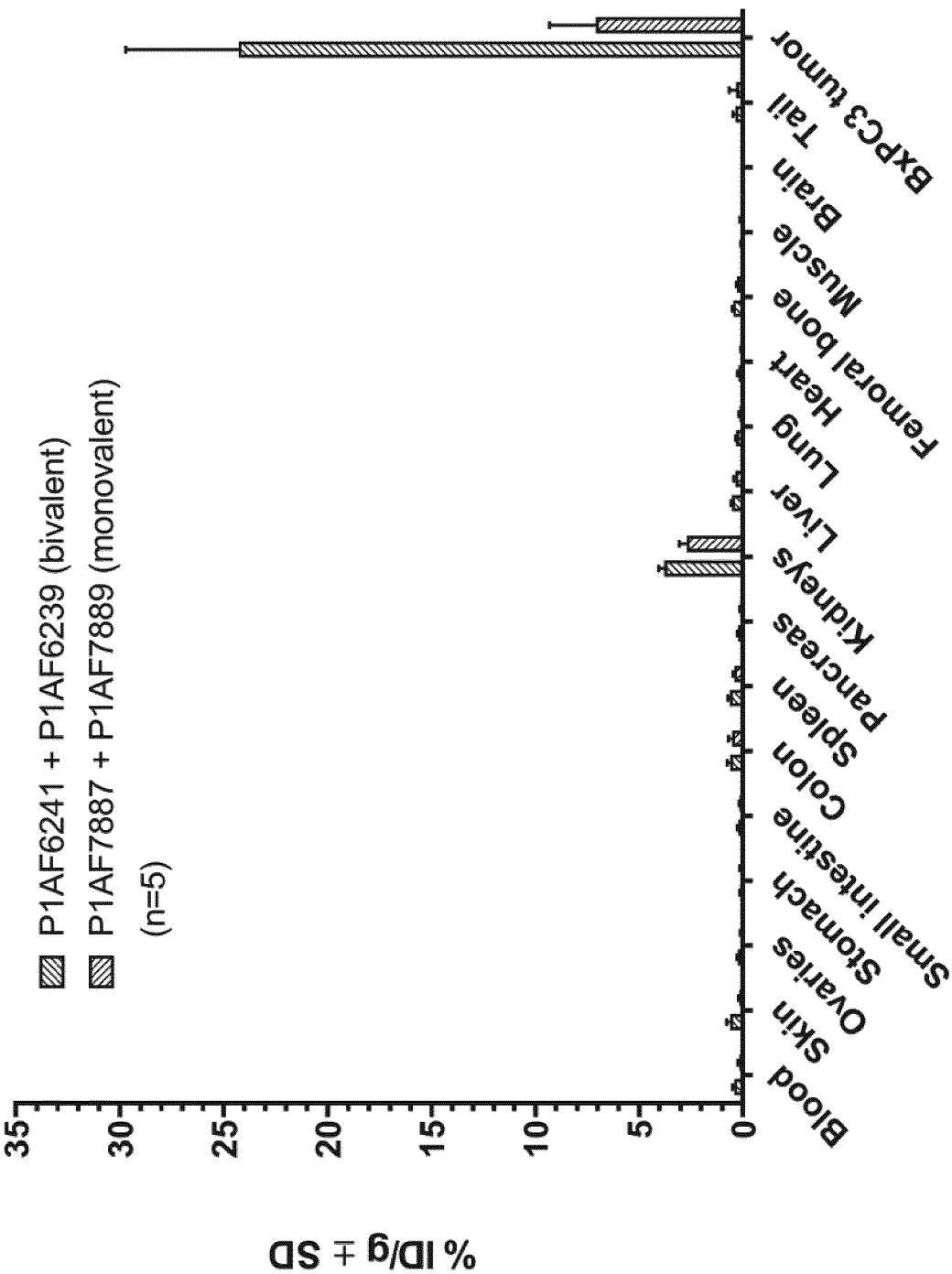


FIG. 31

INTERNATIONAL SEARCH REPORT

International application No.

