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(56) Fremdragne publikationer:
WO-A1-93/05789
US-A- 5 583 112
US-B1- 6 262 029
CHEENU BHARGAVA ET AL: "Targeted dianthin is a powerful toxin to treat pancreatic carcinoma when applied in combination with the glycosylated triterpene SO1861", MOLECULAR ONCOLOGY, vol. 11, no. 11, 15 September 2017 (2017-09-15), pages 1527-1543, XP055677214, ISSN: 1574-7891, DOI: 10.1002/1878-0261.12115
ROGER GILABERT-ORIOL ET AL: "Combinatorial approach to increase efficacy of Cetuximab, Panitumumab and Trastuzumab by dianthin conjugation and co-application of SO1861", BIOCHEMICAL PHARMACOLOGY, vol. 97, no. 3, 1 October 2015 (2015-10-01), pages 247-255, XP055677222, US ISSN: 0006-2952, DOI: 10.1016/j.bcp.2015.07.040
ALEXANDER WENG ET AL: "Saponins modulate the intracellular trafficking of protein toxins", JOURNAL OF CONTROLLED RELEASE, vol. 164, no. 1, 1 November 2012 (2012-11-01), pages 74-86, XP055428873, NL ISSN:

Fortsættes ...

0168-3659, DOI: 10.1016/j.jconrel.2012.10.002
FRØKIÆR H ET AL: "Optimization of hapten-protein conjugation by high-performance capillary electrophoresis", JOURNAL OF CHROMATOGRAPHY A, ELSEVIER, AMSTERDAM, NL, vol. 717, no. 1, 24 November 1995 (1995-11-24), pages 75-81, XP004038529, ISSN: 0021-9673, DOI: 10.1016/0021-9673(95)00642-X
LUIS ALVAREZ-SALAS: "Nucleic Acids as Therapeutic Agents", CURRENT TOPICS IN MEDICINAL CHEMISTRY, vol. 8, no. 15, 1 October 2008 (2008-10-01), pages 1379-1404, XP055302276, NL ISSN: 1568-0266, DOI: 10.2174/156802608786141133
MUNIYAN ANBARASAN ET AL: "Characterization and in vitro antibacterial activity of saponin-conjugated silver nanoparticles against bacteria that cause burn wound infection", WORLD JOURNAL OF MICROBIOLOGY AND BIOTECHNOLOGY, SPRINGER NETHERLANDS, DORDRECHT, vol. 33, no. 7, 20 June 2017 (2017-06-20), pages 1-12, XP036261778, ISSN: 0959-3993, DOI: 10.1007/S11274-017-2309-3 [retrieved on 2017-06-20]
DATABASE EMBASE [Online] ELSEVIER SCIENCE PUBLISHERS, AMSTERDAM, NL; 13 April 2015 (2015-04-13), SAITO M ET AL: "3[beta]-hydroxysterol [Delta]24-reductase on the surface of hepatitis C virus-related hepatocellular carcinoma cells can be a target for molecular targeting therapy", Database accession no. EMB-2015017035 & SAITO M ET AL: "3[beta]-hydroxysterol [Delta]24-reductase on the surface of hepatitis C virus-related hepatocellular carcinoma cells can be a target for molecular targeting therapy", PLOS ONE 20150413 PUBLIC LIBRARY OF SCIENCE USA, vol. 10, no. 4, 13 April 2015 (2015-04-13) , ISSN: 1932-6203
CRAVEN RACHEL A ET AL: "Proteomics and the search for biomarkers for renal cancer", CLINICAL BIOCHEMISTRY, ELSEVIER, AMSTERDAM, NL, vol. 46, no. 6, 7 December 2012 (2012-12-07), pages 456-465, XP028996892, ISSN: 0009-9120, DOI: 10.1016/J.CLINBIOCHEM.2012.11.029
DATABASE MEDLINE [Online] US NATIONAL LIBRARY OF MEDICINE (NLM), BETHESDA, MD, US; 24 November 1995 (1995-11-24), FRØKIAER H ET AL: "Optimization of hapten-protein conjugation by high-performance capillary electrophoresis.", Database accession no. NLM8520687 & FRØKIAER H ET AL: "Optimization of hapten-protein conjugation by high-performance capillary electrophoresis.", JOURNAL OF CHROMATOGRAPHY. A 24 NOV 1995, vol. 717, no. 1-2, 24 November 1995 (1995-11-24), pages 75-81, ISSN: 0021-9673
HILL KENNETH E ET AL: "Cerebellar Purkinje cells incorporate immunoglobulins and immunotoxins in vitro: implications for human neurological disease and immunotherapeutics", JOURNAL OF NEUROINFLAMMATION, BIOMED CENTRAL LTD., LONDON, GB, vol. 6, no. 1, 29 October 2009 (2009-10-29), page 31, XP021063705, ISSN: 1742-2094, DOI: 10.1186/1742-2094-6-31

DESCRIPTION

TECHNICAL FIELD

[0001] The invention relates to a therapeutic combination for use as a medicament, wherein the therapeutic combination comprises: (a) a first pharmaceutical composition comprising a first proteinaceous molecule and at least one saponin covalently bound to said first proteinaceous molecule; and (b) a second pharmaceutical composition comprising a second proteinaceous molecule, comprising an effector moiety, wherein the binding site of the first proteinaceous molecule and the binding site of the second proteinaceous molecule are the same. The invention also relates to the first pharmaceutical composition for use as a medicament. The invention also relates to the first pharmaceutical composition, further comprising the second proteinaceous molecule. The invention also relates to the first pharmaceutical composition, further comprising the second proteinaceous molecule, for use as a medicament. Furthermore, the invention relates to the first pharmaceutical composition, further comprising the second proteinaceous molecule, for use in the treatment or prophylaxis of cancer in a patient in need thereof.

BACKGROUND

[0002] Molecules with a therapeutic biological activity are in many occasions in theory suitable for application as an effective therapeutic drug for the treatment of a disease such as a cancer in human patients in need thereof. A typical example are small-molecule biologically active moieties. However, many if not all potential drug-like molecules and therapeutics currently used in the clinic suffer from at least one of a plethora of shortcomings and drawbacks. When administered to a human body, therapeutically active molecules may exert off-target effects, in addition to the biologically activity directed to an aspect underlying a to-be-treated disease or health problem. Such off-target effects are undesired and bear a risk for induction of health- or even life-threatening side effects of the administered molecule. It is the occurrence of such adverse events that cause many drug-like compounds and therapeutic moieties to fail phase III clinical trials or even phase IV clinical trials (post-market entry follow-up). Therefore, there is a strong desire to provide drug molecules such as small-molecule therapeutics, wherein the therapeutic effect of the drug molecule should, e.g., (1) be highly specific for a biological factor or biological process driving the disease, (2) be sufficiently safe, (3) be sufficiently efficacious, (4) be sufficiently directed to the diseased cell with little to no off-target activity on non-diseased cells, (5) have a sufficiently timely mode of action (e.g. the administered drug molecule should reach the targeted site in the human patient within a certain time frame and should remain at the targeted site for a certain time frame), and/or (6) have sufficiently long lasting therapeutic activity in the patient's body, amongst others. Unfortunately, to date, 'ideal' therapeutics with many or even all of the beneficial characteristics here above outlined, are not available to the patients, despite already long-lasting and intensive research and despite the impressive progress made in several areas of the individually addressed encountered difficulties and drawbacks.

[0003] Chemotherapy is one of the most important therapeutic options for cancer treatment. However, it is often associated with a low therapeutic window because it has no specificity towards cancer cells over dividing cells in healthy tissue. The invention of monoclonal antibodies offered the possibility of exploiting their specific binding properties as a mechanism for the targeted delivery of cytotoxic agents to cancer cells, while sparing normal cells. This can be achieved by chemical conjugation of cytotoxic effectors (also known as payloads or warheads) to antibodies, to create antibody-drug conjugates (ADCs). Typically, very potent payloads such as emtansine (DM1) are used which have a limited therapeutic index (a ratio that compares toxic dose to efficacious dose) in their unconjugated forms. The conjugation of DM1 to trastuzumab (ado-trastuzumab emtansine), also known as Kadcycla, enhances the tolerable dose of DM1 at least two-fold in monkeys. In the

past few decades tremendous efforts and investments have been made to develop therapeutic ADCs. However, it remains challenging to bring ADCs into the clinic, despite promising preclinical data. The first ADC approved for clinical use was gemtuzumab ozogamicin (Mylotarg, CD33 targeted, Pfizer/Wyeth) for relapsed acute myelogenous leukemia (AML) in 2000. Mylotarg was however, withdrawn from the market at the request of the Federal Drug Administration (FDA) due to a number of concerns including its safety profile. Patients treated with Mylotarg were more often found to die than patients treated with conventional chemotherapy. Mylotarg was admitted to the market again in 2017 with a lower recommended dose, a different schedule in combination with chemotherapy or on its own, and a new patient population. To date, only five ADCs have been approved for clinical use, and meanwhile clinical development of approximately fifty-five ADCs has been halted. However, interest remains high and approximately eighty ADCs are still in clinical development in nearly six-hundred clinical trials at present.

[0004] Despite the potential to use toxic payloads that are normally not tolerated by patients, a low therapeutic index (a ratio that compares toxic dose to efficacious dose) is a major problem accounting for the discontinuance of many ADCs in clinical development, which can be caused by several mechanisms such as off-target toxicity on normal cells, development of resistance against the cytotoxic agents and premature release of drugs in the circulation. A systematic review by the FDA of ADCs found that the toxicity profiles of most ADCs could be categorized according to the payload used, but not the antibody used, suggesting that toxicity is mostly determined by premature release of the payload. Of the approximately fifty-five ADCs that were discontinued, it is estimated that at least twenty-three were due to a poor therapeutic index. For example, development of a trastuzumab tesirine conjugate (ADCT-502, HER-2 targeted, ADC therapeutics) was recently discontinued due to a narrow therapeutic index, possibly due to an on-target, off-tissue effect in pulmonary tissue which expresses considerable levels of HER2. In addition, several ADCs in phase 3 trials have been discontinued due to missing primary endpoint. For example, phase 3 trials of a depatuxizumab mafodotin conjugate (ABT-414, EGFR targeted, AbbVie) tested in patients with newly diagnosed glioblastoma, and a mirvetuximab soravtansine conjugate (IMGN853, folate receptor alpha (FRα) targeted, ImmunoGen) tested in patients with platinum-resistant ovarian cancer, were recently stopped, showing no survival benefit. It is important to note that the clinically used dose of some ADCs may not be sufficient for its full anticancer activity. For example, ado-trastuzumab emtansine has an MTD of 3.6 mg/kg in humans. In preclinical models of breast cancer, ado-trastuzumab emtansine induced tumor regression at dose levels at or above 3 mg/kg, but more potent efficacy was observed at 15 mg/kg. This suggests that at the clinically administered dose, ado-trastuzumab emtansine may not exert its maximal potential anti-tumor effect.

[0005] ADCs are mainly composed of an antibody, a cytotoxic moiety such as a payload, and a linker. Several novel strategies have been proposed and carried out in the design and development of new ADCs to overcome the existing problems, targeting each of the components of ADCs. For example, by identification and validation of adequate antigenic targets for the antibody component, by selecting antigens which have high expression levels in tumor and little or no expression in normal tissues, antigens which are present on the cell surface to be accessible to the circulating ADCs, and antigens which allows internalizing of ADCs into the cell after binding; and alternative mechanisms of activity; design and optimize linkers which enhance the solubility and the drug-to-antibody ratio (DAR) of ADCs and overcome resistance induced by proteins that can transport the chemotherapeutic agent out of the cells; enhance the DAR ratio by inclusion of more payloads, select and optimize antibodies to improve antibody homogeneity and developability. In addition to the technological development of ADCs, new clinical and translational strategies are also being deployed to maximize the therapeutic index, such as, change dosing schedules through fractionated dosing; perform biodistribution studies; include biomarkers to optimize patient selection, to capture response signals early and monitor the duration and depth of response, and to inform combination studies.

[0006] An example of ADCs with clinical potential are those ADCs such as brentuximab vedotin, inotuzumab ozogamicin, moxetumomab pasudotox, and polatuzumab vedotin, which are evaluated as a treatment option for lymphoid malignancies and multiple myeloma. Polatuzumab vedotin, binding to CD79b on (malignant) B-

cells, and pinatuzumab vedotin, binding to CD22, are tested in clinical trials wherein the ADCs each were combined with co-administered rituximab, a monoclonal antibody binding to CD20 and not provided with a payload [B. Yu and D. Liu, Antibody-drug conjugates in clinical trials for lymphoid malignancies and multiple myeloma; Journal of Hematology & Oncology (2019) 12:94]. Combinations of monoclonal antibodies such as these examples are yet a further approach and attempt to arrive at the 'magic bullet' which combines many or even all of the aforementioned desired characteristics of ADCs.

[0007] Meanwhile in the past few decades, nucleic acid-based therapeutics are under development. Therapeutic nucleic acids can be based on deoxyribonucleic acid (DNA) or ribonucleic acid (RNA), Anti-sense oligonucleotides (ASOs, AONs), and short interfering RNAs (siRNAs), MicroRNAs, and DNA and RNA aptamers, for approaches such as gene therapy, RNA interference (RNAi). Many of them share the same fundamental basis of action by inhibition of either DNA or RNA expression, thereby preventing expression of disease-related abnormal proteins. The largest number of clinical trials is being carried out in the field of gene therapy, with almost 2600 ongoing or completed clinical trials worldwide but with only about 4% entering phase 3. This is followed by clinical trials with ASOs. Luis M. Alvarez-Salas (Nucleic Acids as Therapeutic Agents. Current Topics in Medicinal Chemistry, 2008; 8:1379-1404) published a review article, describing therapeutic nucleic acids that have progressed to clinical trials with anticancer and antiviral protocols.

[0008] Similarly to ADCs, despite the large number of techniques being explored, therapeutic nucleic acids share two major issues during clinical development: delivery into cells and off-target effects. For instance, ASOs such as peptide nucleic acid (PNA), phosphoramidate morpholino oligomer (PMO), locked nucleic acid (LNA) and bridged nucleic acid (BNA), are being investigated as an attractive strategy to inhibit specifically target genes and especially those genes that are difficult to target with small molecules inhibitors or neutralizing antibodies. Currently, the efficacy of different ASOs is being studied in many neurodegenerative diseases such as Huntington's disease, Parkinson's disease, Alzheimer's disease, and amyotrophic lateral sclerosis and also in several cancer stages. The application of ASOs as potential therapeutic agents requires safe and effective methods for their delivery to the cytoplasm and/or nucleus of the target cells and tissues. Although the clinical relevance of ASOs has been demonstrated, inefficient cellular uptake, both in vitro and in vivo, limit the efficacy of ASOs and has been a barrier to therapeutic development. Cellular uptake can be < 2% of the dose resulting in too low ASO concentration at the active site for an effective and sustained outcome. This consequently requires an increase of the administered dose which induces off-target effects. Most common side-effects are activation of the complement cascade, the inhibition of the clotting cascade and toll-like receptor mediated stimulation of the immune system.

[0009] Chemotherapeutics are most commonly small molecules, however, their efficacy is hampered by the severe off-target side toxicity, as well as their poor solubility, rapid clearance and limited tumor exposure. Scaffold-small-molecule drug conjugates such as polymer-drug conjugates (PDCs) are macromolecular constructs with pharmacologically activity, which comprises one or more molecules of a small-molecule drug bound to a carrier scaffold (e.g. polyethylene glycol (PEG)).

[0010] Such conjugate principle has attracted much attention and has been under investigation for several decades. The majority of conjugates of small-molecule drugs under pre-clinical or clinical development are for oncological indications. However, up-to-date only one drug not related to cancer has been approved (Movantik, a PEG oligomer conjugate of opioid antagonist naloxone, AstraZeneca) for opioid-induced constipation in patients with chronic pain in 2014, which is a non-oncology indication. Translating application of drug-scaffold conjugates into treatment of human subjects provides little clinical success so far. For example, PK1 (N-(2-hydroxypropyl)methacrylamide (HPMA) copolymer doxorubicin; development by Pharmacia, Pfizer) showed great anti-cancer activity in both solid tumors and leukemia in murine models, and was under clinical investigation for oncological indications. Despite that it demonstrated significant reduction of nonspecific toxicity and improved pharmacokinetics in man, improvements in anticancer efficacy turned out to be marginal in patients, and as a consequence further development of PK1 was discontinued.

[0011] The failure of scaffold-small-molecule drug conjugates is at least partially attributed to its poor accumulation at the tumor site. For example, while in murine models PK1 showed 45-250 times higher accumulation in the tumor than in healthy tissues (liver, kidney, lung, spleen, and heart), accumulation in tumor was only observed in a small subset of patients in the clinical trial.

[0012] A potential solution to the aforementioned problems is application of nanoparticle systems for drug delivery such as liposomes. Liposomes are sphere-shaped vesicles consisting of one or more phospholipid bilayers, which are spontaneously formed when phospholipids are dispersed in water. The amphiphilicity characteristics of the phospholipids provide it with the properties of self-assembly, emulsifying and wetting characteristics, and these properties can be employed in the design of new drugs and new drug delivery systems. Drug encapsulated in a liposomal delivery system may convey several advantages over a direct administration of the drug, such as an improvement and control over pharmacokinetics and pharmacodynamics, tissue targeting property, decreased toxicity and enhanced drug activity. An example of such success is liposome-encapsulated form of a small molecule chemotherapy agent doxorubicin (Doxil: a pegylated liposome-encapsulated form of doxorubicin; Myocet: a non-pegylated liposomal doxorubicin), which have been approved for clinical use. Cheenu Bhargava et al. (Targeted dianthin is a powerful toxin to treat pancreatic carcinoma when applied in combination with the glycosylated triterpene SO1861. Molecular Oncology; 2017, 11(11):1527-154) report the results of *in vitro* tests with a fusion protein of the ribosome-inactivating protein dianthin and human epidermal growth factor in combination with SO1861 in the pancreatic carcinoma cell lines BxPC-3 and MIA PaCa-2. Gilabert-Oriol et al. (Combinatorial approach to increase efficacy of Cetuximab, Panitumumab and Trastuzumab by dianthin conjugation and co-application of SO1861. Biochemical Pharmacology; 2015, 97(3): 247-255) describe the chemical conjugation of the therapeutic antibodies Cetuximab (anti-EGFR, Erbitux®), Panitumumab (anti-EGFR, Vectibix®) and Trastuzumab (anti-HER2, Herceptin®) to the toxin dianthin. Cytotoxicity of these conjugates on MCF-7 cells was tested, also in combination with SO1861. Alexander Weng et al. (Saponins modulate the intracellular trafficking of protein toxins; Journal of Controlled Release, 2015; 164(1):74-86) disclose that combining a targeted anti-tumor toxin (i.e. saporin-EGF) with certain saponins increases the therapeutic efficacy of the toxin in an animal model (tumor bearing BALB/c mice). Frokier H et al. (Optimization of hapten-protein conjugation by high-performance capillary electrophoresis. Journal of Chromatography; 1995, 717(1):75-81) describe the conjugation of haptens, including soyasaponin 1, to a carrier protein, Kunitz soybean trypsin inhibitor (KSTI). WO 93/05789 concerns the use of saponins for for enhancement of immune responses against an antigen, i.e. the use of saponins as an adjuvant.

[0013] A solution still needs to be found that allows for drug therapies such as anti-tumor therapies, applicable for non-systemic use when desired, wherein the drug has for example an acceptable safety profile, little off-target activity, sufficient efficacy, sufficiently low clearance rate from the patient's body, etc.

SUMMARY

[0014] For an embodiment of the present invention, it is a first goal to provide an improved biologically active compound or composition comprising such improved biologically active compound.

[0015] It is one of several objectives of embodiments of the current invention to provide a solution to the problem of non-specificity, encountered when administering small-molecule therapeutically active compounds to a human patient in need thereof. It is one of several objectives of embodiments of the current invention to provide a solution to the problem of drugs with non-optimal specificity for a biological factor or biological process driving a disease. It is one of several objectives of embodiments of the current invention to provide a solution to the problem of insufficient safety characteristics of current drugs, when administered to human patients in need thereof. It is one of several objectives of embodiments of the current invention to provide a

solution to the problem of current drugs being less efficacious than desired, when administered to human patients in need thereof. It is one of several objectives of embodiments of the current invention to provide a solution to the problem of current drugs being not sufficiently directed to the diseased cell with little to no off-target activity on non-diseased cells, when administered to human patients in need thereof. It is one of several objectives of embodiments of the current invention to provide a solution to the problem that current drugs do not have a sufficiently timely mode of action (e.g. the administered drug molecule should reach the targeted site in the human patient within a certain time frame and should remain at the targeted site for a certain time frame), when administered to human patients in need thereof. It is one of several objectives of embodiments of the current invention to provide a solution to the problem that current drugs have not sufficiently long lasting therapeutic activity in the patient's body, when administered to human patients in need thereof.

[0016] At least one of the above objectives of embodiments of the invention is achieved by providing a therapeutic combination as defined in the claims for use as a medicament. The therapeutic combination comprises a first molecule comprising covalently bound saponin and comprises a second molecule comprising an effector molecule, wherein both the first and second molecule comprise a binding site for the same epitope exposed on the same cell-surface molecule of a targeted cell.

[0017] The present invention will be described with respect to particular embodiments but the invention is not limited thereto but only by the claims. The embodiments of the invention described herein can operate in combination and cooperation, unless specified otherwise.

[0018] In accordance with the invention, the therapeutic combination comprises: (a) a first pharmaceutical composition comprising a first proteinaceous molecule comprising a binding site for binding to an epitope on a cell-surface molecule and at least one saponin covalently bound to said first proteinaceous molecule preferably to an amino-acid residue of said first proteinaceous molecule, the first pharmaceutical composition optionally further comprising a pharmaceutically acceptable excipient; and (b) a second pharmaceutical composition comprising a second proteinaceous molecule, the second proteinaceous molecule comprising a binding site for binding to the epitope on the cell-surface molecule of (a) and an effector moiety, the second pharmaceutical composition optionally further comprising a pharmaceutically acceptable excipient, wherein the binding site of the first proteinaceous molecule and the binding site of the second proteinaceous molecule are the same, and wherein the cell-surface molecule and the epitope on the cell-surface molecule, to which the first proteinaceous molecule can bind, and the cell-surface molecule and the epitope on the cell-surface molecule, to which the second proteinaceous molecule can bind, are the same. Thus, the first proteinaceous molecule is a conjugate comprising a binding site for binding to an epitope on a cell-surface molecule and comprising at least one saponin covalently bound to said first proteinaceous molecule preferably to an amino-acid residue of said first proteinaceous molecule. Thus, the second proteinaceous molecule is a conjugate comprising the binding site for binding to the epitope on the same cell-surface molecule and comprising an effector moiety.

[0019] An aspect of the invention relates to a therapeutic combination for use as a medicament, wherein the therapeutic combination comprises: (a) a first pharmaceutical composition comprising a first proteinaceous molecule comprising a binding site for binding to an epitope on a cell-surface molecule and at least one saponin covalently bound to said first proteinaceous molecule preferably to an amino-acid residue of said first proteinaceous molecule, the first pharmaceutical composition optionally further comprising a pharmaceutically acceptable excipient; and (b) a second pharmaceutical composition comprising a second proteinaceous molecule, the second proteinaceous molecule comprising a binding site for binding to the epitope on the cell-surface molecule of (a) and an effector moiety, the second pharmaceutical composition optionally further comprising a pharmaceutically acceptable excipient, wherein the binding site of the first proteinaceous molecule and the binding site of the second proteinaceous molecule are the same, and wherein the cell-surface molecule and the epitope on the cell-surface molecule, to which the first proteinaceous molecule can bind, and the cell-surface molecule and the epitope on the cell-surface molecule, to which the second

proteinaceous molecule can bind, are the same.

[0020] An embodiment is the therapeutic combination of the invention for use in the treatment or prevention of a cancer in a human subject, wherein the therapeutic combination comprises: (a) the first pharmaceutical composition of the invention; and (b) the second pharmaceutical composition of the invention, wherein the cell-surface molecule is expressed on a tumor cell surface, and preferably the cell-surface molecule is a tumor cell-specific surface molecule, and wherein preferably the epitope is a tumor-cell specific epitope.

[0021] In accordance with the invention, the binding site of the first proteinaceous molecule and the second proteinaceous molecule comprises or consists of an immunoglobulin or a binding fragment or binding domain of said immunoglobulin, such as any one or more of an antibody, an IgG, a molecule comprising or consisting of a Vhh domain or Vh domain, a Fab, an scFv, an Fv, a dAb, an F(ab)2, Fcab fragment, and/or comprises or consists of at least one ligand, the ligand for binding to a cell-surface molecule such as EGF.

[0022] In an embodiment, the tumor-cell surface molecule is a cell-surface receptor specifically present at a tumor cell, and, preferably, the epitope is a tumor-cell specific epitope.

[0023] In accordance with the invention, the at least one saponin is a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin.

[0024] In an embodiment, the epitope is an epitope on a tumor-cell receptor, preferably a tumor-cell specific epitope, and wherein the receptor is preferably a tumor-cell specific receptor, more preferably a receptor selected from CD71, CA125, EpCAM(17-1A), CD52, CEA, CD44v6, FAP, EGF-IR, integrin, syndecan-1, vascular integrin alpha-V beta-3, HER2, EGFR, CD20, CD22, Folate receptor 1, CD146, CD56, CD19, CD138, CD27L receptor, PSMA, CanAg, integrin-alphaV, CA6, CD33, mesothelin, Cripto, CD3, CD30, CD239, CD70, CD123, CD352, DLL3, CD25, ephrinA4, MUC1, Trop2, CEACAM5, CEACAM6, HER3, CD74, PTK7, Notch3, FGF2, C4.4A, FLT3, CD38, FGFR3, CD7, PD-L1, CTLA4, CD52, PDGFRA, VEGFR1, VEGFR2, most preferably selected from CD71, HER2 and EGFR.

[0025] In an embodiment, the effector moiety comprised by the second proteinaceous molecule comprises or consists of at least one proteinaceous molecule, preferably selected from any one or more of a peptide, a protein, an enzyme such as urease and Cre-recombinase, a proteinaceous toxin, a ribosome-inactivating protein, a protein toxin selected from Table A5 and/or a bacterial toxin, a plant toxin; preferably the protein toxin is dianthin and/or saporin.

[0026] In an embodiment, the first proteinaceous molecule comprises more than one covalently bound saponin, preferably 2, 3, 4, 5, 6, 8, 10, 16, 32, 64, 128 or 1-100 saponins, or any number of saponins therein between, such as 7, 9, 12 saponins.

[0027] In an embodiment, the first pharmaceutical composition and the second pharmaceutical composition are administered to the patient in need thereof.

[0028] Specifically envisaged, within the context of the invention, are any of the following first proteinaceous molecules or any of the following second proteinaceous molecules, or any of the following semi-finished products consisting of any of the first proteinaceous molecules or any of the second proteinaceous molecules, comprising the binding site of the invention and either comprising at least one effector moiety of the invention when the second proteinaceous molecule is considered or comprising at least one saponin of the invention, when the first proteinaceous molecule is considered, wherein the semi-finished products are suitable for application in the manufacture of an antibody-drug conjugate conjugated with a saponin of the invention or an

antibody-oligonucleotide conjugate conjugated with a saponin of the invention:

Anti-EGFR antibody - saponin;

Anti-EGFR antibody - triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin;

Anti-EGFR antibody - SO1861;

Anti-EGFR antibody - GE1741;

Anti-EGFR antibody - SA1641;

Anti-EGFR antibody - Quil-A;

Anti-EGFR antibody - QS-21;

Anti-EGFR antibody - saponins in water soluble saponin fraction of Quillaja saponaria;

Cetuximab - saponin;

Cetuximab - triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin;

Cetuximab - SO1861;

Cetuximab - GE1741;

Cetuximab - SA1641;

Cetuximab - Quil-A;

Cetuximab - QS-21;

Cetuximab - saponins in water soluble saponin fraction of Quillaja saponaria;

Anti-HER2 antibody - saponin;

Anti-HER2 antibody - triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin;

Anti-HER2 antibody - SO1861;

Anti-HER2 antibody - GE1741;

Anti-HER2 antibody - SA1641;

Anti-HER2 antibody - Quil-A;

Anti-HER2 antibody - QS-21;

Anti-HER2 antibody - saponins in water soluble saponin fraction of Quillaja saponaria;

Trastuzumab - saponin;

Trastuzumab - triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid

function in a carbohydrate substituent at the C-3beta-OH group of the saponin;

Trastuzumab - SO1861;

Trastuzumab - GE1741;

Trastuzumab - SA1641;

Trastuzumab - Quil-A;

Trastuzumab - QS-21;

Trastuzumab - saponins in water soluble saponin fraction of Quillaja saponaria;

Anti-CD71 antibody - saponin;

Anti-CD71 antibody - triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin;

Anti-CD71 antibody - SO1861;

Anti-CD71 antibody - GE1741;

Anti-CD71 antibody - SA1641;

Anti-CD71 antibody - Quil-A;

Anti-CD71 antibody - QS-21;

Anti-CD71 antibody - saponins in water soluble saponin fraction of Quillaja saponaria;

OKT-9 - saponin;

OKT-9 -triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin;

OKT-9 - SO1861;

OKT-9 - GE1741;

OKT-9 - SA1641;

OKT-9 - Quil-A;

OKT-9 - QS-21;

OKT-9 - saponins in water soluble saponin fraction of Quillaja saponaria;

Anti-EGFR antibody - oligonucleotide;

Anti-EGFR antibody - antisense oligonucleotide;

Anti-EGFR antibody - siRNA;

Anti-EGFR antibody - antisense BNA;

Anti-EGFR antibody - antisense BNA(HSP27);

Anti-EGFR antibody - proteinaceous toxin;

Anti-EGFR antibody - ribosome inactivating protein;

Anti-EGFR antibody - dianthin;

Anti-EGFR antibody - saporin;

Cetuximab - oligonucleotide;

Cetuximab - antisense oligonucleotide;

Cetuximab - siRNA;

Cetuximab - antisense BNA;

Cetuximab - antisense BNA(HSP27);

Cetuximab - proteinaceous toxin;

Cetuximab - ribosome inactivating protein;

Cetuximab - dianthin;

Cetuximab - saporin;

Anti-HER2 antibody - oligonucleotide;

Anti-HER2 antibody - antisense oligonucleotide;

Anti-HER2 antibody - siRNA;

Anti-HER2 antibody - antisense BNA;

Anti-HER2 antibody - antisense BNA(HSP27);

Anti-HER2 antibody - proteinaceous toxin;

Anti-HER2 antibody - ribosome inactivating protein;

Anti-HER2 antibody - dianthin;

Anti-HER2 antibody - saporin;

Trastuzumab - oligonucleotide;

Trastuzumab - antisense oligonucleotide;

Trastuzumab - siRNA;

Trastuzumab - antisense BNA;

Trastuzumab - antisense BNA(HSP27);

Trastuzumab - proteinaceous toxin;

Trastuzumab - ribosome inactivating protein;

Trastuzumab - dianthin;

Trastuzumab - saporin;

Anti-CD71 antibody - oligonucleotide;

Anti-CD71 antibody - antisense oligonucleotide;

Anti-CD71 antibody - siRNA;

Anti-CD71 antibody - antisense BNA;

Anti-CD71 antibody - antisense BNA(HSP27);

Anti-CD71 antibody - proteinaceous toxin;

Anti-CD71 antibody - ribosome inactivating protein;

Anti-CD71 antibody - dianthin;

Anti-CD71 antibody - saporin;

OKT-9 - oligonucleotide;

OKT-9 - antisense oligonucleotide;

OKT-9 - siRNA;

OKT-9 - antisense BNA;

OKT-9 - antisense BNA(HSP27);

OKT-9 - proteinaceous toxin;

OKT-9 - ribosome inactivating protein;

OKT-9 - dianthin; and

OKT-9 - saporin.

[0029] In an embodiment, the semi-finished product consists of any one of the first proteinaceous molecules or second proteinaceous molecules, covalently coupled to at least one oligonucleotide when the first proteinaceous molecule is considered or covalently coupled to at least one effector moiety when the second proteinaceous molecules is considered, preferably selected from:

anti-EGFR antibody (- oligonucleotide)(- saponin) conjugate, wherein the oligonucleotide is any one or more of antisense oligonucleotide, siRNA, antisense BNA, and antisense BNA(HSP27), and wherein the saponin is any one or more of a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin, SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria, wherein the anti-EGFR antibody preferably is cetuximab;

anti-EGFR antibody (- proteinaceous toxin)(- saponin) conjugate, wherein the proteinaceous toxin is any one or more of a ribosome inactivating protein, dianthin and saporin, and wherein the saponin is any one or more of a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin, SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria, wherein the anti-EGFR antibody preferably is cetuximab;

anti-HER2 antibody (- oligonucleotide)(- saponin) conjugate, wherein the oligonucleotide is any one or more of antisense oligonucleotide, siRNA, antisense BNA, and antisense BNA(HSP27), and wherein the saponin is any one or more of a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin, SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria, wherein the anti-HER2 antibody preferably is trastuzumab;

anti-HER2 antibody (- proteinaceous toxin)(- saponin) conjugate, wherein the proteinaceous toxin is any one or more of a ribosome inactivating protein, dianthin and saporin, and wherein the saponin is any one or more of a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin, SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria, wherein the anti-HER2 antibody preferably is trastuzumab;

anti-CD71 antibody (- oligonucleotide)(- saponin) conjugate, wherein the oligonucleotide is any one or more of antisense oligonucleotide, siRNA, antisense BNA, and antisense BNA(HSP27), and wherein the saponin is any one or more of a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin, SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria, wherein the anti-CD71 antibody preferably is OKT-9; and

anti-CD71 antibody (- proteinaceous toxin)(- saponin) conjugate, wherein the proteinaceous toxin is any one or more of a ribosome inactivating protein, dianthin and saporin, and wherein the saponin is any one or more of a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin, SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria, wherein the anti-CD71 antibody preferably is OKT-9.

[0030] In an embodiment, the binding site is selected from cetuximab, trastuzumab, OKT-9, and/or wherein the effector moiety is selected from dianthin, saporin and antisense BNA(HSP27), and/or wherein the saponin is selected from SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria.

[0031] In an embodiment, the binding site is selected from cetuximab, trastuzumab, OKT-9, and/or wherein the effector moiety is selected from dianthin, saporin and antisense BNA(HSP27), and/or wherein the saponin is selected from SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria.

[0032] Specifically envisaged, within the context of the invention, is an ADC or an AOCs or a semi-finished ADC conjugate or a semi-finished AOC conjugate comprising the binding site of the invention and comprising at least one effector moiety of the invention and/or comprising at least one saponin of the invention, of Structure C:

A (- S)b (- E)c Structure C,

wherein A is the binding site;

S is the saponin;

E is the effector moiety;

b = 0 - 64, preferably 0, 1, 2, 3, 4, 8, 16, 32, 64 or any whole number or fraction therein between;

c = 0 - 8, preferably 0, 1, 2, 3, 4, 6, 8 or any whole number or fraction therein between,

wherein S is coupled to A and/or E, E is coupled to A and/or S, preferably S is coupled to A and E is coupled to A.

[0033] In an embodiment, A is an anti-EGFR antibody such as cetuximab, an anti-HER2 antibody such as trastuzumab, an anti-CD71 antibody such as OKT-9, and/or wherein S is any one or more of a saponin, a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprising a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin, SO1861, GE1741, SA1641, Quil-A, QS-21, and saponins in water soluble saponin fraction of Quillaja saponaria, and/or wherein E is any one or more of an oligonucleotide, an antisense oligonucleotide, an siRNA, an antisense BNA, and an antisense BNA(HSP27), and/or any one or more of a proteinaceous toxin, a ribosome inactivating protein, dianthin and saporin.

[0034] In an embodiment, the saponin, if present, and/or the effector moiety, if present, is covalently coupled via at least one linker, such as a cleavable linker, and/or via at least one oligomeric or polymeric scaffold, such as a linker based on N-ε-maleimidocaproic acid hydrazide (EMCH) succinimidyl 3-(2-pyridyldithio)propionate or 3-(2-Pyridyldithio)propionic acid N-hydroxysuccinimide ester (SPDP), and 1-[Bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate (HATU), and such as a scaffold based on a Dendron such as a G4-Dendron or a tri-functional linker such as the tri-functional linker of Scheme II, and/or at least a lysine side chain and/or a cysteine side chain of the binding site, preferably a monoclonal antibody or fragments or domains thereof, is involved in the covalent bond with the saponin and/or the effector moiety and/or the linker and/or the cleavable linker and/or the scaffold, wherein preferably the saponin and/or the effector moiety is covalently linked to the binding site, preferably an antibody, wherein the covalent link comprises or consists of an amide bond, a hydrazone bond, a disulphide bond.

DEFINITIONS

[0035] The term "linker" has its regular scientific meaning, and here refers to a chemical moiety or a linear stretch of amino-acid residues complexed through peptide bonds, which attaches a molecule or an atom to another molecule, e.g. to a ligand or to an effector molecule or to a scaffold. Typically, the linker comprises a chain of atoms linked by chemical bonds. Any linker molecule or linker technology known in the art can be used in the present disclosure. Where indicated, the linker is a linker for covalently binding of molecules through a chemical group on such a molecule suitable for forming a covalent linkage or bond with the linker. The linker may be a non-cleavable linker, e.g., the linker is stable in physiological conditions. The linker may be a cleavable linker, e.g. a linker that is cleavable, in the presence of an enzyme or at a particular pH range or value, or under physiological conditions such as intracellular conditions in the endosomes such as the late endosomes and the lysosomes of mammalian cells such as human cells. Exemplary linkers that can be used in the context of the present disclosure includes, but is not limited to, N-ε-maleimidocaproic acid hydrazide (EMCH), succinimidyl 3-(2-pyridyldithio)propionate or 3-(2-Pyridyldithio)propionic acid N-hydroxysuccinimide ester (SPDP), and 1-[Bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate (HATU).

[0036] The term "tri-functional linker" has its regular scientific meaning, and here refers to a linker which attaches three molecules via a chemical group on each of the three molecules. The skilled person is able to design such tri-functional linkers, based on the present disclosure and the common general knowledge. Such tri-functional linker can exhibit, for instance, a maleimido group that can be used for conjugation to targeting ligands that exhibit thiol groups to perform a thiol-ene reaction. In addition, the tri-functional linker could exhibit a dibenzocyclooctyne (DBCO) group to perform the so-called strain-promoted alkyne-azide cycloaddition (SPAAC, click chemistry) with an azido bearing saponin. Finally, the tri-functional linker could obtain a third functional group such as a trans-cyclooctene (TCO) group to perform the so-called inverse electron demand Diels-Alder (IEDDA) reaction with a tetrazine (Tz) bearing effector molecule. The skilled person will appreciate that the chemical groups of the tri-functional linker can be all three the same, or different, or the linker may comprise two of the same chemical groups for linking a molecule to the tri-functional linker. The formed bonds between the tri-functional linker can be covalent or non-covalent, and covalent bonds are preferred. The formed bonds between the tri-functional linker and the one or two or three bound molecules via respective chemical groups, can be cleavable (labile) bonds, such as cleavable under acidic conditions inside cells such as endosomes and lysosomes of mammalian cells such as human cells, or can be non-cleavable bonds. Of course, the tri-functional linker may encompass one or two chemical groups for forming covalent bonds while the further two or one chemical group(s), respectively, are/is for forming a non-covalent bond. Of course, the tri-functional linker may encompass one or two chemical groups for forming cleavable bonds while the further two or one chemical group(s), respectively, are/is for forming a non-cleavable bond.

[0037] The term "cleavable", such as used in the term "cleavable linker" or "cleavable bond" has its regular scientific meaning, and here refers to being subject to cleavage under acidic conditions, reductive conditions, enzymatic conditions or light-induced conditions. For example, a cleavable linker may be subject to cleavage under acidic conditions, preferably said cleavable linker is subject to cleavage in vivo under acidic conditions as present in endosomes and/or lysosomes of mammalian cells, preferably human cells, preferably at pH 4.0 - 6.5, and more preferably at pH ≤ 5.5. As another example, a cleavable linker may be subject to cleavage by an enzyme, e.g. by cathepsin. Furthermore, an example of a covalent bond cleavable under reductive conditions is a disulphide bond.

[0038] The terms "oligomer" and "polymer" in the context of an oligomeric or polymeric scaffold has its regular scientific meaning. A polymer here refers to a substance which has a molecular structure built up chiefly or completely from a large number of equal or similar units bonded together; an oligomer here refers to a polymer whose molecules consist of relatively few repeating units. For example, a structure comprising 5-10 or less equal or similar units, may be called an oligomeric structure, whereas a structure comprising 10-50 monomeric units or more may be called a polymeric structure, whereas a structure of 10 monomeric units may be called either oligomeric or polymeric.

[0039] The term "binding site" has its regular scientific meaning, and here refers to a region or an epitope on a molecule, e.g. a protein, DNA or RNA, to which another molecule can bind.

[0040] The term "scaffold" has its regular scientific meaning, and here refers to an oligomeric or polymeric template or a carrier or a base (base molecule or base structure), to which one or more molecules, e.g. ligand molecule, glycoside, saponin of the invention, effector molecule, can be covalently bound, either directly, or via a linker, such as a cleavable linker. A scaffold may have a structurally ordered formation such as a polymer, oligomer, dendrimer, dendronized polymer, or dendronized oligomer or have an assembled polymeric structure such as a hydrogel, microgel, nanogel, stabilized polymeric micelle or liposome, but excludes structures that are composed of non-covalent assemblies of monomers such as cholesterol/phospholipid mixtures. A scaffold may comprise a polymeric or oligomeric structure, such as poly- or oligo(amines), e.g., polyethylenimine and poly(amidoamine); or structures such as polyethylene glycol, poly- or oligo(esters), such as poly(lactids), poly(lactams), polylactide-co-glycolide copolymers; or poly(dextrin), poly- or oligosaccharides, such as cyclodextrin or polydextrose; or structures such as natural and/or artificial poly- or oligoamino acids

such as poly-lysine or a peptide or a protein, DNA oligo- or polymers, stabilized RNA polymers or PNA (peptide nucleic acid) polymers. Preferably, the polymeric or oligomeric structures are biocompatible, wherein biocompatible means that the polymeric or oligomeric structure does not show substantial acute or chronic toxicity in organisms and can be either excreted as it is or fully degraded to excretable and/or physiological compounds by the body's metabolism.

[0041] The term "ligand" has its regular scientific meaning, and here refers to any molecule or molecules which may selectively bind to a target cell-surface molecule or target cell-surface receptor expressed at target cells, e.g. target cancer cells or target auto-immune cells. The ligand may bind to an epitope comprised by receptors or other antigens on the target cells. Preferably, the cell-binding ligands are antibodies.

[0042] The term "antibody" as used herein is used in the broadest sense, which may refer to an immunoglobulin (Ig) defined as a protein belonging to the class IgG, IgM, IgE, IgA, or IgD (or any subclass thereof), or a functional binding fragment or binding domain of an immunoglobulin. In the context of the present invention, a "binding fragment" or a "binding domain" of an immunoglobulin is defined as antigen-binding fragment or -domain or other derivative of a parental immunoglobulin that essentially maintains the antigen binding activity of such parental immunoglobulin. Functional fragments and functional domains are antibodies in the sense of the present invention even if their affinity to the antigen is lower than that of the parental immunoglobulin. "Functional fragments and -domains" in accordance with the invention include, but are not limited to, F(ab')2 fragments, Fab' fragments, Fab fragments, scFv, dsFv, single-domain antibody (sdAb), monovalent IgG, scFv-Fc, reduced IgG (rIgG), minibody, diabodies, triabodies, tetrabodies, Fc fusion proteins, nanobodies, variable V domains such as VHH, Vh, and other types of antigen recognizing immunoglobulin fragments and domains. The fragments and domains may be engineered to minimize or completely remove the intermolecular disulphide interactions that occur between the CH1 and CL domains. Functional fragment and -domains offer the advantage of greater tumor penetration because of their smaller size. In addition, the functional fragment or -domain can be more evenly distributed throughout the tumor mass as compared to whole immunoglobulin.

[0043] The antibodies (immunoglobulins) of the present invention may be bi- or multifunctional. For example, a bifunctional antibody has one arm having a specificity for one receptor or antigen, while the other arm recognizes a different receptor or antigen. Alternatively, each arm of the bifunctional antibody may have specificity for a different epitope of the same receptor or antigen of the target cell.

[0044] The antibodies (immunoglobulins) of the present invention may be, but are not limited to, polyclonal antibodies, monoclonal antibodies, human antibodies, humanized antibodies, chimeric antibodies, resurfaced antibodies, anti-idiotypic antibodies, mouse antibodies, rat antibodies, rat/mouse hybrid antibodies, llama antibodies, llama heavy-chain only antibodies, heavy-chain only antibodies, and veterinary antibodies. Preferably, the antibody (immunoglobulin) of the present invention is a monoclonal antibody. The resurfaced, chimeric, humanized and fully human antibodies are also more preferred because they are less likely to cause immunogenicity in humans. The antibodies of the ADC of the present invention preferably specifically binds to an antigen expressed on the surface of a cancer cell, an autoimmune cell, a diseased cell, an aberrant cell, while leaving any healthy cell essentially unaltered (e.g. by not binding to such normal cell, or by binding to a lesser extent in number and/or affinity to such healthy cell).

[0045] Specific antibodies that can be used for the ADCs of the present invention include, but are not limited to, anti-HER2 monoclonal antibody such as trastuzumab and pertuzumab, anti-CD20 monoclonal antibody such as rituximab, ofatumumab, tositumomab and ibritumomab, anti-CA125 monoclonal antibody such as oregovomab, anti-EpCAM (17-1A) monoclonal antibody such as edrecolomab, anti-EGFR monoclonal antibody such as cetuximab, panitumumab and nimotuzumab, anti-CD30 monoclonal antibody such brentuximab, anti-CD33 monoclonal antibody such as gemtuzumab and huMy9-6, anti-vascular integrin alpha-v beta-3 monoclonal antibody such as etaracizumab, anti-CD52 monoclonal antibody such as alemtuzumab,

anti-CD22 monoclonal antibody such as epratuzumab, anti-CEA monoclonal antibody such as labetuzumab, anti-CD44v6 monoclonal antibody such as bivatuzumab, anti-FAP monoclonal antibody such as sibrotuzumab, anti-CD19 monoclonal antibody such as huB4, anti-CanAg monoclonal antibody such as huC242, anti-CD56 monoclonal antibody such huN901, anti-CD38 monoclonal antibody such as daratumumab, anti-CA6 monoclonal antibody such as DS6, anti-IGF-IR monoclonal antibody such as cixutumumab and 3B7, anti-integrin monoclonal antibody such as CNTO 95, and anti-syndecan-1 monoclonal antibody such as B-B4. An embodiment is the protein-toxin conjugate of Stemline: ELZONRIS™ (tagraxofusp, SL-401) - ELZONRIS is a novel targeted therapy directed to the interleukin-3 (IL-3) receptor-α (CD123), a target present on a wide range of malignancies.

[0046] Any other molecules than antibodies that bind to a cell receptor or antigen of a target cell can also be used as the cell-binding ligand for the ligand-drug conjugates of the present invention and the ligands provided with covalently bound saponin according to the invention. These ligands include, but are not limited to, proteins, polypeptides, peptides, small molecules. Examples of these non-antibody ligands are interferons (e.g. IFN-α, IFN-β, and IFN-γ), transferrins, lectins, epidermal growth factors (EGF) and EGF-like domains, gastrin-releasing peptides (GRP), platelet-derived growth factors (PDGF), transforming growth factors (TGF), vaccinia growth factor (VGF), insulin and insulin-like growth factors (IGF, e.g. IGF-1 and IGF-2), other suitable hormones such as thyrotropin releasing hormones (TRH), melanocyte-stimulating hormones (MSH), steroid hormones (e.g. estrogen and androgen), somatostatin, lymphokines (e.g. IL-2, IL-3, IL-4, and IL-6), colony-stimulating factors (CSF, e.g. G-CSF, M-CSF and GM-CSF), bombesin, gastrin, Arg-Gly-Asp or RGD, aptamers (e.g. AS-1411, GBI-10, RNA aptamers against HIV glycoprotein), small molecules (e.g. folate, anisamide phenylboronic acid), vitamins (e.g., vitamin D), carbohydrates (e.g. hyaluronic acid, galactose).

[0047] An "effector molecule" or "effector moiety" or "payload" has its regular scientific meaning and in the context of this invention is any substance that affects the metabolism of a cell by interaction with an intracellular effector molecule target, wherein this effector molecule target is any molecule or structure inside cells excluding the lumen of compartments and vesicles of the endocytic and recycling pathway but including the membranes of these compartments and vesicles. Said structures inside cells thus include the nucleus, mitochondria, chloroplasts, endoplasmic reticulum, Golgi apparatus, other transport vesicles, the inner part of the plasma membrane and the cytosol.

[0048] The effector molecule or -moiety is a pharmaceutically active substance, such as a toxin such as a proteinaceous toxin, a drug, a polypeptide or a polynucleotide. A pharmaceutically active substance in this invention is an effector molecule or -moiety that is used to achieve a beneficial outcome in an organism, preferably a vertebrate, more preferably a mammal such as non-human subjects or a human being/subject. Benefits include diagnosis, prognosis, treatment, cure and prevention (prophylaxis) of diseases and/or symptoms and/or health problems. The pharmaceutically active substance may also lead to undesired and sometimes even harmful side effects (adverse events such as observed during clinical trials). In this case, pros and cons must be weighed to decide whether the pharmaceutically active substance is suitable in the particular case. If the effect of the pharmaceutically active substance inside a cell is predominantly beneficial for the organism as a whole, the cell is called a target cell. If the effect inside a cell is predominantly harmful for the organism as a whole, the cell is called an off-target cell. In artificial systems such as cell cultures and bioreactors, target cells and off-target cells depend on the purpose and are defined by the user. Examples of effector molecules and -moieties are a drug, a toxin, a polypeptide (such as an enzyme), a polynucleotide (including polypeptides and polynucleotides that comprise non-natural amino acids or nucleic acids), and any combination thereof.

[0049] An effector molecule or effector moiety that is a drug may include anti-cancer agents, antiinflammatory agents, and anti-infective (e.g., anti-fungal, antibacterial, anti-parasitic, anti-viral) agents. Preferably, the drug molecule of the present invention is an anti-cancer agent or an anti-auto-immune agent. Suitable anti-cancer agents include, but are not limited to, alkylating agents, antimetabolites, spindle poison plant alkaloids,

cytotoxic/antitumor antibiotics, topoisomerase inhibitors, photosensitizers, and kinase inhibitors. Also included in the definition of "anti-cancer agent" are: e.g. (i) anti-hormonal agents that act to regulate or inhibit hormone action on tumors such as anti-estrogens and selective estrogen receptor modulators; (ii) aromatase inhibitors that inhibit the enzyme aromatase, which regulates estrogen production in the adrenal glands; (iii) anti-androgens; (iv) protein kinase inhibitors; (v) lipid kinase inhibitors; (vi) antisense oligonucleotides, particularly those which inhibit expression of genes in signaling pathways implicated in aberrant cell proliferation; (vii) ribozymes such as VEGF expression inhibitors and HER2 expression inhibitors; (viii) vaccines such as gene therapy vaccines; topoisomerase 1 inhibitors; (ix) anti-angiogenic agents; and pharmaceutically acceptable salts, acids, solvates and derivatives of any of the above.

[0050] An effector molecule or -moiety that is a toxin may include, but is not limited to, proteinaceous toxins (e.g. bacterial-derived toxins, and plant-derived toxins), toxins targeting tubulin filaments, toxins targeting DNA, toxins targeting RNA. Examples of proteinaceous toxins are saporin, dianthin, ricin, modeccin, abrin, volkensin, viscumin, shiga toxin, shiga-like toxin, pseudomonas exotoxin (PE, also known as exotoxin A), diphtheria toxin (DT), and cholera toxin. Examples of tubulin filaments-targeting toxins are maytansinoids (e.g. DM1 and DM4), auristatins (e.g. Monomethyl auristatin E (MMAE) and Monomethyl auristatin F (MMAF)), toxoids, tubulysins, cryptophycins, rhizoxin. Examples of DNA-targeting toxins are calicheamicins: N-Acetyl- γ-calicheamicin, CC-1065 analogs, duocarmycins, doxorubicin, methotrexate, benzodiazepines, camptothecin analogues, and anthracyclines. Examples of DNA-targeting toxins are amanitins, spliceostatins, and thailanstatins. A toxin, as used in this invention, is defined as a pharmaceutically active substance that is able to kill or inactivate a cell. Preferably, a targeted toxin is a toxin that is only, or at least predominantly, toxic for target cells but not for off-target cells. The net effect of the targeted toxin is preferably beneficial for the organism as a whole.

[0051] An effector molecule or -moiety that is a polypeptide may be, e.g., a polypeptide that recover a lost function, such as for instance enzyme replacement, gene regulating functions, or a toxin. Examples of polypeptides as effector molecules are, e.g., Cas9; toxins (e.g. saporin, dianthin, gelonin, (de)bouganin, agrostin, ricin (toxin A chain); pokeweed antiviral protein, apoptin, diphtheria toxin, pseudomonas exotoxin) metabolic enzymes (e.g. argininosuccinate lyase, argininosuccinate synthetase), enzymes of the coagulation cascade, repairing enzymes; enzymes for cell signaling; cell cycle regulation factors; gene regulating factors (transcription factors such as NF-κB or gene repressors such as methionine repressor).

[0052] An effector molecule or an effector moiety that is a polynucleotide may, e.g., be a polynucleotide that comprises coding information, such as a gene or an open reading frame encoding a protein. It may also comprise regulatory information, e.g. promotor or regulatory element binding regions, or sequences coding for micro RNAs. Such polynucleotide may comprise natural and artificial nucleic acids. Artificial nucleic acids include, e.g. peptide nucleic acid (PNA), Morpholino and locked nucleic acid (LNA), as well as glycol nucleic acid (GNA) and threose nucleic acid (TNA). Each of these is distinguished from naturally occurring DNA or RNA by changes to the backbone of the molecule. Examples of nucleotides as effector molecules are, e.g., DNA: single stranded DNA (e.g. DNA for adenine phosphoribosyltransferase); linear double stranded DNA (e.g. clotting factor IX gene); circular double stranded DNA (e.g. plasmids); RNA: mRNA (e.g. TAL effector molecule nucleases), tRNA, rRNA, siRNA, miRNA, antisense RNA; anti-sense oligonucleotides (ASOs, AONs e.g. PNA, PMO, LNA and BNA).

[0053] The term "proteinaceous", used in e.g. "proteinaceous molecule" and "proteinaceous toxin", are molecules and toxins comprising at least a string of amino acid residues that can be obtained as an expression product from a single mRNA. Such a molecule or toxin may further comprise any post-translational modifications, a carbohydrate such as an N- or O-linked carbohydrate, disulphide bonds, phosphorylations, sulphatations, etc., as a result of any post-translational modification, and/or may further comprise any other modification such as those resulting from chemical modifications (e.g., linking of effector moieties, saponin, scaffolds, ligands, etc., either directly to e.g. an amino-acid side chain, or via at least one linker (covalently) bound to the molecule for chemically modifying the proteinaceous molecule, and chemically bound (covalently)

to the proteinaceous molecule). The term "proteinaceous" also encompasses and includes assemblies of such molecules, e.g. homodimers, heterotrimers, heterohexamers or complex assemblies such as ribosomes.

[0054] The terms "specific" and "specifically", in the context of for example "specific binding" and "receptor or molecular target specifically present or expressed at the surface of a tumor cell", have their normal scientific meaning known in the art, and here refer to e.g. a binding interaction of a first molecule with a second molecule which occurs with a higher affinity relative to any putative binding of the first molecule to a further molecule different from the second molecule, or e.g. to the expression or expression to a higher extent when e.g. the number of receptors or molecular targets is considered, of a cell-surface receptor or molecular target on the surface of a first type of cell such as a tumor cell, autoimmune cell, diseased cell, aberrant cell, relative to the extent of expression of the same receptor or molecular target at a second type of cell such as a healthy cell, etc., wherein expression at the second type of cell can be fully absent or very low, relative to any extent of expression on the tumor cell, etc. Furthermore, the term "specific", for example in "specific binding", has its normal scientific meaning known in the art, and here has the meaning of indicating a molecule that can have an interaction with another molecule with higher binding affinity than background interactions between molecules. Similarly, the term "specificity" refers to an interaction, for example, between two molecules or between a cell and a molecule, which has higher binding affinity than background interactions between molecules. Binding molecules such as immunoglobulins bind via their binding site such as immunoglobulin variable regions of the immunoglobulin, to binding sites on molecules, such as epitopes, cell-surface receptors, etc., with a higher binding affinity than background interactions between molecules. In the context of the invention, background interactions are typically interactions with an affinity lower than a K_D of 10E-4 M. Similarly, "specific binding domains" are domains that preferentially bind to binding sites on molecules, such as epitopes, cell-surface receptors, etc., with a higher binding affinity than background interactions between molecules. In the context of the invention, "background interactions" are typically interactions with an affinity lower than a K_D of 10E-4 M. Preferably, specific binding domains bind with an affinity higher than a K_D of about 10E-5 M.

[0055] The term "binding" is defined as interactions between molecules that can be distinguished from background interactions.

[0056] Throughout the specification, the term "fragment" refers to an amino acid sequence which is part of a protein domain or which builds up an intact protein domain. Binding fragments according to the invention must have binding specificity for the respective target such as a cell-surface receptor, e.g. on the surface of a diseased cell such as a tumor cell.

[0057] The term "ADC" or "antibody-drug conjugate" has its regular scientific meaning known to the skilled person, and here refers to a class of biopharmaceutical drugs designed as a targeted therapy for treating e.g. cancer. Unlike chemotherapy, ADCs are intended to target and kill tumor cells while sparing healthy cells. ADCs are composed of an antibody linked to a biologically active cytotoxic (anticancer) payload or drug. ADCs combine the targeting capabilities of monoclonal antibodies with the cancer-killing ability of cytotoxic drugs. They are designed with the intention to discriminate between healthy cells and diseased tissue such as tumor cells in a tumor.

[0058] The term "Saponinum album" has its normal meaning and here refers to a mixture of saponins produced by Merck KGaA (Darmstadt, Germany) containing saponins from *Gypsophila paniculata* and *Gypsophila arostii,* containing SA1657 and mainly SA1641.

[0059] The term "Quillajasaponin" has its normal meaning and here refers to the saponin fraction of Quillaja saponaria and thus the source for all other QS saponins, mainly containing QS-18 and QS-21.

[0060] "QS-21" or "QS21" has its regular scientific meaning and here refers to a mixture of QS-21 A-apio

(-63%), QS-21 A-xylo (-32%), QS-21 B-apio (-3.3%), and QS-21 B-xylo (-1.7%).

[0061] Similarly, "QS-21A" has its regular scientific meaning and here refers to a mixture of QS-21 A-apio (-65%) and QS-21 A-xylo (-35%).

[0062] Similarly, "QS-21B" has its regular scientific meaning and here refers to a mixture of QS-21 B-apio (-65%) and QS-21 B-xylo (-35%).

[0063] The term "Quil-A" refers to a commercially available semi-purified extract from Quillaja saponaria and contains variable quantities of more than 50 distinct saponins, many of which incorporate the triterpene-trisaccharide substructure Gal-(1→2)-[Xyl-(1→3)]-GlcA- at the C-3beta-OH group found in QS-7, QS-17, QS18, and QS-21. The saponins found in Quil-A are listed in van Setten (1995), Table 2 *[*Dirk C. van Setten, Gerrit van de Werken, Gijsbert Zomer and Gideon F. A. Kersten, Glycosyl Compositions and Structural Characteristics of the Potential Immuno-adjuvant Active Saponins in the Quillaja saponaria Molina Extract Quil A, RAPID COMMUNICATIONS IN MASS SPECTROMETRY, VOL. 9,660-666 (1995)]. Quil-A and also Quillajasaponin are fractions of saponins from Quillaja saponaria and both contain a large variety of different saponins with largely overlapping content. The two fractions differ in their specific composition as the two fractions are gained by different purification procedures.

[0064] The term "QS1861" and the term "QS1862" refer to QS-7 and QS-7 api. QS1861 has a molecular mass of 1861 Dalton, QS1862 has a molecular mass of 1862 Dalton. QS1862 is described in Fleck et al. (2019) in Table 1, row no. 28 [Juliane Deise Fleck, Andresa Heemann Betti, Francini Pereira da Silva, Eduardo Artur Troian, Cristina Olivaro, Fernando Ferreira and Simone Gasparin Verza, Saponins from Quillaja saponaria and Quillaja brasiliensis: Particular Chemical Characteristics and Biological Activities, Molecules 2019, 24, 171; doi:10.3390/molecules24010171]. The described structure is the api-variant QS1862 of QS-7. The molecular mass is 1862 Dalton as this mass is the formal mass including proton at the glucuronic acid. At neutral pH, the molecule is deprotonated. When measuring in mass spectrometry in negative ion mode, the measured mass is 1861 Dalton.

[0065] The terms first, second, third in the description and in the claims, are used for distinguishing between similar elements and not necessarily for describing a sequential or chronological order. The terms are interchangeable under appropriate circumstances. The embodiments of the invention can operate in other sequences than described or illustrated herein.

[0066] Furthermore, the various embodiments, although referred to as "preferred" or "*e.g.*" or "for example" or "in particular" are to be construed as exemplary manners in which the invention may be implemented rather than as limiting the scope of the invention.

[0067] The term "comprising", used in the claims, should not be interpreted as being restricted to the elements or steps listed thereafter; it does not exclude other elements or steps. It needs to be interpreted as specifying the presence of the stated features, integers, steps or components as referred to, but does not preclude the presence or addition of one or more other features, integers, steps or components, or groups thereof. Thus, the scope of the expression "a pharmaceutical composition comprising A and B" should not be limited to a pharmaceutical composition consisting only of components A and B, rather with respect to the present invention, the only enumerated components of the pharmaceutical composition are A and B, and further the claim should be interpreted as including equivalents of those components. Similarly, the scope of the expression "a method comprising step A and step B" should not be limited to a method consisting only of steps A and B, rather with respect to the present invention, the only enumerated steps of the method are A and B, and further the claim should be interpreted as including equivalents of those steps.

[0068] In addition, reference to a feature by the indefinite article "a" or "an" does not exclude the possibility

that more than one of the features such as for example a component, excipient, saponin, etc. are present, unless the context clearly requires that there is one and only one of the features. The indefinite article "a" or "an" thus usually means "at least one".

BRIEF DESCRIPTION OF THE DRAWINGS

[0069]

Figure 1. 1T2C in vivo activity. The 1T2C combination of 50mg/kg cetuximab-(Cys-L-SO1861)^4 + 25 mg/kg cetuximab-(-L-HSP27BNA)^4 in A431 tumor bearing mice reveals strong tumor targeted gene silencing, compared to the controls.

Figure 2. 1T2C in vivo activity. The 1T2C combination of 40 mg/kg trastuzumab-(Cys-L-SO1861)^4 + 0.02/0.03 mg/kg trastuzumab-saporin in a PDX tumor mouse model (high HER2 expression) shows effective tumor growth inhibition.

Figure 3. 1-target 2-component. EGFR targeted cell killing in A431 cells (EGFR^{++}) (A, C) and CaSKi cells (EGFR^{+}) (B, D) by a therapeutic combination according to the invention. A, B) Cetuximab-(Cys-L-SO1861)^{3,7} titration + fixed concentration 10 pM cetuximab-saporin and controls on A431 (A) and CaSKi (B) cells. C, D) Cetuximab-saporin titration + fixed concentration of 75nM cetuximab-(Cys-L-SO1861)^{3,7} and controls on A431 (C) and CaSKi (D) cells. Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 4. 1-target 2-component. EGFR targeted cell killing in HeLa cells (EGFR$^{+/-}$) (A, C) and A2058 cells (EGFR^{-}) (B, D) by a therapeutic combination according to the invention. A, B) Cetuximab-(Cys-L-SO1861)^{3,7} titration + fixed concentration 10 pM cetuximab-saporin and controls on HeLa (A) and CaSKi (B) cells. C, D) Cetuximab-saporin titration + fixed concentration of 75nM cetuximab-(Cys-L-SO1861)^{3,7} and controls on Hela (C) and A2058 (D) cells. Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 5: 1-target 2-component. HER2 targeted cell killing in SKBR3 cells (HER2^{++}) (A, B) by a therapeutic combination according to the invention. A) Trastuzumab-(Cys-L-SO1861)^4 titration + fixed concentration 50 pM trastuzumab-saporin and controls on SKBR3 cells. B) Trastuzumab-saporin titration + fixed concentration of 2.5nM trastuzumab-(Cys-L-SO1861)^4 and controls on SKBR3 cells. Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 6. 1-target 2-component. HER2 targeted cell killing in JIMT-1 cells (HER2$^{+/-}$) (A, C) and MDA-MB-468 cells (HER2) (B, D) by a therapeutic combination according to the invention. A, B) Trastuzumab-(Cys-L-SO1861)^4 titration + fixed concentration of 50 pM trastuzumab-saporin and controls on JIMT-1 (A) and MDA-MB-468 (B) cells. C, D) Trastuzumab-saporin titration + fixed concentration of 2.5nM trastuzumab-(Cys-L-SO1861)^4 and controls on JIMT-1 (C) and MDA-MB-468 (D) cells. Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 8: *Chloroquine inhibits the 1-target 2-component.* HER2 and EGFR targeted cell killing in SK-BR-3 (HER2^{++}) and A431 cells (EGFR^{++}), by a therapeutic combination according to the invention + chloroquine. A) Trastuzumab-saporin titration + fixed concentration of 5nM trastuzumab-(Cys-L-SO1861)^4 + 0.5 µM chloroquine and control on SK-BR-3 cells. B) Cetuximab-saporin titration + fixed concentration of 5 nM cetuximab-(Cys-L-SO1861)^{3,8} + 0.5 µM chloroquine and control on A431 cells. Remark: For target receptor

expression data of each cell line (determined by FACS analysis) see table 19.

Figure 9: 1-target 2-component. EGFR targeted gene silencing in A431 cells ($EGFR^{++}$) and A2058 cells (EGFR) by a therapeutic combination according to the invention. A,B) Cetuximab-$(Cys\text{-}L\text{-}SO1861)^{3,8}$ titration + fixed concentration of 100 nM Cetuximab-$(Lys\text{-}L\text{-}HSP27BNA)^{4}$ and control on A431 cells (A) and A2058 cells (B). C, D) Cetuximab-$(Lys\text{-}L\text{-}HSP27BNA)^{4}$ titration + fixed concentration of 77 nM Cetuximab-$(Cys\text{-}L\text{-}SO1861)^{3,8}$ and control on A431 cells (C) and A2058 cells (D). Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 10: 2-target 2-component. A) EGFR and HER2 targeted cell killing in MDA-MB-468 cells ($EGFR^{++}$) and HeLa cells ($EGFR^{+/-}$) and HER2 targeted cell killing in SK-BR-3 cells ($HER2^{++}$) and JIMT-1 cells ($HER2^{+/-}$) by a therapeutic combination according to the invention. A) Cetuximab-Cys-$(dendron(\text{-}L\text{-}SO1861)^{4})^{3,9}$ titration + fixed concentration 10 pM cetuximab-saporin and controls on MDA-MB-468 cells (A) and HeLa cells (B). C,D) Trastuzumab-Cys-$(dendron(\text{-}L\text{-}SO1861)^{4})^{4}$ titration + fixed concentration 50 pM trastuzumab-saporin and controls on SK-BR-3 cells (C) and JIMT-1 cells (D). Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 11. 1-target 2-component. SK-BR-3 cells ($HER2^{+/-}$) can efficiently be killed with the therapeutic combination according to the invention, Tratuzumab-saporin + 2.5 nM trastuzumab-$(Cys\text{-}L\text{-}SO1861)^{4}$, however titration of T-DM1 + 2.5 nM trastuzumab-$(Cys\text{-}L\text{-}SO1861)^{4}$ is not effective at such low toxin concentrations. T-DM1 is Trastuzumab-emtansine (Kadcyla®), carrying ~3.5 emtansine (DM1) toxin molecules per antibody (DAR3.5). Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 12. *Control treatments on all cell lines.* A-D) Cell viability when trastuzumab (A), cetuximab (B), T-DM1, (C) free toxins: saporin and dianthin (D) or saporin coupled to a non-cell binding IgG (D) are treated with the indicated cell lines SK-BR-3, JIMT-1, MDA-MB-468, A431, CaSki, HeLa, A2058, BT-474. Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 13. 1-target 2-component. EGFR targeted cell killing in A431 cells ($EGFR^{++}$) (A) and CaSKi cells ($EGFR^{+}$) (B) and A2058 cells ($EGFR^{-}$) by a therapeutic combination according to the invention. A, B, C) Cetuximab-$(Cys\text{-}L\text{-}QSmix)^{4,1}$ titration + fixed concentration 10 pM cetuximab-saporin or 10pM cetuximab-dianthin and controls in A431 cells (A), CaSKi cells (B) and A2058 cells (C). QSmix is a mixture of saponins from an extract Quillaja Saponaria. Remark: For target receptor expression data of each cell line (determined by FACS analysis) see table 19.

Figure 14: 1-target 2-component concept: mAb1-SO1861 + *mAb1-protein toxin.* SO1861 and toxin (ribosomal inactivating protein) are each, independently, conjugated to an antibody (mAb1) for delivery and internalization into target cells. 1) mAb1-SO1861 and mAb1-protein toxin bind to the cell surface receptor, 2) receptor-mediated endocytosis of both conjugates occurs, 3) at low endolysosomal pH and appropriate concentration, SO1861 becomes active to enable endolysosomal escape, 4) release of toxin into cytoplasm occurs and 5) toxin induces cell death

Figure 15: 1-target 2-component concept: mAb1-SO1861 + *mAb2-BNA oligo.* SO1861 and antisense BNA oligo nucleotide are each, independently, conjugated to an antibody (mAb1) for delivery and internalization into target cells. 1) mAb1-SO1861 and mAb1-BNAoligo bind to the cell surface receptor, 2) receptor-mediated endocytosis of both conjugates occurs, 3) at low endolysosomal pH and appropriate concentration, SO1861 becomes active to enable endolysosomal escape, 4) release of BNA oligo into cytoplasm occurs and 5) target gene silencing.

Figure 16: 1-target 2-component concept: mAb1-(scaffold(-SO1861)^{n})^{n} + *mAb1-protein toxin.* Dendron(-SO1861)^{n} and protein toxin (ribosomal inactivating protein) are each, independently, conjugated to an antibody (mAb1) for delivery and internalization into target cells. 1) mAb1-dendron(-SO1861)^{4} and mAb1-protein toxin bind to the cell surface receptor, 2) receptor-mediated endocytosis of both conjugates occurs, 3) at low endolysosomal pH and appropriate concentration, SO1861 becomes active to enable endolysosomal escape, 4) release of toxin into cytoplasm occurs and 5) toxin induces cell death

Figure 17. *Antibody-SO1861 conjugation procedure.* Shown is the coupling reaction of the linking of four moieties of a plant-derived saponin SO1861 to the four cysteines in the light chain of an antibody. First, the disulphide bonds in the IgG are disrupted under influence of exposure to TCEP (Tris(2-carboxyethyl)phosphine); second, the saponin SO1861 comprising a chemical linker bound to it, is added together with trifluoro acetic acid, and four saponin moieties are linked to the IgG. For producing cleavable 'ready to conjugate' saponins the aldehyde group of SO1861 was reacted with an **EMCH** (ε-maleimidocaproic acid hydrazide) linker. The hydrazide group of EMCH forms an acid cleavable hydrazone bond with the aldehyde of SO1861. At the same time the EMCH linker presents a maleimide group that is thiol (sulfhydryl group) reactive and thus can be conjugated to thiols of the IgG, i.e. the ligand moiety. Herewith, an endosomal escape enhancing conjugate of the invention is provided, and/or a first binding molecule of the invention is provided.

Figure 18. SO1861-EMCH synthesis

Figure 19. Dendron-(-L-SO1861)^{4} synthesis

Figure 20. Dendron-(-L-SO1861)^{8} synthesis

Figure 21. SG181-L-trifcuntional linker-L-HSP27BNA synthesis

Figure 22. HSP27BNA-dendron-(-L-SO1861)^{4} synthesis

Figure 23. Dendron(NEM)^{4} synthesis synthesis

Figure 24. SO1861-EMCH synthesis

Figure 25: Scaffold precursor with four amino groups for saponin linkage and an azide group for click chemistry.

Figure 26: Evidence for the coupling of saponins to the model scaffold. The inset shows the theoretically expected peeks and intensity distribution for coupled saponins. The experimental data obtained by LC-MS/ESI-MS show almost exactly the same peaks at *m/z* 758-760 Da proving successful saponin coupling.

Figure 27: Cytotoxicity assays using the targeted toxin dianthin-Epidermal Growth Factor (dianthin-EGF). Untreated cells were normalized to 1. The polymeric structure (Pentrimer) has no influence on cell viability neither in the presence nor in the absence of Dianthin-EGF and saponin (SA1641) indicating no intrinsic cytotoxicity of the polymeric structure. The clickable targeted toxin (Dianthin-EGF-Alkyne) has a markedly reduced activity, which is a result of the toxin modification but does not have any relation to the scaffold. The functionalized polymeric structure has the same activity as the unclicked targeted toxin, indicating that the functionalization of the scaffold does not impair effector molecule activity. The effect of saponins is identical in the presence and absence of the polymeric structure showing that the polymeric structure does not impair the efficacy of the saponins in the two-component system.

Figure 28: H-NMR spectrum of (A) SO1861 and (B) SO1861-EMCH (EMCH = N-ε-maleimidocaproic acid hydrazide). (A) The peak at 9.43 ppm (H^{a}) corresponds to the aldehyde proton of SO1861. (B) The peak at 6.79 ppm (H^{c}) corresponds to the maleimide protons of SO1861-EMCH, while the peak at 7.68 ppm (H^{b})

corresponds to the hydrazone proton. The absence of the signal at 9.43 ppm indicates a quantitative conversion of the aldehyde group.

Figure 29: (A) MALDI-TOF-MS spectrum of SO1861-EMCH and (B) SO1861-EMCH-mercaptoethanol. (A) RP mode: *m/z* 2124 Da ($[M+K]^+$, saponin-EMCH), *m/z* 2109 Da ($[M+K]^+$, SO1861-EMCH), *m/z 2094* Da ($[M+Na]^+$, SO1861-EMCH). (B) RP mode: *m/z* 2193 Da ($[M+K]^+$, saponin-EMCH-mercaptoethanol), *m/z* 2185 Da ($[M+K]^+$, SO1861-EMCH-mercaptoethanol), *m/z* 2170 Da ($[M+Na]^+$, SO1861-EMCH-mercaptoethanol).

Figure 30: SO1861 structure with highlighted chemical groups for conjugation of endosomal escape enhancing saponins to a polymeric structure. Highlighted groups are aldehyde (black circle), carboxylic acid (dashed circle), alkene (dashed pentagon), and alcohol (dashed box). The aldehyde group (arrow) is most suitable group for chemoselective and reversible conjugation reactions.

Figure 31: Strategy for producing (A) stable and (B) cleavable 'ready-to conjugate' endosomal escape enhancer saponins.

Figure 32: Hydrolysis of the hydrazone bond of SO1861-EMCH under acidic conditions.

Figure 33: SO1861-EMCH structure. (A) Standard molecular structure, and (B) 3D model. Maleimide group is marked with a circle.

Figure 34: (A) SO1861-EMCH synthesis scheme. (B) MALDI-TOF-MS spectra of SO1861 (*m/z* 1861 Da) and (C) SO1861-EMCH (*m/z*2068 Da) in negative reflector mode. TFA: trifluoroacetic acid, r.t: room temperature, h: hours, and MW: molecular weight.

Figure 35: MALDI-TOF-MS spectra of SO1861-EMCH (A) before and (B) after hydrolysis in HCl solution at pH 3.

Figure 36: Reaction scheme of SO1861-EMCH conjugation to any amine-bearing polymeric structure.

Figure 37: MALDI-TOF-MS spectra of (A) BSA-SO1861 (*m/z* 70.0 kDa, 72.1 kDa, 74.2 kDa), and (B) BSA (*m/z* 66.6 kDa).

Figure 38: Reaction scheme of (A) SO1861-EMCH and (B) SO1861-HATU (HATU = 1-[Bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate) conjugation to a cyanine 3 dye labeled polyamidoamine (PAMAM) G5 dendrimer.

Figure 39: MALDI-TOF-MS spectra of (A) Cy3-PAMAM, (B-D) Cy3-PAMAM-SO1861 with increasing SO1861-EMCH feed equivalents from (B) up to bottom (D). (B) corresponds to Cy3-PAMAM-SO1861 with 5 SO1861 attached per PAMAM, (C) corresponds to Cy3-PAMAM-SO1861 with 13 SO1861 attached per PAMAM, and (D) corresponds to Cy3-PAMAM-SO1861 with 51 SO1861 attached per PAMAM.

Figure 40: MALDI-TOF-MS spectra of (A) Cy3-PAMAM-SO1861 with 5 equivalents feed SO1861-EMCH and (B) Cy3-PAMAM-SO1861 with 30 equivalents feed SO1861-EMCH.

Figure 41: MALDI-TOF-MS spectra of Cy3-PAMAM-NC-SO1861 (NC = stable bond ("non-cleavable").

Figure 42: (A) Reaction scheme and MALDI-TOF-MS spectra of (B) Cy3-PAMAM-NC-SO1861-Dibenzocyclooctyne (DBCO), (C) Cy3-PAMAM-$(SO1861)_5$-DBCO, and (D) Cy3-PAMAM-$(SO1861)_{27}$-DBCO.

Figure 43: Reaction scheme of (A) dianthin-EGF-Alexa488 and (B) dianthin-EGF-Alexa488-SS-PEG-N_3. MALDI-TOF-MS spectra of (C) dianthin-EGF, (D) dianthin-EGF-Alexa488, and (E) dianthin-EGF-Alexa488-SS-PEG-N_3; Alexa488: Alexa Fluor 488 dye.

Figure 44: Reaction scheme of (A) dianthin-Alexa488 and (B) dianthin-Alexa488-SS-PEG-N_3. MALDI-TOF-MS

spectra of (C) dianthin, (D) dianthin-Alexa488, and (E) dianthin-Alexa488-SS-PEG-N_3; Alexa488: Alexa Fluor 488 dye.

Figure 45: Fluorescence images of SDS-PAGE gel performed on a VersaDoc imaging system. M = marker, P = Cy3-PAMAM-(SO1861)$_{27}$-DBCO, D = dianthin-EGF-Alexa488-SS-PEG-N_3, C1 = Cy3-PAMAM-(SO1861)$_5$-Dianthin-EGF-Alexa488, C2 = Cy3-PAMAM-NC-SO1861-Dianthin-EGF-Alexa488, and C3 = Cy3-PAMAM-(SO1861)$_{27}$-Dianthin-EGF-Alexa488.

Figure 46: (A) Synthesis scheme of Cy3-PAMAM-NC-SO1861 via reductive amination. (B, and C) Respective MALDI-TOF-MS spectra.

Figure 47: Reaction scheme for the generation of poly(SO1861) using SO1861-EMCH as monomer, the APS / TMEDA system as polymerization initiator, and aminopropanethiol as radical quencher.

Figure 48: MALDI-TOF-MS spectra of poly(SO1861) reaction batches. (A) SO1861-EMCH at 60 °C, (B) SO1861-EMCH + 11^{-3} equivalents APS at 60 °C, (C) SO1861-EMCH + 11^{-3} equivalents APS / TMEDA at 60 °C.

Figure 49: DNA approach. Usage of the principle of DNA-origami to generate a DNA based scaffold that is able to conjugate and release glycoside molecules. In addition, one of the DNA strands obtains a click chemistry moiety that can be used for conjugation to a targeted toxin to form a functionalized scaffold. bp: base pair.

Figure 50: Poly(peptide-SO1861) approach. Usage of a peptide sequence that can conjugate and release glycoside molecules and which can react with itself to form a poly(peptide-SO1861) construct. The poly(peptide) chain endings can be further modified with click chemistry moieties (e.g., BCN-NHS linker) that can be used for conjugation to a toxin.

Figure 51. MALDI-TOF-MS spectra of (A) native peptide, (B) peptide-SO1861 conjugate.

Figure 52. Molecular structure of G4-dendron with protected amino groups.

Figure 53. Synthesis scheme for the generation of dendron based scaffolds and functional scaffolds.

Figure 54. (A) Reaction scheme for partial dye labeling and deprotection of the G4-dendron. (B) MALDI-TOF-MS spectrum of deprotected and partially dye labeled G4-dendron.

Figure 55. MALDI-TOF-MS spectra of G4-dendron-SO1861 scaffolds with (A) 22 feed equivalents of SO1861-EMCH, (B) 10 feed equivalents of SO1861-EMCH, and (C) 3 feed equivalents of SO1861-EMCH.

Figure 56. Cell viability curves of HeLa cells treated with (A) EGFR cell surface expression as determined by FACS analyses of HeLa cells (B, see Table 19), cell viability of HeLa cells treated with SO1861 + dianthin-EGF (Dia-EGF), SO1861 + dianthin-EGF + 500 nM chloroquine, SO1861 + dianthin-EGF + 500 nM PAMAM, SO1861 + dianthin-EGF + 667 nM dendron (C) cell viability of HeLa cells treated with SO1861 + dianthin-EGF, SO1861 + dianthin-EGF + 500 nM chloroquine, SO1861 + dianthin-EGF + 500 nM PAMAM, SO1861 + dianthin-EGF + 500 nM PAMAM-(SH)$_{16}$, SO1861 + dianthin-EGF + 500 nM PAMAM-(SH)$_{65}$, SO1861 + dianthin-EGF + 500 nM PAMAM-(SH)$_{108}$ (D) cell viability of HeLa cells treated with SO1861 + dianthin-EGF, SO1861 + dianthin-EGF + 500 nM chloroquine, SO1861 + dianthin-EGF + 500 nM PAMAM, SO1861 + dianthin-EGF + 500 nM PAMAM-(mPEG)$_3$, SO1861 + dianthin-EGF + 500 nM PAMAM-(mPEG)$_8$, SO1861 + dianthin-EGF + 500 nM PAMAM-(mPEG)$_{18}$.

Figure 57. (A) Reaction scheme of the thiolation of PAMAM using the thiolation reagent 2-iminothiolane. MALDI-TOF-MS spectra of (B) native PAMAM, (C) thiolated PAMAM-(SH)$_{16}$, (D) thiolated PAMAM-(SH)$_{65}$, and (E) thiolated PAMAM-(SH)$_{108}$.

Figure 58. (A) Reaction scheme of the PEGylation of PAMAM using the PEGylating reagent $mPEG_{2k}$-NHS. MALDI-TOF-MS spectra of (B) native PAMAM, (C) PEGylated PAMAM-$(mPEG_{2k})_3$, (D) PEGylated PAMAM-$(mPEG_{2k})_8$, and (E) PEGylated PAMAM-$(mPEG_{2k})_{18}$.

Figure 59: Basic scaffold with click chemistry function to link any desired effector molecule. The user determines the position of the click chemistry position in the effector molecule and all further properties of the effector molecule, e.g. choice and position of an optional ligand.

Figure 60: Functionalized scaffold with pre-bound effector molecule and click chemistry function to link any desired ligand. Optionally, a pH-sensitive linkage can be provided to release the effector molecule from the scaffold after reaching the endosomes.

DETAILED DESCRIPTION

[0070] In order for a bioactive molecule to work, the molecule must be able to engage with its target, e.g. in the blood serum, on the outside of the cell surface or inside a cell or an organelle. The active moiety of almost all protein-based targeted toxins, e.g., must enter the cytosol of the target cell to mediate its target modulatory effect. In many constellations the toxin remains ineffective since (1) the targeting moiety is poorly internalized and remains bound to the outside of the cells, (2) is recycled back to the cell surface after internalization or (3) transported to the endolysosomes where it is degraded. Although these fundamental issues are known for decades and more than 500 targeted toxins have been investigated in the past decades, the problems have not been solved yet and only one antibody-targeted protein toxin, moxetumomab pasudotox-tdfk (LUMOXITI®, AstraZeneca Pharmaceuticals LP), has been approved for relapsed or refractory hairy cell leukemia by the FDA to date. Further approved ADCs are Elzonris, Ontak.

[0071] To overcome these problems, many strategies have been described including approaches to redirect the toxins to endogenous cellular membrane transport complexes of the biosynthetic pathway in the endoplasmic reticulum and techniques to disrupt or weaken the membrane integrity of endosomes, i.e. the compartments of the endocytic pathway in a cell, and thus facilitating the endosomal escape. This comprises the use of lysosomotropic amines, carboxylic ionophores, calcium channel antagonists, various cell-penetrating peptides of viral, bacterial, plant, animal, human and synthetic origin, other organic molecules and light-induced techniques. Although the efficacy of the targeted toxins was typically augmented in cell culture hundred- or thousand-fold, in exceptional cases more than million-fold, the requirement to co-administer endosomal escape enhancers with other substances harbors new problems including additional side effects, loss of target specificity, difficulties to determine the therapeutic window and cell type-dependent variations.

[0072] All strategies, including physicochemical techniques, require enhancer molecules that interact more or less directly with membranes and comprise essentially small chemical molecules, secondary metabolites, peptides and proteins. A common feature of all these substances is that they are *per se* not target cell-specific and distribute with other kinetics than the targeted toxins. This is one major drawback of the current approaches.

[0073] The present invention will be described with respect to particular embodiments but the invention is not limited thereto but only by the claims. The embodiments of the invention described herein can operate in combination and cooperation, unless specified otherwise.

[0074] While the invention has been described in terms of several embodiments, it is contemplated that alternatives, modifications, permutations and equivalents thereof will become apparent to one having ordinary

skill in the art upon reading the specification and upon study of the drawings and graphs. The invention is not limited in any way to the illustrated embodiments. Changes can be made without departing from the scope which is defined by the appended claims.

[0075] An aspect of the invention relates to a therapeutic combination for use as a medicament as defined in the claims, wherein the therapeutic combination comprises: (a) a first pharmaceutical composition comprising a first proteinaceous molecule comprising a binding site for binding to an epitope on a cell-surface molecule and at least one saponin covalently bound to said first proteinaceous molecule preferably to an amino-acid residue of said first proteinaceous molecule, the first pharmaceutical composition optionally further comprising a pharmaceutically acceptable excipient; and (b) a second pharmaceutical composition comprising a second proteinaceous molecule, the second proteinaceous molecule comprising a binding site for binding to the epitope on the cell-surface molecule of (a) and an effector moiety, the second pharmaceutical composition optionally further comprising a pharmaceutically acceptable excipient, wherein the binding site of the first proteinaceous molecule and the binding site of the second proteinaceous molecule are the same, and wherein the cell-surface molecule and the epitope on the cell-surface molecule, to which the first proteinaceous molecule can bind, and the cell-surface molecule and the epitope on the cell-surface molecule, to which the second proteinaceous molecule can bind, are the same.

[0076] The inventors established that the therapeutic window of an antibody drug conjugate, such as a second proteinaceous molecule in the second pharmaceutical composition of the invention, increases when administered to a tumor-bearing mammal (mouse) to whom also the first pharmaceutical composition is administered. The first proteinaceous protein has at least one saponin bound thereto, preferably covalently, more preferably via a cleavable linker. The saponin augments the therapeutic efficacy of the effector moiety bound to the second proteinaceous molecule, likely by enhancing the endosomal escape of the effector moiety into the cytosol where the activity of the effector moiety is desired. This way, already at a lower dose than the conventional dose of the ADC, i.e. the second proteinaceous molecule, therapeutic effect is established under influence of the presence of the first proteinaceous molecule comprising the saponin near, at and inside the targeted cell. The targeted cell is for example a diseased cell such as a tumor cell or an auto-immune cell or a B-cell disease related B-cell, etc. The effector moiety is for example a toxin as part of an ADC or an oligonucleotide such as a BNA according to the invention.

[0077] By targeting the same cell-surface molecule with the first and second proteinaceous molecule, the delivery of the saponin and the effector molecule at and inside the cytosol of the very same targeted cell is improved and more specific. An aberrant cell selected for targeting by the binding site of the first and second proteinaceous molecule ideally bears the cell-surface molecule to a high extent and/or specifically, when (neighboring) healthy cells in a patient are considered. Thus, the epitope on the targeted cell-surface molecule is ideally unique to the targeted diseases cells, and is at least specifically present and exposed at the surface of the targeted cells. Binding of the first and second proteinaceous molecules is followed by endocytosis of the complexes of the first proteinaceous molecule and the target cell-surface molecule and the second proteinaceous molecule and the target cell-surface molecule. Since the first and second proteinaceous molecules have to enter the same target cell through binding interaction with the very same cell-surface molecules, accumulation of a therapeutically active amount of first and second proteinaceous molecules inside the target cells is only possible and occurring if expression levels of the targeted cell-surface molecule is above a certain minimal expression threshold. At the same time, the fact that the effector moiety bound to the second proteinaceous molecule is only capable of exerting its intracellular (e.g. cytotoxic or gene silencing) activity in the presence of the first proteinaceous molecule bearing the covalently bound saponin, when both the first and second proteinaceous molecules were capable to enter the target cell in sufficient amounts by binding to sufficiently exposed and expressed cell-surface molecule, also provides a safeguard against negative and undesired side effects of the effector moiety towards e.g. healthy cells and healthy tissue not meant to be targeted and affected by the effector moiety, when expression of the targeted cell-surface molecule is sufficiently low at the healthy cells. That is to say, low expression of the cell-surface molecule bound by the

binding site of the first and second proteinaceous molecules, does not allow entrance of both the first and second proteinaceous molecules to amounts that would in concert result in endosomal escape of the effector moiety under influence of the saponin bound to the first proteinaceous molecule. Since the ADC can be used at lower dose compared to when the first proteinaceous molecule was not added to the therapeutic regimen, ADC entrance in healthy cells to low extent already bears a lower risk for occurrence of unwanted side effects when for example the targeting and killing of target diseased cells such as tumor cells and auto-immune cells is considered.

[0078] An aspect of the invention relates to a therapeutic combination as defined in the claims for use in the treatment or prevention of a cancer in a human subject, wherein the therapeutic combination comprises: (a) the first pharmaceutical composition of the invention; and (b) the second pharmaceutical composition of the invention, wherein the cell-surface molecule is expressed on a tumor cell surface, and preferably the cell-surface molecule is a tumor cell-specific surface molecule, and wherein preferably the epitope is a tumor-cell specific epitope.

[0079] An aspect of the invention relates to the therapeutic combination for use of the invention, wherein the second pharmaceutical composition of the invention and the first pharmaceutical composition are administered to the patient in need thereof.

[0080] Before the present invention one of the major hurdles of introducing ADCs on the market was the small therapeutic window: a therapeutically effective dose of an ADC is accompanied with (unacceptable) side effects, hampering development and implication in treatment of patients with the ADCs. By the application of the first proteinaceous molecule of the invention it has now become possible to guide one or multiple glycoside molecules (saponin) to a (target) cell, together with the ADC carrying a payload or together with a (monoclonal) antibody conjugated with an oligonucleotide such as a BNA according to the invention. In particular, it was not possible to specifically guide an effector moiety of a second proteinaceous molecule and a (predefined, controllable) particular number or range of glycoside molecules (saponins) per effector moiety at the same time to the cytosol of cells, such as via the endocytic pathway of a cell.

[0081] A solution provided for by the invention comprises the covalent binding of at least one saponin to the first proteinaceous molecule. A further solution provided for by the invention comprises (first) polymerizing the glycoside molecules (saponins) using an oligomeric or polymeric scaffold, and providing the first proteinaceous molecule with a cluster of covalently bound saponins, enabling re-monomerization of the one or more saponins at the intracellular site where the mode of action of the saponin is desired, e.g. after endocytosis. "Polymerizes" in this context means the reversible and/or irreversible multiple conjugation of saponin molecules to the first proteinaceous molecule, either via linker, or directly or via a polymeric or oligomeric structure to form a scaffold or the reversible and/or irreversible multiple conjugation of (modified) saponins thereby forming a polymeric or oligomeric structure to form a scaffold. "Re-monomerization" in this context means the cleavage of the saponins from the first proteinaceous molecule, from the linker linking the saponin(s) to the first proteinaceous molecule or from the scaffold, for example after endocytosis, and regaining the (native) chemical state of the unbound saponins, which unbound saponins may or may not comprise additional chemical groups such as a chemical group for linking the saponin to a linker, an amino-acid residue of the first proteinaceous molecule or to the scaffold, and/or a (chemical) linker bound to a chemical group of the saponin such as an aldehyde group or carboxylic acid group. Due to the complex chemistry of the saponins for example the 'polymerization' of saponins at a scaffold or other linking linker and their 're-monomerization' at a desired location such as intracellularly e.g. after endocytosis, was a challenging task. In particular, the chemical reactions used for providing the linkers and the scaffold comprising covalently linked glycosides for covalent binding to the first proteinaceous molecule, e.g. triterpenoid saponins (polymerization of the glycosides), normally occur in water-free organic solvents, but saponins and for example biocompatible polymers applied as a scaffold for bearing bound saponins, are water-soluble molecules. The chemical properties of the unmodified saponin further prohibited polymerization by itself and, one other

possible solution, to bind multiple saponins (directly) to the effector molecule was estimated not to be very promising, as an effector molecule (drug, toxin, polypeptide or polynucleotide) does typically not provide sufficient binding sites and because the coupling product would become quite heterogeneous and/or coupling biologically active molecules such as a saponin and e.g. a peptide, a toxin, a nucleic acid together bears the risk for influencing and hampering the activity of one or even both molecules bound together in such saponin-comprising conjugate. Further, there was a considerable risk that the effector moiety comprised by the second proteinaceous molecule loses its function after coupling of a saponin to the e.g. ADC or antibody-oligonucleotide conjugate (AOC). Embodiments of the present invention solves at least one of these drawbacks.

[0082] In accordance with the invention, the binding site of the first proteinaceous molecule and the second proteinaceous molecule comprises or consists of an immunoglobulin or a binding fragment or binding domain of said immunoglobulin, such as any one or more of an antibody, an IgG, a molecule comprising or consisting of a Vhh domain or Vh domain, a Fab, an scFv, an Fv, a dAb, an F(ab)2, Fcab fragment, and/or comprises or consists of at least one ligand, the ligand for binding to a cell-surface molecule such as EGF or a cytokine. The inventors established that such immunoglobulins, domains thereof, ligands, etc., are particularly suitable for application as the binding site of the first (and second) proteinaceous molecule comprising the binding site. For example, antibodies and binding domains of antibodies are suitable for targeting an epitope on the exposed surface of a selected cell-surface molecule, resulting in targeting the first (and second) proteinaceous molecule to cells expressing the cell-surface molecule and having said cell-surface molecule on their cell surface. Similarly, ligands such as EGF, targeting the EGFR on target cells, are suitable for application as the binding site in the first and second proteinaceous molecules. Preferred are binding sites for the epitope, which are specific for the binding to the cell-surface molecule. Binding sites based on antibodies or domains or binding fragments thereof for example provide for such desired specificity for a selected epitope on a selected cell-surface molecule of a selected cell for targeting such as a diseased cell, a tumor cell, an auto-immune cell, etc. Therefore, binding sites based on antibodies or binding molecules (fragments, domains) are preferred for the first and second proteinaceous molecules.

[0083] In an embodiment of the invention, the tumor-cell surface molecule is a cell-surface receptor specifically present at a tumor cell, and, preferably, the epitope is a tumor-cell specific epitope. Tables A2, A3 and A4 list such preferred examples of cell-surface molecules comprising the epitope for the binding site of the first and second proteinaceous molecule. When the cell-surface molecule is specifically expressed on the target cell, and when the epitope on the cell-surface molecule, to which the binding site can bind, is specifically present in the cell-surface molecule, specific targeting of the first and second proteinaceous molecule to the desired cell such as a tumor cell exposing the tumor-cell surface molecule, is facilitated, whereas other cells such as healthy cells, which do not express the cell-surface molecule or do express the cell-surface molecule to a lower extent, are not targeted by the first and second proteinaceous molecule or are targeted to a lower extent.

[0084] In accordance with the invention, the at least one saponin is a triterpenoid saponin and/or a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and optionally comprises a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin.

[0085] In an embodiment of the invention, the at least one saponin is a single specific saponin or is a mixture of two or more different saponins, such as one or more of the saponins in Table A1 or Scheme I, SO1861, SA1657, GE1741, SA1641, QS-21, QS-21A, QS-21 A-api, QS-21 A-xyl, QS-21B, QS-21 B-api, QS-21 B-xyl, QS-7-xyl, QS-7-api, QS-17-api, QS-17-xyl, QS1861, QS1862, Quillajasaponin, Saponinum album, QS-18, Quil-A, Gyp1, gypsoside A, AG1,AG2, SO1542, SO1584, SO1658, SO1674, SO1832, or any of their stereomers and/or any combinations thereof, preferably the saponin is SO1861 and/or GE1741 and/or SA1641 and/or QS-21 and/or saponin with a quillaic acid aglycon core, a Gal-(1→2)-[Xyl-(1→3)]-GlcA carbohydrate substituent at

the C-3beta-OH group and a Glc-(1→3)-Xyl-(1→4)-Rha-(1→2)-[Xyl-(1→3)-4-OAc-Qui-(1→4)]-Fuc carbohydrate substituent at the C-28-OH group, and/or is 3-O-beta-D-galactopyranosyl-(1→2)-[beta-D-xylopyranosyl-(1->3)]-beta-D-glucuronopyranosyl quillaic acid 28-O-beta-D-glucopyranosyl-(1→3)-beta-D-xylopyranosyl-(1→4)- alpha-L-rhamnopyranosyl-(1→2)-[beta-D-xylopyranosyl-(1→3)-4-OAc-beta-D-quinovopyranosyl-(1→4)]-beta-D-fucopyranoside, preferably the saponin is SO1861 and/or QS-21.

[0086] In an embodiment of the invention, the at least one saponin is a bisdesmosidic saponin having a molecular mass of at least 1.500 Dalton and comprising an oleanan-type triterpene containing an aldehyde group at the C-23 position and optionally a hydroxyl group at the C-16 position, with a first branched carbohydrate side chain at the C-3 position which first branched carbohydrate side chain optionally contains glucuronic acid, wherein the saponin contains an ester group with a second branched carbohydrate side chain at the C-28 position which second branched carbohydrate chain preferably comprises at least four carbohydrate units, optionally containing at least one acetyl residue such as two acetyl residues and/or optionally at least one deoxy carbohydrate and/or a quinovose and/or a glucose and/or 4-methoxycinnamic acid and/or optionally comprising 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid and/or optionally comprising 5-O-[5-O-Rha-(1→2)-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid bound to a carbohydrate via an ester bond, or wherein the at least one saponin is QS-21 or any one or more of QS-21A, QS-21 A-api, QS-21 A-xyl, QS-21B, QS-21 B-api, QS-21 B-xyl, QS-7-xyl, QS-7-api, QS-17-api, QS-17-xyl, QS-18, QS1861, protonated QS-1861 (QS1862), Quil-A.

[0087] In an embodiment of the invention, the at least one saponin is a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23, wherein the saponin is covalently coupled to the first proteinaceous molecule, preferably covalently coupled to an amino-acid residue of the first proteinaceous molecule, via an aldehyde function in the saponin, preferably said aldehyde function in position C-23, preferably via at least one linker, and/or via at least one cleavable linker, wherein the amino-acid residue preferably is selected from cysteine and lysine.