PCT/EP2022/050359

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

1. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international search was carried out on the basis of a sequence listing:
 - a. ☒ forming part of the international application as filed:
 - ☒ in the form of an Annex C/ST.25 text file.
 - ☐ on paper or in the form of an image file.
 - b. ☐ furnished together with the international application under PCT Rule 13ter.1(a) for the purposes of international search only in the form of an Annex C/ST.25 text file.
 - c. ☐ furnished subsequent to the international filing date for the purposes of international search only:
 - ☐ in the form of an Annex C/ST.25 text file (Rule 13ter.1(a)).
 - ☐ on paper or in the form of an image file (Rule 13ter.1(b) and Administrative Instructions, Section 713).
2. ☐ In addition, in the case that more than one version or copy of a sequence listing has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that forming part of the application as filed or does not go beyond the application as filed, as appropriate, were furnished.
3. Additional comments:

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2022/050359

A. CLASSIFICATION OF SUBJECT MATTER

INV. A61P35/00 C07K16/30 A61K51/10
ADD.

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

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C07K A61K

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

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

EPO-Internal, BIOSIS, EMBASE, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2019/122350 A1 (STUHLER GERNOT [DE]) 27 June 2019 (2019-06-27) page 1, paragraph 2; last; figure 1 -----	1-88
X	AGNES BANASZEK ET AL: "On-target restoration of a split T cell-engaging antibody for precision immunotherapy", NATURE COMMUNICATIONS, vol. 10, no. 1, 26 November 2019 (2019-11-26), XP055730520, DOI: 10.1038/s41467-019-13196-0 the whole document ----- -/-	1-88



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents :

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

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

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

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

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

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

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

"&" document member of the same patent family

Date of the actual completion of the international search

7 April 2022

Date of mailing of the international search report

20/04/2022

Name and mailing address of the ISA/

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040,
Fax: (+31-70) 340-3016

Authorized officer

Covone-van Hees, M

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2022/050359

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	Kip Dudgeon ET AL: "General strategy for the generation of human antibody variable domains with increased aggregation resistance", Proceedings of the National Academy of Sciences of the United States of America, 3 July 2012 (2012-07-03), pages 10879-10884, XP055084733, United States DOI: 10.1073/pnas.1202866109 Retrieved from the Internet: URL:http://search.proquest.com/docview/1023506243 the whole document -----	1-88
A	WO 2017/087789 A1 (REVITOPÉ LTD [GB]; COBBOLD MARK [US]) 26 May 2017 (2017-05-26) paragraphs [0005], [0006], [0098], [0099], [0113], [0014], [0126], [0129] figures 3,4,5B -----	1-88
A	YAZAKI PAUL J ET AL: "A series of anti-CEA/anti-DOTA bispecific antibody formats evaluated for pre-targeting: comparison of tumor uptake and blood clearance", REDUCED ELIMINATION OF IGG ANTIBODIES BY ENGINEERING THE VARIABLE REGION,, vol. 26, no. 3, 1 March 2013 (2013-03-01), pages 187-193, XP002792685, ISSN: 1741-0134, DOI: 10.1093/PROTEIN/GZS096 the whole document -----	1-88
L	WO 2022/008688 A1 (HOFFMANN LA ROCHE [CH]; HOFFMANN LA ROCHE [US]) 13 January 2022 (2022-01-13) same applicant; the whole document -----	1-88
L	WO 2021/009047 A1 (HOFFMANN LA ROCHE [CH]) 21 January 2021 (2021-01-21) same applicant; the whole document -----	1-88

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2022/050359

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2019122350 A1	27-06-2019	EP 3759135 A1 US 2021070846 A1 WO 2019122350 A1	06-01-2021 11-03-2021 27-06-2019
WO 2017087789 A1	26-05-2017	AU 2016358296 A1 AU 2020220048 A1 CA 3003482 A1 CN 108289952 A CN 112807429 A DK 3377103 T3 EP 3377103 A1 EP 3903818 A1 ES 2873846 T3 HU E054131 T2 JP 2018538262 A PL 3377103 T3 PT 3377103 T US 2017152316 A1 US 2018346568 A1 WO 2017087789 A1	17-05-2018 03-09-2020 26-05-2017 17-07-2018 18-05-2021 19-04-2021 26-09-2018 03-11-2021 04-11-2021 30-08-2021 27-12-2018 04-10-2021 26-05-2021 01-06-2017 06-12-2018 26-05-2017
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