[0088] Table A1 and Scheme I and the above embodiments summarize a series of saponins that have been identified for their endosomal escape enhancing activity when contacted to mammalian cells, in particular human tumor cells, in free form together with a second molecule (e.g. an effector moiety or effector molecule, such as a toxin, an oligonucleotide). Indeed, in cell-based bioassays using human tumor cells it was established for the saponins tabulated in Table A1 and those in Scheme I and in the various embodiments of the invention described herein, that under influence of these saponins, when bound to the first proteinaceous molecule, a second molecule (effector moiety) such as a nucleic acid and/or a toxin such as a protein toxin (e.g. one or more of the protein toxins listed in Table A5), bound to the second proteinaceous molecule, is delivered into the cytosol with increased efficiency and/or efficacy, presumably through intracellular release from the (late) endosomes and lysosomes. That is to say, endosomal and/or lysosomal escape of such second molecules (effector moieties bound to a second proteinaceous molecule of the invention), e.g. nucleic acids and/or toxins, is less efficient in the absence of the saponin.

[0089] Surprisingly, the inventors now demonstrate that a water-soluble saponin fraction from Quillaja saponaria, comprising QS-21 and its family members QS-21A, QS-21 A-api, QS-21 A-xyl, QS-21B, QS-21 B-api, QS-21 B-xyl, QS-7-xyl, QS-7-api, QS-17-api, QS-17-xyl, QS1861, QS1862, QS-18 and Quil-A, also exhibits the ability to potentiate a biological effect in vitro of e.g. a nucleic acid bound to a monoclonal antibody or a protein toxin bound to a monoclonal antibody (examples of a second proteinaceous molecule of the invention comprising covalently bound oligonucleotide or payload such as a (protein) toxin), when administered to tumor cells of a mammalian species (human) in the form of a covalent conjugate comprising a monoclonal antibody (first proteinaceous molecule of the invention), together with the second proteinaceous molecule comprising the effector moiety (the aforementioned second proteinaceous molecule) and the at least one glycoside such as the QS-21 and its family member saponins encompassed by such QS-21 preparation (e.g. water soluble fraction of Quillaja saponaria), comprised by the first proteinaceous molecule as a covalent

conjugate, wherein the effector molecule and the glycoside, e.g. saponin fraction of Quillaja saponaria, QS-21, SO1861, SA1641, GE1741, are covalently bound to for example the proteinaceous molecules directly or via a linker or via a polymeric or oligomeric scaffold, either directly or via at least one linker. Without wishing to be bound by any theory, the observed stimulation or potentiation of for example antisense BNA mediated reduction of tumor-cell HSP27 expression (HSP27 gene silencing) in the presence of saponins derived from Quillaja saponaria in vitro may (also) relate to activation of the inflammasome in the tumor cell by the saponins, for example resulting in tumor cell pyroptosis. The inventors established that second proteinaceous molecules conjugated to for example antisense BNA or dianthin or saporin, exerted any anti-tumor cell activity in vitro at all or improved anti-tumor cell activity when contacted with cells in bio-based cell assays, when in the presence of the first proteinaceous molecule of the invention, comprising the saponin, and targeted to the same binding site on the same tumor cells as the cell surface molecule targeted by the second proteinaceous molecule.

[0090] QS-21, and also the water-soluble saponins fraction comprising QS-21 from Quillaja saponaria is already for a long time known and previously intensively applied for its immune-potentiating abilities, e.g. as an adjuvant in e.g. sub-unit vaccines. For example, QS-21 is applied in two phase III clinical trials with human patients, who were vaccinated with a sub-unit vaccine mixed with an adjuvant comprising QS-21 (Glaxo-Smith-Kline, MAGRIT trial, DERMA study), wherein the sub-unit was MAGE-A3 protein, which is specifically expressed and presented by tumor cells. The anti-tumor vaccinations, potentiated with QS-21, aimed for extension of disease-free survival of the cancer patients (melanoma; non-small cell lung cancer). In addition, QS-21 has been tested as an adjuvant in clinical trials for developing anti-cancer vaccine treatment, for vaccines for HIV-1 infection, in development of a vaccine against hepatitis B, and for anti-malaria vaccine development using QS-21 comprising adjuvants AS01 and AS02 of Glaxo-Smith-Kline. Previous studies revealed an immune response elicited against MAGE-A3 peptides presented at the cancer cell surface, under influence of the QS-21 saponin comprising adjuvant (AS15; GSK). To the surprise of the inventors, the saponin fraction of Quillaja saponaria, and thus likely QS-21 (as part of the water soluble saponin fraction of Quillaja saponaria) potentiates the anti-tumor cell activity of e.g. a payload such as a protein toxin (dianthin), bound to the second proteinaceous molecule (e.g. the ligand EGF).

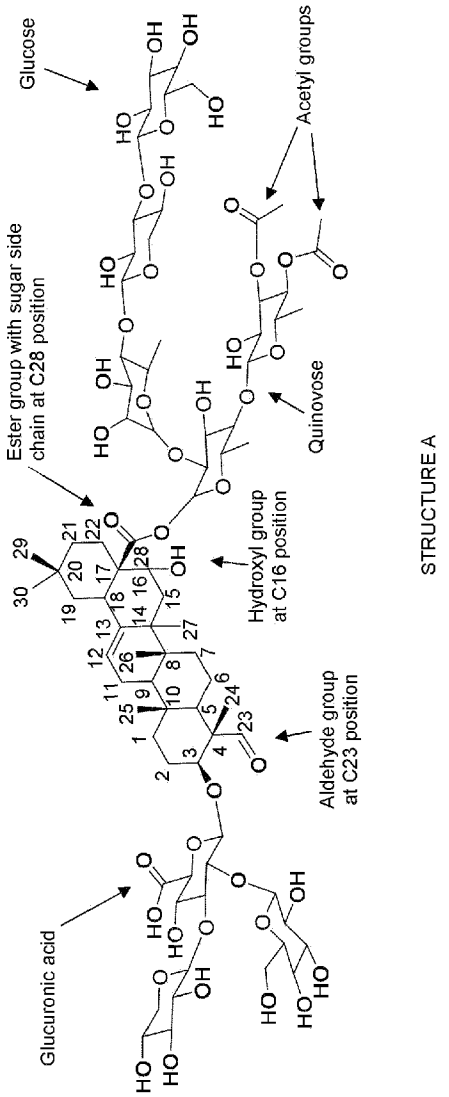
[0091] The inventors show that a tumor-cell targeting monoclonal antibody provided with covalently coupled antisense BNA such as BNA(HSP27), and contacted with the tumor cells together with a first proteinaceous molecule of the invention with covalently coupled saponin (e.g. SO1861, QS-21), both the BNA and the saponin coupled to the respective antibody (e.g. cetuximab) of the first and second proteinaceous molecule via a cleavable bond is capable of silencing HSP27 in vivo in tumors, compared to control and compared to ADC only, without presence of first proteinaceous molecule with coupled saponin. Co-administering an ADC or an antibody-oligonucleotide conjugate (AOC), such as an antibody-BNA conjugate, with a first proteinaceous molecule with a saponin thus endows the ADC or AOC with anti-tumor cell activity not seen with only the ADC or only the AOC at the same dose. Noteworthy, the AOC (the second proteinaceous molecule) and the monoclonal antibody with covalently coupled saponin (first proteinaceous molecule) increase HSP27 expression in tumor cells, when administered to tumor-bearing mice separately in separate groups of mice, compared to a control group (vehicle administered, only). Only co-administration of the AOC comprising the effector moiety of the invention (second proteinaceous molecule) and the first proteinaceous molecule with covalently coupled saponin, displays reduced HSP27 expression when compared to controls. The antisense BNA (HSP27) was BNA with oligo nucleic acid sequence 5'-GGCacagccagtgGCG-3' according to Zhang et al. (2011) [Y Zhang, Z Qu, S Kim, V Shi, B Liao1, P Kraft, R Bandaru, Y Wu, LM Greenberger and ID Horak, Down-modulation of cancer targets using locked nucleic acid (LNA)-based antisense oligonucleotides without transfection, Gene Therapy (2011) 18, 326-333]. Noteworthy, to the best of the knowledge of the inventors, BNA is designed for application as a free nucleic acid. The inventors are now the first to demonstrate that the antisense BNA can be covalently coupled through a (non-)cleavable linker with a ligand or an antibody, in a way that gene-silencing activity is retained in vitro and more importantly in vivo in the tumor cells of a tumor-bearing animal. This approach of providing BNA based AOCs opens new ways to administer targeted BNA to

human (cancer) patients in need thereof.

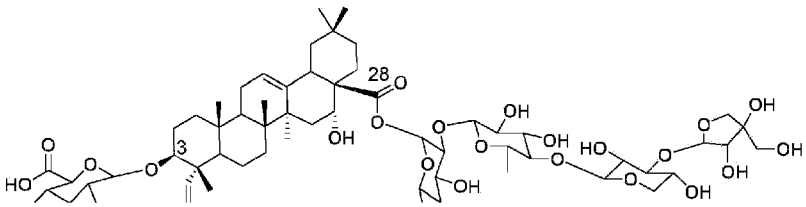
[0092] The inventors demonstrate here that covalently coupling saponins such as saponins in the water-soluble fraction of Quillaja saponaria, QS-21, SA1641, SO1861, Table A1, Scheme I, to a first proteinaceous molecule, such as via a tri-functional linker, e.g. the tri-functional linker of Scheme II and Structure B, or via an oligomeric or polymeric structure of a scaffold comprising covalently bound saponins, results in improved cell toxicity exerted by the effector moiety such as a toxin, comprised by the second proteinaceous molecule, under influence of the covalently coupled saponin in the first proteinaceous molecule.

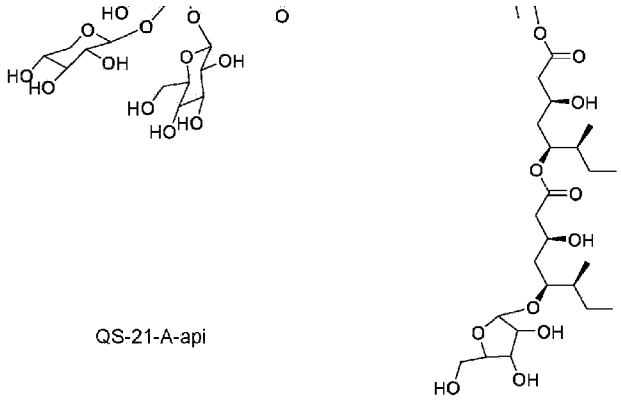
[0093] In an embodiment, the first proteinaceous molecule comprising a saponin comprises one or several or all of the indicated structural features of the saponin of Structure A in Scheme I, and/or a saponin selected from any one or more of the further saponins in Scheme I:

SCHEME I

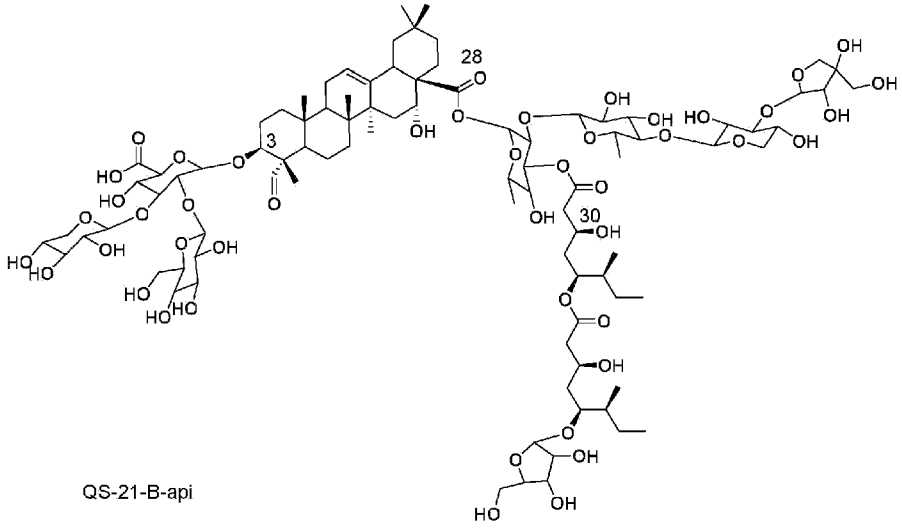


SCHEME I (continued)



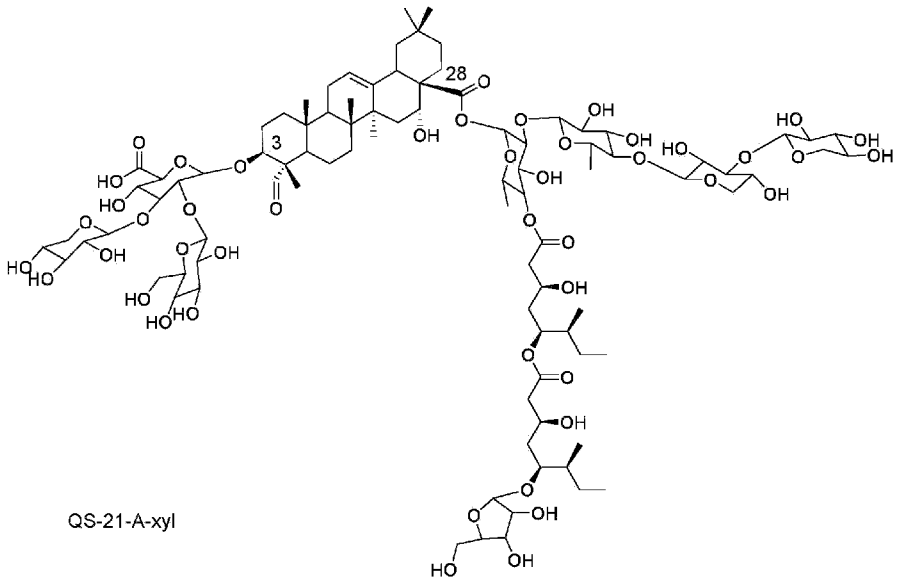


QS-21-A-api

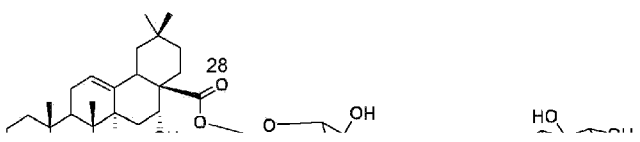


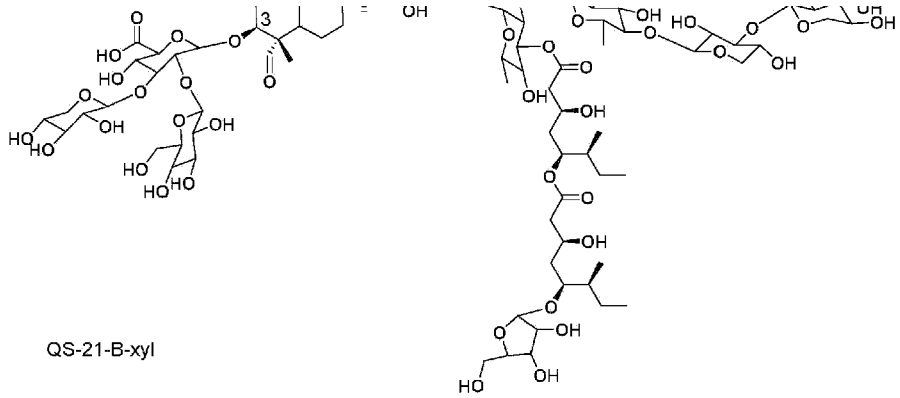
QS-21-B-api

SCHEME I (continued)



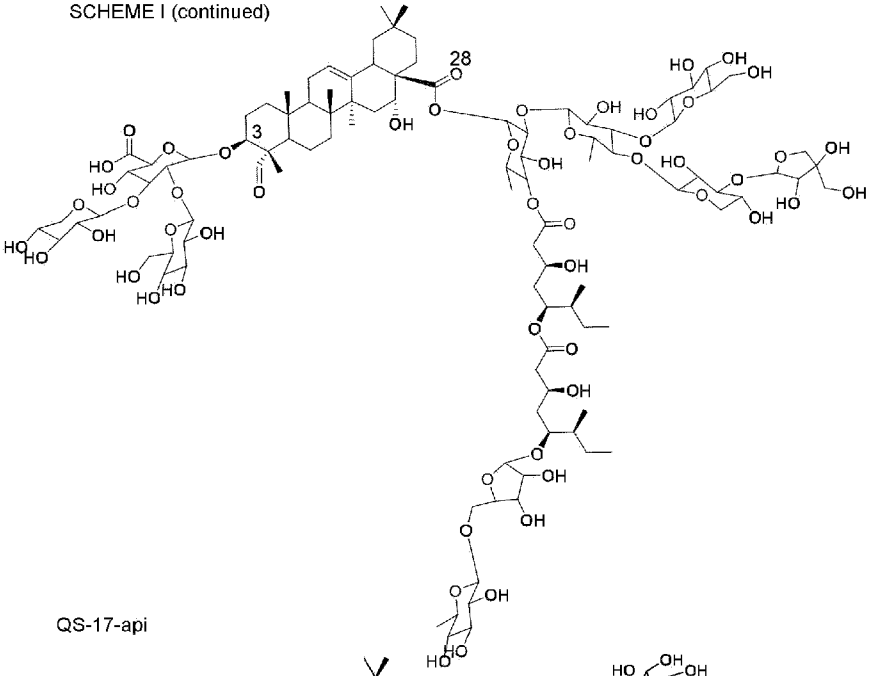
QS-21-A-xyl



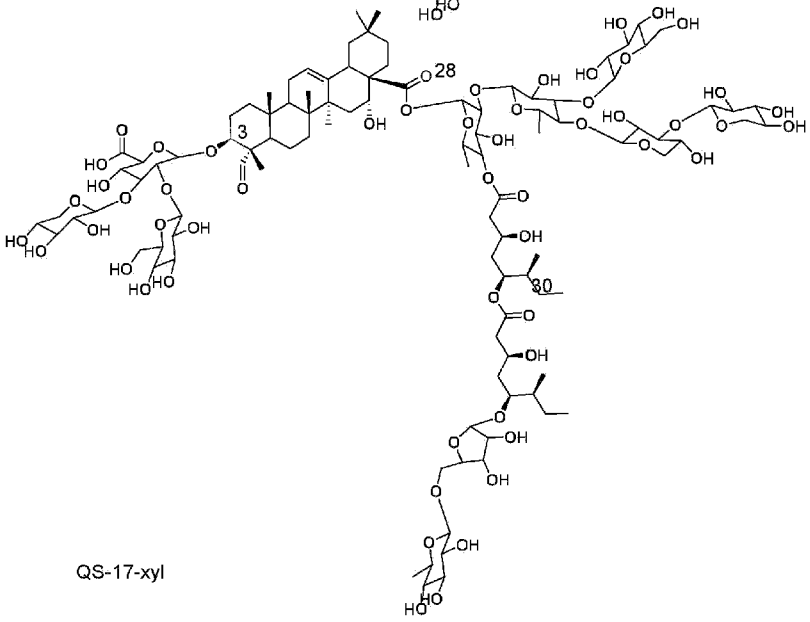


QS-21-B-xyl

SCHEME I (continued)

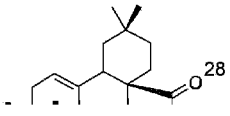


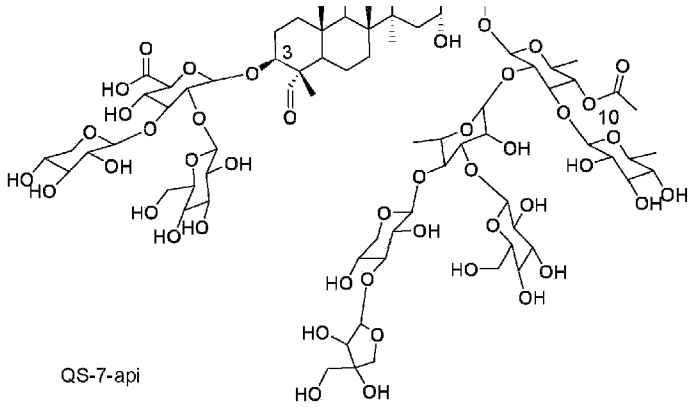
QS-17-api



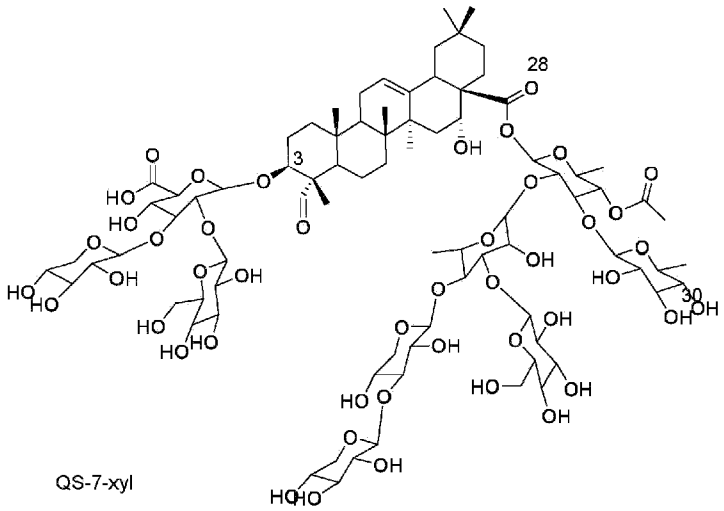
QS-17-xyl

SCHEME I (continued)





QS-7-api



QS-7-xyl

According to the invention, a glycoside, such as a saponin according to the invention, bound to the first proteinaceous molecule of the invention, which has the 'ideal' structure for the purpose of enhancing endosomal escape of an effector molecule bound to the second proteinaceous molecule of the invention is a bisdesmosidic saponin according to Structure A of Scheme I, having a molecular mass of at least 1.500 Dalton and comprising an oleanan-type triterpene containing an aldehyde group at the C-23 position and optionally a hydroxyl group at the C-16 position, with a first branched carbohydrate side chain at the C-3 position which first branched carbohydrate side chain optionally contains glucuronic acid, wherein the saponin contains an ester group with a second branched carbohydrate side chain at the C-28 position which second branched carbohydrate chain preferably comprises at least four carbohydrate units, optionally containing at least one acetyl residue such as two acetyl residues and/or optionally comprising deoxy carbohydrates and/or optionally comprising quinovose and/or optionally comprising glucose and/or optionally comprising 4-methoxycinnamic acid and/or optionally comprising 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid and/or optionally comprising 5-O-[5-O-Rha-(1→2)-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid bound to a carbohydrate via an ester bond.

[0094] SO1861 is different from the *"ideal structure"* displayed in Scheme I, Structure A, only in having only one acetyl residue at the quinovose and having an additional xylose. The "ideal structure" of a saponin for enhancing endosomal escape of an effector molecule or effector moiety, is a saponin which preferably has the Structure A of Scheme I, and saponins which display the endosomal escape enhancing activity have one or more of the structural features displayed in Structure A of Scheme I. Without wishing to be bound by any theory, the inventors belief that the Structure A of Scheme I represents an "ideal saponin" (and not a minimum requirement saponin) for endosomal escape enhancing activity, which means that not all of the structures can or must be present in each saponin with at least sufficient endosomal escape enhancing activity to promote

accumulation of the effector moiety in the cytosol, and which means that some saponins might have other structure elements such as acyl chains, and/or for yet other saponins that display endosomal escape enhancing activity, the sugars can be different than the sugars displayed in Scheme I. For example, the QS-21 saponin and some of the saponins in the water soluble fraction of Quillaja saponaria (Quillaja saponins; Quil-A) differ in the carbohydrate modification at C-28 when the ideal structure of Structure A in Scheme I is considered: presence of an acyl chain in QS-21 for example. In the water soluble fraction of Quillaja saponaria, saponins such as QS-7, QS1862, are similar to the ideal Structure A, and are similar to SO1861.

[0095] In an embodiment of the invention, the aldehyde function in position C-23 of the at least one saponin is covalently coupled to linker N-ε-maleimidocaproic acid hydrazide, which linker is covalently coupled via a thio-ether bond to a sulfhydryl group in the first proteinaceous molecule, such as a sulfhydryl group of a cysteine.

[0096] In an embodiment of the invention, the at least one saponin is a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position C-23 and comprises a glucuronic acid function in a carbohydrate substituent at the C-3beta-OH group of the saponin, wherein the saponin is covalently coupled to an amino-acid residue of the first proteinaceous molecule via the glucuronic acid function in the carbohydrate substituent at the C-3beta-OH group of the saponin, preferably via at least one linker, wherein the amino-acid residue preferably is selected from cysteine and lysine.

[0097] In an embodiment of the invention, the glucuronic acid function in the carbohydrate substituent at the C-3beta-OH group of the at least one saponin is covalently coupled to linker 1-[Bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate, which linker is covalently coupled via an amide bond to an amine group in the first proteinaceous molecule, such as an amine group of a lysine or an N-terminus of the first proteinaceous molecule.

[0098] The inventors established that covalent coupling, preferably via cleavable bonds or linkers, of the saponin to the first proteinaceous molecule, according to any of the embodiments here above, provides efficient and cell-targeted potentiation of the activity of an effector moiety bound to the second proteinaceous molecule, wherein the first and second proteinaceous molecules comprise the same binding site. Coupling saponin to a cysteine side chain or a lysine side chain of the first proteinaceous molecule such as a monoclonal antibody, directly or via a linker proved to be a beneficial way of specific and efficient delivery of effector-moiety potentiating activity inside the target cell, when also the effector moiety is delivered in the same target cell by using the second proteinaceous molecule comprising the same binding site as the first proteinaceous molecule.

[0099] To explain the invention in more detail, the process of cellular uptake of substances (although the inventors do not wish to be bound by any theory) and the used terminology in this invention is described. The uptake of extracellular substances into a cell by vesicle budding is called endocytosis. Said vesicle budding can be characterized by (1) receptor-dependent ligand uptake mediated by the cytosolic protein clathrin, (2) lipid-raft uptake mediated by the cholesterol-binding protein caveolin, (3) unspecific fluid uptake (pinocytosis), or (4) unspecific particle uptake (phagocytosis). All types of endocytosis run into the following cellular processes of vesicle transport and substance sorting called the endocytic pathways. The endocytic pathways are complex and not fully understood. Without wishing to be bound by any theory, organelles may be formed *de novo* and mature into the next organelle along the endocytic pathway. It is however, now hypothesized that the endocytic pathways involve stable compartments that are connected by vesicular traffic. A compartment is a complex, multifunctional membrane organelle that is specialized for a particular set of essential functions for the cell. Vesicles are considered to be transient organelles, simpler in composition, and are defined as membrane-enclosed containers that form *de novo* by budding from a preexisting compartment. In contrast to compartments, vesicles can undergo maturation, which is a physiologically irreversible series of biochemical changes. Early endosomes and late endosomes represent stable compartments in the endocytic pathway while primary endocytic vesicles, phagosomes, multivesicular bodies (also called endosome carrier vesicles),

secretory granules, and even lysosomes represent vesicles. The endocytic vesicle, which arises at the plasma membrane, most prominently from clathrin-coated pits, first fuses with the early endosome, which is a major sorting compartment of approximately pH 6.5. A large part of the cargo and membranes internalized are recycled back to the plasma membrane through recycling vesicles (recycling pathway). Components that should be degraded are transported to the acidic late endosome (pH lower than 6) via multivesicular bodies. Lysosomes are vesicles that can store mature lysosomal enzymes and deliver them to a late endosomal compartment when needed. The resulting organelle is called the hybrid organelle or endolysosome. Lysosomes bud off the hybrid organelle in a process referred to as lysosome reformation. Late endosomes, lysosomes, and hybrid organelles are extremely dynamic organelles, and distinction between them is often difficult. Degradation of an endocytosed molecule occurs inside an endolysosome or lysosome. Endosomal escape is the active or passive release of a substance from the inner lumen of any kind of compartment or vesicle from the endocytic pathway, preferably from clathrin-mediated endocytosis, or recycling pathway into the cytosol. Endosomal escape thus includes but is not limited to release from endosomes, endolysosomes or lysosomes, including their intermediate and hybrid organelles.

[0100] Unless specifically indicated otherwise and in particular when relating to the endosomal escape mechanism of the glycoside molecule such as the saponin of the invention, whenever the word "endosome" or "endosomal escape" is used herein, it also includes the endolysosome and lysosome, and escape from the endolysosome and lysosome, respectively. After entering the cytosol, said substance might move to other cell units such as the nucleus.

[0101] In formal terms, a glycoside is any molecule in which a sugar group is bound through its anomeric carbon to another group via a glycosidic bond. Glycoside molecules, such as saponins, in the context of the invention are such molecules that are further able to enhance the effect of an effector moiety, without wishing to be bound by any theory, in particular by facilitating the endosomal escape of the effector moiety. Without wishing to be bound by any theory, the glycoside molecules (saponins, such as those listed in Table A1) interact with the membranes of compartments and vesicles of the endocytic and recycling pathway and make them leaky for said effector moieties resulting in augmented endosomal escape. With the term "the scaffold is able to augment endosomal escape of the effector moiety" is meant that the at least one saponin (glycoside molecule), which is coupled to the polymeric or oligomeric structure of the scaffold, is able to enhance endosomal escape of an effector moiety when both molecules are within an endosome, e.g. a late endosome, optionally and preferably after the at least one glycoside such as a saponin is released from the first proteinaceous molecule such as from a linker or polymeric or oligomeric structure comprised by said first proteinaceous molecule, e.g., by cleavage of a cleavable bond between the at least one glycoside (saponin) and the first proteinaceous molecule (for example via a polymeric or oligomeric structure of a scaffold and/or via a linker). Even though a bond between the at least one glycoside such as a saponin according to the invention and the first proteinaceous molecule, optionally via a linker or a scaffold, may be a "stable bond", that does not mean that such bond cannot be cleaved in the endosomes by, e.g., enzymes. For instance, the glycoside or saponin, optionally together with a linker or a part of the oligomeric or polymeric structure of a scaffold, may be cleaved off from the remaining linker fragment or oligomeric or polymeric structure. It could, for instance be that a protease cuts a (proteinaceous) linker or proteinaceous polymeric structure, e.g., albumin, thereby releasing the at least one glycoside, saponin. It is, however, preferred that the glycoside molecule (preferably saponin) is released in an active form, preferably in the original form that it had before it was (prepared to be) coupled to the first proteinaceous molecule optionally via a linker and/or an oligomeric or polymeric scaffold; thus the glycoside (saponin) has its natural structure after such cleavage or the glycoside (saponin) has (part of) a chemical group or linker bound thereto, after such cleavage, while glycoside biological activity (saponin biological activity), e.g. endosomal/lysosomal escape enhancing activity towards an effector moiety present in the same endosome or lysosome, is maintained or restored upon said cleavage of the bond between the glycoside (saponin) and the carrier molecule, i.e. the first proteinaceous molecule optionally comprising a linker and/or a scaffold of the invention. With regard to the present invention the term "stable" with respect to bonds between e.g. saponins and amino-acid residues of the first proteinaceous

molecule, a linker, a polymeric or oligomeric structures (of the scaffold), ligands, (monoclonal) immunoglobulins or binding domains or -fragments thereof, and/or effectors (effector moieties, effector molecules), is meant that the bond is not readily broken or at least not designed to be readily broken by, e.g., pH differences, salt concentrations, or UV-light, reductive conditions. With regard to the present invention the term "cleavable" with respect to bonds between e.g. saponins and the first proteinaceous molecule, linkers, amino-acid residues, polymeric or oligomeric structures of the scaffold, ligands, antibodies and/or effectors, is meant that the bond is designed to be readily broken by, e.g., pH differences, salt concentrations, under reductive conditions. The skilled person is well aware of such cleavable bonds and how to prepare them.

[0102] In an embodiment of the invention, the epitope is an epitope on a tumor-cell receptor, preferably a tumor-cell specific epitope, and the receptor is preferably a tumor-cell specific receptor, more preferably a receptor selected from CD71, CA125, EpCAM(17-1A), CD52, CEA, CD44v6, FAP, EGF-IR, integrin, syndecan-1, vascular integrin alpha-V beta-3, HER2, EGFR, CD20, CD22, Folate receptor 1, CD146, CD56, CD19, CD138, CD27L receptor, PSMA, CanAg, integrin-alphaV, CA6, CD33, mesothelin, Cripto, CD3, CD30, CD239, CD70, CD123, CD352, DLL3, CD25, ephrinA4, MUC1, Trop2, CEACAM5, CEACAM6, HER3, CD74, PTK7, Notch3, FGF2, C4.4A, FLT3, CD38, FGFR3, CD7, PD-L1, CTLA4, CD52, PDGFRA, VEGFR1, VEGFR2, most preferably selected from CD71, HER2 and EGFR.

[0103] In an embodiment of the invention, the tumor-cell receptor, preferably the tumor-cell specific receptor, is internalized by the tumor cell after binding to the first proteinaceous molecule of the invention and/or after binding to the second proteinaceous molecule of the invention, and, preferably, binding of the first proteinaceous molecule and/or the second proteinaceous molecule to the tumor-cell receptor, such as the tumor-cell specific receptor, is followed by tumor-cell receptor-mediated internalization, e.g. via endocytosis, of a complex of the first proteinaceous molecule and the tumor-cell receptor and of a complex of the second proteinaceous molecule and the tumor-cell receptor.

[0104] In an embodiment of the invention, the binding site of the first proteinaceous molecule and the second proteinaceous molecule comprises a monoclonal antibody or at least one of a cell-surface molecule binding domain and/or -fragment thereof, and, preferably, comprises or consists of any one of cetuximab, daratumumab, gemtuzumab, trastuzumab, panitumumab, brentuximab, inotuzumab, moxetumomab, polatuzumab, obinutuzumab, OKT-9 anti-CD71 monoclonal antibody of the IgG type, pertuzumab, rituximab, ofatumumab, Herceptin, alemtuzumab, pinatuzumab, OKT-10 anti-CD38 monoclonal antibody, an antibody of Table A2 or Table A3 or Table A4, preferably cetuximab or trastuzumab or OKT-9, or at least one cell-surface molecule binding fragment and/or -domain thereof, with the proviso that the first binding site is the same as the second binding site.

[0105] An effector molecule, or effector moiety, in the context of this invention is any substance that affects the metabolism of a cell by interaction with an intracellular effector molecule target, wherein this effector molecule target is any molecule or structure inside cells excluding the lumen of compartments and vesicles of the endocytic and recycling pathway but including the membranes of these compartments and vesicles. Said structures inside cells thus include the nucleus, mitochondria, chloroplasts, endoplasmic reticulum, Golgi apparatus, other transport vesicles, the inner part of the plasma membrane and the cytosol. Cytosolic delivery of an effector moiety in the context of the invention preferably means that the effector moiety is able to escape the endosome (and/or lysosome), which, as defined previously, also includes escaping the endolysosome and the lysosome, and is preferably able to reach the effector moiety target as described herein. The invention also encompasses a new type of molecule, referred to as scaffold that serves to bring both an effector moiety and at least one glycoside molecule such as a saponin of the invention in an endosome at the same time in a pre-defined ratio, when the effector moiety is comprised by the second proteinaceous molecule of the invention and the saponin is comprised by the first proteinaceous molecule. Within the context of the present invention, the polymeric or oligomeric structure of the scaffold is a structurally ordered formation such as a polymer, oligomer, dendrimer, dendronized polymer, or dendronized oligomer or it is an assembled polymeric

structure such as a hydrogel, microgel, nanogel, stabilized polymeric micelle or liposome, but excludes structures that are composed of non-covalent assemblies of monomers such as cholesterol/phospholipid mixtures. The terms "polymer, oligomer, dendrimer, dendronized polymer, or dendronized oligomer" have their ordinary meaning. In particular a polymer is a substance which has a molecular structure built up chiefly or completely from a large number of equal or similar units bonded together and an oligomer is a polymer whose molecules consist of relatively few repeating units. There is no consensus about one specific cut-off for "many" and "a few" as used in the above definition of polymer and oligomer, respectively. However, as the scaffold may comprise a polymeric or an oligomeric structure, or both, the full range of numbers of similar units bonded together applies to such structure. i.e. from 2 monomeric units to 100 monomeric units, 1000 monomeric units, and more. A structure comprising 5 or less, for instance maybe called an oligomeric structure, whereas a structure comprising 50 monomeric units maybe called a polymeric structure. A structure of 10 monomeric units maybe called either oligomeric or polymeric. A scaffold as defined herein, further comprises at least one glycoside molecule such as a saponin of the invention. A scaffold preferably includes a polymeric or oligomeric structure such as poly- or oligo(amines), e.g., polyethylenimine and poly(amidoamine), and biocompatible structures such as polyethylene glycol, poly- or oligo(esters), such as poly(lactids), poly(lactams), polylactide-co-glycolide copolymers, and poly(dextrin), poly- or oligosaccharides, such as cyclodextrin or polydextrose, and poly- or oligoamino acids, such as poly-lysine or a peptide or a protein, or DNA oligo- or polymers. An assembled polymeric structure as defined herein comprises at least one scaffold and, optionally, other individual polymeric or oligomeric structures. Other individual polymeric or oligomeric structures of said assembly may be (a) scaffolds (thus comprising at least one glycoside molecule such as a saponin of the invention), (b) functionalized scaffolds (thus comprising at least one glycoside molecule such as a saponin, and a ligand, antibody, etc. as the first proteinaceous molecule, (c) polymeric or oligomeric structures without a glycoside molecule such as a saponin of the invention (See Table A1 for example), without a ligand, antibody, etc., as the first proteinaceous molecule. A functionalized assembled polymeric structure is an assembled polymeric structure that contains (a) at least one functionalized scaffold or (b) at least one scaffold and at least one polymeric structure comprising at least one ligand, antibody, etc. as the first proteinaceous molecule. Polymeric or oligomeric structures within an assembled polymeric structure that do not comprise any of the above mentioned molecules (i.e. no glycosides such as saponins, no first proteinaceous molecule such as ligands, antibodies) are in particular added as structural components of the assembled structures, which help to build up or to stabilize the assembled structure ("glue-like").Without wishing to be bound by any theory, the acidic environment seems to be a prerequisite for the synergistic action between glycoside (saponin) and effector moiety.

[0106] Whether or not a first proteinaceous molecule comprising saponins, either or not further comprising one or more (cleavable) linkers and/or optionally a scaffold, is able to disturb the acidic environment and inhibit the endosomal escape function of the at least one glycoside (saponin) can be easily determined with an assay as described in Example 3 and as known in the art. The inhibition is described as "fold amount increases of glycoside necessary to induced 50% cell killing". It is preferred that the scaffold does not lead to an increase that is at least the increase in glycoside molecules (saponins) necessary to obtain 50% cell killing observed when using Chloroquine as a positive control. Alternatively, and preferably, the first proteinaceous molecule comprising saponins, either or not further comprising one or more (cleavable) linkers and/or optionally a scaffold does not lead to an at least 4-fold increase of glycoside molecules to induce 50% cell killing, more preferably does not lead to an at least 2-fold increase. The fold increase is to be measured in assay, essentially as described in Example 4, wherein Chloroquine, as a positive control, induces a 2-fold increase in glycoside amount, preferably saponin amount wherein the saponin is any one or more of the saponins of the invention (see Table A1, Scheme I, previous embodiments) to observe 50% cell killing.

[0107] With the term "improving or enhancing an effect of an effector moiety" is meant that the glycoside molecule, preferably a saponin of the invention, increases the functional efficacy of that effector moiety (e.g. the therapeutic index of a toxin or a drug or an oligonucleotide such as a BNA; the metabolic efficacy of a modifier in biotechnological processes; the transfection efficacy of genes in cell culture research experiments),

preferably by enabling or improving its target engagement. Acceleration, prolongation, or enhancement of antigen-specific immune responses are preferably not included. Therapeutic efficacy includes but is not limited to a stronger therapeutic effect, preferably with lower dosing and/or with less side effects. "Improving an effect of an effector moiety" can also mean that an effector moiety, which could not be used because of lack of effect (and was e.g. not known as being an effector moiety), becomes effective when used in combination with the present invention. Any other effect, which is beneficial or desired and can be attributed to the combination of effector moiety and the second proteinaceous molecule, as provided by the invention is considered to be "an improved effect". In an embodiment, the scaffold comprising bound saponin(s) and comprised by the first proteinaceous molecule enhances an effect of the effector moiety comprised by the second proteinaceous molecule which effect is intended and/or desired. In case of a first proteinaceous molecule comprising saponin bound to a proteinaceous scaffold, the proteinaceous polymeric structure of the scaffold as such may have, for instance, an effect on colloid osmotic pressure in the blood stream. If such effect is not the intended or desired effect of such a functionalized scaffold comprised by the first proteinaceous molecule, the proteinaceous structure of the scaffold is not an effector moiety as defined in the invention. Or, for instance in case of a DNA- or RNA-based scaffold carrying bound saponins and comprised by the first proteinaceous molecule, parts of that DNA or RNA may have an (unintended) function, e.g., by interfering with expression. If such interference is not the intended or desired effect of the ultimate functionalized scaffold, the DNA- or RNA polymeric structure of the scaffold is not the effector moiety as defined in the invention.

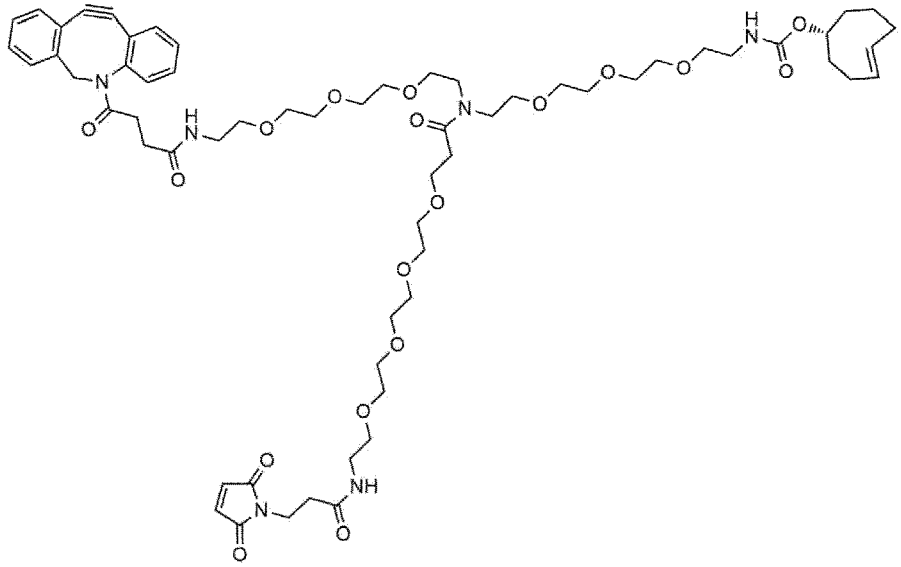
[0108] A number of preferred features can be formulated for endosomal escape enhancers comprised by the first proteinaceous molecule, i.e. a glycoside, preferably a saponin according to the invention: (1) they are preferably not toxic and do not invoke an immune response, (2) they preferably do not mediate the cytosolic uptake of the effector moiety into off-target cells, (3) their presence at the site of action is preferably synchronized with the presence of the effector moiety, (4) they are preferably biodegradable or excretable, and (5) they preferably do not substantially interfere with biological processes of the organism unrelated to the biological activity of the effector molecule with which the endosomal escape enhancer is combined with, e.g. interact with hormones. Examples of glycoside molecules such as saponins of the invention that fulfill the before mentioned criteria, at least to some extent, are bisdesmosidic triterpenes, preferably bisdesmosidic triterpene saponins, such as SO1861, SA1641, QS-21, GE1741, and the saponins in Table A1, Scheme I.

[0109] In an embodiment, the at least one saponin is covalently bound to the first proteinaceous molecule via a linker and/or via a polymeric or oligomeric structure, via a non-cleavable bond or via a cleavable bond, wherein preferably said cleavable bond is subject to cleavage under acidic conditions, reductive conditions, enzymatic conditions or light-induced conditions, more preferably the cleavable bond is a hydrazone bond or a hydrazide bond subject to cleavage under acidic conditions, and/or is a bond susceptible to proteolysis, for example proteolysis by Cathepsin B, and/or is a bond susceptible for cleavage under reductive conditions such as a disulphide bond. According to the invention, typically the saponin is a saponin listed in Table A1, Scheme I. It has been proven beneficial for the activity of the saponin, e.g. the endosomal escape enhancing activity inside cells when the entry into the cell and the accumulation inside the cytosol of an effector moiety covalently coupled to the second proteinaceous molecule, is considered, when the saponin is covalently coupled to the first proteinaceous molecule involving a hydrazone bond, and/or a hydrazide bond, and/or a disulphide bond. Such bond types readily cleave under the acidic conditions inside (late) endosomes and lysosomes of mammalian cells, e.g. human cells, and/or under the reductive conditions. Alternatively, the inventors also demonstrate that covalent coupling of saponin to the first proteinaceous molecule via a bond that is not readily cleavable under the physiological conditions inside cells, e.g. (late) endosomes, lysosomes, cytosol, is also beneficial to the potentiating activity of the saponin on the biological effect of e.g. an effector moiety such as a nucleic acid (e.g. BNA silencing HSP27) and a proteinaceous toxin such as saporin. Throughout the application, including the claims, the term 'cleavable linker', 'cleavable bond', etc., is also referred to as 'labile linker' ('L') and 'labile bond', for example in the context of cleavage of such a bond or linker in the (late) endosome and/or lysosome when a conjugate of the invention, e.g. a first proteinaceous molecule optionally comprising a scaffold with saponins coupled to the first proteinaceous molecule through a linker and/or via the

scaffold via hydrazone bonds or disulphide bonds, is referred to. For example, Figure 1 shows the in vivo HSP27 gene silencing in human tumors in mice. The tumor-bearing mice were treated with a first proteinaceous molecule consisting of monoclonal antibody with saponin bound thereto via a labile linker (hydrazone bond) according to the invention, whereas the second proteinaceous molecule comprised bound antisense BNA for silencing the HSP27 gene in the tumor cells, covalently coupled to the monoclonal antibody (same type as the first monoclonal antibody) via a a disulphide bond. That is to say, without wishing to be bound by any theory, the hydrazone bond and the disulphide bond are cleaved in the (late) endosomes and/or lysosomes of the targeted tumor cells that express the epitope on the targeted cell-surface molecule, here the EGFR, at the cell surface, once the therapeutic combination of the invention is internalized by e.g. endocytosis. Cleavage of the bonds likely contributes to the endosomal escape enhancing activity of the saponin when the entry of the BNA from the endosome and/or lysosome into the cytosol is considered, although such cleavage is not a necessity for observing the gene silencing effect of the combination of the cetuximab-SO1861 conjugate and the cetuximab-BNA conjugate of the invention.

[0110] The skilled person will appreciate that a tri-functional linker is a scaffold of the invention suitable for covalently coupling one, two or three saponin moieties. For the tri-functional linker covalent coupling of one or two saponin moieties is preferred. The second and/or third binding site is for example suitable for covalent coupling a proteinaceous ligand such as the first proteinaceous molecule. Typical proteinaceous ligands are EGF for targeting (tumor) cells expressing EGFR at the cell surface, and cytokines for targeting tumor cells or autoimmune cells. Moreover, the second or third binding site of the tri-functional linker is suitable for covalent coupling of an immunoglobulin such as a monoclonal antibody, i.e. the first proteinaceous molecule for binding to a cell surface molecule such as a tumor cell surface molecule, preferably a tumor-cell specific molecule, more preferably a tumor cell receptor that is specifically (over-)expressed at the surface of the tumor cell. Similarly, the immunoglobulin, or any fragment(s) and/or domain(s) thereof which encompass the binding specificity of the immunoglobulin, is suitable for binding to a cell surface molecule such as a receptor, expressed at the surface of an autoimmune cell. Thus, in an embodiment, the first proteinaceous molecule comprises the tri-functional linker, said linker comprises or consists of a covalently bound saponin, e.g. QS-21, SO1861, and the covalently bound binding site such as a cell targeting moiety such as a ligand or an antibody for (specific) binding to a tumor cell, an auto-immune cell, a diseased cell, an aberrant cell, a non-healthy cell, a B-cell disease.

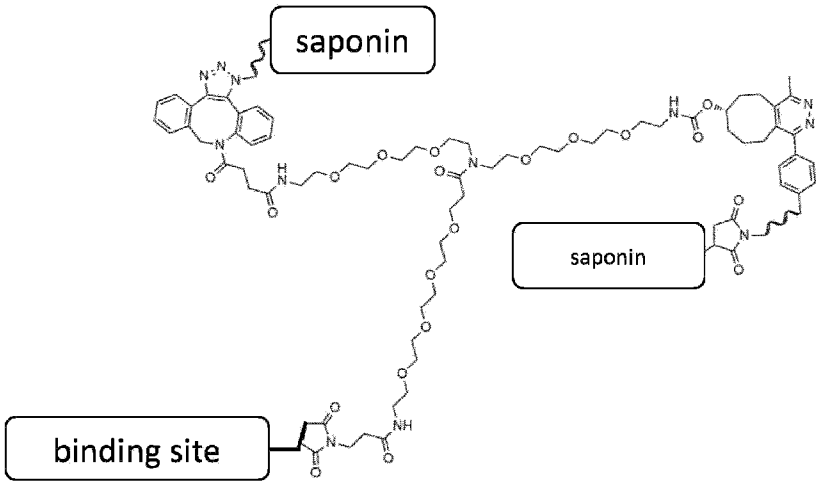
[0111] In an embodiment, the first proteinaceous molecule comprises the oligomeric tri-functional linker as the scaffold core structure, according to Scheme II:



SCHEME II,

wherein the saponins are covalently bound to the tri-functional linker scaffold via labile, cleavable hydrazone

linkers (acid sensitive) and/or via a maleimide comprising bond, whereas the binding of the scaffold to the binding site such as an antibody is established via labile, cleavable hydrazone linkers (acid sensitive) and/or via a maleimide comprising bond with cysteines in the binding site, therewith forming Structure B:



Structure B,

such as 1, 2, 3 or 4 cysteines such that 1-4 scaffolds are covalently bound to a single e.g. antibody such as a monoclonal antibody.

[0112] In an embodiment,the linkage between saponin and the first proteinaceous molecule preferably occurs via an acid-labile bond that is stable at pH 7.4 and, preferably releases the saponin below pH 6.5, more preferably between pH 6.5 and 5.0. This is, e.g., realized via an imine formed by an amino group of a linker linking the saponin and the first proteinaceous molecule and the aldehyde group of the saponin. Other chemical bonds that fulfill the pH-condition can also be used for aldehyde coupling, e.g. particular hydrazones or acetals, requiring hydrazides and hydroxyl groups as the functional group of the linker, respectively. If the bond is a cleavable bond, a saponin is preferably attached to the polymeric or oligomeric structure of a scaffold via an aldehyde function or via one of the carboxyl groups in saponin, more preferably through the aldehyde function, preferably an aldehyde function in position 23. Alternatively, a saponin is preferably attached to the first proteinaceous molecule via the polymeric or oligomeric structure of the scaffold via a linker that connects the polymeric or oligomeric structure of the scaffold either via the aldehyde function or via the carboxylic acid function of the glycoside molecule, i.e. the saponin.

[0113] In an embodiment, the at least one saponin is bound to the first proteinaceous molecule via a stable bond. In a more preferred embodiment, the stable bond between saponin first proteinaceous molecule preferably occurs via an amide coupling or amine formation. This is, e.g., realized via carbodiimide mediated amide bond formation by an amino group of a polymeric or oligomeric scaffold structure linking the saponin and the first proteinaceous molecule together, and the activated glucuronic acid group of the saponin. Chemical bonds that fulfill the stable bond definition can also be used for aldehyde coupling, e.g. particular amines derived after reductive amination, requiring primary amino groups as the functional group of a polymeric or oligomeric structure of a scaffold or a linker. If the bond is a stable bond, the saponin is preferably attached to a linker or a scaffold via one of the carboxyl groups of the saponin, the linker or scaffold further linked to the first proteinaceous molecule.

[0114] In an embodiment, the saponin is coupled to the binding site via a scaffold according to the invention, wherein the chemical group for covalently coupling of the scaffold to the binding site is a click chemistry group.

[0115] In an embodiment, the saponin is coupled to the binding site via a scaffold according to the invention, wherein the click chemistry group is a tetrazine, an azide, an alkene or an alkyne, or a cyclic derivative of any of these groups, preferably an azide. A click chemistry group is a functional chemical group suitable for click

chemistry, which is defined as a reaction that is modular, wide in scope, gives very high yields, generates only inoffensive byproducts, offers high selectivity, and high tolerance over different functional groups, and is stereospecific. The required process characteristics include simple reaction conditions, readily available starting materials and reagents, the use of no solvent or a solvent that is benign (such as water) or easily removed, and simple product isolation. The click chemistry group for coupling the saponin to the binding site in the first proteinaceous molecule optionally via a scaffold or a linker, is preferably a tetrazine, azide, alkene, or alkyne, or reactive derivates of them such as methyl-tetrazine or maleimide (alkene), more preferably an alkyne, or a cyclic derivative of these groups, such as cyclooctyne (e.g. aza-dibenzocyclooctyne, difluorocyclooctyne, bicyclo[6.1.0]non-4-yne, dibenzocyclooctyne).

[0116] A first proteinaceous molecule, according to the invention, thus comprises at least one saponin. With "at least one" in this context is meant that the first proteinaceous molecule comprises one saponin molecule but may also comprise a couple (e.g. two, three or four) of saponins or a multitude (e.g. 10, 20 or 100) of saponins. Depending on the application, the first proteinaceous molecule may comprise a covalently bound scaffold with covalently bound saponins, wherein the scaffold may be designed such that it comprises a defined number of saponins. Preferably, a first proteinaceous molecule according to the invention comprises a defined number or range of saponins, rather than a random number. This is especially advantageous for drug development in relation to marketing authorization. A defined number in this respect means that a first proteinaceous molecule preferably comprises a previously defined number of saponins. This is, e.g., achieved by designing a scaffold comprising a polymeric structure with a certain number of possible moieties for the saponin(s) to attach. Under ideal circumstances, all of these moieties are coupled to a saponin and the scaffold than comprises the prior defined number of saponins. It is envisaged to offer a standard set of scaffolds, comprising, e.g., two, four, eight, sixteen, thirty-two, sixty-four, etc., saponins so that the optimal number can be easily tested by the user according to his needs. An embodiment is the first proteinaceous molecule of the invention comprising the scaffold of the invention, wherein the saponin is present in a defined range as, e.g., under non-ideal circumstances, not all moieties present in a polymeric structure bind a saponin. Such ranges may for instance be 2 - 4 saponin molecules per scaffold, 3 - 6 saponin molecules per scaffold, 4 - 8 saponin molecules per scaffold, 6 - 8 saponin molecules per scaffold, 6 - 12 saponin molecules per scaffold and so on. In such case, a first proteinaceous molecule comprising a scaffold according to the invention thus comprises 2, 3 or 4 saponins if the range is defined as 2 - 4.

[0117] The scaffold is fundamentally independent of the type of saponin covalently bound to the scaffold, the scaffold subsequently (in sequential order) covalently coupled to the first proteinaceous molecule. Thus, first proteinaceous molecule comprising the scaffold is the basis product for a new platform technology. Since the at least one covalently bound saponin mediates intracellular delivery of the effector moiety bound to the second proteinaceous molecule, the scaffold technology according to the invention is the first system known that mediates controlled intracellular effector moiety delivery by saponins. The scaffold provides an optimized and functionally active unit that can be linked to the saponin(s) and to the binding site comprised by the first proteinaceous molecule, e.g. a ligand, an antibody, etc., at a single and defined position.

[0118] In an embodiment, the first proteinaceous molecule comprises a scaffold according to the invention, wherein the number of monomers of the polymeric or oligomeric structure is an exactly defined number or range. Preferably, the polymeric or oligomeric structure comprises structures such as poly(amines), e.g., polyethylenimine and poly(amidoamine), or structures such as polyethylene glycol, poly(esters), such as poly(lactides), poly(lactams), polylactide-co-glycolide copolymers, poly(dextrin), or a peptide or a protein, or structures such as natural and/or artificial polyamino acids, e.g. poly-lysine, DNA polymers, stabilized RNA polymers or PNA (peptide nucleic acid) polymers, either appearing as linear, branched or cyclic polymer, oligomer, dendrimer, dendron, dendronized polymer, dendronized oligomer or assemblies of these structures, either sheer or mixed. Preferably, the polymeric or oligomeric structures are biocompatible, wherein biocompatible means that the polymeric or oligomeric structure does not show substantial acute or chronic toxicity in organisms and can be either excreted as it is or fully degraded to excretable and/or physiological

compounds by the body's metabolism. Assemblies can be built up by covalent cross-linking or non-covalent bonds and/or attraction. They can therefore also form nanogels, microgels, or hydrogels, or they can be attached to carriers such as inorganic nanoparticles, colloids, liposomes, micelles or particle-like structures comprising cholesterol and/or phospholipids. Said polymeric or oligomeric structures preferably bear an exactly defined number or range of coupling moieties for the coupling of glycoside molecules (and/or effector molecules and/or carrier molecules such as a ligand, monoclonal antibody or a fragment thereof). Preferably at least 50%, more preferably at least 75%, more preferably at least 85%, more preferably at least 90%, more preferably at least 95%, more preferably at least 98%, more preferably at least 99%, most preferably 100% of the exactly defined number or range of coupling moieties in the polymeric or oligomeric structure is occupied by a glycoside molecule in a scaffold according to the invention.

[0119] Preferably, a dendron is a branched, clearly defined tree-like polymer with a single chemically addressable group at the origin of the tree, called the focal point. A dendrimer is a connection of two or more dendrons at their focal point. A dendronized polymer is a connection of the focal point of one or more dendrons to a polymer. In a preferred embodiment, a scaffold according to the invention is provided, wherein the polymeric or oligomeric structure comprises a linear, branched or cyclic polymer, oligomer, dendrimer, dendron, dendronized polymer, dendronized oligomer or assemblies of these structures, either sheer or mixed, wherein assemblies can be built up by covalent cross-linking or non-covalent attraction and can form nanogels, microgels, or hydrogels, and wherein, preferably, the polymer is a derivative of a poly(amine), e.g., polyethylenimine and poly(amidoamine), and structures such as polyethylene glycol, poly(esters), such as poly(lactids), poly(lactams), polylactide-co-glycolide copolymers, and poly(dextrin), and structures such as natural and/or artificial polyamino acids such as poly-lysine, or a peptide or a protein or DNA polymers, stabilized RNA polymers or PNA (peptide nucleic acid) polymers. Preferably, the polymeric or oligomeric structures are biocompatible.

[0120] An embodiment is the therapeutic combination for use according to the invention, wherein the first proteinaceous molecule comprises more than one covalently bound saponin, preferably 2, 3, 4, 5, 6, 8, 10, 16, 32, 64, 128 or 1-100 saponins, or any number of saponins therein between, such as 7, 9, 12 saponins.

[0121] An embodiment is the therapeutic combination for use according to the invention, wherein the second proteinaceous molecule comprises the effector moiety, wherein said effector moiety is a pharmaceutically active substance, such as a toxin, a drug, a polypeptide and/or a polynucleotide. An embodiment is the second proteinaceous molecule of the invention wherein the effector moiety is a toxin, a micro RNA, or a polynucleotide encoding a protein. Typically, the first and second proteinaceous molecule comprises the binding site for targeting the saponin or the effector moiety respectively, to e.g. a tumor cell or an auto-immune cell or a cell related to a B-cell disease. Preferably the binding site of the first and second proteinaceous molecule comprises an immunoglobulin or at least one or more binding domains and/or binding fragments thereof, such immunoglobulin preferably selected from any of the monoclonal antibodies of any of Tables A2-A4, such as cetuximab, OKT-9, trastuzumab.

[0122] A pharmaceutically active substance in this invention is an effector moiety that is used to achieve a beneficial outcome in an organism, preferably a vertebrate, more preferably a human being such as a cancer patient or an auto-immune patient. Benefit includes diagnosis, prognosis, treatment, cure and/or prevention of diseases and/or symptoms. The pharmaceutically active substance may also lead to undesired harmful side effects. In this case, pros and cons must be weighed to decide whether the pharmaceutically active substance is suitable in the particular case. If the effect of the pharmaceutically active substance inside a cell is predominantly beneficial for the whole organism, the cell is called a target cell. If the effect inside a cell is predominantly harmful for the whole organism, the cell is called an off-target cell. In artificial systems such as cell cultures and bioreactors, target cells and off-target cells depend on the purpose and are defined by the user.

[0123] An effector moiety that is a polypeptide may be, e.g., a polypeptide that recover a lost function, such as for instance enzyme replacement, gene regulating functions, or a toxin.

[0124] An embodiment is the therapeutic combination for use of the invention, wherein the effector moiety comprised by the second proteinaceous molecule comprises or consists of any one or more of an oligonucleotide, a nucleic acid and a xeno nucleic acid, preferably selected from any one or more of a vector, a gene, a cell suicide inducing transgene, deoxyribonucleic acid (DNA), ribonucleic acid (RNA), anti-sense oligonucleotide (ASO, AON), short interfering RNA (siRNA), microRNA (miRNA), DNA aptamer, RNA aptamer, mRNA, mini-circle DNA, peptide nucleic acid (PNA), phosphoramidate morpholino oligomer (PMO), locked nucleic acid (LNA), bridged nucleic acid (BNA), 2'-deoxy-2'-fluoroarabino nucleic acid (FANA), 2'-O-methoxyethyl-RNA (MOE), 2'-O,4'-aminoethylene bridged nucleic acid, 3'-fluoro hexitol nucleic acid (FHNA), a plasmid, glycol nucleic acid (GNA) and threose nucleic acid (TNA), or a derivative thereof, more preferably a BNA, for example a BNA for silencing HSP27 protein expression.

[0125] An embodiment is the therapeutic combination for use of the invention, wherein the effector moiety comprised by the second proteinaceous molecule comprises or consists of at least one proteinaceous molecule, preferably selected from any one or more of a peptide, a protein, an enzyme such as urease and Cre-recombinase, a proteinaceous toxin, a ribosome-inactivating protein, a protein toxin selected from Table A5 and/or a bacterial toxin, a plant toxin, more preferably selected from any one or more of a viral toxin such as apoptin; a bacterial toxin such as Shiga toxin, Shiga-like toxin, Pseudomonas aeruginosa exotoxin (PE) or exotoxin A of PE, full-length or truncated diphtheria toxin (DT), cholera toxin; a fungal toxin such as alpha-sarcin; a plant toxin including ribosome-inactivating proteins and the A chain of type 2 ribosome-inactivating proteins such as dianthin e.g. dianthin-30 or dianthin-32, saporin e.g. saporin-S3 or saporin-S6, bouganin or de-immunized derivative debouganin of bouganin, shiga-like toxin A, pokeweed antiviral protein, ricin, ricin A chain, modeccin, modeccin A chain, abrin, abrin A chain, volkensin, volkensin A chain, viscumin, viscumin A chain; or an animal or human toxin such as frog RNase, or granzyme B or angiogenin from humans, or any fragment or derivative thereof; preferably the protein toxin is dianthin and/or saporin.

[0126] An embodiment is the therapeutic combination for use of the invention, wherein the effector moiety comprised by the second proteinaceous molecule comprises or consists of at least one payload, preferably selected from any one or more of a toxin targeting ribosomes, a toxin targeting elongation factors, a toxin targeting tubulin, a toxin targeting DNA and a toxin targeting RNA, more preferably any one or more of emtansine, pasudotox, maytansinoid derivative DM1, maytansinoid derivative DM4, monomethyl auristatin E (MMAE, vedotin), monomethyl auristatin F (MMAF, mafodotin), a Calicheamicin, N-Acetyl-γ-calicheamicin, a pyrrolobenzodiazepine (PBD) dimer, a benzodiazepine, a CC-1065 analogue, a duocarmycin, Doxorubicin, paclitaxel, docetaxel, cisplatin, cyclophosphamide, etoposide, docetaxel, 5-fluorouracyl (5-FU), mitoxantrone, a tubulysin, an indolinobenzodiazepine, AZ13599185, a cryptophycin, rhizoxin, methotrexate, an anthracycline, a camptothecin analogue, SN-38, DX-8951f, exatecan mesylate, truncated form of *Pseudomonas aeruginosa* exotoxin (PE38), a Duocarmycin derivative, an amanitin, α-amanitin, a spliceostatin, a thailanstatin, ozogamicin, tesirine, Amberstatin269 and soravtansine, or a derivative thereof.

[0127] An embodiment is the therapeutic combination for use of the invention, wherein the second proteinaceous molecule comprises or consists of any one of Gemtuzumab ozogamicin, Brentuximab vedotin, Trastuzumab emtansine, Inotuzumab ozogamicin, Moxetumomab pasudotox and Polatuzumab vedotin and an antibody-drug conjugate of Table A2 and Table A3, or at least one tumor-cell receptor binding-domain thereof and/or at least one tumor-cell receptor binding-fragment thereof, wherein said domain(s) or fragment(s) comprise(s) the effector moiety and are preferably (a) tumor-cell specific receptor binding-domain(s) and/or (a) tumor-cell specific receptor binding-fragment(s).

[0128] Synchronization is the missing link between a very successful delivery strategy for mice and its application in humans. Indeed, the inventors established in a series of in vivo mouse tumor models that

separately administering to the mice a dose of free saponin and a dose of ADC (second proteinaceous molecule according to the invention) did not result in any desired anti-tumor activity such as delayed tumor growth, tumor regression, diminished and slower tumor growth, compared to control animals not treated with the ADC and free saponin. The free saponin was administered using various routes of administration and using various time points of administering the free saponin compared to the moment of administering the ADC (administering free saponin before, during and after administering the ADC). The ADC tested in in vivo tumor models was cetuximab-dianthin (with free SO1861), or trastuzumab-saporin (with free SO1861). Varying the dose of free saponin did not provide for an efficacious anti-tumor activity. The ADCs referred to were administered at a dose that in itself did not inflict any beneficial anti-tumor effect on the tumor-bearing animals. Surprisingly, the inventors now established that beneficial anti-tumor activity in various in vitro mammalian cell-based bioassays and/or in various in vivo animal tumor models can be achieved by treating the animals with conjugates according to the invention, comprising a scaffold according to the invention. The scaffold for example being a tri-functional linker with a covalently bound saponin (e.g. SO1861, QS-21) via a cleavable or non-cleavable linkage, and with a covalently bound effector moiety (e.g. dianthin, silencing BNA (HSP27) via a non-cleavable bond or a cleavable bond, and with a covalently bound monoclonal antibody such as cetuximab, trastuzumab, OKT-9, or the scaffold being a dendron, such as a dendron to which for example four moieties can bind such as four saponin molecules, or a dendron for binding for example two saponins and two effector molecules, the dendron comprising a chemical group for (covalent) coupling to a ligand or an antibody or fragment or domain thereof. Reference is made to the Examples section, exemplifying various of these scaffolds according to the invention, showing in vivo and/or in vitro anti-tumor cell activity when cell toxicity exerted by e.g. a proteinaceous toxin is considered or when gene silencing in the tumor cell is considered.

[0129] Without wishing to be bound by any theory, in view of the failures observed when treatment of tumor-bearing animals with an ADC together with free saponin is considered, it is preferred to synchronize the presence of both, the at least one saponin, and the effector moiety, preferably a toxin, in compartments or vesicles of the endocytic pathway of the target cell, e.g. a tumor cell or an auto-immune cell. With ADC and free saponin, synchronizing the presence of the molecules in the late endosomes, in order to obtain the synergistic effects *in vivo* was not beneficially obtainable according to attempts of the inventors. In one aspect, the invention preferably solves at least the following problem with respect to combining the effector moiety comprised by the second proteinaceous molecule and the saponins comprised by the first proteinaceous molecule: without wishing to be bound by any theory the only reasonable chemical group within, e.g., the saponins that can be used for (covalent), in particular single and cleavable, retainable coupling is required for the endosomal escape activity. Known restrictions are most likely the reason why saponins have not been used in combination with pharmaceutically active substances in clinical investigations other than the application of saponins in vaccination regimes wherein the use of an immune-potentiating adjuvant substance was implied, although the striking endosomal escape enhancer effect of, e.g., saponins listed in Table A1 and Scheme I is known for more than 10 years. For example providing a first proteinaceous molecule of the invention with a covalently conjugated scaffold solves these difficulties, at least in part. Surprisingly, the saponins previously applied for their immune-potentiating activity in the vaccination context involving saponins as adjuvant component, are now also suitably for (covalent) coupling to the first proteinaceous molecule of the invention, for anti-tumor activity in vitro and in vivo.

[0130] An effector molecule useful in the present invention preferably relies on late endosomal escape for exerting its effect. Some effectors, such as, e.g., a pseudomonas exotoxin, are rerouted to other organelles prior to the "late endosomal stage" and, thus, would normally not benefit from coupling to the second proteinaceous molecule according to the present invention. However, such toxin may be adapted for use with the present invention, e.g., by deleting the signal peptide responsible rerouting. In particular toxins that are highly toxic and would require only one molecule to escape the endosomes to kill a cell maybe modified to be less potent. It is preferred to use a toxin that kills a cell if at least 2, more preferably at least 5, more preferably at least 10, more preferably at least 20, more preferably at least 50, most preferably at least 100 toxin molecules escape the endosome. It is further preferred that a second proteinaceous molecule of the invention

comprises a covalently conjugated functionalized scaffold, i.e. a scaffold comprising covalently bound effector moietie(s) for targeting the scaffold comprising the bound effector moietie(s) at a target cell such as a tumor cell or an auto-immune cell. Further, in order to reduce off-target toxicity, cell membrane non-permeable small molecule toxins are preferred effector molecules over cell membrane permeable toxins.

[0131] The term "ligand" as used in this invention has its ordinary meaning and preferably means a molecule or structure that is able to bind another molecule or structure on the cell surface of a target cell, wherein said molecule or structure on the cell surface can be endocytosed and is preferably absent or less prominent on off-target cells. Preferably, said molecule or structure on the cell surface is constitutively endocytosed. More preferably a ligand in this invention induces endocytosis of said molecule or structure on the cell surface of target cells after binding to said molecule or structure. This is for instance the case for Epidermal Growth Factor Receptor (EGFR), present on the surface of a variety of cancer cells. Examples of molecules or structures on the cell surface of target cells that are constitutively endocytosed, are for instance Claudin-1 or major histocompatibility complex class II glycoproteins. A ligand can, e.g., be an antibody, a growth factor or a cytokine. Combining in a carrier molecule a toxin with a ligand is one possibility to create a targeted toxin. A toxin that is only toxic in a target cell because it interferes with processes that occur in target cells only can also be seen as a targeted toxin (as in off-target cells it cannot exert its toxic action, e.g. apoptin). Preferably, a targeted toxin is a toxin that is combined with a ligand or e.g. a monoclonal antibody in order to be active in target cells and not in off-target cells (as it is only bound to and endocytosed by target cells). In a functionalized scaffold comprising a carrier molecule comprising a ligand and an effector molecule, the ligand or the monoclonal antibody guides the effector molecule and scaffold to the target cells. After internalization, the at least one glycoside, preferably a saponin, mediates the endosomal escape of the effector molecule. The saponin is typically a saponin listed in Table A1 and Scheme I, and preferably the saponin is SO1861 and/or QS-21, and/or SA1641 and/or GE1741.

[0132] Preferably, the effector moiety bound to the second proteinaceous molecule, which effect is enhanced by the saponins bound to the first proteinaceous molecule, detaches from the second proteinaceous molecule, e.g. an antibody, when endocytosed. This can be achieved by a cleavable bond that breaks, e.g. under acidic, reductive, enzymatic or light-induced conditions.

[0133] An embodiment is the therapeutic combination for use of the invention, wherein the first proteinaceous molecule comprises more than one covalently bound saponin, preferably 2, 3, 4, 5, 6, 8, 10, 16, 32, 64, 128 or 1-100 saponins, or any number of saponins therein between, such as 7, 9, 12 saponins.

[0134] An embodiment is the therapeutic combination for use of the invention, wherein the at least one saponin is covalently bound directly to an amino-acid residue of the first proteinaceous molecule, preferably to a cysteine and/or to a lysine, and/or is covalently bound via at least one linker and/or via at least one cleavable linker and/or via at least one oligomeric or polymeric scaffold, preferably 1-8 of such scaffolds or 2-4 of such scaffolds, wherein the at least one scaffold is optionally based on a dendron, wherein 1-32 saponins, such as 2, 3, 4, 5, 6, 8, 10, 16, 32 saponins, or any number of saponins therein between, such as 7, 9, 12 saponins, are covalently bound to the at least one scaffold.

[0135] An embodiment is the therapeutic combination for use of the invention, wherein the cleavable linker is subject to cleavage under acidic conditions, reductive conditions, enzymatic conditions or light-induced conditions, and preferably the cleavable linker comprises a cleavable bond selected from a hydrazone bond and a hydrazide bond subject to cleavage under acidic conditions, and/or a bond susceptible to proteolysis, for example proteolysis by Cathepsin B, and/or a bond susceptible for cleavage under reductive conditions such as a disulphide bond.

[0136] An embodiment is the therapeutic combination for use of the invention, wherein the cleavable linker is subject to cleavage *in vivo* under acidic conditions as present in endosomes and/or lysosomes of mammalian

cells, preferably human cells, preferably at pH 4.0 - 6.5, and more preferably at pH ≤ 5.5.

[0137] An embodiment is the therapeutic combination for use of the invention, wherein the oligomeric or polymeric scaffold comprises a polymeric or oligomeric structure and comprises at least one chemical group, the at least one chemical group for covalently coupling of the scaffold to the amino-acid residue of said first proteinaceous molecule.

[0138] An embodiment is the therapeutic combination for use of the invention, wherein the at least one saponin is covalently bound to the polymeric or oligomeric structure of the scaffold via a cleavable linker according to the invention.

[0139] An embodiment is the therapeutic combination for use of the invention, wherein the at least one saponin is covalently bound to the polymeric or oligomeric structure of the scaffold via any one or more of an imine bond, a hydrazone bond, a hydrazide bond, an oxime bond, a 1,3-dioxolane bond, a disulphide bond, a thio-ether bond, an amide bond, a peptide bond or an ester bond, preferably via at least one linker.

[0140] An embodiment is the therapeutic combination for use of the invention, wherein the at least one saponin comprises an aldehyde function in position C-23 and optionally a glucuronic acid function in the carbohydrate substituent at the C-3beta-OH group of the at least one saponin, which aldehyde function is involved in the covalent bonding to the polymeric or oligomeric structure of the scaffold, and/or, if present, the glucuronic acid function is involved in the covalent bonding to the polymeric or oligomeric structure of the scaffold, either via direct binding or via at least one linker.

[0141] An embodiment is the therapeutic combination for use of the invention, wherein the aldehyde function in position C-23 of the at least one saponin is covalently coupled to linker N-ε-maleimidocaproic acid hydrazide, which linker is covalently coupled via a thio-ether bond to a sulfhydryl group in the polymeric or oligomeric structure of the scaffold, such as a sulfhydryl group of a cysteine.

[0142] An embodiment is the therapeutic combination for use of the invention, wherein the glucuronic acid function in the carbohydrate substituent at the C-3beta-OH group of the at least one saponin is covalently coupled to linker 1-[Bis(dimethylamino)methylene]-1H-1 ,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate, which linker is covalently coupled via an amide bond to an amine group in the polymeric or oligomeric structure of the scaffold, such as an amine group of a lysine or an N-terminus of a proteinaceous molecule.

[0143] An embodiment is the therapeutic combination for use of the invention, wherein the at least one chemical group of the scaffold, for covalently coupling of the oligomeric or polymeric scaffold to the amino-acid residue of said first proteinaceous molecule, is a click chemistry group, preferably selected from a tetrazine, an azide, an alkene or an alkyne, or a cyclic derivative of these groups, more preferably the click chemistry group is an azide.

[0144] An embodiment is the therapeutic combination for use of the invention, wherein the polymeric or oligomeric structure of the scaffold comprises a linear, branched and/or cyclic polymer, oligomer, dendrimer, dendron, dendronized polymer, dendronized oligomer, a DNA, a polypeptide, poly-lysine, a poly-ethylene glycol, or an assembly of these polymeric or oligomeric structures which assembly is preferably built up by covalent cross-linking.

[0145] An embodiment is the therapeutic combination for use of the invention, wherein the first pharmaceutical composition and the second pharmaceutical composition are administered to the patient in need thereof.

[0146] As said before, the at least one saponin that is comprised by the first proteinaceous molecule according

to the invention increases the efficacy of at least current and new effector moieties as defined in this invention. Potential side-effects will be decreased due to lowering of dosing of the effector moiety comprised by the second proteinaceous molecule, without lowering the efficacy. Especially cancer medicines, and in particular the classical chemotherapy medicaments, are notorious for their side effects. Because of targeting and synchronization in time and place of both the pharmaceutically active substance comprised by the second proteinaceous molecule and the saponin comprised by the first proteinaceous molecule, since the first and second proteinaceous molecule bear the same binding site for the same epitope on the same cell-surface molecule, a therapeutic combination according to the invention is especially valuable for use as a medicament, in particular for use in a method of treating cancer. The invention thus provides a therapeutic combination according to the invention for use in a method of treating cancer. The invention also provides a therapeutic combination according to the invention for use in a method of treating acquired or hereditary disorders, in particular monogenic deficiency disorders. Thus, an aspect of the invention relates to a therapeutic combination according to the invention, wherein the second proteinaceous molecule comprises a covalently bound effector moiety, for use in a method for the treatment of a cancer or an auto-immune disease.

[0147] An aspect of the invention relates to the therapeutic combination for use according to the invention, wherein the first and second proteinaceous molecule comprise an antibody as the binding site for targeting a tumor cell, for use in the treatment or prophylaxis of a cancer, wherein the therapeutic combination is administered to a human subject in need thereof.

[0148] An aspect of the invention relates to the therapeutic combination for use in the treatment of a human subject who suffers from a cancer or who is at risk of developing a cancer and who is in need of said treatment, the treatment comprising the step of administering to the human subject an effective dose of the therapeutic combination of the invention, wherein the first proteinaceous molecule comprises a saponin and the second proteinaceous molecule comprises an effector moiety. An aspect of the invention relates to the therapeutic combination of the invention for use as a medicament. An aspect of the invention relates to the therapeutic combination of the invention for use in a method for treatment or prophylaxis of a cancer or an auto-immune disease, in a human subject in need thereof..

[0149] A further application in medicine is the substitution of intracellular enzymes in target cells that produce these enzymes in insufficient amount or insufficient functionality. The resulting disease might be hereditary or acquired. In most cases, only symptomatic treatment is possible and for a number of rare diseases, insufficient treatment options lead to a shortened life span of concerned patients. An example for such a disease is phenylketonuria, which is an inborn error of metabolism that results in decreased metabolism of the amino acid phenylalanine. The disease is characterized by mutations in the gene for the hepatic enzyme phenylalanine hydroxylase. Phenylketonuria is not curable to date. The incidence is approximately 1:10,000 with the highest known incidence in Turkey with 1:2,600. A second proteinaceous molecule, preferably an antibody, with bound phenylalanine hydroxylase or with a bound polynucleotide that encodes phenylalanine hydroxylase can be used to target liver cells by use of a suitable specific antibody, and to substitute the defect enzyme in hepatocytes. This is one example of use of the therapeutic combination of the invention comprising a first proteinaceous molecule with a saponin bound thereto ad a second proteinaceous molecule with the enzyme or the oligonucleotide bound thereto according to the invention for substitution or gene therapy. In a preferred embodiment, a therapeutic combination according to the invention for use in a method of gene therapy or substitution therapy is provided.

[0150] The present invention also provides the therapeutic combination for use in a method of treating cancer, the method comprising administering a medicament comprising a therapeutic combination according to the invention to a patient in need thereof, preferably administering an effective dose of said medicament to a patient in need thereof, preferably a human cancer patient.

[0151] Considerations concerning forms suitable for administration are known in the art and include toxic

effects, solubility, route of administration, and maintaining activity. For example, pharmacological compositions injected into the bloodstream should be soluble.

[0152] Suitable dosage forms, in part depend upon the use or the route of entry, for example transdermal or by injection. Such dosage forms should allow the compound to reach a target cell whether the target cell is present in a multicellular host. Other factors are known in the art, and include considerations such as toxicity and dosage form which retard the compound or composition from exerting its effect.

[0153] An embodiment is the combination of an endosomal escape enhancing conjugate according to the invention, comprising the first proteinaceous molecule comprising at least one covalently bound saponin, and a binding moiety, wherein the binding moiety comprises at least one effector moiety, the binding moiety being the second proteinaceous molecule comprising the bound effector moiety, wherein the endosomal escape enhancing conjugate and the binding moiety are, independently from one another, able to specifically bind to a target cell-specific surface molecule or structure, thereby inducing receptor-mediated endocytosis of a complex of the endosomal escape enhancing conjugate and the target cell-specific surface molecule, and of the complex of the binding moiety and the target cell-specific surface molecule, wherein the endosomal escape enhancing conjugate and the binding moiety can bind to the same target cell-specific surface molecule via their same binding site. An embodiment is the combination according to the invention, wherein the endosomal escape enhancing conjugate is able to compete with the binding moiety for binding to the target cell-specific surface molecule or structure. An embodiment is the combination according to the invention, wherein the endosomal escape enhancing conjugate and the binding moiety are, independently from one another, able to specifically bind to the same epitope. An embodiment is the combination for use in a method for the treatment of an aberrancy such as a cancer according to the invention, wherein said endosomal escape enhancing conjugate and said binding moiety are to be administered concomitant or sequentially, preferably concomitant.

[0154] An aspect of the disclosure relates to a kit comprising a first container containing an endosomal escape enhancing conjugate according to the invention (i.e. the first proteinaceous molecule) and a second container containing a binding moiety according to the invention (i.e. the second proteinaceous molecule), the kit further comprising instructions for using the binding molecules (i.e. the therapeutic combination comprising the first and second pharmaceutical compositions).

TABLE A1. Saponins displaying (late) endosomal/lysosomal escape enhancing activity, and saponins comprising a structure reminiscent to such saponins displaying (late) endosomal/lysosomal escape enhancing activity

Saponin Name	**Aglycon core**	**Carbohydrate substituent at the C-3beta-OH group**	**Carbohydrate substituent at the C-28-OH group**
NP-005236	2alpha-Hydroxyoleanolic acid	GlcA-	Glc/Gal-
AMA-1	16alpha-Hydroxyoleanolic acid	Glc-	Rha-(1→2)-[Xyl-(1→4)]-Rha-
AMR	16alpha-Hydroxyoleanolic acid	Glc-	Rha-(1→2)-[Ara-(1→3)-Xyl-(1→4)]-Rha-
alpha-Hederin	Hederagenin (23-Hydroxyoleanolic acid)	Rha-(1→2)-Ara-	-
NP-012672	16alpha,23-Dihydroxyoleanolic acid	Ara/Xyl-(1→4)-Rha/Fuc-(1→2)-Glc/Gal-(1→2)-Rha/Fuc-(1→2)-GlcA-	Ara/Xyl-
NP-017777	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→4)-Rha-(1→2)-[R-(→4)]-Fuc- (R = 4E-Methoxycinnamic acid)
NP-017778	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→4)-Rha-(1→2)-[R-(→4)]-Fuc- (R = 4Z-

			Methoxycinnamic acid)
NP-017774	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→4)-[Gal-(1→3)]-Rha-(1→2)-4-OAc-Fuc-
NP-018110[c], NP-017772[d]	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-3,4-di-OAc-Fuc-
NP-018109	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-[R-(→4)]-3-OAc-Fuc- (R = 4E-Methoxycinnamic acid)
NP-017888	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Glc-(1→3)-Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-4-OAc-Fuc-
NP-017889	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Glc-(1→3)-Xyl-(1→4)-Rha-(1→2)-4-OAc-Fuc-
NP-018108	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Ara/Xyl-(1→3)-Ara/Xyl-(1→4)-Rha/Fuc-(1→2)-[4-OAc-Rha/Fuc-(1→4)]-Rha/Fuc-
SA1641[a], AE X55[b]	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→3)-Xyl-(1→4)-Rha-(1→2)-[Qui-(1→4)]-Fuc-
NP-017674	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api-(1→3)-Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-Fuc-
NP-017810	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→4)-[Gal-(1→3)]-Rha-(1→2)-Fuc-
AG1	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-Fuc-
NP-003881	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Ara/Xyl-(1→4)-Rha/Fuc-(1→4)-[Glc/Gal-(1→2)]-Fuc-
NP-017676	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api-(1→3)-Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-[R-(→4)]-Fuc- (R = 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid)
NP-017677	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api-(1→3)-Xyl-(1→4)-Rha-(1→2)-[R-(→4)]-Fuc- (R = 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid)
NP-017706	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api-(1→3)-Xyl-(1→4)-Rha-(1→2)-[Rha-(1→3)]-4-OAc-Fuc-
NP-017705	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api-(1→3)-Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-[Rha-(1→3)]-4-OAc-Fuc-
NP-017773	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	6-OAc-Glc-(1→3)-Xyl-(1→4)-Rha-(1→2)-[3-OAc-Rha-(1→3)]-Fuc-
NP-017775	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Glc-(1→3)-Xyl-(1→4)-Rha-(1→2)-[3-OAc-Rha-(1→3)]-Fuc-
SA1657	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→3)-Xyl-(1→4)-Rha-(1→2)-[Qui-(1→4)]-Fuc-
AG2	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Glc-(1→3)-[Xyl-(1→4)]-Rha-(1→2)-[Qui-(1→4)]-Fuc-

SO1861	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Glc-(1→3)-Xyl-(1→4)-Rha-(1→2)-[Xyl-(1→3)-4-OAc-Qui-(1→4)]-Fuc-
GE1741	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→3)-Xyl-(1→4)-Rha-(1→2)-[3,4-di-OAc-Qui-(1→4)]-Fuc-
SO1542	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Glc-(1→3)-[Xyl-(1→4)]-Rha-(1→2)-Fuc-
SO1584	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	6-OAc-Glc-(1→3)-[Xyl-(1→4)]-Rha-(1→2)-Fuc-
SO1658	Gypsogenin	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Glc-(1→3)-[Xyl-(1→3)-Xyl-(1→4)]-Rha-(1→2)-Fuc-
SO1674	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Glc-(1→3)-[Xyl-(1→3)-Xyl-(1→4)]-Rha-(1→2)-Fuc-
SO1832	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→3)-Xyl-(1→4)-Rha-(1→2)-[Xyl-(1→3)-4-OAc-Qui-(1→4)]-Fuc-
QS-7 (also referred to as QS1861)	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api/Xyl-(1→3)-Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-[Rha-(1→3)]-4OAc-Fuc-
QS-7 api (also referred to as QS1862)	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api-(1→3)-Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-[Rha-(1→3)]-4OAc-Fuc-
QS-17	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api/Xyl-(1→3)-Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-[R-(→4)]-Fuc- (R = 5-O-[5-O-Rha-(1→2)-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-

			dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid)
QS-18	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api/Xyl-(1→3)-Xyl-(1→4)-[Glc-(1→3)]-Rha-(1→2)-[R-(→4)]-Fuc- (R = 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid)
QS-21 A-apio	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api-(1→3)-Xyl-(1→4)-Rha-(1→2)-[R-(→4)]-Fuc- (R = 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid)
QS-21 A-xylo	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→3)-Xyl-(1→4)-Rha-(1→2)-[R-(→4)]-Fuc- (R = 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid)
QS-21 B-apio	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Api-(1→3)-Xyl-(1→4)-Rha-(1→2)-[R-(→3)]-Fuc- (R = 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid)
QS-21 B-xylo	Quillaic acid	Gal-(1→2)-[Xyl-(1→3)]-GlcA-	Xyl-(1→3)-Xyl-(1→4)-Rha-(1→2)-[R-(→3)]-Fuc- (R = 5-O-[5-O-Ara/Api-3,5-dihydroxy-6-methyl-octanoyl]-3,5-dihydroxy-6-methyl-octanoic acid)

beta-Aescin (described: Aescin Ia)	Protoaescigenin-21(2-methylbut-2-enoate)-22-acetat	Glc-(1→2)-[Glc-(1→4)]-GlcA-	-
Teaseed saponin I	23-Oxo-barringtogenol C - 21,22-bis(2-methylbut-2-enoate)	Glc-(1→2)-Ara-(1→3)-[Gal-(1→2)]-GlcA-	-
Teaseedsaponin J	23-Oxo-barringtogenol C - 21,22-bis(2-methylbut-2-enoate)	Xyl-(1→2)-Ara-(1→3)-[Gal-(1→2)]-GlcA-	-
Assamsaponin F	23-Oxo-barringtogenol C - 21(2-methylbut-2-enoate)-16,22-diacetat	Glc-(1→2)-Ara-(1→3)-[Gal-(1→2)]-GlcA-	-
Digitonin	Digitogenin	Glc-(1→3)-Gal-(1→2)-[Xyl-(1→3)]-Glc-(1→4)-Gal-	-
Primula acid 1	3,16,28-Trihydroxyoleanan-12-en	Rha-(1→2)-Gal-(1→3)-[Glc-(1→2)]-GlcA-	-
AS64R	Gypsogenic acid	-	Glc-(1→3)-[Glc-(1→6)]-Gal-
		Carbohydrate substituent at the C-23-OH group	
AS6.2	Gypsogenic acid	Gal-	Glc-(1→3)-[Glc-(1→6)]-Gal-
a, b: Different names refer to different isolates of the same structure c, d: Different names refer to different isolates of the same structure			

TABLE A2 - ADCs which were previously investigated in the human clinical setting, and subsequently retracted from further clinical investigation

Drug Name	Indication	Target	Last Development Stage	
Monoclonal Antibody Conjugate to Target EGFR for Oncology	Oncology	Cells Expressing Epidermal Growth Factor Receptor (Proto Oncogene c ErbB 1 or Receptor Tyrosine Protein Kinase erbB 1 or HER1 or ERBB1 or EGFR or EC 2.7.10.1)	Discovery	
Affilutin	Multiple Myeloma (Kahler Disease)		Discovery	
IMGN-779	Myelodys-plastic Syndrome	Cells Expressing Myeloid Cell Surface Antigen CD33 (Sialic Acid Binding Ig Like Lectin 3 or gp67 or CD33)	IND/CTA Filed	
Neuradiab	Non-Hodgkin Lymphoma	Cells Expressing Tenascin (Cytotactin or GMEM or GP 150-225 or Glioma Associated Extracellular Matrix Antigen or Hexabrachion or JI or Myotendinous Antigen or Neuronectin or Tenascin C or TNC)	Phase I	
IMGN-779	Refractory Acute Myeloid Leukemia; Relapsed Acute Myeloid Leukemia	Cells Expressing Myeloid Cell Surface Antigen CD33 (Sialic Acid Binding Ig Like Lectin 3 or gp67 or CD33)	Phase I	
AGS-67E	Acute Myelocytic Leukemia (AML, Acute Myeloblas-tic Leukemia)	Cells Expressing Leukocyte Antigen CD37 (Tetraspanin 26 or CD37)	Phase I	
AGS-67E	Hairy Cell Leukemia; Non-Hodgkin Lymphoma; Refractory Chronic Lymphocy-tic Leukemia (CLL); Relapsed Chronic Lymphocy-tic Leukemia (CLL); T-Cell Leukemia	Cells Expressing Leukocyte Antigen CD37 (Tetraspanin 26 or CD37)	Phase I	
ASG-15ME	Metastatic Transitional (Urothelial) Tract Cancer	Cells Expressing SLIT And NTRK Like Protein 6 (SLITRK6)	Phase I	
vandortuzumab vedotin	Metastatic Hormone Refractory (Castration Resistant, Androgen-Indepen-dent) Prostate Cancer	Cells Expressing Metalloreductase STEAP1 (Six Transmembrane	Phase I	

Drug Name	Indication	Target	Last Development Stage	
		Epithelial Antigen Of The Prostate 1 or STEAP1 or EC 1.16.1.)		
CDX-014	Ovarian Cancer	Cells Expressing Hepatitis A Virus Cellular Receptor 1 (Kidney Injury Molecule 1 or T Cell Immunoglobulin And Mucin Domain Containing Protein 1 or T-Cell Immunoglobulin Mucin Receptor 1 or T Cell Membrane Protein 1 or CD365 or HAVCR1)	Phase I	
AGS-16M18	Liver Cancer; Renal Cell Carcinoma		Phase I	
vorsetuzumab mafodotin	Non-Hodgkin Lymphoma; Renal Cell Carcinoma	Cells Expressing CD70 Antigen (CD27 Ligand or Tumor Necrosis Factor Ligand Superfamily Member 7 or CD70)	Phase	
denintuzumab mafodotin	Acute Lymphocy-tic Leukemia (ALL, Acute Lympho-blastic Leukemia); B-Cell Non-Hodgkin Lymphoma; Burkitt Lymphoma; Lympho-blastic Lymphoma; Mantle Cell Lymphoma	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T Cell Surface Antigen Leu 12 or CD19)	Phase I	
SGN-CD70A	Diffuse Large B-Cell Lymphoma; Follicular Lymphoma; Mantle Cell Lymphoma; Metastatic Renal Cell Carcinoma; Non-Hodgkin Lymphoma	Cells Expressing CD70 Antigen (CD27 Ligand or Tumor Necrosis Factor Ligand Superfamily Member 7 or CD70)	Phase I	
RG-7636	Metastatic Melanoma	Endothelin B Receptor (Endothelin Receptor Non Selective Type or EDNRB)	Phase I	
SC-006	Metastatic Colorectal Cancer		Phase I	
MM-310	Breast Cancer; Endome-trial Cancer; Esophageal Cancer; Gastric Cancer; Gastroeso-phageal (GE) Junction Carcino-mas; Head And Neck Cancer Squamous Cell Carcinoma; Non-Small Cell Lung Cancer; Ovarian	Ephrin Type A Receptor 2 (Epithelial Cell Kinase or Tyrosine Protein Kinase Receptor ECK or EPHA2 or EC 2.7.10.1)	Phase I	

Drug Name	Indication	Target	Last Development Stage	
	Cancer; Pancreatic Ductal Adenocar-cinoma; Prostate Cancer; Small-Cell Lung Cancer; Soft Tissue Sarcoma; Solid Tumor; Transitional Cell Carcinoma (Urothelial Cell Carcinoma)			
PF-06647263	Metastatic Breast Cancer; Ovarian Cancer	Cells Expressing Ephrin A4 (EPH Related Receptor Tyrosine Kinase Ligand 4 or EFNA4)	Phase I	
PF-06263507	Solid Tumor	Cells Expressing Trophoblast Glycoprotein (M6P1 or 5T4 Oncofetal Antigen or 5T4 Oncofetal Trophoblast Glycoprotein or Wnt Activated Inhibitory Factor 1 or TPBG)	Phase I	
PF-06650808	Metastatic Breast Cancer; Non-Small Cell Lung Cancer; Ovarian Cancer	Cells Expressing Neurogenic Locus Notch Homolog Protein 3 (NOTCH3)	Phase I	
XMT-1522	Breast Cancer; Gastric Cancer; Non-Small Cell Lung Cancer	Receptor Tyrosine Protein Kinase ERBB 2 (Metastatic Lymph Node Gene 19 Protein or Proto Oncogene Neu or Proto Oncogene C ErbB 2 or Tyrosine Kinase Type Cell Surface Receptor HER2 or p185erbB2 or HER2 or CD340 or ERBB2 or EC 2.7.10.1); Tubulin	Phase I	
AMG-595	Anaplastic Astrocyto-ma; Recurrent Glioblasto-ma Multiforme (GBM)	Cells Expressing Epidermal Growth Factor Receptor (Proto Oncogene c ErbB 1 or Receptor Tyrosine Protein Kinase erbB 1 or HER1 or ERBB1 or EGFR or EC 2.7.10.1)	Phase I	
pinatuzumab vedotin	Chronic Lymphocytic Leukemia (CLL)	Cells Expressing B Cell Receptor CD22 (B Lymphocyte Cell Adhesion Molecule or Sialic Acid Binding Ig Like Lectin 2 or T Cell	Phase I	

Drug Name	Indication	Target	Last Development Stage	
		Surface Antigen Leu 14 or CD22)		
cantuzumab ravtansine	Colorectal Cancer; Non-Small Cell Lung Cancer; Pancreatic Cancer; Solid Tumor		Phase I	
AVE-9633	Acute Myelocytic Leukemia (AML, Acute Myeloblas-tic Leukemia)	Cells Expressing Myeloid Cell Surface Antigen CD33 (Sialic Acid Binding Ig Like Lectin 3 or gp67 or CD33)	Phase I	
BIWI-1[(1)]	Breast Cancer; Carcino-mas; Esophageal Cancer; Head And Neck Cancer Squamous Cell Carcinoma	Cells Expressing CD44 Antigen (CDw44 or Epican or Extracellular Matrix Receptor III or GP90 Lymphocyte Homing/Adhesion Receptor or HUTCH I or Heparan Sulfate Proteoglycan or Hermes Antigen or Hyaluronate Receptor or Phagocytic Glycoprotein 1 or CD44)	Phase I	
RG-7882	Epithelial Ovarian Cancer; Fallopian Tube Cancer; Pancreatic Cancer; Peritoneal Cancer	Cells Expressing Mucin 16 (Ovarian Cancer Related Tumor Marker CA125 or Ovarian Carcinoma Antigen CA125 or MUC16)	Phase I	
ASG-5ME	Adenocar-cinoma; Hormone Refractory (Castration Resistant, Androgen-Indepen-dent) Prostate Cancer; Metastatic Adenocar-cinoma of The Pancreas	Cells Expressing Choline Transporter Like Protein 4 (Solute Carrier Family 44 Member 4 or SLC44A4)	Phase I	
DCDS-0780A	B-Cell Non-Hodgkin Lymphoma		Phase I	
SC-004	Endome-trial Cancer; Epithelial Ovarian Cancer; Fallopian Tube Cancer; Peritoneal Cancer		Phase I	
RG-7600	Ovarian Cancer; Pancreatic Ductal Adenocar-cinoma		Phase I	
sofituzumab vedotin	Epithelial Ovarian Cancer; Fallopian Tube Cancer; Ovarian Cancer; Pancreatic Cancer; Peritoneal Cancer	Cells Expressing Mucin 16 (Ovarian Cancer Related Tumor Marker CA125 or Ovarian Carcinoma Antigen CA125 or MUC16)	Phase I	

Drug Name	Indication	Target	Last Development Stage	
IMGN-289	Breast Cancer; Esophageal Cancer; Gastric Cancer; Head And Neck Cancer Squamous Cell Carcinoma; Non-Small Cell Lung Cancer; Solid Tumor	Cells Expressing Epidermal Growth Factor Receptor (Proto Oncogene c ErbB 1 or Receptor Tyrosine Protein Kinase erbB 1 or HER1 or ERBB1 or EGFR or EC 2.7.10.1)	Phase I	
SAR-428926	Breast Cancer; Colorectal Cancer; Gastric Cancer; Non-Small Cell Lung Cancer; Ovarian Cancer; Prostate Cancer; Solid Tumor	Cells Expressing Lysosome Associated Membrane Glycoprotein 1 (CD107 Antigen Like Family Member A or CD107a or LAMP1)	Phase I	
SGNCD-19B	B-Cell Non-Hodgkin Lymphoma; Diffuse Large B-Cell Lymphoma; Follicular Lymphoma	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T Cell Surface Antigen Leu 12 or CD19)	P Phase I	
SGNCD-123A	Refractory Acute Myeloid Leukemia; Relapsed Acute Myeloid Leukemia	Cells Expressing Interleukin 3 Receptor Subunit Alpha (CD123 or IL3RA)	Phase I	
SGNCD-352A	Refractory Multiple Myeloma; Relapsed Multiple Myeloma	Cells Expressing SLAM Family Member 6 (Activating NK Receptor or NK T B Antigen or CD352 or SLAMF6)	Phase I	
RG-7841	Breast Cancer; Non-Small Cell Lung Cancer; Solid Tumor	Cells Expressing Lymphocyte Antigen 6E (Retinoic Acid Induced Gene E Protein or Stem Cell Antigen 2 or Thymic Shared Antigen 1 or LY6E)	Phase I	
IMGN-388	Solid Tumor	Cells Expressing Integrin Alpha V (Vitronectin Receptor Subunit Alpha or CD51 or ITGAV)	Phase I	
lorvotuzumab mertansine	Refractory Multiple Myeloma; Relapsed Multiple Myeloma	Cells Expressing Neural Cell Adhesion Molecule 1 (Antigen Recognized By Monoclonal Antibody	Phase I	

Drug Name	Indication	Target	Last Development Stage	
		5.1H11 or CD56 or NCAM1)		
lorvotuzumab mertansine	Neuroendo-crine Carcinoma; Neuroendo-crine Tumors; Non-Small Cell Lung Cancer; Ovarian Cancer; Skin Cancer	Cells Expressing Neural Cell Adhesion Molecule 1 (Antigen Recognized By Monoclonal Antibody 5.1H11 or CD56 or NCAM1)	Phase I	
BAY-794620	Lung Cancer; Solid Tumor	Cells Expressing Carbonic Anhydrase 9 (Carbonate Dehydratase IX or pMW1 or Membrane Antigen MN or P54/58N or Renal Cell Carcinoma Associated Antigen G250 or CA9 or EC 4.2.1.1)	Phase I	
RG-7598	Refractory Multiple Myeloma; Relapsed Multiple Myeloma		Phase I	
Oncolysin B	B-Cell Leukemia; Lymphoma	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T Cell Surface Antigen Leu 12 or CD19)	Phase I	
ADCT-502[(1)]	Bladder Cancer; Breast Cancer; Esophageal Cancer; Gastric Cancer; Non-Small Cell Lung Cancer	Cells Expressing Receptor Tyrosine Protein Kinase ERBB 2 (Metastatic Lymph Node Gene 19 Protein or Proto Oncogene Neu or Proto Oncogene C ErbB 2 or Tyrosine Kinase Type Cell Surface Receptor HER2 or p185erbB2 or HER2 or CD340 or ERBB2 or EC 2.7.10.1)	Phase I	
AMG-172	Renal Cell Carcinoma	Cells Expressing CD70 Antigen (CD27 Ligand or Tumor Necrosis Factor Ligand Superfamily Member 7 or CD70)	Phase I	
ImmuRAIT-LL2	B-Cell Non-Hodgkin Lymphoma	Cells Expressing B Cell Receptor CD22	Phase I/II	

Drug Name	Indication	Target	Last Development Stage	
		(B Lymphocyte Cell Adhesion Molecule or Sialic Acid Binding Ig Like Lectin 2 or T Cell Surface Antigen Leu 14 or CD22)		
indusatumab vedotin	Adenocar-cinoma Of The Gastroe-sophageal Junction; Gastric Cancer	Cells Expressing Heat Stable Enterotoxin Receptor (Guanylyl Cyclase C or or Intestinal Guanylate Cyclase or GUCY2C or EC 4.6.1.2)	Phase I/II	
clivatuzumab tetraxetan	Pancreatic Cancer	Cells Expressing Mucin 1 (Breast Carcinoma Associated Antigen DF3 or Episialin or H23AG or Krebs Von Den Lungen 6 or PEMT or Peanut Reactive Urinary Mucin or Polymorphic Epithelial Mucin or Tumor Associated Epithelial Membrane Antigen or Tumor Associated Mucin or CD227 or MUC1)	Phase I/II	
depatuxizumab mafodotin[(2)]	Recurrent Malignant Glioma	Epidermal Growth Factor Receptor (Proto Oncogene c ErbB 1 or Receptor Tyrosine Protein Kinase erbB 1 or HER1 or ERBB1 or EGFR or EC 2.7.10.1)	Phase I/II	
CDX-014	Metastatic Renal Cell Carcinoma; Papillary Renal Cell Carcinoma	Cells Expressing Hepatitis A Virus Cellular Receptor 1 (Kidney Injury Molecule 1 or T Cell Immunoglobulin And Mucin Domain Containing Protein 1 or T-Cell Immunoglobulin Mucin Receptor 1 or T Cell Membrane Protein 1 or CD365 or HAVCR1)	Phase I/II	
vadastuximab	Refractory Acute Myeloid	Cells Expressing	Phase I/II	

Drug Name	Indication	Target	Last Development Stage	
talirine[(1)]	Leukemia; Relapsed Acute Myeloid Leukemia	Myeloid Cell Surface Antigen CD33 (Sialic Acid Binding Ig Like Lectin 3 or gp67 or CD33)		
vadastuximab talirine	Myelodys-plastic Syndrome	Cells Expressing Myeloid Cell Surface Antigen CD33 (Sialic Acid Binding Ig Like Lectin 3 or gp67 or CD33)	Phase I/II	
MLN-2704	Metastatic Hormone Refractory (Castration Resistant, Androgen-Indepen-dent) Prostate Cancer	Cells Expressing Glutamate Carboxypeptidase 2 (Folate Hydrolase 1 or Prostate Specific Membrane Antigen or PSMA or Pteroylpoly Gamma Glutamate Carboxypeptidase or Cell Growth Inhibiting Gene 27 Protein or FOLH1 or EC 3.4.17.21)	Phase I/II	
Oncolysin B	AIDS - Related Lymphoma	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T Cell Surface Antigen Leu 12 or CD19)	Phase I/II	
coltuximab ravtansine	Diffuse Large B-Cell Lymphoma	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T Cell Surface Antigen Leu 12 or CD19)	Phase II	
coltuximab ravtansine	Acute Lymphocy-tic Leukemia (ALL, Acute Lympho-blastic Leukemia)	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T Cell Surface Antigen Leu 12 or CD19)	Phase II	
coltuximab ravtansine	Diffuse Large B-Cell Lymphoma	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T	Phase II	

Drug Name	Indication	Target	Last Development Stage	
		Cell Surface Antigen Leu 12 or CD19)		
indusatumab vedotin[2]	Adenocar-cinoma Of The Gastroe-sophageal Junction; Gastric Cancer; Metastatic Adenocar-cinoma of The Pancreas	Cells Expressing Heat Stable Enterotoxin Receptor (Guanylyl Cyclase C or or Intestinal Guanylate Cyclase or GUCY2C or EC 4.6.1.2)	Phase II	
depatuxizumab mafodotin	Squamous Non-Small Cell Lung Cancer	Epidermal Growth Factor Receptor (Proto Oncogene c ErbB 1 or Receptor Tyrosine Protein Kinase erbB 1 or HER1 or ERBB1 or EGFR or EC 2.7.10.1)	Phase II	
depatuxizumab mafodotin[2]	Anaplastic Astrocyto-ma; Anaplastic Oligoastro-cytoma; Gliosar-coma; High-Grade Glioma; Oligoden-droglioma; Pediatric Diffuse Intrinsic Pontine Glioma; Recurrent Glioblasto-ma Multiforme (GBM)	Epidermal Growth Factor Receptor (Proto Oncogene c ErbB 1 or Receptor Tyrosine Protein Kinase erbB 1 or HER1 or ERBB1 or EGFR or EC 2.7.10.1)	Phase II	
lifastuzumab vedotin	Non-Small Cell Lung Cancer	Sodium Dependent Phosphate Transport Protein 2B (Sodium Phosphate Transport Protein 2B or NaPi3b or Sodium/Phosphate Cotransporter 2B or NaPi 2b or Solute Carrier Family 34 Member 2 or SLC34A2)	Phase II	
lifastuzumab vedotin	Ovarian Cancer	Sodium Dependent Phosphate Transport Protein 2B (Sodium Phosphate Transport Protein 2B or NaPi3b or Sodium/Phosphate Cotransporter 2B or NaPi 2b or Solute Carrier Family 34 Member 2 or SLC34A2)	Phase II	
Bismab-A	Acute Myelocytic Leukemia (AML, Acute Myeloblas-tic Leukemia)	Cells Expressing Myeloid Cell Surface Antigen CD33 (Sialic Acid Binding Ig Like	Phase II	

Drug Name	Indication	Target	Last Development Stage	
		Lectin 3 or gp67 or CD33)		
denintuzumab mafodotin	Diffuse Large B-Cell Lymphoma; Follicular Lymphoma	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T Cell Surface Antigen Leu 12 or CD19)	Phase II	
Avicidin[1]	Colorectal Cancer; Prostate Cancer	Cells Expressing Epithelial Cell Adhesion Molecule (Adenocarcinoma Associated Antigen or Cell Surface Glycoprotein Trop 1 or Epithelial Cell Surface Antigen or Epithelial Glycoprotein 314 or KS 1/4 Antigen or KSA or Tumor Associated Calcium Signal Transducer 1 or CD326 or EPCAM)	Phase II	
pinatuzumab vedotin	Diffuse Large B-Cell Lymphoma; Follicular Lymphoma	Cells Expressing B Cell Receptor CD22 (B Lymphocyte Cell Adhesion Molecule or Sialic Acid Binding Ig Like Lectin 2 or T Cell Surface Antigen Leu 14 or CD22)	Phase II	
SGN-15	Metastatic Breast Cancer; Non-Small Cell Lung Cancer; Ovarian Cancer; Prostate Cancer	Cells Expressing Lewis Y Antigen (CD174)	Phase II	
cantuzumab ravtansine	Gastric Cancer; Gastroe-sophageal (GE) Junction Carcino-mas		Phase II	
ASP-6183	Ovarian Cancer		Phase II	
SAR-566658	Metastatic Breast Cancer	Cells Expressing Sialoglycotope CA6 Antigen	Phase II	
Oncolysin S	Small-Cell Lung Cancer	Cells Expressing Neural Cell Adhesion Molecule 1 (Antigen Recognized By Monoclonal Antibody 5.1 H11 or CD56 or NCAM1)	Phase II	
lorvotuzumab	Small-Cell Lung Cancer	Cells Expressing	Phase II	

Drug Name	Indication	Target	Last Development Stage	
mertansine		Neural Cell Adhesion Molecule 1 (Antigen Recognized By Monoclonal Antibody 5.1H11 or CD56 or NCAM1)		
glembatumumab vedotin	Metastatic Melanoma; Metastatic Uveal Melanoma; Osteosar-coma; Squamous Non-Small Cell Lung Cancer	Cells Expressing Transmembrane Glycoprotein NMB (Transmembrane Glycoprotein HGFIN or GPNMB)	Phase II	
MM-302	Metastatic Breast Cancer	Cells Expressing Receptor Tyrosine Protein Kinase ERBB 2 (Metastatic Lymph Node Gene 19 Protein or Proto Oncogene Neu or Proto Oncogene C ErbB 2 or Tyrosine Kinase Type Cell Surface Receptor HER2 or p185erbB2 or HER2 or CD340 or ERBB2 or EC 2.7.10.1)	Phase II/III	
Neuradiab	Brain Cancer; Glioblasto-ma Multiforme (GBM)	Cells Expressing Tenascin (Cytotactin or GMEM or GP 150-225 or Glioma Associated Extracellular Matrix Antigen or Hexabrachion or JI or Myotendinous Antigen or Neuronectin or Tenascin C or TNC)	Phase III	
clivatuzumab tetraxetan	Metastatic Adenocar-cinoma of The Pancreas	Cells Expressing Mucin 1 (Breast Carcinoma Associated Antigen DF3 or Episialin or H23AG or Krebs Von Den Lungen 6 or PEMT or Peanut Reactive Urinary Mucin or Polymorphic Epithelial Mucin or Tumor Associated Epithelial Membrane Antigen or Tumor Associated Mucin or	Phase III	

Drug Name	Indication	Target	Last Development Stage	
		CD227 or MUC1)		
depatuxizumab mafodotin[2]	Glioblasto-ma Multiforme (GBM)	Epidermal Growth Factor Receptor (Proto Oncogene c ErbB 1 or Receptor Tyrosine Protein Kinase erbB 1 or HER1 or ERBB1 or EGFR or EC 2.7.10.1)	Phase III	
vadastuximab talirine[1]	Acute Myelocytic Leukemia (AML, Acute Myeloblas-tic Leukemia)	Cells Expressing Myeloid Cell Surface Antigen CD33 (Sialic Acid Binding Ig Like Lectin 3 or gp67 or CD33)	Phase III	
glembatumumab vedotin[2]	Metastatic Breast Cancer	Cells Expressing Transmembrane Glycoprotein NMB (Transmembrane Glycoprotein HGFIN or GPNMB)	Phase III	
Oncolysin B	B-Cell Leukemia; Lymphoma	Cells Expressing B Lymphocyte Antigen CD19 (B Lymphocyte Surface Antigen B4 or Differentiation Antigen CD19 or T Cell Surface Antigen Leu 12 or CD19)	Phase III	
ImmuRAIT-LL2	B-Cell Leukemia	Cells Expressing B Cell Receptor CD22 (B Lymphocyte Cell Adhesion Molecule or Sialic Acid Binding Ig Like Lectin 2 or T Cell Surface Antigen Leu 14 or CD22)	Preclinical	
indusatumab vedotin	Metastatic Colorectal Cancer	Cells Expressing Heat Stable Enterotoxin Receptor (Guanylyl Cyclase C or or Intestinal Guanylate Cyclase or GUCY2C or EC 4.6.1.2)	Preclinical	
ASG-15ME	Lung Cancer	Cells Expressing SLIT And NTRK Like Protein 6 (SLITRK6)	Preclinical	
HTI-1511	Bile Duct Cancer (Cholangiocarcinoma) ; Breast Cancer; Colorectal Cancer; Non-Small Cell Lung Cancer	Cells Expressing Epidermal Growth Factor Receptor (Proto Oncogene c	Preclinical	

Drug Name	Indication	Target	Last Development Stage	
		ErbB 1 or Receptor Tyrosine Protein Kinase erbB 1 or HER1 or ERBB1 or EGFR or EC 2.7.10.1)		
ZW-33	Gastric Cancer; Metastatic Breast Cancer	Cells Expressing Receptor Tyrosine Protein Kinase ERBB 2 (Metastatic Lymph Node Gene 19 Protein or Proto Oncogene Neu or Proto Oncogene C ErbB 2 or Tyrosine Kinase Type Cell Surface Receptor HER2 or p185erbB2 or HER2 or CD340 or ERBB2 or EC 2.7.10.1)	Preclinical	
ZW-33	Ovarian Cancer	Cells Expressing Receptor Tyrosine Protein Kinase ERBB 2 (Metastatic Lymph Node Gene 19 Protein or Proto Oncogene Neu or Proto Oncogene C ErbB 2 or Tyrosine Kinase Type Cell Surface Receptor HER2 or p185erbB2 or HER2 or CD340 or ERBB2 or EC 2.7.10.1)	Preclinical	
SGNCD-352A	Non-Hodgkin Lymphoma	Cells Expressing SLAM Family Member 6 (Activating NK Receptor or NK T B Antigen or CD352 or SLAMF6)	Preclinical	
HuMax-CD74-ADC	Oncology	Cells Expressing HLA Class II Histocompatibility Antigen Gamma Chain (HLA DR Antigens Associated Invariant Chain or Ia Antigen Associated Invariant Chain or p33 or CD74)	Preclinical	
sacituzumab govitecan	Pancreatic Ductal Adenocar-cinoma	Cells Expressing Tumor Associated		

Drug Name	Indication	Target	Last Development Stage	
		Calcium Signal Transducer 2 (Cell Surface Glycoprotein Trop 2 or Membrane Component Chromosome 1 Surface Marker 1 or Pancreatic Carcinoma Marker Protein GA733-1 or TACSTD2)		
sacituzumab govitecan	Adenocar-cinoma; Cervical Cancer; Colorectal Cancer; Endome-trial Cancer; Epithelial Ovarian Cancer; Esophageal Cancer; Follicular Thyroid Cancer; Gastric Cancer; Glioblasto-ma Multiforme (GBM); Head And Neck Cancer Squamous Cell Carcinoma; Hepato-cellular Carcinoma; Kidney Cancer (Renal Cell Cancer); Metastatic Hormone Refractory (Castration Resistant, Androgen-Indepen-dent) Prostate Cancer; Metastatic Transitional (Urothelial) Tract Cancer; Transitional Cell Cancer (Urothelial Cell Cancer)	Cells Expressing Tumor Associated Calcium Signal Transducer 2 (Cell Surface Glycoprotein Trop 2 or Membrane Component Chromosome 1 Surface Marker 1 or Pancreatic Carcinoma Marker Protein GA733-1 or TACSTD2)		
sacituzumab govitecan	Hepato-cellular Carcinoma	Cells Expressing Tumor Associated Calcium Signal Transducer 2 (Cell Surface Glycoprotein Trop 2 or Membrane Component Chromosome 1 Surface Marker 1 or Pancreatic Carcinoma Marker Protein GA733-1 or TACSTD2)		
sacituzumab govitecan	Metastatic Breast Cancer; Transitional Cell Cancer (Urothelial Cell Cancer)	Cells Expressing Tumor Associated Calcium Signal Transducer 2 (Cell Surface Glycoprotein Trop 2 or Membrane Component Chromosome 1 Surface Marker 1 or Pancreatic Carcinoma Marker Protein GA733-1 or TACSTD2)		

Drug Name	Indication	Target	Last Development Stage	
sacituzumab govitecan	Non-Small Cell Lung Cancer; Small-Cell Lung Cancer	Cells Expressing Tumor Associated Calcium Signal Transducer 2 (Cell Surface Glycoprotein Trop 2 or Membrane Component Chromosome 1 Surface Marker 1 or Pancreatic Carcinoma Marker Protein GA733-1 or TACSTD2)		
sacituzumab govitecan	Metastatic Breast Cancer	Cells Expressing Tumor Associated Calcium Signal Transducer 2 (Cell Surface Glycoprotein Trop 2 or Membrane Component Chromosome 1 Surface Marker 1 or Pancreatic Carcinoma Marker Protein GA733-1 or TACSTD2)		
(1) Discontinued due to adverse events				
(2) Discontinued due to lack of efficacy				

TABLE A3 - ADCs that reached phase III clinical development

Drug Name	Indication	Development Stage	Last Development Stage	Reason for Discontinuation
trastuzumab emtansine	Gastric Cancer	Marketed	Phase II/III	Unspecified
MM-302	Metastatic Breast Cancer	Discontinued	Phase II/III	Business/Strategic Decision
trastuzumab emtansine	Metastatic Breast Cancer	Marketed	Phase III	Unspecified
trastuzumab emtansine	Gastric Cancer	Marketed	Phase III	Unspecified
ibritumomab tiuxetan	Diffuse Large B-Cell Lymphoma	Marketed	Phase III	
inotuzumab ozogamicin	Follicular Lymphoma	Marketed	Phase III	
inotuzumab ozogamicin	Diffuse Large B-Cell Lymphoma; Non-Hodgkin Lymphoma	Marketed	Phase III	Lack of Efficacy
rovalpituzumab tesirine	Small-Cell Lung Cancer	Phase III	Phase III	
rovalpituzumab	Small-Cell Lung Cancer	Phase III	Phase III	

Drug Name	Indication	Development Stage	Last Development Stage	Reason for Discontinuation
tesirine				
Neuradiab	Brain Cancer; Glioblastoma Multiforme (GBM)	Inactive	Phase III	Unspecified
clivatuzumab tetraxetan	Metastatic Adenocarcinoma of The Pancreas	Inactive	Phase III	Unspecified
depatuxizumab mafodotin	Glioblastoma Multiforme (GBM)	Inactive	Phase III	Lack of Efficacy
vadastuximab talirine	Acute Myelocytic Leukemia (AML, Acute Myeloblastic Leukemia)	Discontinued	Phase III	Adverse Events
glembatumumab vedotin	Metastatic Breast Cancer	Discontinued	Phase III	Lack of Efficacy
Oncolysin B	B-Cell Leukemia; Lymphoma	Discontinued	Phase III	Business/Strategic Decision

TABLE A4. Tumor-specific cell-surface receptor targets which can be targeted by immunoglobulins according to the invention, and antibodies that can be used for the ADCs and the antibodies provided with a saponin, and the ADCs provided with a saponin, of the present invention (not presented as a limitation; further immunoglobulins are equally suitable for the invention)

Target cell-surface receptor	Example monoclonal antibodies
HER2	anti-HER2 monoclonal antibody such as trastuzumab and pertuzumab
CD20	anti-CD20 monoclonal antibody such as rituximab, ofatumumab, tositumomab and ibritumomab
CA125	anti-CA125 monoclonal antibody such as oregovomab
EpCAM (17-1A)	anti-EpCAM (17-1A) monoclonal antibody such as edrecolomab
EGFR	anti-EGFR monoclonal antibody such as cetuximab, panitumumab and nimotuzumab
CD30	anti-CD30 monoclonal antibody such brentuximab
CD33	anti-CD33 monoclonal antibody such as gemtuzumab and huMy9-6
vascular integrin alpha-v beta-3	anti-vascular integrin alpha-v beta-3 monoclonal antibody such as etaracizumab
CD52	anti-CD52 monoclonal antibody such as alemtuzumab
CD22	anti-CD22 monoclonal antibody such as epratuzumab
CEA	anti-CEA monoclonal antibody such as labetuzumab
CD44v6	anti-CD44v6 monoclonal antibody such as bivatuzumab
FAP	anti- FAP monoclonal antibody such as sibrotuzumab

TABLE A4. Tumor-specific cell-surface receptor targets which can be targeted by immunoglobulins according to the invention, and antibodies that can be used for the ADCs and the antibodies provided with a saponin, and the ADCs provided with a saponin, of the present invention (not presented as a limitation; further immunoglobulins are equally suitable for the invention)	
Target cell-surface receptor	**Example monoclonal antibodies**
CD19	anti-CD19 monoclonal antibody such as huB4
CanAg	anti-CanAg monoclonal antibody such as huC242
CD56	anti-CD56 monoclonal antibody such huN901
CD38	anti-CD38 monoclonal antibody such as daratumumab
CA6	anti-CA6 monoclonal antibody such as DS6
IGF-IR	anti-IGF-IR monoclonal antibody such as cixutumumab and 3B7
integrin	anti-integrin monoclonal antibody such as CNTO 95
syndecan-1	anti-syndecan-1 monoclonal antibody such as B-B4

Table A5: RIPs from plants*

Plant Family	**Plant Species**	**Proteins**	**Classification**
	Sambucus ebulus L.	Ebulitin α, Ebulitin β, Ebulitin γ	RIP 1
		Ebulin f, Ebulin l, Ebulin r1, Ebulin r2, SEA	RIP 2
		SEAII, SELfd, SELId, SELIm	lectin
	Sambucus nigra L.	α-Nigntin, β-Nigntin, γ-Nigritin, Nigritin f1, Nigritin f2	RIP 1
		basic Nigrin b, Nigrin b = SNA-V, Nigrin f = SNA-Vf, Nigrin 11, Nigrin l2, Nigrin s, SNA-I, SNA-I', SNA-If, SNAflu-I, SNLRP1, SNLRP2	RIP 2
Adoxaceae		SNA-Id, SNA-Im, SNA-II, SNA-III, SNA-IV = SNA-IVf, SNA-IVI, SNApol-I, SNApol-II, TrSNA-I, TrSNA-If	lectin
	Sambucus racemosa L.	basic racemosin b, SRA	RIP 2
		SRLbm = SRAbm	lectin
	Sambucus sieboldiana (Miq.) Blume ex Graebn.	SSA = SSA-b-1, Sieboldin-b = SSA-b-2	RIP 2
		SSA-b-3, SSA-b-4	lectin
Aizoaceae	Mesembryanthe-mum crystallinum L.	RIP1	RIP 1
	Amaranthus caudatus L.	Amaranthin = ACA	lectin
	Amaranthus cruentus L.	ACL	lectin
	Amaranthus hypochondriacus L. [Syn.: Amaranthus leucocarpus S. Watson]	A. leucocarpus lectin	lectin

Plant Family	Plant Species	Proteins	Classification
	Amaranthus mangostanus L.	Amaramangin	RIP 1
Amaranthaceae	Amaranthus tricolor L.	AAP-27	RIP 1
	Amaranthus viridis L.	Amaranthin	RIP 1
	Beta vulgaris L.	Beetin-27 = BE27, Beetin-29 = BE29, Betavulgin	RIP 1
	Celosia argentea L. [Syn.: Celosia cristata L.]	CCP-25, CCP-27	RIP 1
	Chenopodium album L.	CAP30	RIP 1
	Spinacia oleracea L.	SoRIP1 = BP31	RIP 1
		SoRIP2	RIP 1 candidate
	Aralia elata (Miq.) Seem.	Aralin	RIP 2
Araliaceae	Panax ginseng C.A.Mey	Panaxagin	peculiar RIP 1 candidate/RNase
	Panax quinquefolius L.	Quinqueginsin	peculiar RIP 1 candidate/RNase
	Asparagus officinalis L.	Asparin 1, Asparin 2	RIP 1
	Drimia maritima (L.) Steam [Syn.: Charybdis maritima (L.) Speta]	Charybdin	RIP 1
Asparagaceae	Muscari armeniacum Leichtlin ex Baker	Musarmin 1, Musarmin 2, Musarmin 3, Musarmin 4	RIP 1
	Polygonatum multiflorum (L.) All.	PMRIPm, PMRIPt	RIP 2
	Yucca gloriosa var. tristis Carrière [Syn.: Yucca recurvifolia Salisb.]	Yucca leaf protein = YLP	RIP 1
Basellaceae	Basella rubra L.	Basella RIP 2a, Basella RIP 2b, Basella RIP 3	RIP 1
	Agrostemma githago L.	Agrostin 2, Agrostin 5, Agrostin 6, Agrostin	RIP 1
	Dianthus barbatus L.	Dianthin 29	RIP 1
	Dianthus caryophyllus L.	Dianthin 30, Dianthin 32	RIP 1
	Dianthus chinensis L. [Syn.: Dianthus sinensis Link]	D. sinensis RIP	RIP 1
	Gypsophila elegans M.Bieb.	Gypsophilin	RIP 1

Plant Family	Plant Species	Proteins	Classification
	Silene chalcedonica (L.) E.H.L.Krause [Syn.: Lychnis chalcedonica L.]	Lychnin	RIP 1
	Silene glaucifolia Lag. [Syn.: Petrocoptis glaucifolia (Lag.) Boiss]	Petroglaucin 1, Petroglaucin 2	RIP 1
Caryophyllaceae	Silene laxipruinosa Mayol & Rosselló [Syn.: Petrocoptis grandiflora Rothm.]	Petrograndin	RIP 1
	Saponaria ocymoides L.	Ocymoidin	RIP 1
	Saponaria officinalis L.	Saporin-L1 = SO-L1, Saporin-L2 = SO-L2, Saporin-L3 = SO-L3, Saporin-I = SO-I = SO-4, Saporin-R1 = SO-R1, Saporin-R2 = SO-R2, Saporin-R3 = SO-R3, SO3a, SO3b, Saporin-S5 = Saporin 5 = SO-S5, Saporin-S6 = Saporin 6 = SO-6 = SO-S6, Saporin-S8 = SO-S8, Saporin-S9 = Saporin 9 = SO-S9, SAP-C, SAP-S	RIP 1
	Myosoton aquaticum (L.) Moench [Syn.: Stellaria aquatica (L.) Scop.]	Stellarin	RIP 1
	Stellaria media (L.) Vill.	RIP Q3	RIP 1
	Vaccaria hispanica (Mill.) Rauschert [Syn.: Vaccaria pyramidata Medik.]	Pyramidatin	RIP 1
	Benincasa hispida (Thunb.) Cogn.	Hispin	RIP 1
		α-benincasin, β-benincasin	sRIP 1
	Bryonia cretica subsp. dioica (Jacq.) Tutin. [Syn.: Bryonia dioica L.]	Bryodin 1 = BD1, Bryodin 2, Bryodin-L, Bryodin-R	RIP 1
		BDA	lectin/ RIP 2 like
Cucurbitaceae	Citrullus colocynthis (L.) Schrad .	Colocin 1, Colocin 2	RIP 1
		Foetidissimin	peculiar RIP 2
	Cucurbita foetidissima Kunth	Foetidissimin II	RIP 2
	Cucumis ficifolius A.Rich. [Syn.: Cucumis figarei Delile ex Naudin]	Cucumis figarei RIP = CF-RIP	RIP 1 candidate

Plant Family	Plant Species	Proteins	Classification
	Cucurbita maxima Duchesne	Cucurmoschin	sRIP 1 candidate
	Cucurbita moschata Duchesne [Syn.: Cucurbita moschata (Duchesne ex Lam.) Duchesne ex Poir.]	Cucurmosin, Cucurmosin 2, C. moschata RIP, Moschatin, PRIP 1, PRIP 2	RIP 1
		α-moschin, β-moschin	sRIP 1 candidate
	Cucurbita pepo L.	Pepocin	RIP 1
	Cucurbita pepo var. texana (Scheele) D.S.Decker [Syn.: Cucurbita texana (Scheele) A. Gray]	Texanin	RIP 1
	Gynostemma pentaphyllum (Thunb.) Makino	Gynostemmin	RIP 1
	Lagenaria siceraria (Molina) Standi .	Lagenin	RIP 1 candidate
	Luffa acutangula (L.) Roxb.	Luffaculin-1, Luffaculin-2	RIP 1
		Luffangulin	sRIP 1
		Luffa acutangula fruit lectin	lectin
	Luffa cylindrica (L.) M.Roem [Syn.: Luffa aegyptiaca Mill.]	Luffin, Luffin-a, Luffin-b, α-luffin, β-luffin, LRIP	RIP 1
		Luffacylin, Luffin P1	sRIP 1
		Luffin-S, LuffinS(1), LuffinS(2) = luffin S2, LuffinS(3)	sRIP 1 candidate
	Marah oreganus (Torr. & A. Gray) Howell	MOR-I, MOR-II	RIP 1
	Momordica balsamina L.	Balsamin, MbRIP-1, Momordin II	RIP 1
	Momordica charantia L.	MAP 30, α-momorcharin = α-MC = α-MMC, β-momorcharin = β-MC = β-MMC, δ-momorcharin = δ-MMC, Momordin, Momordin = Momordica charantia inhibitor, Momordin II, Momordin-a, Momordin-b	RIP 1
		γ-momorcharin = γ-MMC, Charantin	sRIP 1
		RIP 1 candidate	RIP 1 candidate
		MCL = M. charantia lectin, anti-H Lectin, Momordica agglutinin, Momordin, protein fraction 1, protein fraction 2	lectin
		MCL = Momordica charantia seed lectin = Momordica charantia lectin, MCL1	RIP 2
	Momordica cochinchinensis Spreng .	Cochinin B, Momorcochin, Momorcochin-S	RIP 1

Plant Family	Plant Species	Proteins	Classification
	Siraitia grosvenorii (Swingle) C.Jeffrey ex A.M.Lu & Zhi Y.Zhang [Syn.: Momordica grosvenorii Swingle]	Momorgrosvin	RIP 1
	Sechium edule (Jacq.) Sw.	Sechiumin	RIP 1
		Sechium edule fruit lectin	lectin
	Trichosanthes anguina L.	Trichoanguin	RIP 1
		SGSL	lectin/ RIP 2 like
	Trichosanthes cordata Roxb.	TCA-I, TCA-II	lectin
	Trichosanthes cucumerina L.	TCSL	lectin/ RIP 2 candidate
	Trichosanthes cucumeroides (Ser.) Maxim.	β-trichosanthin = β-TCS	RIP 1
	Trichosanthes kirilowii Maxim.	α-kirilowin, β-kirilowin, TAP 29, TK-35, Trichobitacin, Trichokirin, Trichomislin = TCM, Trichosanthin = Trichosanthes antiviral protein = TAP = TCS = α-trichosanthin = α-TCS = GLQ223, Trichosanthin, β-trichosanthin = β-TCS, γ-trichosenthin = γ-TCS	RIP 1
		Trichokirin S1, S-Trichokirin, Trichosanthrip	sRIP 1
		TKL-1 = Trichosanthes kirilowii lectin-1	lectin/ RIP 2 candidate
		TK-I, TK-II, TK-III, Trichosanthes kirilowii lectin	lectin
	Trichosanthes kirilowii Maximovicz var. japonica (Miquel) Kitamura	Karasurin-A, Karasurin-B, Karasurin-C	RIP 1
	Trichosanthes lepiniate	Trichomaglin	RIP 1
	Trichosanthes dioica Roxb.	TDSL	lectin/ RIP 2 candidate
	Trichosanthes sp. Bac Kan 8-98	Trichobakin	RIP 1
Cupressaceae	Thuja occidentalis L.	Arborvitae RIP	RIP candidate
	Croton tiglium L.	Crotin I	RIP 1 candidate
		Crotin 2	RIP 1
	Euphorbia characias L.	E. characias lectin	lectin
	Suregada multiflora (A.Juss.) Baill. [Syn.: Gelonium multiflorum A.Juss.]	Gelonin = GAP 31	RIP 1

Plant Family	Plant Species	Proteins	Classification
	Hura Crepitans L.	Hura crepitans RIP, Hura crepitans RIP-5	RIP 1
		Hura crepitans latex lectin	RIP 2
		Crepitin, Hurin, Hura crepitans seed lectin	lectin
Euphorbiaceae	Jatropha curcas L.	Curcin, Curcin 2, Curcin-L, Jc-SCRIP	RIP 1
	Manihot palmata Müll. Arg.	Mapaimin	RIP 1
	Manihot esculenta Crantz. [Syn.: Manihot utilissima Pohl]	Manutin 1, Manutin 2	RIP 1
	Ricinus communis L.	Ricin = crystalline Ricin = Ricin D, Ricin E, RCA = Ricinus communis agglutinin = RCAI = RCA120 = R. communis hemagglutinin = RCB-PHA I, RCAII = RCA60 = RCB-PHA II	RIP 2
	Ricinus communis, USA	Ricin 1, Ricin 2, Ricin 3	RIP 2
	Ricinus communis, India	Ricin I, Ricin II, Ricin III	RIP 2
	Ricinus sanguienus, France	$Ricin_{11}$, $Ricin_{12}$, $Ricin_2$	RIP 2
	Abrus precatorius L.	Abrin, Abrin-a = Abrin C = Abrin-III, Abrin-b, Abrin-c = Abrin A = Abrin-I, Abrin-d, Abrin-II, APA = Abrus precatorius agglutinin = Abrus lectin = AAG, APA-I, APA-II	RIP 2
	Abrus pulchellus Thwaites	Pulchellin, Pulchellin P!, Pulchellin PII, Pulchellin PIII	RIP 2
Fabaceae	Pisum sativum subsp. sativum L. [Syn.: Pisum sativum var. arvense (L.) Poir.]	α-pisavin, β-pisavin	RIP 1
	Pisum sativum var. macrocarpon	Sativin	RIP 1 candidate
Iridaceae	Iris hollandica var. Professor Blaauw	IrisRIP = IRIP, IrisRIP.A1, IrisRIP.A2, IrisRIP.A3	RIP 1
		IRA, IRAb, IRAr	RIP 2
	Clerodendrum aculeatum (L.) Schltdl.	CA-SRI	RIP 1 candidate
Lamiaceae	Clerodendrum inerme (L.) Gaertn .	CIP-29	RIP 1
		CIP-34	RIP 1 candidate
	Leonurus japonicus Houtt.	Leonurin	RIP candidate
	Cinnamomum bodinieri H. Lév	Bodinierin	RIP 2

Plant Family	Plant Species	Proteins	Classification
	Cinnamomum camphora (L.) J.Presl	Camphorin	RIP 1
		Cinnamomin, Cinnamomin 1, Cinnamomin 2, Cinnamomin 3	RIP 2
Lauraceae		Cinphorin	sRIP 2
	Cinnamomum parthenoxylon (Jack) Meisn. [Syn.: Cinnamomum porrectum (Roxb.) Kosterm.]	Porrectin	RIP 2
Malvaceae	Abelmoschus esculentus (L.) Moench	Abelesculin	RIP 1
	Boerhaavia diffusa L.	Boerhaavia inhibitor	RIP 1 candidate
	Bougainvillea spectabilis Willd.	BAP I, Bouganin = Bougainvillea RIP I	RIP 1
	Bougainvillea × buttiana cv. Enid Lancester	BBP-24, BBP-28	RIP 1
Nyctaginaceae	Bougainvillea × buttiana cv. Mahara	BBAP1	RIP 1
	Mirabilis expansa (Ruiz & Pav.) Standl.	ME1, ME2	RIP 1
	Mirabilis jalapa L.	MAP, MAP-2, MAP-3, MAP-4, MAP-S	RIP 1
Olacaceae	Malania oleifera Chun & S. K. Lee	Malanin	lectin/ RIP 2 candidate
	Ximenia americana L.	Riproximin = Rpx, Rpx-I, Rpx-II	RIP 2
Passifloraceae	Adenia digitata (Harv.) Engl.	Modeccin = Modeccin 4B, Modeccin 6B	RIP 2
	Adenia ellenbeckii Harms	A. ellenbeckii lectin	RIP 2 candidate
	Adenia fruticosa Burtt Davy	A. fruticosa lectin	lectin
	Adenia glauca Schinz	A. glauca lectin	RIP 2 candidate
	Adenia goetzei Harms (unresolved name)	A. goetzei lectin	RIP 2
	Adenia keramanthus Harms	A. keramanthus lectin	RIP 2 candidate
	Adenia lanceolata Engl.	Lanceolin	RIP 2
	Adenia racemosa W. J. de Wilde	A. racemosa lectin	lectin
	Adenia spinosa Burtt Davy	A. spinosa lectin	RIP 2 candidate

Plant Family	Plant Species	Proteins	Classification
	Adenia stenodactyla Harms	Stenodactylin	RIP 2
	Adenia venenata Forssk.	A. venenata lectin	RIP 2 candidate
	Adenia volkensii Harms	Volkensin	RIP 2
	Phytolacca americana L.	α-PAP, PAP = Phytolacca americana protein = pokeweed antiviral protein, PAP-I, PAP-II, PAP-III, PAP-C, PAP-H, PAP-R, PAP-S, PAP-S1, PAP-S2	RIP 1
	Phytolacca dioica L.	Diocin 1, Diocin 2, PD-L1, PD-L2, PD-L3, PD-L4, PD-S1, PD-S2, PD-S3	RIP 1
Phytolaccaceae	Phytolacca dodecandra L'Hér,	Dodecandrin, Dodecandrin C	RIP 1
	Phytolacca heterotepala H. Walter	Heterotepalin 4, Heterotepalin 5b	RIP 1
	Phytolacca insularis Nakai	Insularin = PIP = Phytolacca insularis antiviral protein, PIP2 = P. insularis antiviral protein 2	RIP 1
	Hordeum vulgare L.	Barley toxin = Barley translation inhibitor = Barley Protein Synthesis Inhibitor = BPSI = RIP 30, Barley toxin I = Barley translation inhibitor I, Barley toxin II = Barley translation inhibitor II = Barley Protein Synthesis Inhibitor II = BPSI II, Barley toxin III = Barley translation inhibitor III, JIP60	RIP 1
Poaceae	Oryza sativa L.	Oryza sativa RIP	RIP 1
	Secale cereale L.	RPSI	RIP 1
	Triticum aestivum L.	Tritin, Tritin 1, Tritin 2, Tritin 3, Tritin-S, Tritin-L	RIP 1
	Zea mays L.	b-32 = maize RIP = maize proRiP1, Maize proRIP2	RIP 3/ peculiar RIP 1
Ranunculaceae	Eranthis hyemalis (L.) Salisb .	EHL	RIP 2
	Phoradendron californicum Nutt.	PCL	RIP 2
	Viscum album L. (Himalayan mistletoe)	HmRip, HmRip 1, HmRip 2, HmRip 3, HmRip 4	RIP 2
Santalaceae	Viscum album L. (European mistletoe)	ML-I = Mistletoe lectin I = Viscumin = Eu-ML = EML-1 = VAA-I, ML-II = Mistletoe lectin II = VAA-II, ML-III = Mistletoe lectin III = VAA-III	RIP 2
	Viscum articulatum Burm. f.	Articulatin-D	RIP 2
	Viscum coloratum (Kom.) Nakai [Syn.: Viscum album	KML, KML-C, KML-IIL, KML-IIU, VCA	RIP 2

Plant Family	Plant Species	Proteins	Classification
	subsp. coloratum Kom.]		
Solanaceae	Nicotiana tabacum L.	CIP31	RIP-like protein
		TRIP	RIP 1 candidate
Thymelaeaceae	Phaleria macrocarpa (Scheff.) Boerl.	P. macrocarpa RIP	RIP candidate
*Schrot J, Weng A, Melzig MF, et al. Ribosome-inactivating and related proteins. Toxins (Basel). 2015 May 8;7(5):1556-615 .			

[0155] It is part of the invention that the therapeutic combination, the first pharmaceutical composition, the first proteinaceous molecule, the second pharmaceutical composition or the second proteinaceous molecule of the invention is further combined with a covalent conjugate (complex) of a binding molecule or a binding moiety and a saponin, or is further combined with a pharmaceutical compound, an antibody, etc., therewith providing a composition comprising three or more enhancers, pharmaceutically active ingredients, etc., e.g. a conjugate of the invention (e.g. a first proteinaceous molecule and/or a second proteinaceous molecule) combined with a binding moiety complexed with an effector molecule, further combined with a pharmaceutical, which is either or not linked to a saponin, and which is either or not coupled to a ligand such as a targeting immunoglobulin, a domain or a fragment thereof. Furthermore, in an embodiment, the second proteinaceous molecule is provided with two or more effector moieties such as a toxin or immunotoxin, wherein the two or more effector moieties are the same or different.

Exemplary embodiments

[0156] In an embodiment of the invention,the saponin is a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function, in position 23, and wherein the saponin is preferably a saponin that can be isolated from *Gypsophila* or *Saponaria* species, more preferably the saponin is the saponin SO1861 or any of its diastereomers.

[0157] In an embodimentof the invention, the binding site is at least a ligand, such as an immunoglobulin, with at least an effector moiety bound thereto.

[0158] In an embodiment, of the invention, the binding site is an immunoglobulin or at least a binding domain thereof for binding to a cell surface molecule, wherein preferably the cell surface molecule is selected from any of HER2, EGFR, CD20, CD22, Folate receptor 1, CD146, CD56, CD19, CD138, CD27L, PSMA, CanAg, integrin-alphaV, CA6, CD33, mesothelin, Cripto, CD3, CD30, CD33, CD239, CD70, CD123, CD352, DLL3, CD25, ephrinA4, MUC1, Trop2, CEACAM5, HER3, CD74, PTK7, Notch3, FGF2, C4.4A, FLT3, CD71.

[0159] In an embodiment of the invention, a linker is coupled to the glycoside via a cleavable bond, and the ligand is an immunoglobulin, wherein preferably said cleavable bond is subject to cleavage under acidic, reductive, enzymatic or light-induced conditions, and preferably the cleavable bond is a covalent bond, preferably an imine bond, a hydrazone bond, an oxime bond, a 1,3-dioxolane bond or an ester bond, wherein preferably the cleavable bond is a disulfide bond or a peptide bond.

[0160] In an embodiment of the invention, the saponin moiety is a terminal saponin, preferably the saponin SO1861, the linker is a chemical linker covalently linking the saponin to the binding site of the first proteinaceous molecule, and the binding site of the second and first proteinaceous molecule is an

immunoglobulin such as trastuzumab or cetuximab, the linker preferably providing a cleavable bond between the terminal saponin moiety and the binding site comprised by the first proteinaceous molecule.

[0161] An embodiment is the combination for use according to the invention, comprising an endosomal escape enhancing conjugate (i.e. the first proteinaceous molecule) according to the invention and a binding moiety (i.e. the second proteinaceous molecule), wherein the binding moiety comprises at least one effector moiety, wherein the endosomal escape enhancing conjugate and the binding moiety are, independently from one another, able to specifically bind to a target cell-specific surface molecule or structure, thereby inducing receptor-mediated endocytosis of a complex of the endosomal escape enhancing conjugate and the target cell-specific surface molecule, and of the complex of the binding moiety and the target cell-specific surface molecule.

[0162] An embodiment is the combination for use according to the invention, wherein the endosomal escape enhancing conjugate and the binding moiety are able to specifically bind to the same target cell-specific surface molecule or structure.

[0163] An embodiment is the combination for use according to the invention, wherein the endosomal escape enhancing conjugate is able to compete with the binding moiety for binding to the target cell-specific surface molecule or structure.

[0164] An embodiment is the combination for use according to the invention, wherein the endosomal escape enhancing conjugate and the binding moiety are, independently from one another, able to specifically bind to the same epitope.

[0165] An embodiment is the combination for use according to the invention, wherein the endosomal escape enhancing conjugate is able to specifically bind to a first epitope, which is the same as the first epitope to which the binding moiety is able to specifically bind.

[0166] An embodiment is the combination for use according to the invention, wherein the target cell-specific surface molecule or structure is selected from HER2, EGFR, CD20, CD22, Folate receptor 1, CD146, CD56, CD19, CD138, CD27L, PSMA, CanAg, integrin-alphaV, CA6, CD33, mesothelin, Cripto, CD3, CD30, CD33, CD239, CD70, CD123, CD352, DLL3, CD25, ephrinA4, MUC1, Trop2, CEACAM5, HER3, CD74, PTK7, Notch3, FGF2, C4.4A, FLT3, CD71.

[0167] An embodiment is the combination for use according to the invention, wherein the glycoside molecule is a bisdesmosidic triterpene, preferably a saponin.

[0168] An embodiment is the combination for use according to the invention, wherein the glycoside molecule is a bisdesmosidic triterpene saponin.

[0169] An embodiment is the combination for use according to the invention, wherein the saponin is a bisdesmosidic triterpene saponin belonging to the type of a 12,13-dehydrooleanane with an aldehyde function in position 23.

[0170] An embodiment is the combination for use according to the invention, wherein the saponin is a saponin that can be isolated from *Gypsophila* or *Saponaria* species.

[0171] An embodiment is the combination according to the invention, wherein the saponin is a SO1861 or any of its diastereomers.

[0172] An embodiment is the combination for use according to the invention, wherein the at least one

glycoside is bound to the ligand (binding site for the epitope on the cell-surface molecule) via a cleavable bond, wherein preferably said cleavable bond is subject to cleavage under acidic, reductive, enzymatic or light-induced conditions, and wherein the cleavable bond preferably is a disulfide bond or a peptide bond.

[0173] An embodiment is the combination for use according to the invention, wherein the cleavable bond is a covalent bond, preferably an imine bond, a hydrazone bond, an oxime bond, a 1,3-dioxolane bond or an ester bond.

[0174] An embodiment is the combination for use according to the invention, wherein the endosomal escape enhancing conjugate comprises a defined number of glycosides or a defined range.

[0175] An embodiment is the combination for use according to the invention, wherein the defined range is between 1 - 30 glycoside(s), preferably between 1 - 20, more preferably between 1 - 10, more preferably between 1 - 6, more preferably between 2-6, more preferably between 2 - 5, more preferably between 3-5, more preferably between 3-4 glycosides.

[0176] An embodiment is the combination for use according to the invention, wherein the effector moiety is a pharmaceutically active substance, such as a toxin such as a proteinaceous toxin, a drug, a polypeptide or a polynucleotide.

[0177] An embodiment is the combination for use according to the invention, wherein the target cell is a diseased cell or a disease-related cell, preferably a tumor cell or a tumor-associated cell (e.g. tumor vascular cell), or an immune cell (e.g. a T regulatory cell), or an autoimmune cell.

[0178] An embodiment is the combination for use according to the invention, wherein the at least one effector moiety is bound to the binding moiety (second proteinaceous molecule) via a cleavable bond, wherein preferably said cleavable bond is subject to cleavage under acidic, reductive, enzymatic or light-induced conditions, and/or wherein the cleavable bond is a disulfide bond or a peptide bond.

[0179] An embodiment is the combination for use according to the invention, wherein the glycoside (saponin) is capable of augmenting endosomal escape of the effector molecule.

[0180] An embodiment is the combination according to the invention, for use as a medicament.

[0181] An embodiment is the combination for use in a method of treating cancer or an autoimmune disease.

[0182] An embodiment is the combination for use according to the invention, wherein the endosomal escape enhancing conjugate (first proteinaceous molecule) and the binding moiety (second proteinaceous molecule) are to be administered concomitant or sequentially, preferably concomitant.

[0183] The invention is further illustrated by the following examples, which should not be interpreted as limiting the present invention in any way.

EXAMPLES

EXAMPLE A - TREATING A MAMMALIAN TUMOR-BEARING ANIMAL WITH A CONJUGATE OF THE INVENTION IN COMBINATION WITH AN ADC RESULTS IN SURVIVAL AND TUMOR REGRESSION

[0184] Female Balb/c nude mice were injected subcutaneously with a suspension of human A431 tumor cells.

Under the skin of the mice, a human epidermal carcinoma developed in the xenograft animal tumor model. After injection of the tumor cells, the xenograft tumor was allowed to develop to a size of approximately 170-180 mm^3. The A431 tumor cells have the following characteristics: high EGFR expressors, medium CD71 expressors, low HER2 expressors.

[0185] In Table A, the results of the treatment of control mice and tumor-bearing mice are presented. Tumor-bearing mice were treated with the indicated antibodies directed to either human Her2/neu, human EGFR, or human CD71, which are cell-surface receptors on the xenograft tumor. Cetuximab was covalently conjugated with saponin SO1861. The SO1861 was first provided with the linker EMCH (N-ε-maleimidocaproic acid hydrazide), which EMCH is a maleimide-and-hydrazide crosslinker for covalently conjugating sulfhydryls (reduced cysteines of the antibody)) to carbonyls (aldehyde or ketones; here the carbonyl of the aldehyde at position C-23 of the saponin). The saponin-EMCH was covalently coupled to reduced cysteines of the Cetuximab, forming a covalent thio-ether bond between the EMCH and the cysteine side chain. The ADCs trastuzumab-saporin (covalent conjugate) and anti-CD71 mAb (OKT-9, IgG) - saporin (covalent conjugate) were tested for their tumor-attacking efficacy in the mice, measured as tumor volume in time after start of the treatment with the ADCs. The dose of the ADCs was sub-optimal in the tumor model. That is to say, from previous experiments, it was established at which sub-optimal dose of the ADCs no tumor-regression or arrest of tumor growth would be observable.

TABLE A: RESULTS OF TREATING A MAMMALIAN TUMOR-BEARING ANIMAL WITH A CONJUGATE OF THE INVENTION IN COMBINATION WITH AN ADC RESULTS IN SURVIVAL AND TUMOR REGRESSION

Treatment group	**Patient / healthy animal**	**treatment**	**tumor size (volume in mm^3 or '+' for growth, '-' for regression, and 'stable' for growth nor regression)**
1	xenograft	vehicle	2000 mm^3 (death/euthanasia)
2	xenograft	Trastuzumab-saporin	2000 mm^3 (death/euthanasia)
3	xenograft	Anti-CD71 mAb OKT-9 – saporin (covalent conjugate)	2000 mm^3 (death/euthanasia)
4	xenograft	Cetuximab-SO1861 (covalent conjugate)	2000 mm^3 (death/euthanasia)
5	xenograft	Cetuximab	> 170 mm^3, but < 2000 mm^3 (death/euthanasia)
6	**xenograft**	**Trastuzumab-saporin** (covalent conjugate) **+ Cetuximab-SO1861** (covalent conjugate)	**Tumor regression from 180 mm^3 at the start of treatment back to 80 mm^3 (survival)**
7	**xenograft**	**Anti-CD71 mAb OKT-9 - saporin** (covalent conjugate) **+ Cetuximab-SO1861** (covalent conjugate)	**Tumor regression from 180 mm^3 at the start of treatment back to 40 mm^3 (survival)**

[0186] These results demonstrate that the combination therapy of an ADC at a dose which is ineffective when treatment of tumor-bearing mice with the ADC alone is considered (tumor growths, death of the mice is not prevented (euthanasia)), with a conjugate of the invention consisting of a tumor-cell specific receptor targeting

antibody covalently bound to a saponin, i.e. SO1861, the covalent conjugate administered to the mice suffering from cancer, at a non-effective dose when administered alone (tumor growths, death of the mice is not prevented (euthanasia)), provides an efficient and efficacious treatment regimen, expressed as tumors in regression and prolonged survival of the treated animals (beyond the duration of the experiment). The sub-optimal dose of ADC combined with a covalently bound saponin-comprising conjugate of the invention which has no anti-tumor activity when administered alone, thus provide for an effective treatment option for cancer patients, wherein a relative low dose of the ADC is efficacious. A lower dose of ADC bears the promise of less risk for adverse events, or even no side effects at all. In addition, the stimulatory effect of the saponin-bearing conjugate of the invention when the efficacy of the ADC is considered, shows that ADCs which previously have proven to lack efficacy when tumor patient treatment is concerned, may gain renewed attention and value, since ADC efficacy is improved in combination therapy setting, as the current example demonstrated. Reference is made to Table A2 and Table A3, summarizing ADCs which were previously investigated in the human clinical setting, but then were for some ADCs retracted from further clinical investigation. Especially the ADCs for which clinical development was terminated due to observed lack of efficacy and/or due to occurrence of unacceptable adverse event are ADCs which may gain renewed value for cancer patients when combined with a covalently bound saponin-comprising conjugate of the invention, such as the cetuximab-saponin tested.

EXAMPLE B - saponins mixture of Quillaja saponaria comprising QS-21, with endosomal/lysosomal escape enhancing activity

[0187] Scheme I displays the common molecular structure of a series of QS-21 saponins (in part adapted from: Conrado Pedebos, Laércio Pol-Fachin, Ramon Pons, Cilaine V. Teixeira Hugo Verli, Atomic Model and Micelle Dynamics of QS-21 Saponin, Molecules 2014, 19, 3744-3760). A mixture of water-soluble saponins obtained from Quillaja saponaria (Sigma-Aldrich, product No. S4521; Roth, Item No. 6857; InvivoGen, product 'Quil-A') may be applied in the endosomal/lysosomal escape enhancing conjugate, composition, combination of the invention, based on endosomal/lysosomal escape enhancing properties of at least one individual saponin present in the mixture, e.g. QS-21, or based on a combination of two or more of the saponins comprised by the mixture, such as QS-21 and QS-7.

[0188] The inventors demonstrated that the mixture of saponins from Quillaja saponaria at 2,5 microgram/ml dose was capable of enhancing endosomal escape of dianthin, as tested with mammalian tumor cells in a cell-based bioassay. The effector moiety exposed to the cells was dianthin covalently coupled to the ligand EGF: EGF-dianthin. Cells tested were tumor cell lines HeLa for free saponins, and A431, MDA-MB-468, CaSki and A2058 for testing the saponins when covalently coupled to cetuximab.

Example 1

[0189] The 1 target 2-components system (1T2C) is the combination treatment of mAb1-protein toxin and mAb1-SO1861, as illustrated in Figure 14. SO1861-EMCH was conjugated via cysteine residues (Cys) and HSP27BNA oligo was conjugated via lysine residues to cetuximab (monoclonal antibody recognizing and binding human EGFR), both with a DAR 4 resulting in the production of 2 conjugates: cetuximab-(Cys-L-SO1861)^4 and cetuximab-(Lys-L-HSP27BNA)^4. The combination of cetuximab-(Cys-L-SO1861)^4 (intraperitoneal administration, (i.p.)) and cetuximab-(Lys-L-HSP27BNA)^4 (intravenous administration, (i.v.)) was tested in a A431 xenograph 'mouse tumor model for EGFR tumor targeted gene silencing activity. Dosings started at day 12 when tumors reached ~150mm^3 in size and tumor samples were collected at 72h after the first dosing and analysed for HSP27 gene expression compared to control gene mRNA expression levels (reference genes). This revealed that 1 dosing of 50 mg/kg cetuximab-(Cys-L-SO1861)^4+ 25 mg/kg

cetuximab-(Lys-L-HSP27BNA)^4 resulted in a 50% reduction in HSP27 gene expression in the A431 tumors compared to single dosing of cetuximab-(Cys-L-SO1861)^4 or cetuximab-(Lys-L-HSP27BNA)^4 mono therapies (Figure 1). Compared to the vehicle control tumors, a reduction of 40% HSP27 gene silencing was observed. This shows and enables that the combination of cetuximab-conjugated SO1861 + cetuximab-conjugated HSP27BNA oligo, according to the 1T2C invention, induces efficient targeted delivery of a therapeutic antisense oligo nucleotide in the cytoplasm of solid tumor cells, thereby inducing tumor targeted gene silencing, *in vivo*.

[0190] Next, SO1861-EMCH was conjugated via cysteine residues (Cys) to trastuzumab (monoclonal antibody recognizing and binding human HER2), with a DAR 4 resulting in the production of trastuzumab-(Cys-L-SO1861)^4. The combination of trastuzumab-(Cys-L-SO1861)^{4}and trastuzumab-saporin (trastuzumab protein toxin conjugate) was tested in a mouse tumor model (patient derived xenograph tumor model, PDX) with high HER2 expression levels and resistant for trastuzumab mono therapy. The combination, according to the 1T2C invention of 40 mg/kg trastuzumab-(Cys-L-SO1861)^4 (intraperitoneal administration, (i.p.)) + 0.03 (Day1, 8)/ 0.02 (Day 15, 22, 30, 36,43) mg/kg trastuzumab-saporin (intravenous administration, (i.v.)) revealed strong tumor growth inhibition compared to the vehicle control and the 40 mg/kg trastuzumab-(Cys-L-SO1861)^4 or 0.03/0.02 mg/kg trastuzumab-saporin mono therapies (Figure 2). Besides, in tumor bearing mice that were treated with a lower dosing combination (40 mg/kg trastuzumab-(Cys-L-SO1861)^4+ 0.01 mg/kg trastuzumab-saporin) no tumor growth inhibiting activity was observed (Figure 2). This shows and enables that the 1T2C combination of trastuzumab conjugated SO1861 + trastuzumab conjugated protein toxin induces efficient targeted delivery of a therapeutic protein toxin in the cytoplasm of solid tumor cells, thereby inducing tumor cell death and tumor growth inhibition, *in vivo.*

Example 2.

[0191] The 1 target 2-components system (1T2C) is the combination treatment of mAb1-SO1861 and mAb1-protein toxin (Figure 14)

[0192] SO1861-EMCH was conjugated via cysteine residues (Cys) to cetuximab (monoclonal antibody recognizing and binding human EGFR), with a DAR 3,7 (cetuximab-(Cys-L-SO1861)^{3,7}). Cetuximab-(Cys-L-SO1861)^{3,7} was titrated on a fixed concentration of 10 pM cetuximab-saporin (cetuximab, conjugated to the protein toxin, saporin) and targeted protein toxin mediated cell killing on EGFR expressing cells (A431, EGFR^{++}; CaSKi, EGFR^{+}) was determined. This revealed strong cell killing at low concentrations of cetuximab-(Cys-L-SO1861)^{3,7} (A431: IC50= 0,6 nM and Caski IC50= 1 nM; Figure 3A, 3B) whereas cetuximab, cetuximab-(Cys-L-SO1861)^{3,7} or cetuximab + 10 pM cetuximab-saporin could not induce any cell killing activity in EGFR expressing cells. This shows that cetuximab conjugated SO1861 efficiently enhances endosomal escape of the cetuximab conjugated protein toxin (at non-effective concentrations), thereby inducing cell killing of EGFR expressing cells. The cell killing activity in A431 is more effective compared to CaSki correlating with EGFR expression levels in these cell lines. EGFR receptor binding competition between both conjugates within the 1T2C is also observed when cetuximab-(Cys-L-SO1861)^{3,7} concentrations increase, cell killing activity declines due to outcompeting receptor binding and internalization of cetuximab-saporin (Figure 3A, 3B).

[0193] Next, cetuximab-saporin was titrated on a fixed concentration of 75 nM cetuximab-(Cys-L-SO1861)^{3,7} and targeted protein toxin mediated cell killing on EGFR expressing cells was determined. This revealed that 75nM cetuximab-(Cys-L-SO1861)^{3,7} in combination with low concentrations cetuximab-saporin induced already efficient cell killing in EGFR expressing cells (A431 : IC50= 0.4 pM; and CaSKi: (IC50= 2 pM; Figure 3C and 3D), whereas cetuximab-saporin alone or cetuximab-saporin + 75 nM cetuximab showed cell killing only at

high concentrations cetuximab-saporin (IC50= 40 pM, IC50= 1000 pM, resp.) in both cell lines (Figure 3C, 3D). All this shows that relatively low concentrations of cetuximab-saporin can be effective and induce cell killing only in combination with low cetuximab-SO1861 concentrations in high EGFR expressing cells. The receptor competition between both conjugates within the 1T2C system is also observed in the cetuximab-toxin titration treatments when the cell killing activity of cetuximab-saporin with and without 75 nM cetuximab was compared (Figure 3C, 3D).

[0194] Next, cetuximab-(Cys-L-SO1861)^{3,7} was titrated on a fixed concentration of 10 pM cetuximab-saporin and targeted protein toxin mediated cell killing on low EGFR expressing cells or cells without EGFR expression (HeLa, EGFR$^{+/-}$; A2058, EGFR^{-}) was determined. Cells with low (HeLa) or no (A2058) EGFR expression were not sensitive at all for any combination of cetuximab-(Cys-L-SO1861)^{3,7}+ 10 pM cetuximab-saporin (HeLa: IC50> 1000 nM; A2058: IC50> 1000 nM; Figure 4A, 4B). This shows that in the absence of sufficient EGFR receptor expression, effective intracellular delivered SO1861 concentrations are not optimal (threshold) to induce endosomal protein toxin escape and toxin-mediated cell killing. Next, cetuximab-saporin was titrated on a fixed concentration of 75 nM cetuximab-(Cys-L-SO1861)^{3,7} and targeted protein toxin mediated cell killing on low (HeLa) or no (A2058) EGFR expressing cells was determined. Low EGFR expressing cells (HeLa) showed cell killing only at high cetuximab-saporin concentrations in combination with 75 nM cetuximab-(Cys-L-SO1861)^{3,7} (HeLa: IC50= 60 pM), Figure 4C), whereas A2058 cells (EGFR^{-}) are not sensitive at any of the tested concentrations (A2058: IC50> 10.000 pM; Figure 4D). All this shows that cells with low or no EGFR receptor expression are not susceptible for the combination of cetuximab-(Cys-L-SO1861)^{3,7} + cetuximab-saporin, due to a lack of sufficient EGFR receptor that facilitates the antibody-mediated delivery of sufficient SO1861 within the endolysosomal compartments, to facilitate the escape of the protein toxin.

[0195] Next, SO1861-EMCH was conjugated via cysteine residues (Cys) to trastuzumab (monoclonal antibody recognizing and binding human HER2, with a DAR 4, (trastuzumab-(Cys-L-SO1861)^{4}). Trastuzumab-(Cys-L-SO1861)^{4} was titrated on a fixed concentration of 50 pM trastuzumab-saporin (trastuzumab, conjugated to the protein toxin, saporin) and targeted protein toxin mediated cell killing on HER2 expressing cells (SK-BR-3, HER2^{++}) was determined. This revealed strong cell killing at low concentrations of trastuzumab-(Cys-L-SO1861)^{4} (SK-BR-3: IC50= 0,8 nM; Figure 5A) whereas equivalent concentrations trastuzumab, trastuzumab-(Cys-L-SO1861)^{4} or trastuzumab + 50 pM trastuzumab-saporin could not induce any cell killing activity in HER2 expressing cells. This shows that trastuzumab conjugated SO1861 efficiently enhances endosomal escape of the trastuzumab conjugated protein toxin (at non-effective concentrations), thereby inducing cell killing of HER2 expressing cells. The receptor competition between both conjugates within the 1T2C is also observed when trastuzumab-(Cys-L-SO1861)^{4} concentrations increase, cell killing activity declines due to outcompeting receptor binding and internalization of trastuzumab-saporin (Figure 5A).

[0196] Next, trastuzumab-saporin was titrated on a fixed concentration of 2,5 nM trastuzumab-(Cys-L-SO1861)^{4}, according to the invention and targeted protein toxin mediated cell killing on HER2 expressing cells was determined. This revealed that 2,5 nM trastuzumab-(Cys-L-SO1861)^{4} in combination with low concentrations trastuzumab-saporin induced already efficient cell killing in HER2 expressing cells (SK-BR-3: IC50= 2 pM; Figure 5B), whereas trastuzumab-saporin alone or trastuzumab-saporin + 2,5 nM trastuzumab showed cell killing only at high concentrations trastuzumab-saporin (Figure 5B). All this shows that relatively low concentrations of trastuzumab-saporin can be effective and induce cell killing only in combination with low trastuzumab-(Cys-L-SO1861)^{4} concentrations in high HER2 expressing cells.

[0197] Next, trastuzumab-(Cys-L-SO1861)^{4} was titrated on a fixed concentration of 50 pM trastuzumab-saporin, according to the invention and targeted protein toxin mediated cell killing on low HER2 expressing

cells (A431 HER2$^{+/-}$) or cells without HER2 expression (JIMT-1: HER2^{+}; MDA-MB-468: HER2^{-}) was determined. Cells with low or no HER2 expression were not sensitive at all for any combination of trastuzumab-(Cys-L-SO1861)^{4}+ 50 pM trastuzumab-saporin (JIMT-1: IC50> 1000 nM; MDA-MB-468: IC50> 1000 nM; Figure 6A, 6B). This shows that in the absence of sufficient HER2 receptor expression, effective intracellular delivered SO1861 concentrations are not optimal (threshold) to induce endosomal protein toxin escape and toxin-mediated cell killing. Next, trastuzumab-saporin was titrated on a fixed concentration of 2,5 nM trastuzumab-(Cys-L-SO1 861)^{4} and targeted protein toxin mediated cell killing on low or no HER2 expressing cells was determined. Low HER2 expressing cells (JIMT-1) showed cell killing only at high trastuzumab-saporin concentrations in combination with 2,5 nM trastuzumab-(Cys-L-SO1861)^{4} (JIMT-1: IC50> 10.000 pM; Figure 6C), whereas MDA-MB-468 cells (HER2) are not sensitive at any of the tested concentrations (MDA-MB-468: IC50> 10.000 pM; Figure 6D).

[0198] All this shows that cells with low or no HER2 receptor expression are not susceptible for the combination of trastuzumab-(Cys-L-SO1861)^{3,7} + trastuzumab-saporin, due to a lack of sufficient HER2 receptor that facilitates the antibody-mediated delivery of sufficient SO1861 within the endolysosomal compartments, to facilitate the escape of the protein toxin.

Example 3

[0199] In order to show that the activity of the 1T2C system is driven by the acidification of the endolysosomal compartments, the 1T2C system , according to the invention was tested in combination with an endosomal acidification inhibitor, chloroquine. Trastuzumab-saporin was titrated in combination with 5 nM trastuzumab-(Cys-L-SO1861)^{4} in combination with or without chloroquine. Trastuzumab-saporin + 5 nM trastuzumab-(Cys-L-SO1861)^{4} showed a strong cell killing activity in high HER2 expressing cells (SK-BR-3, HER2^{++}; IC50= 0.2 pM;), however, trastuzumab-saporin + 5 nM trastuzumab-(Cys-L-SO1861)^{4} + 0.5 µM chloroquine resulted in strong inhibition of the 1T2C cell killing activity in SK-BR-3 (HER2^{++}) cells (IC50= 40pM). This shows that activity of the antibody conjugated SO1861 is reduced/blocked when acidification of endoslysomes is prohibited (Figure 8A). Same results were derived with the 1T2C combination, according to the invention of cetuximab-saporin + 5 nM cetuximab-(Cys-L-SO1861)^{3,8} (IC50= 1 pM) compared with cetuximab-saporin + 5 nM cetuximab-(Cys-L-SO1861)^{3,8}+ 0.5 µM chloroquine (IC50= 200 pM) in EGFR expressing cells (A431, EGFR^{++}; Figure 8B).

Example 4

[0200] The 1 target 2-components system (1T2C) can also be the combination treatment of mAb1-SO1861 and mAb1-antisense BNA oligo nucleotide as illustrated in Figure 15. For this we used an antisense BNA oligonucleotide against the mRNA of a cancer specific target gene (upregulated in cancer cells), heat shock protein 27 (HSP27). Upon release into the cytoplasm the antisense BNA recognizes and binds the mRNA encoding for HSP27, targeting the mRNA for destruction thereby depleting the HSP27 mRNA expression within the cancer cell. HSP27BNA was conjugated to cetuximab with a DAR4 (Cetuximab-(Lys-L-HSP27BNA)^{4}) and tested in combination with cetuximab-(Cys-L-SO1861)^{3,8}for enhanced HSP27 gene silencing activity in EGFR expressing cells (A431, EGFR^{++}) and non-expressing cells (A2058, EGFR), according to the invention (Figure 15). Cetuximab-(Cys-L-SO1861)^{3,8}+ 100 nM cetuximab-(Lys-L-HSP27BNA)^{4} showed strong HSP27 gene silencing in EGFR expressing cells (A431: IC50= nM, Figure 9A), whereas cetuximab-(Cys-L-SO1861)^{3,8} alone did not show any gene silencing activity. In A2058 cells (EGFR) no gene silencing activity was observed in the

1T2C combination (Figure 9B). Next, cetuximab-(Lys-L-HSP27BNA)^4 + 76.9 nM Cetuximab-(Cys-L-SO1861)^{3,8} show strong HSP27 gene silencing activity in EGFR expressing cells (A431: IC50= 4 nM, Figure 9C), whereas cetuximab-(Lys-L-HSP27BNA)^4 or cetuximab-(Cys-L-SO1861)^{3,8} or the combination of Cetuximab-(Lys-L-HSP27BNA)^4 + 77 nM cetuximab did not reveal any significant gene silencing activity (IC50> 100nM). When the experiment was performed in EGFR non-expressing cells (A2058) no gene silencing activity was observed in the 1T2C combination (IC50>100 nM; Figure 9D). All this shows that the 1T2C system efficiently delivers an antisense BNA oligo to the cytoplasm of high EGFR expressing cells, thereby inducing mRNA degradation of the BNA target mRNA resulting in target gene silencing.

Example 5

[0201] The 1 target 2-components system (1T2C) can also be the combination treatment of mAb1-(scaffold(-SO1861)^n)^n and mAb1-protein toxin as illustrated in Figure 16. Dendron(-L-SO1861)^4 was conjugated to cetuximab via cysteine residues (Cys) conjugation with a DAR3,9 and cetuximab-Cys-(dendron(-L-SO1861)^4)^{3,9} was tested for enhanced cell killing activity in combination with an anti-EGFR antibody-protein toxin conjugate (cetuximab-saporin) in EGFR expressing cells (MDA-MB-468). Cetuximab-Cys-(dendron(-L-SO1861)^4)^{3,9} + 10 pM cetuximab-saporin efficiently induces toxin-mediated cell killing in high EGFR expressing cells (IC50= 0.4 nM; Figure 10A), whereas this was not induced by cetuximab-Cys-(dendron(-L-SO1861)^4)^{3,9} or cetuximab + 10 pM cetuximab-saporin or cetuximab (Figure 10A). This shows that according to the 1T2C invention, cetuximab conjugated dendron(-L-SO1861)^4 efficiently enhances endosomal escape of the cetuximab conjugated protein toxin (at non-effective concentrations), thereby inducing cell killing of high HER2 expressing cells.Similar 1T2C experiments were performed in cells that express low levels of EGFR (HeLa, EGFR$^{+/-}$) and this revealed no cell killing activity when the 1T2C combination, according ot the invention was used (IC50> 100pM; Figure 10B) indicating that in the absence of sufficient EGFR receptor expression, effective intracellular SO1861 concentrations are not optimal (threshold) to induce cytoplasmic delivery of the protein toxin that results in toxin-mediated cell killing.

[0202] Next, dendron(-L-SO1861)^4 was conjugated to the anti-HER2 antibody, trastuzumab via cysteine conjugation (Cys) with a DAR4, trastuzumab-Cys-(dendron(-L-SO1861)^4)^4 and tested for enhanced cell killing activity in combination with an anti-HER2 antibody-protein toxin conjugate (trastuzumab-saporin) in HER2 expressing cells (SK-BR-3, HER2^{++}). Trastuzumab-Cys-(dendron(-L-SO1861)^4)^4 + 50 pM trastuzumab-saporin efficiently induces toxin-mediated cell killing (IC50= 2 nM, Figure 10C), whereas this was not induced by trastuzumab-Cys-(dendron(-L-SO1861)^4)^4 or trastuzumab + 50nM trastuzumab-saporin or trastuzumab (Figure 10C). This shows that trastuzumab conjugated dendron(-L-SO1861)^4 efficiently enhances endosomal escape of the trastuzumab conjugated protein toxin (at non-effective concentrations), thereby inducing cell killing of high HER2 expressing cells. Similar experiments in cells that express low levels of HER2 (JIMT-1, HER2$^{+/-}$) revealed no activity of Trastuzumab-Cys-(dendron(-L-SO1861)^4)^4 + 50 pM trastuzumab-saporin (IC50> 200 nM; Figure 10D) indicating that in the absence of sufficient HER2 receptor expression, effective intracellular SO1861 concentrations are not optimal (threshold) to induce endosomal protein toxin escape and toxin-mediated cell killing.

Example 6

[0203] The clinical approved ADC, trastuzuzmab-emtansine (T-DM1) is a conjugate of the anti-Her2 antibody, trastuzumab and the small molecule toxin emtansine (DAR3.5). T-DM1 was titrated in combination with

trastuzumab-(Cys-L-SO1861)^{4} and compared with the antibody protein toxin conjugate, trastuzumab-saporin + Trastuzumab-(Cys-L-SO1861)^{4}, according to the invention. Whereas trastuzumab-saporin + 2.5 nM trastuzumab-(Cys-L-SO1861)^{4} showed enhanced activity compared to Trastuzumab-saporin + 2.5 nM trastuzumab or trastuzumab-saporin alone (IC50= 2 pM, Figure 11), T-DM1 + 25.6 nM trastuzumab-(Cys-L-SO1861)^{4} showed no enhanced cell killing activity (IC50> 100 pM; Figure 11). This shows that the 1T2C system, according to the invention, cannot enhance the delivery of an antibody small molecule conjugate, since small molecules can already passively cross (endolysosomal) membranes.

Example 7

[0204] Figure 12A-D displays the relative cell viability when trastuzumab (Figure 12A), cetuximab (Figure 12B) or T-DM1 (Figure 12C), unconjugated protein toxins, saporin, dianthin and saporin conjugated to a (non-cell binding) IgG antibody (Figure 12D) are administrated to various cancer cell lines SK-BR-3, JIMT-1, MDA-MB-468, A431, CaSki, HeLa, A2058.

[0205] Trastuzumab and cetuximab do not or hardly influence cell viability when exposed to most of the cell lines, with some effect on cell growth inhibition via blocking the function of the HER2 growth factor receptor when trastuzumab is exposed to SK-BR-3 cells at relatively high dose and with some effect on cell growth inhibition via blocking the function of the EGFR growth factor receptor when cetuximab is exposed to MDA-MB-468 cells at relatively high dose.

[0206] TDM-1, or ado-trastuzumab emtansine, is a targeted therapy approved by the U.S. Food and Drug Administration to treat: HER2-positive metastatic breast cancer that has previously been treated with Herceptin (chemical name: trastuzumab) and taxane chemotherapy; early-stage HER2-positive breast cancer after surgery if residual disease was found after neoadjuvant (before surgery) treatment with Herceptin and taxane chemotherapy. The TDM-1 is a combination of Herceptin (Trastuzumab) and the chemotherapy medicine emtansine. Figure 12C shows that the TDM-1 results in decreased cell viability for all cell lines tested at >1000 pM concentrations

[0207] The free toxins saporin and dianthin and the toxin saporin coupled to a control IgG with no affinity for any of the cell surface molecules on the cell lines tested, do not or hardly have any influence on cell viability over a wide range of concentrations toxin tested, up to 100.000 pM (Figure 12D).

Example 8

[0208] The 1 target 2-components system (1T2C) can also be the combination treatment of mAb1-QSmix (mixture of saponins from Quillaja Saponaria) and mAb1-protein toxin.

[0209] QSmix-EMCH was conjugated via cysteine residues (Cys) to cetuximab (monoclonal antibody recognizing and binding human EGFR), with a DAR 4.1 (cetuximab-(Cys-L-QSmix)^{4,1}). Cetuximab-(Cys-L-QSmix)^{4,1} was titrated on a fixed concentration of 10 pM cetuximab-saporin or 10pM cetuximab-dianthin and targeted protein toxin mediated cell killing on A431 (EGFR^{++}), CaSKi (EGFR^{+}) and A2058 (EGFR) cells was determined. This revealed strong cell killing at low concentrations of cetuximab-(Cys-L-QSmix)$^{4.1}$ + 10 pM cetuximab-saporin or 10pM cetuximab-dianthin in A431 (EGFR^{++}) and CaSKi (EGFR^{+}) cells (A431: IC50= 3 nM, Figure 13A; CaSKi: IC50= 1nM, Figure 13B) whereas all control treatments could not induce any cell killing in EGFR expressing cells. In cells that do not express EGFR (A2058; EGFR^{-}) no HSP27 gene silencing is

observed with the combination, according to the invention (IC50> 1000 nM; Figure 13C). This shows that cetuximab conjugated QS21 mix efficiently enhances endosomal escape of the cetuximab conjugated protein toxin (at non-effective concentrations), thereby inducing cell killing only in EGFR expressing cells.

Example 9

dendron(-L-SO1861)^n synthesis (Figure 18, 19, 20)

materials and methods

Abbreviations

[0210]

DCM
: dichloromethane

DIPEA
: *N,N*-diisopropylethylamine

DMF
: *N,N*-dimethylformamide

EDCI.HCl
: 3-((Ethylimino)methyleneamino)-N,N-dimethylpropan-1-aminium chloride

EMCH.TFA
: N-(ε-maleimidocaproic acid) hydrazide, trifluoroacetic acid salt

min
: minutes

r.t.
: retention time

TCEP
: tris(2-carboxyethyl)phosphine hydrochloride

Temp
: temperature

TFA
: trifluoroacetic acid

THF
: tetrahydrofuran

Analytical methods

LC-MS method 1, [1]

[0211] Apparatus: Agilent 1200 Bin. Pump: G1312A, degasser; autosampler, ColCom, DAD: Agilent G1316A,

210, 220 and 220-320 nm, PDA: 210-320 nm, MSD: Agilent LC/MSD G6130B ESI, pos/neg 100-1000; ELSD Alltech 3300 gas flow 1.5 ml/min, gas temp: 40°C; column: Waters XSelect™ CSH C18, 30×2.1mm, 3.5µm, Temp: 35 °C, Flow: 1 mL/min, Gradient: t_0 = 5% A, $t_{1.6min}$ = 98% A, t_{3min} = 98% A, Posttime: 1.3 min, Eluent A: 0.1% formic acid in acetonitrile, Eluent B: 0.1% formic acid in water.

LC-MS method 2, [2]

[0212] Apparatus: Agilent 1260 Bin. Pump: G7112B, Multisampler, Column Comp, DAD: Agilent G7115A, 210, 220 and 220-320 nm, PDA: 210-320 nm, MSD: Agilent LC/MSD G6130B ESI, mass ranges depending on the molecular weight of the product:

[A]pos/neg 100-1000

[B]pos/neg 100-1400

; ELSD Alltech 3300 gas flow 1.5 ml/min, gas temp: 40°C; column: Waters XSelect™ C18, 30×2.1mm, 3.5µm, Temp: 40 °C, Flow: 1 mL/min, Gradient: t_0 = 5% A, $t_{1.6min}$ = 98% A, t_{3min} = 98% A, Posttime: 1.3 min, Eluent A: 0.1% formic acid in acetonitrile, Eluent B: 0.1% formic acid in water.

LC-MS method 3, [3]

[0213] Apparatus: Waters IClass; Bin. Pump: UPIBSM, SM: UPISMFTN with SO; UPCMA, PDA: UPPDATC, 210-320 nm, SQD: ACQ-SQD2 ESI, pos/neg 800-1500; ELSD: gaspressure 40 psi, drift tube temp: 50°C; column: Waters XSelect™ CSH C18, 50×2.1mm, 2.5µm Temp: 25°C, Flow: 0.6 mL/min, Gradient: t_0 = 5% A, $t_{2.0min}$ = 98% A, $t_{2.7min}$ = 98% A, Posttime: 0.3 min, Eluent A: acetonitrile, Eluent B: 10 mM ammonium bicarbonate in water (pH=9.5).

LC-MS method 4, [4]

[0214] Apparatus: Waters IClass; Bin. Pump: UPIBSM, SM: UPISMFTN with SO; UPCMA, PDA: UPPDATC, 210-320 nm, SQD: ACQ-SQD2 ESI, pos/neg 1500-2500; ELSD: gas pressure 40 psi, drift tube temp: 50°C; column: Waters XSelect™ CSH C18, 50×2.1mm, 2.5µm Temp: 25°C, Flow: 0.6 mL/min, Gradient: t_0 = 15% A, $t_{2.0min}$ = 60% A, $t_{2.7min}$ = 98% A, Posttime: 0.3 min, Eluent A: acetonitrile, Eluent B: 10 mM ammonium bicarbonate in water (pH=9.5).

LC-MS method 5, [5]

[0215] Apparatus: Waters IClass; Bin. Pump: UPIBSM, SM: UPISMFTN with SO; UPCMA, PDA: UPPDATC, 210-320 nm, SQD: ACQ-SQD2 ESI, mass ranges depending on the molecular weight of the product:

[A]pos/neg 1500-2500

^{B}neg 2000-3000

; ELSD: gaspressure 40 psi, drift tube temp: 50°C; column: Acquity C18, 50×2.1mm, 1.7µm Temp: 60°C, Flow: 0.6 mL/min, Gradient: t_0 = 2% A, $t_{5.0min}$ = 50% A, $t_{6.0min}$ = 98% A, Posttime: 1.0 min, Eluent A: acetonitrile, Eluent B: 10 mM ammonium bicarbonate in water (pH=9.5).

Preparative methods

Preparative MP-LC method 1, [1]

[0216] Instrument type: Reveleris™ prep MPLC; column: Waters XSelect™ CSH C18 (145×25 mm, 10µ); Flow: 40 mL/min; Column temp: room temperature; Eluent A: 10 mM ammoniumbicarbonate in water pH = 9.0); Eluent B: 99% acetonitrile + 1% 10 mM ammoniumbicarbonate in water; Gradient: t_{0min} = 5% B, t_{1min} = 5% B, t_{2min} = 10% B, t_{17min} = 50% B, t_{18min} = 100% B, t_{23min} = 100% B; Detection UV: 210, 225, 285 nm.

Preparative MP-LC method 2, [2]

[0217] Instrument type: Reveleris™ prep MPLC; Column: Phenomenex LUNA C18(3) (150×25 mm, 10µ); Flow: 40 mL/min; Column temp: room temperature; Eluent A: 0.1% (v/v) Formic acid in water, Eluent B: 0.1% (v/v) Formic acid in acetonitrile; Gradient: t_{0min} = 5% B, t_{1min} = 5% B, t_{2min} = 10% B, t_{17min} = 50% B, t_{18min} = 100% B, t_{23min} = 100% B; Detection UV : 210, 225, 285 nm.

Preparative LC-MS method 1, [3]

[0218] MS instrument type: Agilent Technologies G6130B Quadrupole; HPLC instrument type: Agilent Technologies 1290 preparative LC; Column: Waters XSelect™ CSH (C18, 100×30mm, 10µ); Flow: 25 ml/min; Column temp: room temperature; Eluent A: 100% acetonitrile; Eluent B: 10 mM ammonium bicarbonate in water pH=9.0; lin. gradient depending on the polarity of the product:

$^{A}t_0$ = 20% A, t_{2min} = 20% A, $t_{8.5min}$ = 60% A, t_{10min} = 100% A, t_{13min} = 100% A

$^{B}t_0$ = 5% A, t_{2min} = 5% A, $t_{8.5min}$ = 40% A, t_{10min} = 100% A, t_{13min} = 100% A

$^{C}t_0$ = 10% A, t_{2min} = 10% A, $t_{8.5min}$ = 50% A, t_{10min} = 100% A, t_{13min} = 100% A

; Detection: DAD (220-320 nm); Detection: MSD (ESI pos/neg) mass range: 100 - 800; Fraction collection based on DAD.

Preparative LC-MS method 2, [4]

[0219] MS instrument type: Agilent Technologies G6130B Quadrupole; HPLC instrument type: Agilent Technologies 1290 preparative LC; column: Waters XBridge Shield (C18, 150×19mm, 5µ); Flow: 25 ml/min; Column temp: room temperature; Eluent A: 100% acetonitrile; Eluent B: 10 mM ammonium bicarbonate in water pH=9.0; lin. gradient: t_0= 5% A, $t_{2.5min}$ = 5% A, t_{11min} = 40% A, t_{13min} = 100% A, t_{17min} = 100% A; Detection: DAD (220-320 nm); Detection: MSD (ESI pos/neg) mass range: 100 - 800; Fraction collection based DAD

Flash chromatography

[0220] Grace Reveleris X2® C-815 Flash; Solvent delivery system: 3-piston pump with auto-priming, 4 independent channels with up to 4 solvents in a single run, auto-switches lines when solvent depletes; maximum pump flow rate 250 mL/min; maximum pressure 50bar (725psi); Detection: UV 200-400nm, combination of up to 4 UV signals and scan of entire UV range, ELSD; Column sizes: 4-330g on instrument, luer type, 750g up to 3000g with optional holder.

SO1861-EMCH synthesis (Figure 18 and 24)

[0221] To SO1861 (121 mg, 0.065 mmol) and EMCH.TFA (110 mg, 0.325 mmol) was added methanol (extra dry, 3.00 mL) and TFA (0.020 mL, 0.260 mmol). The reaction mixture stirred at room temperature. After 1.5 hours the reaction mixture was subjected to preparative MP-LC.[1] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (120 mg, 90%) as a white fluffy solid. Purity based on LC-MS 96%.

LRMS (m/z): 2069 $[M-1]^{1-}$

LC-MS r.t. (min): 1.08[4]

Dendron(-L-SO1861)^{4}synthesis (Figure 19)

Intermediate 1:

di-tert-butyl (((6-azidohexanoyl)azanediyl)bis(ethane-2,1-diyl))dicarbamate

[0222] 6-azidohexanoic acid (0.943 g, 6.00 mmol), EDCI.HCl (1.21 g, 6.30 mmol) and Oxyma Pure (0.938 g, 6.60 mmol) were dissolved in DMF (10.0 mL) and the mixture was stirred for 5 min. Next a solution of di-tert-butyl (azanediylbis(ethane-2,1-diyl))dicarbamate (1.82 g, 6.00 mmol) in DMF (5.00 mL) was added and the reaction mixture was stirred at room temperature. After 5 hours the reaction mixture was evaporated *in vacuo* and the residue was dissolved in ethyl acetate (50 mL). The resulting solution was washed with 1N potassium bisulphate solution (50 mL), saturated sodium bicarbonate solution (2 × 50 mL) and brine (50 mL), dried over Na_2SO_4, filtered and evaporated *in vacuo.* The residue was purified by flash chromatography (ethyl acetate - heptane gradient, 10:90 rising to 100:0) to give the title compound (2.67 g, 100%) as a white solid. Purity based on LC-MS 98%.

LRMS (m/z): 287/343/465 $[M-155/M-99/M+23]^{1+}$

LC-MS r.t. (min): 2.02^{2A}

Intermediate 2:

N,N-bis(2-aminoethyl)-6-azidohexanamide dihydrochloride

[0223] To di-tert-butyl (((6-azidohexanoyl)azanediyl)bis(ethane-2,1-diyl))dicarbamate (2.66 g, 6.00 mmol) was added HCl in isopropanol (5-6 N, 20.0 mL, 110 mmol) and the reaction mixture was stirred at room temperature. After 4 hours, the reaction mixture was evaporated *in vacuo* and the resulting crude product was co-evaporated with DCM (3 × 20 mL) to give the crude title product (1.49 g, 79%) as a white solid.

[0224] LRMS (m/z): 243 $[M+1]^{1+}$

Intermediate 3:

tetra-tert-butyl ((5S,5'S)-((((6-azidohexanoyl)azanediyl)bis(ethane-2,1-diyl))bis(azanediyl))bis(6-oxohexane-6,1,5-triyl))tetracarbamate

[0225] To a solution of N,N-bis(2-aminoethyl)-6-azidohexanamide dihydrochloride (1.19 g, 3.76 mmol) in DMF (30.0 mL) and DIPEA (2.62 mL, 15.1 mmol) was added Boc-Lys(Boc)-ONp (3.69 g, 7.90 mmol) and the mixture was stirred at room temperature overnight. The reaction mixture was evaporated *in vacuo* and the residue was dissolved in ethyl acetate (100 mL). The resulting solution was washed with 1N potassium bisulphate solution (100 mL) and saturated sodium bicarbonate solution (5 × 100 mL), dried over Na_2SO_4, filtered and evaporated *in vacuo.* The residue was purified by flash chromatography (DCM - methanol/DCM (1/9, v/v) gradient 100:0 rising to 0:100) to give the give the title product (3.07 g, 91%) as a slightly yellowish solid. Purity based on LC-MS 94%.

LRMS (m/z): 800/900/922 $[M-99/M+1/M+23]^{1+}$

LC-MS r.t. (min): 2.17^{2A}

Intermediate 4:

4-nitrophenyl 3-(acetylthio)propanoate

[0226] 4-Nitrophenyl trifluoroacetate (5.17 g, 22.0 mmol) and 3-(Acetylthio)propionic Acid (2.96 g, 20.0 mmol) were dissolved in DCM (50.0 mL). Next, DIPEA (6.97 mL, 40.0 mmol) was added and the reaction mixture was

stirred at room temperature overnight. The reaction mixture was evaporated *in vacuo* and the residue was dissolved in ethyl acetate (50 mL). The resulting solution was washed with 1N potassium bisulphate solution (50 mL), saturated sodium bicarbonate solution (5 × 50 mL) and brine (50 mL), dried over Na_2SO_4, filtered and evaporated *in vacuo.* The residue was purified by flash chromatography (DCM - methanol/DCM (1/9, v/v) gradient 100:0 rising to 0:100) to give the give the title product (4.90 g, 91%) as a slightly yellowish solid. Purity based on LC-MS 99%.

LRMS (m/z): 292 $[M+23]^{1+}$

LC-MS r.t. (min): 1.94[2A]

Intermediate 5:

(S)-2,6-diamino-N-(2-(6-azido-N-(2-((S)-2,6-diaminohexanamido)ethyl)hexanamido)ethyl)hexanamide tetrahydrochloride

[0227] tetra-tert-butyl ((5S,5'S)-((((6-azidohexanoyl)azanediyl)bis(ethane-2,1-diyl))bis(azanediyl))bis(6-oxohexane-6,1,5-triyl))tetracarbamate (1.80 g, 2.00 mmol) was dissolved in HCl in isopropanol (5-6N, 50.0 ml, 275 mmol) and the reaction mixture was stirred at room temperature overnight. The reaction mixture was evaporated *in vacuo* and the resulting crude product was co-evaporated with DCM (3 × 20 mL) to give the crude title product as a white solid.

LRMS (m/z): 250 $[M+2]^{2+}$, 500 $[M+1]^{1+}$

Intermediate 6:

(2S)-2,6-bis[3-(acetylsulfanyl)propanamido]-N-[2-(6-azido-N-{2-[(2S)-2,6-bis[3-(acetylsulfanyl)propanamido]hexanamido]ethyl}hexanamido)ethyl]hexanamide

[0228] To a solution of (S)-2,6-diamino-N-(2-(6-azido-N-(2-((S)-2,6- diaminohexanamido)ethyl)hexanamido) ethyl)hexanamide tetrahydrochloride (1.29 g, 2.00 mmol) in DMF (30 mL) and DIPEA (3.48 mL, 20.0 mmol) was added 4-nitrophenyl 3-(acetylthio)propanoate (2.26 g, 8.40 mmol) and the reaction mixture was stirred at room temperature over the weekend. The reaction mixture was evaporated *in vacuo* and the residue was dissolved in DCM/methanol (95:5, v/v, 100 mL). The resulting solution was washed with 1N potassium bisulphate solution (100 mL), 1 N sodium hydroxide solution (3 × 100 mL) and brine (100 mL), dried over Na_2SO_4, filtered and evaporated *in vacuo.* The residue was purified by flash chromatography (DCM - methanol/DCM (1/9, v/v) gradient 100:0 rising to 0:100) to give the title product (1.33 g, 65%) as a white solid. On LC-MS an impurity (15%) was found with m/z values corresponding to the product with one deprotected thioacetate group. The impurity was formed during or after work-up. Purity based on LC-MS 85%.

LRMS (m/z): 510 $[M+2]^{2+}$, 1019/1041 $[M+1/M+23]^{1+}$

LC-MS r.t. (min): 1.86[2B]

Intermediate 7:

N,N'-((9S,19S)-14-(6-aminohexanoyl)-1-mercapto-9-(3-mercaptopropanamido)-3,10,18-trioxo-4,11,14,17-tetraazatricosane-19,23-diyl)bis(3-mercaptopropanamide) formate

[0229] Scaffold 2 (102 mg, 0.100 mmol) was dissolved in methanol (1.00 ml). Next, a freshly prepared 1 N Sodium hydroxide solution (0.440 ml, 0.440 mmol) was added and the reaction mixture was stirred at room temperature. After 30 min a 1.0 M trimethylphosphine solution in THF (0.500 ml, 0.500 mmol) was added and the resulting mixture was stirred at room temperature. After 30 min the reaction mixture was evaporated *in vacuo* and co-evaporated with methanol (2 × 10 mL). The residue was dissolved in a mixture of methanol/water (9:1, v/v, 1.00 mL) and the resulting solution was subjected to preparative MP-LC.[2] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (75.6 mg, 87%) as a colorless sticky oil. Purity based on LC-MS 96%.

LRMS (m/z): 513 $[M+2]^{2+}$, 825 $[M+1]^{1+}$

LC-MS r.t. (min): 1.42^{2A}

Intermediate 8:

dendron(-L-SO1861)$_4$-amine

[0230] N, N'-((9S,19S)-14-(6-aminohexanoyl)-1-mercapto-9-(3-mercaptopropanamido)-3,10,18-trioxo-4,11,14,17-tetraazatricosane-19,23-diyl)bis(3-mercaptopropanamide) formate (2.73 mg, 3.13 µmol) was dissolved in a mixture of 20 mM NH_4HCO_3 with 0.5 mM TCEP/acetonitrile (3:1, v/v, 3.00 mL). Next, SO1861-EMCH (29.2 mg, 0.014 mmol) was added and the reaction mixture was stirred at room temperature. After 1.5 hours the reaction mixture was subjected to preparative LC-MS.[3B] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (12.3 mg, 43%) as a white fluffy solid. Purity based on LC-MS 97%.

LRMS (m/z): 1517 $[M-6]^{6-}$, 1821 $[M-5]^{5-}$, 2276 $[M-4]^{4-}$

LC-MS r.t. (min): 4.39^{5A}

Intermediate 9:

dendron(-L-SO1861)$_4$-azide

[0231] Dendron(SO1861)$_4$-amine (6.81 mg, 0.748 µmol) and 2,5-dioxopyrrolidin-1-yl 1-azido-3,6,9,12-

tetraoxapentadecan-15-oate (2.90 mg, 7.48 µmol) were dissolved in DMF(1.00 mL). Next, DIPEA (1.302 µL, 7.48 µmol) was added and the mixture was shaken for 1 min and left standing at room temperature. After 2 hours the reaction mixture was subjected to preparative LC-MS.[3C] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (5.86 mg, 84%) as a white fluffy solid. Purity based on LC-MS 90%.

LRMS (m/z): 2344 $[M-4]^{4-}$

LC-MS r.t. (min): 4.78[5B]

Intermediate 10:

dendron(-L-SO1861)^4-maleimide1

[0232] Dendron$(SO1861)_4$-amine (8.12 mg, 0.891 µmol) and 2,5-dioxopyrrolidin-1-yl 1-(2,5-dioxo-2,5-dihydro-1H-pyrrol-1-yl)-3,6,9,12-tetraoxapentadecan-15-oate (3.94 mg, 8.91 µmol) were dissolved in DMF(1.00 mL). Next, DIPEA (1.55 µL, 8.91 µmol) was added and the mixture was shaken for 1 min and left standing at room temperature. After 3 hours the reaction mixture was subjected to preparative LC-MS.[3C] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (6.76 mg, 80%) as a white fluffy solid. Purity based on LC-MS 66%.

LRMS (m/z): 2358 $[M-4]^{4-}$

LC-MS r.t. (min): 2.13[6C]

Intermediate 11:

dendron(-L-SO1861)^4-maleimide2

[0233] Scaffold 2 (5.10 mg, 5.00 µmol) was dissolved in methanol (100 µL). Next, a freshly prepared 1 N Sodium hydroxide solution (22.0 µL, 22.0 µmol) was added and the mixture was shaken for 1 min and left standing at room temperature. After 30 min a 1.0 M trimethylphosphine solution in THF (25.0 µL, 25.0 µmol) was added and the resulting mixture was shaken for 1 min and left standing at room temperature. After 30 min the reaction mixture was evaporated *in vacuo* and co-evaporated with methanol (2 × 5 mL). The resulting residue was dissolved in a mixture of 20 mM NH_4HCO_3 with 0.5 mM TCEP/acetonitrile (3:1, v/v, 3.242 mL). From this solution, directly, 1000 µL was added to SO1861-EMCH (14.4 mg, 6.94 µmol, 4.5 equiv. compared to the scaffold) and the mixture was shaken for 1 min and left standing at room temperature. After 10 min the reaction mixture was lyophilized overnight. To the resulting residue 2,5-Dioxopyrrolidin-1-yl 3-(2-(2-(3-(2,5-dioxo-2h-pyrrol-1(5h)-yl)propanamido)ethoxy)ethoxy)propanoate (5.84 mg, 0.014 mmol) and DMF (1.00 mL) were added. Next, DIPEA (2.39 µL, 0.014 mmol) was added and the suspension was shaken for 1 min and left standing at room temperature. After 2 hours the reaction mixture was subjected to preparative LC-MS.[3C] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to

give the title compound (10.9 mg, 85%) as a white fluffy solid. Purity based on LC-MS 80%.

LRMS (m/z): 2354 $[M-4]^{4-}$

LC-MS r.t. (min): 4.16^{5B}

Dendron(-L-SO1861)^8 synthesis (Figure 20)

Intermediate 1:

tert-butyl N-[(1S)-1-{[(1S)-1-{[2-(6-azido-N-{2-[(2S)-2,6-bis[(2S)-2,6-bis({[(tert-butoxy)carbonyl]amino})hexanamido]hexanamido]ethyl}hexanamido)ethyl]carbamoyl}-5-[(2S)-2,6-bis({[(tert-butoxy)carbonyl]amino})hexanamido]pentyl]carbamoyl}-5-{[(tert-butoxy)carbonyl]amino)pentyl]carbamate

[0234] (S)-2,6-diamino-N-(2-(6-azido-N-(2-((S)-2,6-diaminohexanamido)ethyl)hexanamido)ethyl)hexanamide tetrahydrochloride (964 mg, 1.50 mmol) was dissolved in DMF (25.0 mL) and triethylamine (2.08 mL, 15.0 mmol). Next, Boc-Lys(Boc)-ONp (3.36 g, 7.18 mmol) was added and the reaction mixture was stirred at room temperature overnight. The reaction mixture was evaporated *in vacuo* and the residue was purified by flash chromatography (DCM - methanol/DCM (1/9, v/v) gradient 100:0 rising to 0:100) to give the title product (2.71 g, 100%) as a white solid. Purity based on LC-MS 97%.

LRMS (m/z): 807 $[M-198]^{2+}$

LC-MS r.t. (min): 2.35^{2B}

Intermediate 2:

(2S,2'S)-N,N'-((5S,15S,22S)-22,26-diamino-10-(6-azidohexanoyl)-15-((S)-2,6-diaminohexanamido)-6,14,21-trioxo-7,10,13,20-tetraazahexacosane-1,5-diyl)bis(2,6-diaminohexanamide) octahydrochloride

[0235] Intermediate 1 (2.71 g, 1.50 mmol) was dissolved in HCl in isopropanol (5-6N, 25.0 ml, 138 mmol) and the reaction mixture was stirred at room temperature overnight. Next, the reaction mixture was evaporated *in vacuo* and the resulting crude product was co-evaporated with DCM (3 × 20 mL) to give the crude title product as a white solid.

LRMS (m/z): 203/254 $[M-200/M+4]^{4+}$, 338 $[M+3]^{3+}$, 507 $[M+2]^{2+}$, 1012 $[M+1]^{1+}$

Intermediate 3:

(2S)-2,6-bis[3-(acetylsulfanyl)propanamido]-N-[(1S)-1-{[2-(6-azido-N-{2-[(2S)-2,6-bis[(2S)-2,6-bis[3-(acetylsulfanyl)propanamido]hexanamido]hexanamido]ethyl}hexanamido)ethyl]carbamoyl}-5-[(2S)-2,6-bis[3-(acetylsulfanyl)propanamido]hexanamido]pentyl]hexanamide

[0236] To (2S,2'S)-N,N'-((5S,15S,22S)-22,26-diamino-10-(6-azidohexanoyl)-15-((S)-2,6-diaminohexanamido)-6,14,21-trioxo-7,10,13,20-tetraazahexacosane-1,5-diyl)bis(2,6-diaminohexanamide) octahydrochloride (300 mg, 0.230 mmol) was added DMF (20.0 mL), triethylamine (320 µl, 2.30 mmol) and 4-nitrophenyl 3-(acetylthio)propanoate (595 mg, 2.21 mmol). The resulting suspension was sonicated at 60 °C for 30 min and left stirring at room temperature overnight. The reaction mixture was evaporated *in vacuo* and the residue was purified by first performing flash chromatography (DCM - methanol/DCM (1/9, v/v) gradient 100:0 rising to 0:100), followed by preparative MP-LC[2] to give the title product (70 mg, 15%) as a white solid. Purity based on LC-MS 100%.

LRMS (m/z): 685 $[M+3]^{3+}$

LC-MS r.t. (min): 1.91[2A]

Intermediate 4:

(2S)-N-[(1S)-1-{[2-(6-amino-N-{2-[(2S)-2,6-bis[(2S)-2,6-bis(3-sulfanylpropanamido)hexanamido]hexanamido]ethyl}hexanamido)ethyl]carbamoyl}-5-[(2S)-2,6-bis(3-sulfanylpropanamido)hexanamido]pentyl]-2,6-bis(3-sulfanylpropanamido)hexanamide formate

[0237] Scaffold 4 (10.0 mg, 4.87 µmol) was dissolved methanol (200 µL). Next, a freshly prepared 1 N Sodium hydroxide solution (42.9 µL, 0.043 mmol) was added and the resulting mixture was shaken for 1 min and left standing at room temperature. After 30 min a 1.0 M trimethylphosphine solution in THF (24.4 µL, 0.024 mmol) was added and the resulting mixture was shaken for 1 min and left standing at room temperature. After 30 min the reaction mixture was diluted with water (1 mL) and directly subjected to preparative MP-LC.[2] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (4.02 mg, 48%) as a white fluffy solid.

LRMS (m/z): 564 $[M+3]^{3+}$, 846 $[M+2]^{2+}$

LC-MS r.t. (min): 1.54[2C]

Intermediate 5:

dendron(-L-SO1861)^{8}-amine

[0238] Scaffold 5 (0.52 mg, 0.299 µmol) and SO1861-EMCH (29.2 mg, 0.014 mmol) were dissolved in a mixture of 20 mM NH_4HCO_3 with 0.5 mM TCEP/acetonitrile (3:1, v/v, 1.00 mL) and the resulting mixture was

shaken for 1 min and left standing at room temperature. After 30 min TCEP (0.30 mg, 1.05 µmol) was added and the reaction mixture was shaken for 1 min. Next, the mixture was directly subjected to preparative LC-MS.[3B] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (2.17 mg, 40%) as a white fluffy solid. Purity based on LC-MS 97%.

LRMS (m/z): 2282 $[M-8]^{8-}$, 2607 $[M-7]^{7-}$

LC-MS r.t. (min): 4.41[5A]

Example 10

SO1861-trifunctional linker-BNAoligo synthesis (Figure 21)

Materials and methods

Trifunctional linker

[0239] Trifunctional linker (DBCO, TCO, maleimide) was ordered at Bio-Synthesis Inc. (Lewisville, Texas).

HSP27BNA oligo

[0240] HSP27BNA(-thiol) oligos (sequence 5'-GGCacagccagtgGCG-3')[39] were ordered at Bio-synthesis Inc. (Lewisville, Texas)

Intermediate 1:

SO1861-azide

[0241] To SO1861 60 mg, 0.032 mmol)) and 1-azido-3,6,9,12-tetraoxapentadecane-15-hydrazide (39.3 mg, 0.129 mmol) was added methanol (extra dry, 1.00 mL) and TFA (9.86 µl, 0.129 mmol) and the reaction mixture was shaken for 1 min and left standing at room temperature. After 2 hours the reaction mixture was subjected to preparative MP-LC.[1] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (58.4 mg, 84%) as a white fluffy solid. Purity based on LC-MS 100%.

LRMS (m/z): 2150 $[M-1]^{1-}$

LC-MS r.t. (min): 1.10[3B]

Intermediate 2:

SO1861-trifunctional linker

[0242] SO1861-azide (45 mg, 0.021 mmol) and trifunctional linker (26.5 mg, 0.022 mmol) were dissolved in DMF (2.50 mL) and the resulting mixture was shaken for 1 min and left standing at room temperature. After 30 min the reaction mixture was subjected to preparative LC-MS.[3C] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (58.4 mg, 84%) as a white fluffy solid. Purity based on LC-MS 89%.

LRMS (m/z): 1677 $[M-2]^{2-}$

LC-MS r.t. (min): 2.54[6A]

Intermediate 3:

(E)-1-(4-((2-(6-(2,5-dioxo-2,5-dihydro-1H-pyrrol-1-yl)hexanoyl)hydrazineylidene)methyl)benzamido)-N-(4-(6-methyl-1,2,4,5-tetrazin-3-yl)benzyl)-3,6,9,12-tetraoxapentadecan-15-amide

[0243] To 1-(4-formylbenzamido)-N-(4-(6-methyl-1,2,4,5-tetrazin-3-yl)benzyl)-3,6,9,12-tetraoxapentadecan-15-amide (28.0 mg, 0.048 mmol) and EMCH.TFA (24.5 mg, 0.072 mmol) was added methanol (extra dry, 2.00 mL) and TFA (11.1 µL, 0.145 mmol) and the reaction mixture stirred at 50 °C. After 30 min the reaction mixture was evaporated *in vacuo* and the resulting residue was purified by MP-LC.[1] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (33.4 mg, 88%) as a bright purple fluffy solid. Purity based on LC-MS 92%.

LRMS (m/z): 394 $[M+2]^{2+}$, 789 $[M+1]^{1+}$

LC-MS r.t. (min): 1.28[7A]

Intermediate 4:

methyltetrazine-BNA oligo

[0244] To HSP27 BNA oligo disulfide (70.0 mg, 0.012 mmol) was dissolved in 20 mM NH_4HCO_3 (20.0 mL). Next, TCEP (14.3 mg, 0.050 mmol) was added and the reaction mixture was shaken for 1 min and left standing at room temperature. The reaction mixture was filtered by using a centrifugal filter with a molecular weight cut-off of 3000 Da (5000 × g for 30 min). Next, a solution of 20 mM NH_4HCO_3 with 2.5 mM TCEP (20.0 mL) was added to the residue solution and the resulting mixture was filtered again under the same conditions

described above. The residue solution was diluted with 20 mM NH_4HCO_3 (30.0 mL) and the resulting mixture was added to a solution of (E)-1-(4-((2-(6-(2,5-dioxo-2,5-dihydro-1H-pyrrol-1-yl)hexanoyl)hydrazineylidene)methyl)benzamido)-N-(4-(6-methyl-1,2,4,5-tetrazin-3-yl)benzyl)-3,6,9,12-tetraoxapentadecan-15-amide (14.8 mg, 18.8 µmol) in acetonitrile (10.0 mL). The reaction mixture was shaken for 1 min and left standing at room temperature. After 30 min the reaction mixture was frozen and lyophilized over the weekend to yield the crude title product as a pink fluffy solid. To the crude product was added a solution of 20 mM NH_4HCO_3 (20.0 mL) and the resulting suspension was filtered over a 0.45 µm syringe filter. The filtrate was filtered using a centrifugal filter with the same conditions as described above. Next, again a solution of 20 mM NH_4HCO_3 (20.0 mL) was added to the residue solution and the resulting mixture was again filtered by using a centrifugal filter with the same conditions described above. The residue solution was diluted with 20 mM NH_4HCO_3 (20.0 mL) and the resulting mixture was lyophilized overnight to yield the title product (90.0 mg, 115%) as a pink fluffy solid. Purity based on LC-MS 91%.

LRMS (m/z): 1631 $[M-4]^{4-}$; 2174 $[M-3]^{3-}$

LC-MS r.t. (min): 0.73[78]

Intermediate 5:

SO1861-trifunctional linker-BNA oligo

[0245] Methyltetrazine-BNA oligo (90.0 mg, 0.014 mmol) and SO1861-trifunctional linker (48.6 mg, 0.014 mmol) were dissolved in a mixture of water/acetonitrile (4:1, v/v, 12.0 mL). The reaction mixture was shaken for 1 min and left standing at room temperature. After 15 min the mixture was subjected to preparative LC-MS.[4A] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (82.0 mg, 60%) as a white fluffy solid. Purity based on LC-MS 92% (2 peaks with both m/z values corresponding to the title compound).

LRMS (m/z): 1641 $[M-6]^{6-}$, 1970 $[M-5]^{5-}$

LC-MS r.t. (min): 3.24 and 3.40[6B]

Intermediate 1:

SO1861-azide

[0246] To SO1861 60 mg, 0.032 mmol)) and 1-azido-3,6,9,12-tetraoxapentadecane-15-hydrazide (39.3 mg, 0.129 mmol) was added methanol (extra dry, 1.00 mL) and TFA (9.86 µl, 0.129 mmol) and the reaction mixture was shaken for 1 min and left standing at room temperature. After 2 hours the reaction mixture was subjected to preparative MP-LC.[1] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (58.4 mg, 84%) as a white fluffy solid. Purity based on LC-MS 100%.

LRMS (m/z): 2150 $[M-1]^{1-}$

LC-MS r.t. (min): 1.10[3B]

Intermediate 2:

SO1861-trifunctional linker

[0247] SO1861-azide (45 mg, 0.021 mmol) and trifunctional linker (26.5 mg, 0.022 mmol) were dissolved in DMF (2.50 mL) and the resulting mixture was shaken for 1 min and left standing at room temperature. After 30 min the reaction mixture was subjected to preparative LC-MS.[3C] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (58.4 mg, 84%) as a white fluffy solid. Purity based on LC-MS 89%.

LRMS (m/z): 1677 $[M-2]^{2-}$

LC-MS r.t. (min): 2.54[6A]

Intermediate 3:

(E)-1-(4-((2-(6-(2,5-dioxo-2,5-dihydro-1H-pyrrol-1-yl)hexanoyl)hydrazineylidene)methyl)benzamido)-N-(4-(6-methyl-1,2,4,5-tetrazin-3-yl)benzyl)-3,6,9,12-tetraoxapentadecan-15-amide

[0248] To 1-(4-formylbenzamido)-N-(4-(6-methyl-1,2,4,5-tetrazin-3-yl)benzyl)-3,6,9,12-tetraoxapentadecan-15-amide (28.0 mg, 0.048 mmol) and EMCH.TFA (24.5 mg, 0.072 mmol) was added methanol (extra dry, 2.00 mL) and TFA (11.1 µL, 0.145 mmol) and the reaction mixture stirred at 50 °C. After 30 min the reaction mixture was evaporated *in vacuo* and the resulting residue was purified by MP-LC.[1] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (33.4 mg, 88%) as a bright purple fluffy solid. Purity based on LC-MS 92%.

LRMS (m/z): 394 $[M+2]^{2+}$, 789 $[M+1]^{1+}$

LC-MS r.t. (min): 1.28[7A]

Intermediate 4:

methyltetrazine-BNA oligo

[0249] To HSP27 BNA oligo disulfide (70.0 mg, 0.012 mmol) was dissolved in 20 mM NH_4HCO_3 (20.0 mL). Next, TCEP (14.3 mg, 0.050 mmol) was added and the reaction mixture was shaken for 1 min and left standing at room temperature. The reaction mixture was filtered by using a centrifugal filter with a molecular weight cut-off of 3000 Da (5000 × g for 30 min). Next, a solution of 20 mM NH_4HCO_3 with 2.5 mM TCEP (20.0 mL) was added to the residue solution and the resulting mixture was filtered again under the same conditions described above. The residue solution was diluted with 20 mM NH_4HCO_3 (30.0 mL) and the resulting mixture was added to a solution of (E)-1-(4-((2-(6-(2,5-dioxo-2,5-dihydro-1H-pyrrol-1-yl)hexanoyl)hydrazineylidene)methyl)benzamido)-N-(4-(6-methyl-1,2,4,5-tetrazin-3-yl)benzyl)-3,6,9,12-tetraoxapentadecan-15-amide (14.8 mg, 18.8 µmol) in acetonitrile (10.0 mL). The reaction mixture was shaken for 1 min and left standing at room temperature. After 30 min the reaction mixture was frozen and lyophilized over the weekend to yield the crude title product as a pink fluffy solid. To the crude product was added a solution of 20 mM NH_4HCO_3 (20.0 mL) and the resulting suspension was filtered over a 0.45 µm syringe filter. The filtrate was filtered using a centrifugal filter with the same conditions as described above. Next, again a solution of 20 mM NH_4HCO_3 (20.0 mL) was added to the residue solution and the resulting mixture was again filtered by using a centrifugal filter with the same conditions described above. The residue solution was diluted with 20 mM NH_4HCO_3 (20.0 mL) and the resulting mixture was lyophilized overnight to yield the title product (90.0 mg, 115%) as a pink fluffy solid. Purity based on LC-MS 91%.

LRMS (m/z): 1631 $[M-4]^{4-}$; 2174 $[M-3]^{3-}$

LC-MS r.t. (min): 0.73[78]

Intermediate 5:

SO1861-trifunctional linker-BNA oligo

[0250] Methyltetrazine-BNA oligo (90.0 mg, 0.014 mmol) and SO1861-trifunctional linker (48.6 mg, 0.014 mmol) were dissolved in a mixture of water/acetonitrile (4:1, v/v, 12.0 mL). The reaction mixture was shaken for 1 min and left standing at room temperature. After 15 min the mixture was subjected to preparative LC-MS.[4A] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (82.0 mg, 60%) as a white fluffy solid. Purity based on LC-MS 92% (2 peaks with both m/z values corresponding to the title compound).

LRMS (m/z): 1641 $[M-6]^{6-}$, 1970 $[M-5]^{5-}$

LC-MS r.t. (min): 3.24 and 3.40[6B]

Example 11

SO1861-BNA oligo conjugation

[0251] HSP27 BNA oligo disulfide (1.10 mg, 0.187 µmol) was dissolved in 20 mM NH_4HCO_3 with 1.0 mM

TCEP (500 µL) and the mixture was shaken for 1 min and left standing at room temperature. After 1 hour the reaction mixture was filtered by using a centrifugal filter with a molecular weight cut-off of 3000 Da (14000 × g for 30 min). The residue solution was diluted with 20 mM NH_4HCO_3 with 1.0 mM TCEP (500 µL) and the resulting mixture was filtered again under the same conditions described above. The residue solution was diluted with 20 mM NH_4HCO_3/acetonitrile (3:1, v/v, 1.00 mL) and the resulting mixture was added to SO1861-EMCH (3.54 mg, 0.375 µmol). The reaction mixture was shaken for 1 min and left standing at room temperature. After 10 min the reaction mixture was subjected to preparative LC-MS.[4A] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (1.25 mg, 85%) as a white fluffy solid. Purity based on LC-MS 100%.

LRMS (m/z): 1561 $[M-5]^{5-}$, 1951 $[M-4]^{4-}$

LC-MS r.t. (min): 2.46[6B]

Dendron(SO1861)^4-BNA oligo conjugation (Figure 22)

[0252] HSP27 BNA oligo disulfide (1.1 mg, 0.187 µmol) was dissolved in 20 mM NH_4HCO_3 with 1.0 mM TCEP (500 µL) and the mixture was shaken for 1 min and left standing at room temperature. After 1 hour the reaction mixture was filtered by using a centrifugal filter with a molecular weight cut-off of 3000 Da (14000 × g for 30 min). The residue solution was diluted with 20 mM NH_4HCO_3 with 1.0 mM TCEP (500 µL) and the resulting mixture was filtered again under the same conditions described above. The residue solution was diluted with 20 mM NH_4HCO_3/acetonitrile (3:1, v/v, 1.0 mL) and the resulting mixture was added to dendron(SO1861)4-maleimide1 (3.54 mg, 0.375 µmol). The reaction mixture was shaken for 1 min and left standing at room temperature. After 10 min the reaction mixture was subjected to preparative LC-MS.[4A] Fractions corresponding to the product were immediately pooled together, frozen and lyophilized overnight to give the title compound (1.25 mg, 85%) as a white fluffy solid. Purity based on LC-MS 94%

LRMS (m/z): 1896 $[M-8]^{8-}$, 2167 $[M-7]^{7-}$

LC-MS r.t. (min): 3.77[6B]

Dendron(NEM)^4 synthesis (Figure 23)

[0253] To benzyl bis(2-((S)-2,6-bis(3-mercaptopropanamido)hexanamido)ethyl)carbamate (1.69 mg, 2.00 µmol) and N-Ethylmaleimide (1.05 mg, 8.40 µmol) was added a mixture of 20 mM NH_4HCO_3/acetonitrile (3:1, v/v, 2.00 mL) and the reaction mixture was stirred at room temperature. After 2 hours, the reaction mixture was lyophilized overnight. The resulting residue was purified by using preparative LC-MS[3A] to give the title compound (1.53 mg, 57%) as a white fluffy solid. Purity based on LC-MS 98%.

LRMS (m/z): 1346 $[M+1]^{1+}$

LC-MS r.t. (min): 1.43[3A]

Example 12

A431 mouse tumor mouse model and vitro and vivo gene silencing studies

Materials and methods

[0254] HSP27BNA with linkers oligos (sequence 5'-GGCacagccagtgGCG-3')[39] were ordered at Bio-Synthesis Inc. (Lewisville, Texas)

***Vitro* RNA isolation and gene expression analyses**

[0255] RNA from cells was isolated and analysed according to standard protocols (Biorad)

***Vivo* mouse tumor model**

[0256] The mouse study was performed at CrownBio (China) according to standard protocols and procedures. Model: Subcutaneous A431 Human Epidermoid Carcinoma Xenograft Model in female BALB/c nude Mice. Tumor volume was monitored and tumor samples were collected for gene silencing analysis (see below)

RNA isolation and gene expression analyses of tumor samples from mice

[0257] Total RNA was isolated from tumors using TriZol (Thermo Fisher) according to the manufacturer's instructions. Isolated RNA was resuspended in nuclease-free water (NFW). RNA samples were checked for their RNA integrity on the gel. To prepare cDNA, first 100 ng total RNA was mixed with Random Hexamers (Qiagen; final concentration 2 µM) in a final volume of 12.5 µl in NFW, denatured for 5min at 70°C and immediately cooled on ice. Next, 7.5 µl of a cDNA synthesis mix was added, consisting of 4 µl 5xRT Buffer (Promega), 0.4 µl 25mM dNTPs (Promega), 1 µl 200 U/µL MMLV RT-enzyme (Promega), 0.5 µL 40 U/µL RNAse Inhibitor (Promega) and 1.6 µL NFW. The following cDNA synthesis protocol was used: 1) 10 minutes 25°C 2) 60 minutes 37°C 3) 5 minutes 85°C 4) ∞ 4°C

[0258] For a single qPCR reaction the following mix was prepared: 1 µL cDNA, 0.05 µL forward primer (250 µM), 0.05 µL reverse primer (250 µM), 8.9 µl LNFW, 10 µl SYBR Green (Bio-Rad). The following qPCR protocol was used: 1 cycle: 5 minutes 95°C, 40 cycles: 15s 95°C + 30s 60°C.

[0259] HSP27 gene expression was calculated using $2^{-(CtHSP27- GEOMEAN(Ctref1;Ctref2 ;Ctref3;Ctref4))}$, where ref1, ref2, ref3 and ref4 are the reference genes IMMT, EIF2S2, GUSB and UBC for the analysis of the tumors. Four reference genes were chosen based on the performance of a GeNORM analysis among nine reference genes tested to choose the most ideal and stable reference gene for this tumor samples. To do so, qPCR results were imported in Qbase+ software program by which two quality measures are calculated: the coefficient of variation of the normalized reference gene expression levels (V); and the geNorm stability M-value (M)1. A reference gene with an M<0.2 and a V<0.15 is considered very stable. Based on this analysis IMMT and EIF2S2 were chosen as the most stable reference genes. However, UBC and GUSB were also

added to the group of reference genes to further enhance the accuracy of the normalization. Each sample was analyzed as technical triplicate on a CFX96 Real-Time qPCR machine (Bio-Rad).

Table 1. Primers used in qPCR are shown below:

Gene	Primer	Sequence (5'-3')
UBC	Forward	CAGCCGGGATTTGGGTCG
	Reverse	CACGAAGATCTGCATTGTCAAGT
GUSB	Forward	TGCGTAGGGACAAGAACCAC
	Reverse	GGGAGGGGTCCAAGGATTTG
IMMT	Forward	AACCCACACCTGCACTTTCA
	Reverse	TTTTCCTGTTGTGCAAGGCG
EIF2S2	Forward	CCACAAGTCGTCCGAGTAGG
	Reverse	GGAGATGTTTGGGCTGACGA
HSP27	Forward	GCAGTCCAACGAGATCACCA
	Reverse	TAAGGCTTTACTTGGCGGCA

Example 13

Conjugate synthesis

Monoclonal antibodies, SO1861, QS saponins

[0260] Trastuzumab (Herceptin®), cetuximab (Erbitux®)and T-DM1 (Kadcyla®) were purchased from the pharmacy (Charite, Berlin). CD71 monoclonal antibody was purchased from BioCell (Okt9, #BE0023). SO1861 was isolated and purified by Analyticon Discovery GmbH from raw plant extract obtained from *Saponaria officinalis L. Quillaja Saponaria* saponin extract (QSmix) was purchased (Sigma Aldrich, #S4521)

materials

[0261] Trastuzumab (Tras, Herceptin®, Roche), Cetuximab (Cet, Erbitux®, Merck KGaA), Tris(2-carboxyethyl)phosphine hydrochloride (TCEP, 98 %, Sigma-Aldrich), 5,5-Dithiobis(2-nitrobenzoic acid) (DTNB, Ellman's reagent, 99%, Sigma-Aldrich), Zeba™ Spin Desalting Columns (2 mL, Thermo-Fisher), NuPAGE™ 4-12% Bis-Tris Protein Gels (Thermo-Fisher), NuPAGE™ MES SDS Running Buffer (Thermo-Fisher), Novex™ Sharp Pre-stained Protein Standard (Thermo-Fisher), PageBlue™ Protein Staining Solution (Thermo-Fischer), Pierce™ BCA Protein Assay Kit (Thermo-Fisher), N-Ethylmaleimide (NEM, 98 %, Sigma-Aldrich), 1 ,4-Dithiothreitol (DTT, 98 %, Sigma-Aldrich), Sephadex G25 (GE Healthcare), Sephadex G50 M (GE Healthcare), Superdex 200P (GE Healthcare), Isopropyl alcohol (IPA, 99.6 %, VWR), Tris(hydroxymethyl)aminomethane (Tris, 99%, Sigma-Aldrich), Tris(hydroxymethyl)aminomethane hydrochloride (Tris.HCL, Sigma-Aldrich), L-Histidine (99%, Sigma-Aldrich), D-(+)-Trehalose dehydrate (99%, Sigma-Aldrich), Polyethylene glycol sorbitan monolaurate (TWEEN 20, Sigma-Aldrich), Dulbecco's Phosphate-Buffered Saline (DPBS, Thermo-Fisher), Guanidine hydrochloride (99%, Sigma-Aldrich), Ethylenediaminetetraacetic acid disodium salt dihydrate (EDTA-Na_2, 99 %, Sigma-Aldrich), sterile filters 0.2 µm and 0.45 µm (Sartorius), Succinimidyl 4-(N-

maleimidomethyl)cyclohexane-1-carboxylate (SMCC, Thermo-Fisher), Dianthin-Cys (Dia-Cys, Dianthin mutant with a single C-terminal cysteine function, Proteogenix), Vivaspin T4 and T15 concentrator (Sartorius), Superdex 200PG (GE Healthcare), Tetra(ethylene glycol) succinimidyl 3-(2-pyridyldithio)propionate (PEG_4-SPDP, Thermo-Fisher), HSP27 BNA disulfide oligonucleotide (Biosynthesis), [O-(7-Azabenzotriazol-1-yl)-N,N,N,N-tetramethyluronium-hexafluorphosphat] (HATU, 97%, Sigma-Aldrich), Dimethyl sulfoxide (DMSO, 99%, Sigma-Aldrich), N-(2-Aminoethyl)maleimide trifluoroacetate salt (AEM, 98 %, Sigma-Aldrich), L-Cysteine (98.5 %, Sigma-Aldrich), deionized water (DI) was freshly taken from Ultrapure Lab Water Systems (MilliQ, Merck), Nickel-nitrilotriacetic acid agarose (Ni-NTA agarose, Protino), Glycine (99.5 %, VWR), 5,5-Dithiobis(2-nitrobenzoic acid (Ellman's reagent, DTNB, 98 %, Sigma-Aldrich), S-Acetylmercaptosuccinic anhydride Fluorescein (SAMSA reagent, Invitrogen) Sodium bicarbonate (99.7 %, Sigma-Aldrich), Sodium carbonate (99.9 %, Sigma-Aldrich), PD MiniTrap desalting columns with Sephadex G-25 resin (GE Healthcare), PD10 G25 desalting column (GE Healthcare), Zeba Spin Desalting Columns in 0.5, 2, 5, and 10 mL (Thermo-Fisher), Vivaspin Centrifugal Filters T4 10 kDa MWCO, T4 100 kDa MWCO, and T15 (Sartorius), Biosep s3000 aSEC column (Phenomenex), Vivacell Ultrafiltration Units 10 and 30 kDa MWCO (Sartorius), Nalgene Rapid-Flow filter (Thermo-Fisher),

Methods

MALDI-TOF-MS

[0262] Matrix-assisted laser desorption/ionization time of flight (MALDI-TOF) spectra were recorded on a MALDI-Mass Spectrometer (Bruker Ultrafex III). Typically, the sample dissolved in MilliQ water in nanomole to micromole range was spotted on the target (MTP 384 target plate polished steel T F, Bruker Daltons) using either super-DHB (99%, Fluka) or sinapinic acid (SA, 99%, Sigma-Aldrich) as the matrix that was dissolved in acetonitrile (MADLI-TOF-MS tested, Sigma) / 0.1 % TFA (7:3 v/v) via the dried-droplet-method. PepMix (Peptide Calibration Standard, Bruker Daltons) or ProteoMass (Protein Calibration Standard, Sigma-Aldrich) served as calibration standards. RP mode refers to reflector positive mode. RN mode refers to reflector negative mode. LP mode refers to linear positive mode.

Dialysis

[0263] Regenerated cellulose membranes: MWCO = 1 and 2 kDa (Spectra/Por), and MWCO = 12-14 kDa (Carl Roth) were used to perform dialysis. Typically, dialysis was carried out for 24 h with 1 L of solvent that was exchanged after first 6 h of the process.

Lyophilization

[0264] Freeze-drying was performed on an Alpha 1-2 LD plus (Martin Christ Gefriertrocknungsanlagen GmbH). Typically, samples were frozen with liquid nitrogen and placed into the freeze-dryer at high vacuum.

UV-Vis

[0265] Absorbance measurements were performed on a Perkin Elmer Lambda 25 UV-Vis or on a Thermo NanoDrop ND-2000 spectrophotometer in the spectral range of 200-750 nm.

[0266] Concentrations were determined using a Thermo Nanodrop 2000 or Lambda 25 spectrometer using the following parameters:

Trastuzumab OD_{280} = 1.5 $(mg/ml)^{-1}$ cm^{-1}

Cetuximab OD_{280} = 1.4 $(mg/ml)^{-1}$ cm^{-1}

HSP27 Oligonucleotide OD_{260} = 153,000 M^{-1} cm^{-1}; $Rz_{260:280}$ = 1.819

Dia-Cys OD_{280} = 0.57 $(mg/ml)^{-1}$ cm^{-1}

PEG_4-SPDP (PDT) OD_{343} = 8,080 M^{-1} cm^{-1}

SAMSA-Fluorescein OD_{495} = 64,500 M^{-1} cm^{-1}; $Rz_{280:495}$ = 0.232

Ellmans (TNB) OD_{412} = 14,150 M^{-1} cm^{-1}

Immobilized metal ion affinity chromatography (IMCA)

[0267] Nickel-nitrilotriacetic acid (Ni-NTA) chromatography was performed to purify histidine-tagged protein and protein-conjugates. Briefly, Ni-NTA agarose (10 mL) was pipetted into a gravity flow column for 5 mL bed volume. The resin was washed with 20 mL deionized water and recharged with 5 mL of 100 mM NiSO4 solution. The resin was washed again five times with 5 mL deionized water and equilibrated five times with DPBS. The protein solution was incubated with the washed Ni-NTA agarose rotating at 4 °C for 30 min. Afterwards, the Ni-NTA protein solution was pipetted back into the gravity flow column. The flow through was collected and the resin was washed three times with 5 mL DPBS. The immobilized sample was then eluted by increasing the imidazole concentration from 50 mL of 125 mM, pH 8 to 50 mL of 250 mM, pH 8. Elution fractions were dialyzed against PBS pH 7.4 to obtain the purified sample.

Size Exclusion Chromatography

[0268] Size exclusion chromatography (SEC) was carried out on an AKTA purifier. Samples were analyzed by SEC using either a Biosep SEC-S3000 column or an Sephadex G50M column (10 × 40 cm) eluting with TBS/ isopropyl alcohol solution (85:15 v/v). Sample purities were determined by integration of the antibody sample peak with respect to the trace aggregate peaks.

Ellman's assay

[0269] Ellman's test (or Ellman's assay) was carried out to quantitatively determine thiol concentration in a sample via spectrophotometry. Presence of thiols was indicated via the stoichiometric release of the 2-nitro-5-thiobenzoate (TNB) from Ellman's reagent in the presence of thiols. TNB obtains an absorption maximum at 412 nm and an extinction coefficient of OD_{412} = 14,150 M^{-1} cm^{-1} in buffered systems. Briefly, 2 µL of a 0.5 mg / mL solution of the Ellman's reagent (5,5-Dithiobis(2-nitrobenzoic acid), DTNB) in phosphate buffer (0.1 M, 1 mM EDTA, pH 7.4) was mixed with 20 µL of a thiol containing sample in buffer. The mixture was vortexed for 5 sec. Then, UV-Vis absorbance at 412 nm was measured on a Thermo Nanodrop 2000 to determine TNB

concentration and thus thiol content of the sample.

Dianthin production

[0270] Dianthin was expressed in a bacterium culture and the protein was purified following conventional cell culturing and protein purification steps known in the art.

Production of Saporin conjugates

[0271] Custom trastuzumab-saporin cetuximab-saporin, CD71mab-saporin conjugates were produced and purchased from Advanced Targeting Systems (San Diego, CA). IgG-saporin and saporin was purchased from Advanced Targeting Systems

Antibody-(Cys-dendron-(L-SO1861)^n)^n synthesis

[0272] Trastuzumab and Cetuximab are referred hereafter as "Ab". Ab was conjugated to dendritic saponin [dendron-(L-SO1861)$_4$-maleimide] via a tetra(ethylene glycol) succinimidyl 3-(2-pyridyldithio)propionate (PEG$_4$-SPDP) linker conducting a thiole-ene Michael-type reaction between Ab and dendritic saponin. The procedure is exemplary described for Cetuximab-(S-dendron-(L-SO1861)$_4$)$_4$:

[0273] Cetuximab was desalted into DPBS pH 7.5 buffer and then normalized to 2.50 mg/ml. To an aliquot of Ab (9.19 mg, 61 nmol) was added an aliquot of freshly prepared PEG$_4$-SPDP solution (5.0 mg/ml, 6.70 mole equivalents, 411 nmol), the mixture vortexed briefly then incubated for 60 minutes at 20 °C with roller-mixing. After incubation, the reaction was quenched with the addition of glycine (20 mg/ml, 7.7 µl), then the SPDP moiety reduced *in situ* by the addition of TCEP (5.0 mg/ml, 4.0 mole equivalents per SPDP, 1.64 µmol). This mixture was roller-mixed for 15 minutes at 20 °C with roller-mixing. The resulting Ab-SH was purified by gel filtration using a zeba spin desalting column into TBS pH 7.5. The Ab-SH was characterized by UV-vis analysis and Ellman's assay (thiol to Ab ratio = 5.4). To the bulk Ab-SH (7.41 mg, 1.93 mg/ml, 49 nmol) was added an aliquot of freshly prepared dendron-(L-SO1861)$_4$-maleimide solution in DMSO (10 mg/ml, 8.0 mole equivalents per Ab, 0.4 µmol, 3.16 mg, 0.32 ml), the mixture vortexed briefly then incubated overnight at 20 °C. Besides the conjugation reaction, two aliquots of the desalted Ab-SH (0.25 mg, 1.67 nmol) were removed prior to conjugation, and were reacted with NEM (8.0 mole equivalents per Ab, 13.3 nmol, 6.7 µl of a 0.25 mg/ml solution) or TBS pH 7.5 buffer (6.7 µl) overnight at 20 °C, as positive and negative controls, respectively. After incubation for 18 hours (prior to addition of NEM), the crude conjugate mixture was centrifuged briefly and 100 µl aliquot removed for analysis by UV-vis and alongside positive and negative controls were characterized by Ellman's assay to obtain dendron-(L-SO1861)$_4$ incorporation. To the bulk Ab - dendron-(L-SO1861)$_4$ mixture was added an aliquot of freshly prepared NEM solution (2.5 mg/ml, 5.0 mole equivalents, 0.25 µmol) and the mixture purified by 1.6 × 30 cm Sephadex G50 column eluting with DPBS pH 7.5 to give purified Cetuximab-(S-dendron-(L-SO1861)$_4$)$_4$ conjugate. The product was filtered to 0.2 µm to clarify and then concentrated carefully to *ca.* 3 mg/ml using a vivaspin T15 concentrator (3,000 g, 5 minute intervals, 5 °C) to give the final Cetuximab-(S-dendron(L-SO1861)$_4$)$_4$ conjugate. Yield: 4.41 mg, 48%. Dendron-(L-SO1861)$_4$ to Ab ratio = 4.4.

Table 2. Summarized reaction outcomes

Batch	Ab feed (mg)	PEG_4-SPDP mol equivalents	Dendron$(SO1861)^4$-maleimide mol equivalents	Obtained DAR	Purity by analytical SEC	Yield
Trastuzumab-(Cys-dendron-(L-$SO1861)^4)^4$	9.0	6.81	8	4.7	99.2 %	2.34 mg (26 %)
Cetuximab-(Cys-dendron-(L-$SO1861)^4)^4$	9.2	6.7	8	4.4	96.7 %	4.41 mg (48%)

Antibody-(L-$SO1861)^4$ (as illustrated in figure 17)

[0274] Trastuzumab, Cetuximab, are referred hereafter as "Ab". Ab was conjugated to the saponin SO18161-EMCH via Michael-type thiol-ene conjugation reaction at DARs of 1, 2, 3, 4, 5, and 6. The SO1861-EMCH molecule obtains a labil (L) hydrazone bond between its structure and its maleimide function generating a labil bond between the saponin and Ab. The procedure is exemplary described for Trastuzumab-$(L\text{-}SO1861)_4$:
To a solution of Cetuximab (40 mg, 8.0 ml) was added 10 µl/ml each of Tris concentrate (127 mg/ml, 1.05M), Tris.HCl concentrate (623 mg/ml, 3.95M) and EDTA-Na_2 concentrate (95 mg/ml, 0.26M) to give a 50mM TBS, 2.5mM EDTA buffer pH 7.5.

[0275] To Cetuximab divided into four portions (each of 9.73 mg, 4.864 mg/ml, 65 nmol) was added an aliquot of freshly prepared TCEP solution (0.5 - 2.0 mg/ml, 1.15 - 7.02 mole equivalents, 75 - 455 nmol), the mixtures vortexed briefly then incubated for 300 minutes at 20 °C with roller-mixing. After incubation (prior to addition of SO1861-EMCH), a *ca.* 1 mg (0.210 ml) aliquot of Ab-SH was removed from each mixture and purified by gel filtration using a zeba spin desalting column into TBS pH 7.5. These aliquots were characterized by UV-vis analysis and Ellman's assay (thiol to Ab ratio = 2.0, 4.2, 5.9 and 6.8 respectively). To each of the bulk Ab-SH was added an aliquot of freshly prepared SO1861-EMCH solution (2 mg/ml, 1.3 mole equivalents per 'thiol', 0.15 - 0.61 µmol, 0.16 - 0.63 ml), the mixtures vortexed briefly then incubated for 120 minutes at 20 °C. Besides each conjugation reaction, two aliquots of desalted Ab-SH (0.25 mg, 1.67 nmol) were reacted with NEM (1.3 mole equivalents per 'thiol', 4.3 - 17.4 nmol, 2.2 - 8.7 µl of a 0.25 mg/ml solution) or TBS pH 7.5 buffer (2.2 - 8.7 µl) for 120 minutes at 20 °C, as positive and negative controls, respectively. After incubation (prior to addition of NEM), a 0.200 ml aliquot of Ab - SO1861-EMCH mixture was removed and purified by gel filtration using zeba spin desalting column into TBS pH 7.5. This aliquot was characterized by UV-vis and alongside positive and negative controls were characterized by Ellman's assay to obtain SO1861-EMCH incorporations. To the bulk Ab - SO1861-EMCH mixture was added an aliquot of freshly prepared NEM solution (2.5 mg/ml, 2.5 - 10 mole equivalents, 0.15 - 0.58 µmol) and the mixtures purified by zeba spin desalting columns eluting with DPBS pH 7.5 to give purified Cetuximab - (L-SO1861) conjugates. The products were normalized to 2.5 mg/ml and filtered to 0.2µm prior to dispensing for biological evaluation.

Table 3. Summarized reaction conditions and results for Trastuzumab-L-SO1861 conjugates

Batch	Ab feed	TCEP feed mole equivalents	SO1861-EMCH feed	Obtained DAR	Purity by analytical SEC (%)	Yield (%)
Tras-$(L\text{-}SO1861)_2$	9.91 mg 66 nmol	1.10	0.15 µmol	1.6	99.2	79

Batch	Ab feed	TCEP feed mole equivalents	SO1861-EMCH feed	Obtained DAR	Purity by analytical SEC (%)	Yield (%)
Tras-(L-SO1861)$_3$	9.91 mg 66 nmol	2.35	0.31 µmol	3.0	99.0	81
Tras-(L-SO1861)$_4$	9.91 mg 66 nmol	3.83	0.46 µmol	4.0	98.4	81
Tras-(L-SO1861)$_3$	9.91 mg 66 nmol	5.77	0.62 µmol	5.3	98.5	79

Table 4. Summarized reaction conditions and results for Cetuximab-L-SO1861 conjugates

Batch	Ab feed	TCEP feed mole equivalents	SO1861-EMCH feed	Obtained DAR	Purity by analytical SEC (%)	Yield (%)
Cet-(L-SO1861)$_1$	9.73 mg 65 nmol	1.15	0.15 µmol	1.4	99.7	74
Cet-(L-SO1861)$_3$	9.73 mg 65 nmol	2.49	0.31 µmol	2.8	99.6	80
Cet-(L-SO1861)$_4$	9.73 mg 65 nmol	4.19	0.46 µmol	4.1	99.0	77
Cet-(L-SO1861)$_6$	9.73 mg 65 nmol	7.02	0.61 µmol	5.6	98.3	80

Antibody-(S-SO1861)^n synthesis

SO1861-S-Mal

[0276] SO1861 from Saponaria officinalis L (15.4 mg, 8.28 µmol) and HATU (140 mg, 368 µmol, 44.5 mole equivalents) were placed as solid into a 20 mL glas vial with magnetic stirrer and 5 mL DMSO was added to dissolve the materials. The dissolved mixture was stirred for 30 min at room temperature. After 30 min, 1 mL of freshly prepared AEM solution (65 mg, 256 µmol, 31 mole equivalents) in DMSO was added to the stirring SO1861-HATU mixture. The reaction mixture was stirred for 17 hours at room temperature. After stirring for 17 hours, the mixture was diluted with deionized water and dialyzed extensively for 24 h against deionized water using regenerated cellulose membrane tubes (Spectra/Por 7) with a MWCO of 1 kDa. After dialysis, the solution was lyophilized to obtain a white powder.

Yield: 7.22 mg (44 %).
Dried aliquots were further used for characterization via MALDI-TOF-MS.

MALDI-TOF-MS (RN mode) (Figure x a): *m/z* 1983 Da ([M - H]$^-$, *SO1861-S-Mal* conjugate), 2136 Da ([M - H]$^-$,

saponin-S-Mal conjugate).

MALDI-TOF-MS (RP mode) (Figure x b): *m/z* 2007 Da ($[M + Na]^+$, *SO1861-S-Mal* conjugate), 2107 Da ($[M - Na]^+$, saponin-S-Mal conjugate).

[0277] Maleimide functionality was verfied by mixing the purified *SO1861-S-Mal* with a freshly prepared L-cysteine solution (1000 mole excess) and observing the expected mass shift in MALDI-TOF-MS (Figure x) yielding a cysteine-conjugate at m/z 2103 Da.

Antibody conjugation

[0278] Trastuzumab and Cetuximab are referred hereafter as "Ab". Ab was conjugated to the saponin SO18161-S-Mal via Michael-type thiol-ene conjugation reaction. The saponin obtains a stable (S) amide bond between its structure and its maleimide function generating a stable bond between the saponin and Ab.

[0279] 5 mg of Ab dissolved in phosphate buffer saline (PBS) was buffer exchanged into tris(hydroxymethyl)-aminomethan buffer saline (TBS) pH 7.5 via zeba spin desalting column and normalized to a concentration of 3 mg/mL. To Ab was added an aliquot of freshly prepared tris(2-carboxyethyl)phosphine (TCEP) solution (0.25 mg/mL, 2.92 mole equivalents), the mixture vortexed briefly then incubated for 90 min at 20 °C. After, the resulting Ab-SH was purified by gel filtration using zeba spin desalting column into TBS pH 7.5. An aliquot was analyzed by Ellman's assay to ascertain sulfhydryl content. The Ab-SH obtained was split into a single portion for conjugation and two aliquots of 0.25 mg for controls. To the main portion of AB-SH was added an aliquot of freshly prepared SO1861-S-Mal solution (2 mg/mL, 2 mol equivalents with respect to sulhydryl content), to the second control aliquot was added buffer TBS pH 7.5. Each mixture was vortexed briefly then incubated for 2 hours at 20 °C. After, an aliquot from each was analyzed by Ellman's assay to ascertain sulfhydryl content remaining in the conjugate with respect to controls. After purification by gel filtration via zeba spin desalting column into Dulbecco's PBS pH 7.1, Ab-S-SO1861 was obtained. The conjugate was further analyzed by SEC to ascertain % purity. For the Trastuzumab-$(S\text{-}SO1861)_4$ sample a purity of 99 % was determined. For the Cetuximab-$(S\text{-}SO1861)_4$ sample a purity of 98.3 % was determined. Retention volumes for Trastuzumab-$(S\text{-}SO1861)_4$ (9.1 mL) and Cetuximab-$(S\text{-}SO1861)_4$ (8.8 mL) were noted to be similar and are consistent with a non-modified Ab (e.g. IgG).

Antibody-(L-HSP27 BNA)^n

Trastuzumab-(L-HSP27)^4,, Cetuximab-(L-HSP27)^4, synthesis via* PEG$_4$-SPDP *with a DAR4 and Cetuximab-(L-HSP27)^2 synthesis via* PEG$_4$-SPDP *with a DAR2

[0280] Trastuzumab, Cetuximab, are referred hereafter as "Ab". Ab was conjugated to HSP27 BNA disulfide via a tetra(ethylene glycol) succinimidyl 3-(2-pyridyldithio)propionate (PEG_4-SPDP) linker forming a labil (L) disulfide bond between Ab and HSP27 BNA . The procedure is exemplary described for Trastuzumab-$(L\text{-}HSP27\ BNA)_4$:

HSP27 BNA disulfide oligo (2.7mg, 470 nmol, 6.10 mg/ml) was reacted with TCEP (10 mole equivalents, 4.7 µmol, 1.34 mg, 50 mg/ml) for 30 minutes at 20 °C with roller mixing. After, the oligo-SH was purified by PD10 G25 desalting column eluting into TBS pH 7.5 and used promptly. Oligo-SH was obtained (2.48 mg, 90%, 1.24

mg/ml, SH to oligo ratio = 0.8)

Trastuzumab (1.5 mg, 10.3 nmol, 2.50 mg/ml) was reacted with an aliquot of freshly prepared PEG_4-SPDP solution (6.81 mole equivalents, 70.1 nmol, 39 µg) in DMSO (1 mg/ml) for 60 minutes at 20 °C with roller mixing. After, the reaction was quenched with glycine (15.1 µl of 2 mg/ml freshly prepared solution in TBS pH 7.5) and then desalted via zeba desalting column eluting with TBS pH 7.5. An aliquot of the resulting Tras-S-PEG_4-SPDP was taken out and tested by UV-Vis analysis. SPDP incorporation was determined using TCEP to liberate pyridiyl-2-thione (PDT) and by UV-vis analysis at 343 nm (SPDP to Ab ratio: 4). The remaining Tras-(S-PEG_4-SPDP)$_4$ was reacted with an aliquot of freshly prepared HSP27 oligonucleotide (oligo-SH) (8 mole equivalents, 82.4 nmol, 1.24 mg/ml) and incubated overnight at 20 °C with roller mixing. After 17 hours, the conjugate was analysed by UV-vis analysis to ascertain incorporation of HSP27 by displacement of pyridiyl-2-thione (PDT) at 343 nm. The crude conjugate was purified using a 1.6 × 33 cm Sephadex G50 column eluting with DPBS pH 7.5. The resulting Trastuzumab-(L-HSP27)$_4$ was obtained as a single fraction. Yield: n.d.. Purity: 96% , HSP27 BNA to Ab ratio = 4.4

Table 5. Summarized reaction conditions and results

Batch	**PEG_4-SPDP mol equivalents**	**HSP27 (oligo-SH) mol equivalents**	**Obtained DAR**	**Purity by analytical SEC (%)**	**Yield (%)**
Tras-(L-HSP27 BNA)^4	6.81	8	4.4	96.0	n.d.
Cet- (L-HSP27 BNA)^4	6.70	8	3.9	93.9	n.d.
Cet- (L-HSP27)^2	2.3	3.6	1.5	94.9	87
n.d. = not determined					

Antibody-(L-HSP27 BNA-L-blocked)^n

Trastuzumab-(L-HSP27-L-blocked)^4,, Cetuximab-(L-HSP27-L-blocked)

[0281] Trastuzumab and Cetuximab are referred hereafter as "Ab". Ab was conjugated a maleimido (Mal) bearing HSP27 derivate which is referred hereafter as "HSP27-Mal". Ab was conjugated to the HSP27-Mal via Michael-type thiol-ene conjugation reaction. The HSP17-Mal obtains a labile (L) hydrazone bond between its structure and its maleimide function generating a labile bond between the HSP27 BNA and Ab. The procedure is exemplary described for Trastuzumab-(L-HSP27 BNA-L-blocked)^n:

Trastuzumab was reconstituted to 21 mg/ml with deionized water (DI), then diluted to 5 mg/ml using histidine buffer pH 6. To a 20 mg (4.0 ml) aliquot was added 10 µl/ml each of Tris concentrate (127 mg/ml, 1.05M), Tris.HCl concentrate (623 mg/ml, 3.95M) and EDTA-Na_2 concentrate (95 mg/ml, 0.26M) to give a 50mM TBS, 2.5mM EDTA buffer pH 7.5. To Trastuzumab (20.30 mg, 4.920 mg/ml, 0.14 µmol) was added an aliquot of freshly prepared TCEP solution (1.00 mg/ml, 2.35 mole equivalents, 0.32 µmol), the mixture vortexed briefly then incubated for 90 minutes at 20 °C with roller-mixing. After incubation (prior to addition of HSP27-Mal), a *ca.* 2 mg (0.439 ml) aliquot of Ab-SH was removed from each mixture and purified by gel filtration using a zeba spin desalting column into TBS pH 7.5. This aliquot was characterized by UV-vis analysis and Ellman's assay (thiol to ab ratio = 4.0). To the bulk Ab-SH (5.1 mg, 35 nmol) was added an aliquot of the HSP27-Mal derivative (freshly prepared in TBS pH 7.5, 2 mg/ml, 1.3 mole equivalents per 'thiol', 182 nmol), the mixture vortexed briefly then incubated for 120 minutes at 20 °C. Besides the Trasuzumab - HSP27 BNA derivative conjugation

reaction, two aliquots of desalted Ab-SH (0.5 mg, 3.3 nmol) were reacted with NEM (1.3 mole equivalents per 'thiol', 17.3 nmol, 6.7 µl of a 0.25 mg/ml solution) or TBS pH 7.5 buffer (6.7 µl) for 120 minutes at 20 °C, as positive and negative controls, respectively. After incubation (prior to addition of NEM), a 0.100 ml aliquot of Ab - HSP27 BNA mixture was removed and purified by gel filtration using zeba spin desalting column into TBS pH 7.5. This aliquot was characterized by UV-vis and alongside positive and negative controls was characterized by Ellman's assay to obtain HSP27 incorporation. To the bulk Ab - HSP27 mixture was added an aliquot of freshly prepared NEM solution (0.25 mg/ml, 2.5 mole equivalents, 89 nmol) and the mixture purified by gel filtration using a 1.6 × 30 cm Sephadex G50M eluting with DPBS pH 7.5 followed by repeated centrifugal filtration and washing using a 100 KDa MWCO concentrator to give purified Trastuzumab-(L-HSP27-L-blocked)$_4$ conjugate. The products were filtered to 0.2µm prior to dispensing for biological evaluation.

Table 6. Summarized reaction conditions and results

Batch	Ab-SH feed	Thiol to Ab ratio	HSP27-Mal feed	Obtained DAR	Purity by analytical SEC (%)	Yield (%)
Tras-(L-HSP27 BNA-L-blocked)^4	5.1 mg 35 nmol	3.98	1.36 mg	3.6	99.4	40
Cet-(L-HSP27 BNA-L-blocked)^4	5.1 mg 35 nmol	4.16	1.36 mg	3.6	97.9	50

Antibody-[Cys-Trifunctional linker-(L-SO1861)-(L-HSP27 BNA)]^n

Trastuzumab-[S-Tri-(L-SO1861)-(L-HSP27)]^4, Trastuzumab-[S-Tri-(blocked)-(L-HSP27)]^4, Cetuximab-[S-Tri-(L-SO1861)-(L-HSP27)]^4, Cetuximab-[S-Tri-(blocked)-(L-HSP27)]^4

[0282] Trastuzumab and Cetuximab are referred hereafter as "Ab". Ab was conjugated via Michael-type thiol-ene reaction to two different maleimide (Mal) bearing HSP27 BNA derivatives which are referred hereafter as "HSP27-Mal". These HSP27-Mal derivatives were namely: 1) Mal-Trifunctional linker-(L-SO1861)-(L-HSP27), 2) Mal-Trifunctional linker-(blocked)-(L-HSP27). The procedure is exemplary described for Trastuzumab-[S-Trifunctional linker-(L-SO1861)-(L-HSP27 BNA)]^4:
Trastuzumab was reconstituted to 21 mg/ml with deionized water (DI), then diluted to 5 mg/ml using histidine buffer pH 6. To a 20 mg (4.0 ml) aliquot was added 10 µl/ml each of Tris concentrate (127 mg/ml, 1.05M), Tris.HCl concentrate (623 mg/ml, 3.95M) and EDTA-Na_2 concentrate (95 mg/ml, 0.26M) to give a 50mM TBS, 2.5mM EDTA buffer pH 7.5.

[0283] To Trastuzumab (20.30 mg, 4.920 mg/ml, 0.14 µmol) was added an aliquot of freshly prepared TCEP solution (1.00 mg/ml, 2.35 mole equivalents, 0.32 µmol), the mixture vortexed briefly then incubated for 90 minutes at 20 °C with roller-mixing. After incubation (prior to addition of construct), a *ca.* 2 mg (0.439 ml) aliquot of Ab-SH was removed from each mixture and purified by gel filtration using a zeba spin desalting column into TBS pH 7.5. These aliquots were characterized by UV-vis analysis and Ellman's assay (thiol to ab ratio = 4.0). The bulk Ab-SH was split into two aliquots (1.1 mg, 7.6 nmol and 1.2 mg, 8.3 nmol), and to each aliquot was added an aliquot of each of the HSP27 BNA-Mal derivatives **1-2** (freshly prepared in TBS pH 7.5, 2 mg/ml, 1.3 mole equivalents per 'thiol', 40 nmol and 43 nmol), the mixtures vortexed briefly then incubated for 120 minutes at 20 °C. Besides the Trastuzumab - HSP27 BNA derivatives **2** conjugation reaction, two aliquots of desalted Ab-SH (0.5 mg, 3.3 nmol) were reacted with NEM (1.3 mole equivalents per 'thiol', 17.3 nmol, 6.7 µl of a 0.25 mg/ml solution) or TBS pH 7.5 buffer (6.7 µl) for 120 minutes at 20 °C, as positive and negative

controls, respectively. After incubation (prior to addition of NEM), a 0.100 ml aliquot of Ab - construct **2** mixture was removed and purified by gel filtration using zeba spin desalting column into TBS pH 7.5. This aliquot was characterized by UV-vis and alongside positive and negative controls was characterized by Ellman's assay to obtain HSP27 BNA derivatives **2** incorporation. To each bulk Ab - construct mixture was added an aliquot of freshly prepared NEM solution (0.25 mg/ml, 2.5 mole equivalents, 19 and 21 nmol) and the mixtures purified by gel filtration using a 1.6 × 30 cm Sephadex G50M eluting with DPBS pH 7.5 followed by repeated centrifugal filtration and washing using a 100 KDa MWCO concentrator to give purified Trastuzumab - construct **1-2** conjugates. The products were filtered to 0.2µm prior to dispensing for biological evaluation.

Table 7. Summarized reaction conditions and results

Batch	Ab-SH feed	TCEP equivalents	Thiol to ab ratio	Purity by analytical SEC (%)	Yield (%)
Tras-[S-Trifunctional linker-(L-SO1861)-(L-HSP27 BNA)]^4	1.1 mg 7.6 nmol	2.35	3.98	97.0	57
Tras-[S-Trifunctional linker-(blocked)-(L-HSP27 BNA)]^4	1.2 mg 8.3 nmol	2.35	3.98	96.6	33
Cet-[S-Trifunctional linker-(L-SO1861)-(L-HSP27 BNA)]^4	1.1 mg 7.6 nmol	2.72	4.16	98.9	47
Cet-[S-Trifunctional linker-(blocked)-(L-HSP27 BNA)]^4	1.2 mg 8.3 nmol	2.72	4.16	99.1	50

Antibody-(L-SO1861)^n-(L-HSP27 BNA)^n

Trastuzumab-(L-SO1861)^4-(L-HSP27)^4, Cetuximab-(L-SO1861)^4-(L-HSP27)^4 synthesis via PEG$_4$-SPDP with a DAR4 Cetuximab-(L-SO1861)^4-(L-HSP27)^2 (FBR703 STB17I7-8) synthesis via PEG$_4$-SPDP with a DAR2

[0284] Trastuzumab-(L-SO1861)$_4$, Cetuximab-(L-SO1861)$_4$ are referred hereafter as "Ab". Ab was conjugated to HSP27 BNA disulfide via a tetra(ethylene glycol) succinimidyl 3-(2 pyridyldithio)propionate (PEG$_4$-SPDP) linker forming a labil (L) disulfide bond between Ab and HSP27 BNA. The procedure is exemplary described for Trastuzumab-(L-SO1861)$_4$-(L-HSP27)$_4$:

HSP27 BNA disulfide oligo (2.7mg, 470 nmol, 6.10 mg/ml) was reacted with TCEP (10 mole equivalents, 4.7 µmol, 1.34 mg, 50 mg/ml) for 30 minutes at 20 °C with roller mixing. After, the oligo-SH was purified by PD10 G25 desalting column eluting into TBS pH 7.5 and used promptly. Oligo-SH was obtained (2.48 mg, 90%, 1.24 mg/ml, SH to oligo ratio = 0.8)

[0285] Trastuzumab-(L-SO1861)$_4$ (1.3 mg, 8.7 nmol, 2.50 mg/ml) was reacted with an aliquot of freshly

prepared PEG_4-SPDP solution (9.26 mole equivalents, 80.3 nmol, 45 µg) in DMSO (1 mg/ml) for 60 minutes at 20 °C with roller mixing. After, the reaction was quenched with glycine (15.1 µl of 2 mg/ml freshly prepared solution in TBS pH 7.5) and then desalted via zeba desalting column eluting with TBS pH 7.5. An aliquot of the resulting Tras-(L-SO1861)-(L-PEG_4-SPDP) was taken out and tested by UV-Vis analysis. SPDP incorporation was determined using TCEP to liberate pyridiyl-2-thione (PDT) and by UV-vis analysis at 343 nm (SPDP to Ab ratio = 4). The remaining Tras-(L-SO1861)-(L-PEG_4-SPDP) was reacted with an aliquot of freshly prepared HSP27 oligonucleotide (oligo-SH) (8 mole equivalents, 54.8 nmol, 0.32 mg, 1.24 mg/ml) and incubated overnight at 20 °C with roller mixing. After 17 hours, the conjugate was analyzed by UV-vis analysis to ascertain incorporation of HSP27 by displacement of pyridiyl-2-thione (PDT) at 343 nm. The crude conjugate was purified using a 1.6 × 33 cm Sephadex G50 column eluting with DPBS pH 7.5. The resulting Trastuzumab-$(L\text{-}SO1861)_4$-$(L\text{-}HSP27\ BNA)_4$ was obtained as a single fraction. Yield: 0.47 mg, 45% (0.49 mg/ml), HSP27 to Ab ratio = 3.5

Table 8. Summarized reaction conditions and results

Batch	PEG_4-SPDP mol equivalents	HSP27 (oligo-SH) mol equivalents	Obtained DAR	Purity by analytical SEC (%)	Yield (%)
Tras-$(L\text{-}SO1861)^4$-$(L\text{-}HSP27\ BNA)^4$	9.26	8	3.5	85.1	45
Cet-$(L\text{-}SO1861)_4$-$(L\text{-}HSP27\ BNA)^4$	7.21	8	3.8	80.8	n.d.
Cet-$(L\text{-}SO1861)_4$-$(L\text{-}HSP27\ BNA)^2$	3.34	3.6	1.8	76.2	81
n.d. = not determined					

Antibody-$(L\text{-}QS\ Mix)^n$

[0286] Trastuzumab, Cetuximab, are referred hereafter as "Ab". Ab was conjugated to the saponin QS Mix-EMCH via Michael-type thiol-ene conjugation reaction. The procedure is exemplary described for Trastuzumab-L-QS Mix:

Trastuzumab ("Ab", 600 mg) was reconstituted to 21 mg/mL with deionized water (DI), then diluted to 5 mg/mL using freshly prepared histidine buffer pH 6 (5mM histidine pH 6, 2% trehalose, 0.01% Tween 20). 10 µL/mL each of Tris concentrate (127 mg/mL, 1.05M), Tris.HCL concentrate (623 mg/mL, 3.95M) and EDTA-Na_2 concentrate (95 mg/ml, 0.26M) was added to give a 50mM TBS, 2.5mM EDTA buffer pH 7.5. To Trastuzumab (603.8 mg, 4.887 mg/mL, 4.0 µmol) was added an aliquot of freshly prepared TCEP solution (1 mg/mL, 2.35 mole equivalents, 9.5 µmol, 2.72 mg), the mixture swirled by hand to mix then incubated for 90 minutes at 20 °C with roller-mixing. After incubation (prior to addition of QS Mix-EMCH), a 2 mg (0.409 mL) aliquot of Ab-SH was removed and purified by gel filtration using zeba spin desalting column into TBS pH 7.5. This aliquot was characterized by UV-vis analysis and Ellman's assay. To the bulk Ab-SH was added an aliquot of freshly prepared QS Mix-EMCH solution (2 mg/mL, 5.2 mole equivalents, 21 µmol, 21.6 mL), the mixtures vortexed briefly then incubated for 120 minutes at 20 °C. Besides the conjugation reaction, two aliquots of desalted Ab-SH (0.5 mg, 0.134 mL, 3.33 nmol) were reacted with NEM (8.00 equivalents, 26.6 nmol, 3.3 µg, 13.3 µL of a 0.25 mg/mL solution) or TBS pH 7.5 buffer (13.3 µL) for 120 minutes at 20 °C, as positive and negative controls, respectively. After incubation (prior to addition of NEM), a ca. 2 mg (0.481 mL) aliquot of Ab - QS Mix-EMCH mixture was removed and purified by gel filtration using zeba spin desalting column into TBS pH 7.5. This aliquot was characterized by UV-vis and alongside positive and negative controls were characterized by

Ellman's assay to obtain QS Mix-EMCH incorporations. To the bulk Ab - QS Mix-EMCH mixture was added an aliquot of freshly prepared NEM solution (2.5 mg/mL, 5 mole equivalents, 20 µmoL, 2.51 mg) and the mixture stored at 2-8 °C overnight. The conjugate was purified by 10 × 40 cm Sephadex G50M column eluting with DPBS pH 7.5 to give purified Trastuzumab - (L-QS Mix) conjugate. The product as a whole was concentrated then normalized to 5 mg/mL using a vivacell 100 concentrator (2,000 g, 4 °C, 200 minutes). The product were filtered to 0.2µm and dispensed for biological evaluation. Yield: n.d. QS Mix to Ab ratio = 4.1.

Table 9. Summarized reaction outcomes

Method	Measure	Result
Analytical SEC	Purity	n.d.
Yield	Mass (percent)	n.d.
Ellman's assay	QS Mix-EMCH incorpoation	4.1

Antibody-(L-HSP27 BNA-L-SO1861)^n

Trastuzumab-(L-HSP27-L-SO1861)^4, Cetiximab-(L-HSP27-L-SO1861)^4 with a DAR4

[0287] Trastuzumab and Cetuximab are referred hereafter as "Ab". Ab was conjugated a saponin SO1861 and maleimido (Mal) bearing HSP27 derivate which is referred hereafter as "HSP27-Mal". Ab was conjugated to the HSP27-Mal via Michael-type thiol-ene conjugation reaction. The HSP17-Mal obtains a labile (L) hydrazone bond between its structure and its maleimide function generating a labile bond between the HSP27 BNA and Ab. The procedure is exemplary described for Trastuzumab-(L-HSP27 BNA-L-SO1861)$_4$:

Trastuzumab was reconstituted to 21 mg/ml with deionized water (DI), then diluted to 5 mg/ml using histidine buffer pH 6. To a 20 mg (4.0 ml) aliquot was added 10 µl/ml each of Tris concentrate (127 mg/ml, 1.05M), Tris.HCl concentrate (623 mg/ml, 3.95M) and EDTA-Na_2 concentrate (95 mg/ml, 0.26M) to give a 50mM TBS, 2.5mM EDTA buffer pH 7.5. To Trastuzumab (20.30 mg, 4.920 mg/ml, 0.14 µmol) was added an aliquot of freshly prepared TCEP solution (1.00 mg/ml, 2.35 mole equivalents, 0.32 µmol), the mixture vortexed briefly then incubated for 90 minutes at 20 °C with roller-mixing. After incubation (prior to addition of HSP27-Mal), a *ca.* 2 mg (0.439 ml) aliquot of Ab-SH was removed from each mixture and purified by gel filtration using a zeba spin desalting column into TBS pH 7.5. This aliquot was characterized by UV-vis analysis and Ellman's assay (thiol to ab ratio = 4.0). To the bulk Ab-SH (4.7 mg, 32 nmol) was added an aliquot of the HSP27-Mal derivative (freshly prepared in TBS pH 7.5, 2 mg/ml, 1.3 mole equivalents per 'thiol', 166 nmol), the mixture vortexed briefly then incubated for 120 minutes at 20 °C. Besides the Trasuzumab - HSP27 BNA derivative conjugation reaction, two aliquots of desalted Ab-SH (0.5 mg, 3.3 nmol) were reacted with NEM (1.3 mole equivalents per 'thiol', 17.3 nmol, 6.7 µl of a 0.25 mg/ml solution) or TBS pH 7.5 buffer (6.7 µl) for 120 minutes at 20 °C, as positive and negative controls, respectively. After incubation (prior to addition of NEM), a 0.100 ml aliquot of Ab - HSP27 BNA mixture was removed and purified by gel filtration using zeba spin desalting column into TBS pH 7.5. This aliquot was characterized by UV-vis and alongside positive and negative controls was characterized by Ellman's assay to obtain HSP27 incorporation. To the bulk Ab - HSP27 mixture was added an aliquot of freshly prepared NEM solution (0.25 mg/ml, 2.5 mole equivalents, 80 nmol) and the mixture purified by gel filtration using a 1.6 × 30 cm Sephadex G50M eluting with DPBS pH 7.5 followed by repeated centrifugal filtration and washing using a 100 KDa MWCO concentrator to give purified Trastuzumab-(L-HSP27 BNA-L-SO1861)$_4$ conjugate. The products were filtered to 0.2µm prior to dispensing for biological evaluation.

Table 10. Summarized reaction conditions and results

Batch	Ab-SH feed	Thiol to Ab ratio	HSP27-Mal feed	Purity by analytical SEC (%)	Yield (%)
Tras-(L-HSP27 BNA-L-SO1861)^4	4.7 mg 32 nmol	3.98	1.57 mg	99.4	40
Cet-(L-HSP27 BNA-L-SO1861)^4	4.7 mg 32 nmol	4.16	1.57 mg	97.9	50

Example 14

Materials and methods

[0288] In our current work, we investigated a model scaffold consisting of four molecular arms for saponin binding via a Schiff base (imine) and one arm for click chemistry. The polymeric structure (Figure 25) is a pentavalent polyethylene glycol-based dendrimer of the first generation (i.e. number of repeated branching cycles) that was purchased from Iris Biotech GmbH (Marktredwitz, Germany). Figure 25 shows a basic scaffold with click chemistry function to link any desired effector molecule. The user determines the position of the click chemistry position in the effector molecule and all further properties of the effector molecule, e.g. choice and position of an optional ligand. The saponin (in this example SA1641) was purified from a saponin composite raw extract from Gypsophila species called Saponinum album obtained from Merck (Darmstadt, Germany). The powdered raw extract (2.5 g) was hydrolyzed in water (100 mL) with sodium hydroxide (0.2 g). The solution was stirred for 20 h at 40 °C and then supplemented with glacial acetic acid until pH 5.0 was reached. To remove tannins, the solution was shaken in a separatory funnel with 30 mL butanol. The aqueous phase was recaptured and butanol extraction repeated two times. The butanol phases were supplemented with anhydrous sodium sulfate, filtered and pooled. Butanol was evaporated and the remaining saponin powder resolved in 20% methanol to a final concentration of 30 mg/mL. After short sonication, different saponins were separated by high performance liquid chromatography (HPLC). Tubes (excluding column) were rinsed with warm water (40 °C) at a flow of 1.5 mL/min and then including Eurospher RP-C18-column (5 µm, 250 × 8 mm) with isopropanol (100%). Saponins were applied to the column and eluted with a methanol gradient (20% methanol to 70% methanol within 30 min at 1.5 mL/min in water supplemented with 0.01% trifluoroacetic acid followed by 70% methanol for further 60 min) [31]. Aliquots of the fractions were analyzed for their SA1641 content by electrospray ionization mass spectrometry (ESI-MS). Fractions containing pure SA1641 were pooled and methanol evaporated. The aqueous solution was frozen as a thin film in a rotating round-bottom flask by use of dry ice. After storage for 16 h at -80 °C, the sample was lyophilized. To produce the scaffold as defined in the invention, the polymeric structure (0.2 mM) and SA1641 (3.2 mM) were solved in water (approx. pH 8) and equal volumes mixed and shaken for 24 h at 26 °C. Then sodium cyanoborohydride ($NaCNBH_3$; 0,1 M) was added in 4-fold molar excess referred to SA1641 and the sample incubated for further 24 h. The structure was then verified by ultra performance liquid chromatography (UPLC)/ESI-MS. The samples were applied to a RP-C4-column and eluted with a methanol gradient (25% methanol to 80% methanol within 15 min in water supplemented with 0.01% trifluoroacetic acid followed by 80% methanol for further 10 min). The fractions were analyzed by use of LockSpray™ that is an ion source designed specifically for exact mass measurement with electrospray ionization using LC-time-of-flight (LC-TOF) mass spectrometers from Waters Corporation.

Results

[0289] Figure 26 relates to a functionalized scaffold with pre-bound effector molecule and click chemistry function to link any desired ligand. Optionally, a pH-sensitive linkage can be provided to release the effector molecule from the scaffold after reaching the endosomes. The inset of Figure 26 shows the theoretically expected mass spectrum obtained from a calculation with the isotope pattern calculator enviPat Web 2.0. The pattern considers the charge of the molecule and the natural occurrence of isotopes, which is the reason that more than one peek is expected for a single substance. The experimental data (Figure 26) obtained by UPLC/ESI-MS show almost exactly the same peaks at m/z 758-760 with same intensity as predicted, thus proving successful SA1641 coupling to the polymeric structure.

Example 15

Materials and methods

[0290] As an example for a pharmaceutical active substance, we used the targeted toxin dianthin-Epidermal Growth Factor (dianthin-EGF). The plasmid His-dianthin-EGF-pET11d [20] (100 ng) was added to 20 µL Escherichia coli Rosetta™ 2 (DE3) pLysS Competent Cells (Novagen, San Diego, CA, USA). Cells were transformed by a heat-shock (30 min on ice, 90 s at 42 °C and 1 min on ice). Thereafter, 300 µL lysogeny broth (LB) was added and the suspension incubated for 1 h at 37 °C while shaking at 200 rpm. A preheated lysogeny broth agar plate with 50 µg/mL ampicillin was inoculated with 100 µl bacteria suspension and the plate incubated overnight at 37 °C. Lysogeny broth (3 mL) with 50 µg/mL ampicillin was inoculated with a colony from the plate and the bacteria were incubated for 8 h at 37 °C and 200 rpm. The suspension (50 µL) was added to 500 mL of lysogeny broth with 50 µg/mL ampicillin and incubated overnight at 37 °C and 200 rpm. Subsequently, the volume was scaled-up to 2.0 L and bacteria grew under the same conditions until an optical density at wavelength 600 nm of 0.9 was reached. Thereafter, protein expression was induced by the addition of isopropyl β-D-1-thiogalactopyranoside (IPTG) at a final concentration of 1 mM. Protein expression lasted for 3 h at 37 °C and 200 rpm. Finally, the bacterial suspension was centrifuged at 5,000 × g and 4 °C for 5 min, resuspended in 20 mL PBS (137 mM NaCl, 2.7 mM KCl, 8.1 mM Na_2HPO_4, 1.47 mM KH_2PO_4) and stored at -20 °C until use. For purification, bacterial suspensions were thawed and lysed by sonication. Lysates were centrifuged (15,800 × g, 4 °C, 30 min) and imidazole added to a final concentration of 20 mM. The supernatant was incubated with 2 mL of Ni-nitrilotriacetic acid agarose under continuous shaking for 30 min at 4 °C in the presence of 20 mM imidazole. Subsequently, the material was poured into a 20-mL-column and washed three times with 10 mL wash buffer (50 mM NaH_2PO_4, 300 mM NaCl, 20 mM imidazole) and dianthin-EGF eluted by 10-mL-portions of increasing concentrations of imidazole (31, 65, 125 and 250 mM) in wash buffer. Eluate fractions (2 mL) were dialyzed overnight at 4 °C against 2.0 L PBS. Desalted dianthin-EGF was concentrated by an Amicon® Ultra-15 (10 kDa) and the protein concentration quantified.

[0291] To introduce a suitable click chemistry group into dianthin-EGF, alkyne-PEGs-N-hydroxysuccinimidyl ester in 8-fold molar excess referred to dianthin-EGF was solved in dimethyl sulfoxide and added to 9 volumes of dianthin-EGF (1 mg in 0.2 M NaH_2PO_4/Na_2HPO_4, pH 8). After incubation at room temperature for 4 h, non-bound alkyne was separated by use of a PD10 column (GE-Healthcare, Freiburg, Germany). Click chemistry with the polymeric structure was conducted by copper(I)-catalyzed alkyne-azide cycloaddition. Alkyne-dianthin-EGF (0.02 mM), dendrimer (0.05 mM), $CuSO_4$ (0.1 mM), tris(3-hydroxypropyltriazolylmethyl)amine (0.5 mM) and sodium ascorbate (5 mM) were incubated under gentle agitation for 1 h at room temperature in 0.1 M NaH_2PO_4/Na_2HPO_4, pH 8. Low molecular mass substances were then separated using a PD10 column.

[0292] To test the efficacy of the invention, we conducted a viability assay with HER14 cells. These cells are fibroblasts stably transfected with the human epidermal growth factor receptor and therefore target cells for

the targeted toxin dianthin-EGF. HER14 cells (2,000 cells/100 µL/well) were seeded into wells of 96-well-cell culture plates and incubated for 24 h in DMEM medium supplemented with 10% fetal calf serum and 1% penicillin/streptomycin at 37 °C, 5% CO_2 and 98% humidity. The different test substances (see results and Figure 27) were then added in triplicates in a volume of 25 µL and supplemented with further 25 µL of medium. After an incubation of 72 h, 30 µL 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (0.5 mg/mL in water) was added per well and incubated for 2 h. Thereafter, the medium was carefully removed and replaced by an aqueous solution containing 10% (v/v) isopropanol, 5% (w/v) sodium dodecyl sulfate and 400 mM HCl, and incubated for 5 min. Solubilized formazan was photometrically quantitated at 570 nM in a microplate reader (Spectra MAX 340 PC, Molecular Devices, Sunnyvale, CA, USA). Untreated cells were normalized to 1 and all samples referred to the untreated control. Significance was determined by unpaired two-sample t-tests.

Results

[0293] The polymeric structure, in the example a pentameric dendrimer (pentrimer), does not have any cytotoxic effect on the target cells, neither in absence nor in presence of SA1641 (Figure 27, column 2 and 3). In the absence of the scaffold, the targeted toxin (dianthin-EGF) shows half maximal toxicity at a concentration of 0.1 nM (column 4). In the presence of SA1641 the same concentration results in death of all cells indicating the general ability of SA1641 to act as an enhancer of the endosomal escape (column 5). The presence of the polymeric structure does not affect the toxicity of dianthin-EGF neither in the presence nor in the absence of SA1641 (columns 6 and 7), indicating that the scaffold does not affect the toxicity of dianthin-EGF. To couple the model polymeric structure via click chemistry to the example pharmaceutically active substance of dianthin-EGF, the substance had to be coupled with an alkyne group before. In consequence of this modification, dianthin-EGF lost some activity (compare columns 8 and 9 with 6 and 7, respectively), however, the undirected alkyne modification does not affect the idea of the invention and is also not required in future applications. We had to introduce the alkyne in an undirected way for test purposes only with the risk to impede the pharmaceutically active center of the toxin. A manufacturer of a pharmaceutically active substance can introduce the click position during synthesis directly into the substance at a position of his choice where the activity of the substance remains unaffected. There was no additional loss of activity when clicking the alkyne-modified pharmaceutically active substance to the polymeric structure indicating that the polymeric structure itself was not toxic (column 10 and 11).

Example 16

Materials

[0294] The following chemicals were used as purchased: methanol (MeOH, LiChrosolv, Merck), N-ε-maleimidocaproic acid hydrazide (EMCH, 95%, TCI Chemicals), trifluoroacetic acid (TFA, 99.8%, Carl Roth), 2-mercaptoethanol (98%, Sigma-Aldrich), poly(amidoamine) (PAMAM dendrimer, ethylenediamine core, generation 5.0 solution, Sigma-Aldrich), cyanine 3 carboxylic acid (Cy3-COOH, 95%, Lumiprobe), 1-[Bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid hexafluorophosphate, N-[(Dimethylamino)-1H-1,2,3-triazolo-[4,5-b]pyridin-1-ylmethylene]-N-methylmethanaminium hexafluorophosphate N-oxide (HATU, 97%, Sigma-Aldrich), bovine serum albumin fraction V (BSA, Carl Roth), dimethylsulfoxide (DMSO, 99%, Carl Roth), 2-Iminothiolane hydrochloride (98%, Sigma-Aldrich), rhodamine b (RhodB, 95%, Merck), Dulbecco's phosphate buffered saline (PBS, Gibco), hydrochloric acid (HCl, 37%, Merck), NHS-PEG_{13}-DBCO (Click Chemistry Tools), Alexa Fluor™ 488 5-TFP (Thermo-Fischer), azido-PEG3-SS-NHS (Conju-Probe), sodium cyanoborohydride ($NaCNBH_3$, 95 %, Sigma-Aldrich), ammonium persulfate

(APS, 98%, Sigma-Aldrich), N,N,N',N'-tetramethylethylenediamine (TMEDA, 99 %, Sigma-Aldrich), customized peptide SESDDAMFCDAMDESDSK (95%, PeptideSynthetics), azido-dPEG$_{12}$-NHS (95%, Quanta Biodesign), PFd-G4-Azide-NH-BOC Dendron (G4-dendron, 95%, Polymer Factory), Cyanin5-DBCO (Cy5-DBCO, 95%, Lumiprobe), Chloroform ($CHCl_3$, 99.5 %, Sigma), Amicon Ultra 0.5 mL centrifugal filters (3 kDa MWCO, Sigma), mPEG-SCM (mPEG$_{2k}$-NHS, 95.6%, Creative PEG Works), Amicon Ultra 15 mL centrifugal filters (10 kDa MWCO, Sigma).

Methods

MALDI-TOF-MS

[0295] MALDI-TOF spectra were recorded on a MALDI-Mass Spectrometer (Bruker Ultrafex III). Typically, the sample dissolved in MilliQ water in nanomolar to micromolar range was spotted on the target (MTP 384 target plate polished steel T F, Bruker Daltons) using either super-DHB (99%, Fluka) or sinapinic acid (SA, 99%, Sigma-Aldrich) as the matrix dissolved in acetonitrile (MADLI-TOF-MS tested, Sigma) / 0.1% TFA (7:3 v/v) via the dried-droplet-method. PepMix (Peptide Calibration Standard, Bruker Daltons) or ProteMass (Protein Calibration Standard, Sigma-Aldrich) served as calibration standards. RP mode refers to reflector positive mode. RN mode refers to reflector negative mode. LP mode refers to linear positive mode.

H-NMR

[0296] ^{1}H NMR analysis was performed using a Bruker 400 MHz NMR spectrometer. The sample preparation, in which 2 mg of sample had been dissolved in 0.8 mL of methanol-D_4 (99%, Deutero), was performed 24 h prior to the measurement.

UV-Vis

[0297] UV-Vis measurements were performed on a NanoDrop ND-1000 spectrophotometer in the spectral range of 200-750 nm.

Size Exclusion Chromatography

[0298] Size exclusion chromatography (SEC) was performed with Sephadex G 25 Superfine from GE Healthcare and on prepacked PD10 columns (GE Healthcare, Sephadex G 25 M). The material was activated by swelling in the respective eluent prior to performing chromatography.

Dialysis

[0299] Regenerated cellulose membranes: MWCO = 1 and 2 kDa (Spectra/Por), and MWCO = 12-14 kDa (Carl Roth) were used to perform dialysis. Typically, dialysis was carried out for 24 h with 1 L of solvent that was exchanged after first 6 h of the process.

Lyophilization

[0300] Freeze-drying was performed on an Alpha 1-2 LD plus (Martin Christ Gefriertrocknungsanlagen GmbH). Typically, samples were frozen with liquid nitrogen and placed into the freeze-dryer at high vacuum.

SO1861-EMCH synthesis (see also Figure 18 and 24)

[0301] SO1861 from Saponaria officinalis L (59 mg, 31.7 µmol) and EMCH (301 mg, 888 µmol) were placed in a round flask with stirrer and dissolved in 13 mL methanol. TFA (400 µL, cat.) was added to the solution and the reaction mixture was stirred for 3 h at 800 rpm and room temperature on a RCT B magnetic stirrer (IKA Labortechnik). After stirring for 3 h, the mix was diluted either with MilliQ water or PBS and dialyzed extensively for 24 h against either with MilliQ water or PBS using regenerated cellulose membrane tubes (Spectra/Por 7) with a MWCO of 1 kDa. After dialysis, the solution was lyophilized to obtain a white powder. Yield 62.4 mg (95 %). Dried aliquots were further used for characterization via ^{1}H NMR and MALDI-TOF-MS.

^{1}H NMR (400 MHz, methanol-D_4) (Figure 28A, SO1861): δ = 0.50-5.50 (m, saponin triterpenoid and sugar backbone protons), 9.43 (1H, s, aldehyde proton of saponin, H^a).

^{1}H NMR (400 MHz, methanol-D_4) (Figure 28B. SO1861-EMCH, PBS workup): δ = 0.50-5.50 (m, saponin triterpenoid and sugar backbone protons), 6.79 (2 H, s, maleimide protons, H^c), 7.62-7.68 (1 H, m, hydrazone proton, H^b).

MALDI-TOF-MS (RP mode) (Figure 29A): *m/z* 2124 Da ($[M+K]^+$, saponin-EMCH), *m/z* 2109 Da ($[M+K]^+$, SO1861-EMCH), *m/z* 2094 Da ($[M+Na]^+$, SO1861-EMCH)

MALDI-TOF-MS (RN mode) (Figure 34C): *m/z* 2275 Da ($[M-H]^-$, saponin-EMCH conjugate), 2244 Da ($[M-H]^-$, saponin-EMCH conjugate), 2222 Da ($[M-H]^-$, saponin-EMCH conjugate), 2178 Da ($[M-H]^-$, saponin-EMCH conjugate), 2144 Da ($[M-H]^-$, saponin-EMCH conjugate), 2122 Da ($[M-H]^-$, saponin-EMCH conjugate), 2092 Da ($[M-H]^-$, saponin-EMCH conjugate), 2070 Da ($[M-H]^-$, SO1861-EMCH), 2038 Da ([M-H]-, SO1832-EMCH), 1936 Da ([M-H]-, SO1730-EMCH), 1861 Da ([M-H]-, SO1861).

SO1861-EMCH-mercaptoethanol

[0302] To SO1861-EMCH (0.1 mg, 48 nmol) 200 µL mercaptoethanol (18 mg, 230 µmol) was added and the solution was shaken for 1 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 1 h, the solution was diluted with methanol and dialyzed extensively for 4 h against methanol using regenerated cellulose membrane tubes (Spectra/Por 7) with a MWCO of 1 kDa. After dialysis, an aliquot was taken out and analyzed via MALDI-TOF-MS.

[0303] MALDI-TOF-MS (Figure 29B) (RP mode): *m/z* 2193 Da ($[M+K]^+$, SO1861-EMCH-mercaptoethanol), *m/z* 2185 Da ($[M+K]^+$, SO1861-EMCH-mercaptoethanol), *m/z* 2170 Da ($[M+Na]^+$, SO1861-EMCH-mercaptoethanol).

BSA-SO1861 synthesis

[0304] 2-iminothiolane (231 µg, 1.1 µmol) dissolved in 47 µL PBS was added to a BSA-RhodB solution (10 mg, 0.15 µmol) in 200 µL PBS and the mix was shaken for 40 min at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 40 min, the reaction mix was immediately run through a Sephadex G25 superfine size exclusion column (16 mL column volume) and SO1861-EMCH (1 mg, 0.5 µmol) dissolved in 100 µL PBS was added to the collected BSA-SH fraction. The reaction mixture was shaken for 12 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 12 h the BSA-SO1861 concentrated using centrifugal filtration at 4,000 rpm (15 °C) via Amicon Ultra 15 filters with a MWCO of 3 kDa. The conjugate was stored as solution in the fridge and aliquots were taken for analysis. Yield: not determined.

[0305] MALDI-TOF-MS (Figure 37A) (LP mode): *m/z* 74.2 kDa ($[M+H]^+$, BSA-SO1861 with 4 SO1861 attached), 72.2 kDa ($[M+H]^+$, BSA-SO1861 with 3 SO1861 attached), 70.2 kDa ($[M+H]^+$, BSA-SO1861 with 2 SO1861 attached), 37.0 kDa ($[M+H]^{2+}$, BSA-SO1861 with 4 SO1861 attached), 35.9 kDa ($[M+H]^{2+}$, BSA-SO1861 with 3 SO1861 attached), 34.7 kDa ($[M+H]^{2+}$, BSA-SO1861 with 2 SO1861 attached).

Cy3-PAMAM

[0306] 720 µL PAMAM dissolved in methanol (30 mg, 1.04 µmol) was placed into a 250 mL round flask and methanol was removed via a rotary evaporator (20 mbar, 60 °C). Remaining PAMAM was dissolved in 9 mL DMSO. HATU (7.6 mg, 20 µmol) dissolved in 0.5 mL DMSO was added to a Cy3-COOH (0.6 mg, 1.2 µmol) solution in DMSO and the mix was shaken for 1 h at 800 rpm at room temperature on a ThermoMixer C (Eppendorf). After shaking for 1 h, the HATU-Cy3 solution was added to the stirring PAMAM solution and the reaction mix was stirred for 12 h at room temperature. After stirring for 12 h, the reaction mix was diluted with MilliQ water and dialyzed extensively for 24 h against MilliQ water using regenerated cellulose membrane tubes (Spectra/Por 6) with a MWCO of 2 kDa. After dialysis, the volume of the conjugate solution was reduced via a rotary evaporator (20 mbar, 60 °C) and the concentrated conjugate solution was run through a Sephadex G25 superfine size exclusion column (16 mL column volume). The first fraction was collected and lyophilized to obtain the viscous pink PAMAM-Cy3 conjugate. PAMAM-Cy3 conjugate formation was confirmed by chromatography on thin layer chromatography (methanol/water, v/v 1:1), and the appearance of a faster band on a Sephadex G 25 superfine column. Yield 21.3 mg (63 %). The dye per PAMAM molar ratio determined by UV-Vis spectrophotometry was 0.43.

[0307] MALDI-TOF-MS (Figure 39A) (LP mode): *m/z* 28.0 kDa ($[M+H]^+$, Cy3-PAMAM).

Cy3-PAMAM-SO1861 synthesis

[0308] Procedure is described exemplary for Cy3-PAMAM-$(SO1861)_5$. 2-iminothiolane (1 mg, 6.7 µmol) dissolved in 250 µL MilliQ water was added to a PAMAM-Cy3 solution (0.5 mg, 17 nmol) in 125 µL MilliQ water and the mix was shaken for 40 min at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 40 min, the reaction mix was immediately run through a Sephadex G25 superfine size exclusion column (16 mL column volume) and SO1861-EMCH (176 µg, 85 nmol) dissolved in 40 µL MilliQ water was added to the collected Cy3-PAMAM-SH fraction. The reaction mixture was shaken for 12 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 12 h, the reaction mix was diluted with MilliQ water and dialyzed extensively for 24 h against MilliQ water using regenerated cellulose membrane tubes (ZelluTrans, Carl Roth) with a MWCO of 12 - 14 kDa. After dialysis, the Cy3-PAMAM-SO1861 solution was concentrated using centrifugal filtration at 4000 rpm (15 °C) via Amicon Ultra 15 filters with a MWCO of 3

kDa. The conjugate was stored as solution in the fridge and aliquots were taken for analysis. Yield: 0.5 mg (75%).

[0309] MALDI-TOF-MS spectra are illustrated in Figures 39B-D, and Figure 40. MALDI-TOF-MS of Cy3-PAMAM-$(SO1861)_6$ (Figure 39B) (LP mode): *m/z* 38.4 kDa ($[M+H]^+$, Cy3-PAMAM-SO1861), 17.9 kDa ($[M+H]^{2+}$, Cy3-PAMAM-SO1861).

[0310] The synthesis of Cy3-PAMAM-$(SO1861)_5$, Cy3-PAMAM-$(SO1861)_{13}$, Cy3-PAMAM-$(SO1861)_{51}$, and Cy3-PAMAM-$(SO1861)_{27}$, has been performed via the above described methodology but differ in the feed equivalents of the starting materials 2-iminothiolane and SO1861-EMCH. The respective feed equivalents of the starting materials and the respective mass of the conjugates are highlighted in Table 11.

Table 11. Reaction parameter for Cy3-PAMAM-SO1861 synthesis.

2-Iminothiolane feed equivalents to Cy3-PAMAM	**SO1861-EMCH feed equivalents to Cy3-PAMAM**	**Mass of conjugate via MALDI-TOF-MS**	**SO1861 molecules attached per PAMAM**	**Resulting conjugate**
384	6	38.7 kDa	~5	Cy3-PAMAM-$(SO1861)_6$, Figure 39 B
384	20	53.9 kDa	~13	Cy3-PAMAM-$(SO1861)_{13}$, Figure 39 C
384	57	133.9 kDa	~51	Cy3-PAMAM-$(SO1861)_{51}$, Figure 39 D
8	5	37.7 kDa	~5	Cy3-PAMAM-$(SO1861)_5$, Figure 40 A
32	30	87.0 kDa	~27	Cy3-PAMAM-$(SO1861)_{27}$, Figure 40 B

Cy3-PAMAM-NC-SO1861 synthesis

[0311] Cy3-PAMAM (0.5 mg, 18 nmol), SO1861 (2.3 mg, 1.24 µmol), and HATU (64.6 mg, 170 µmol) were dissolved separately in 200 µL DMSO. SO1861 and HATU solutions were mixed and shaken for 20 min at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 20 min, Cy3-PAMAM solution was added to the shaking SO1861-HATU solution and the reaction mixture was allowed to shake for 12 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 12 h, the reaction mix was diluted with MilliQ water and dialyzed extensively for 24 h against MilliQ water using regenerated cellulose membrane tubes (ZelluTrans, Carl Roth) with a MWCO of 12-14 kDa. After dialysis, the Cy3-PAMAM-NC-SO1861 solution was concentrated using centrifugal filtration at 4,000 rpm (15 °C) via Amicon Ultra 15 filters with a MWCO of 3 kDa. The Cy3-PAMAM-NC-$(SO1861)_{17}$ conjugate was stored as solution in the fridge and aliquots were taken for analysis. Yield: 0.77 mg (69%).

[0312] MALDI-TOF-MS (Figure 41) (LP mode): *m/z* 62.3 kDa ($[M+H]^+$, Cy3-PAMAM-NC-S01861), 35.7 kDa ($[M+H]^{2+}$, Cy3-PAMAM-NC-SO1861).

G4-dendron dye labeling and deprotection

[0313] PFd-G4-Azide-NH-BOC (G4-dendron) (9.75 mg, 2.11 µmol) was placed into a 2 mL reaction tube (Eppendorf) and dissolved in 200 µL DMSO. 100 µL of a Cy5-DBCO solution in DMSO (1.72 µmol * mL^{-1}, 170 nmol) was added to the G4-dendron solution and the mix was shaken for 12 hours at room temperature and 800 rpm on a ThermoMixer C (Eppendorf). After shaking for 12 h, the reaction mix was diluted with MilliQ water and dialyzed extensively for 24 h against MilliQ water using regenerated cellulose membrane tubes (Spectra/Por 7) with a MWCO of 1 kDa. After dialysis, the solution was lyophilized to obtain a blue powder. The crude product was used as obtained from lyophilization for the deprotection step.

[0314] Partially Cy5 labeled lyophilized G4-dendron was dissolved in 12 mL $CHCl_3$ in 50 mL round flask with stirrer. 12 mL TFA was added and the reaction mix was stirred for 3 h at 800 rpm and room temperature on a RCT B magnetic stirrer (IKA Labortechnik). After stirring for 3 h, the solvent was removed under reduced pressure (50 °C, 30 mbar) on a rotary evaporator (Heidolph WB 2000). After evaporation, the batch was dissolved in MilliQ water and run through a PD10 size exclusion column. G4-dendron conjugate formation was confirmed by chromatography on thin layer chromatography (methanol/water, v/v 1:1), and the appearance of a faster band on a PD10 column. Obtained fraction of size exclusion chromatography was lyophilized to obtain a blue powder.

[0315] Yield 5.7 mg (93 %). The dye per G4-dendron molar ratio determined by UV-Vis spectrophotometry was 0.012.

[0316] MALDI-TOF-MS (Figure 54B) (RP mode): *m/z* 3956 Da ($[M+Na]^+$, Cy5-G4-dendron + PF_6^- counterion), 3820 Da ($[M+Na]^+$, Cy5-G4-dendron - PF_6^- counterion), 3617 Da ($[M+H]^+$, G4-dendron impurity), 3017 ($[M+H]^+$, G4-dendron).

G4-dendron-SO1861 synthesis

[0317] Procedure is described exemplary for the lowest G4-dendron to SO1861-EMCH ratio. 2-iminothiolane (2.65 mg, 19.2 µmol) dissolved in 300 µL MilliQ water was added to a partially Cy5 labeled G4-dendron solution (0.577 mg, 192 nmol) in 252 µL MilliQ water and the mix was shaken for 40 min at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 40 min, the reaction mix was immediately run through a PD10 size exclusion column and SO1861-EMCH (1.19 mg, 575 nmol) dissolved in 100 µL MilliQ water was added to the collected G4-dendron-SH fraction. The reaction mixture was shaken for 12 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 12 h, the reaction mix was concentrated via centrifugal filtration using Amicon Ultra centrifugal filters (3 kDa MWCO). The conjugate was stored as solution in the fridge and aliquots were taken for analysis. Yield: 90 nmol (47%).

[0318] MALDI-TOF-MS spectra are illustrated in Figure 55. MALDI-TOF-MS of G4-dendron-SO1861 (Figure 55 C) (LP mode): *m/z* 10.19 kDa ($[M+H]^+$, Cy5-G4-dendron-$[SO1861]_3$), 9.27 kDa ($[M+H]^+$, G4-dendron-$[SO1861]_3$), 7.92 kDa ($[M+H]^+$, Cy5-G4-dendron-$[SO1861]_2$), 7.14 kDa ($[M+H]^+$, G4-dendron-$[SO1861]_2$), 5.86 kDa ($[M+H]^+$, Cy5-G4-dendron-$[SO1861]_1$), 5.07 kDa ($[M+H]^+$, G4-dendron-$[SO1861]_1$).

[0319] The synthesis of other G4-dendron-$(SO1861)_n$ conjugates has been performed via the above described methodology but differs in the feed equivalents of the starting material SO1861-EMCH. The

respective feed equivalents of the starting materials and the respective mass of the conjugates are highlighted in Table 12.

Table 12. Reaction parameter for G4-dendron-SO1861 synthesis.

2-Iminothiolane feed equivalents to G4-dendron	SO1861-EMCH feed equivalents to G4-dendron	Mass of conjugates via MALDI-TOF-MS	SO1861 molecules attached per G4-dendron	Resulting MS spectrum
100	3	5.07 - 10.18 kDa	~ 1 - 3	Figure 55 C
100	10	5.07 - 11.64 kDa	~ 1 - 4	Figure 55 B
100	22	6.20 - 22.02 kDa	~ 1 - 9	Figure 55 A

PAMAM thiolation

[0320] Procedure is described exemplary for the highest PAMAM to 2-iminothiolane ratio. To a PAMAM (333 µg, 12.8 nmol) solution dissolved in 30 µL methanol 2-iminothiolane (0.53 mg, 3.84 µmol) dissolved in 128 µL MilliQ water was added. The reaction mixture was shaken for 12 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 12 h, the reaction mix was washed 4 times with MilliQ water via centrifugal filtration using Amicon Ultra centrifugal filters (3 kDa MWCO) at 15 °C and 13500 rpm. After washing the sample was lyophilized to obtain a white solid. Yield was not determined.

[0321] MALDI-TOF-MS spectra are illustrated in Figure 57. MALDI-TOF-MS of PAMAM-$(SH)_{108}$ (Figure 57 E) (LP mode): *m/z* 41.5 kDa ($[M+H]^+$, PAMAM-$[SH]_{108}$).

[0322] The synthesis of other PAMAM-iminothiolane conjugates has been performed via the above described methodology but differs in the feed equivalents of the starting material 2-iminothiolane. For the lowest 2-iminothiolane feed reaction Cy3-PAMAM has been used.

[0323] The respective feed equivalents of the starting materials and the respective mass of the conjugates are highlighted in Table 13.

Table 13. Reaction parameter for PAMAM-SH synthesis.

2-Iminothiolane feed equivalents to PAMAM	Mass of conjugates via MALDI-TOF-MS	Iminothiolane molecules attached per PAMAM	Resulting MS spectrum
50	34.4 kDa	~ 16	Fig. 57 C
100	35.9 kDa	~ 65	Fig. 57 D
300	41.5 kDa	~ 108	Fig. 57 E

PAMAM PEGylation

[0324] Procedure is described exemplary for the lowest PAMAM to $mPEG_{2k}$ ratio. To a PAMAM (333 µg, 12.8 nmol) solution dissolved in 10 µL DMSO $mPEG_{2k}$-NHS (0.268 mg, 128 nmol) dissolved in 13 µL DMSO was added. The reaction mixture was shaken for 12 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 12 h, the reaction mix was diluted with MilliQ water and dialyzed extensively for

24 h against MilliQ water using regenerated cellulose membrane tubes (Spectra/Por 6) with a MWCO of 2 kDa. After dialysis, the batch was concentrated via centrifugal filtration using Amicon Ultra 15 mL centrifugal filters (10 kDa MWCO). The concentrated batch was run through a PD10 size exclusion column followed by lyophilization to obtain a white fluffy powder. Yield was not determined.

[0325] MALDI-TOF-MS spectra are illustrated in Figure 58. MALDI-TOF-MS of PAMAM-$(mPEG_{2k})_3$ (Figure 58 C) (LP mode): *m/z* 33.46 kDa ($[M+H]^+$, PAMAM-$[mPEG_{2k}]_3$).

[0326] The synthesis of other PAMAM-$mPEG_{2k}$ conjugates has been performed via the above described methodology but differs in the feed equivalents of the starting material $mPEG_{2k}$-NHS. The respective feed equivalents of the starting materials and the respective mass of the conjugates are highlighted in Table 14.

Table 14. Reaction parameter for PAMAM-$mPEG_{2k}$ synthesis.

$mPEG_{2k}$-NHS feed equivalents to PAMAM	**Mass of conjugates via MALDI-TOF-MS**	**$mPEG_{2k}$ molecules attached per PAMAM**	**Resulting MS spectrum**
10	28.5 kDa	~3	Figure 58 C
20	43.0 kDa	~ 8	Figure 58 D
100	62.8 kDa	~ 18	Figure 58 E

Cy3-PAMAM-SO1861-DBCO synthesis

[0327] Procedure is described exemplary for Cy3-PAMAM-$(SO1861)_{27}$-$(DBCO)_{10}$. Cy3-PAMAM-$(SO1861)_{27}$ (0.41 mg, 4.71 nmol) was freeze-fried and dissolved in 100 µL DMSO. DBCO-PEG_{13}-NHS ester (0.197 mg, 188 nmol) dissolved in DMSO was added to the Cy3-PAMAM-SO1861 solution and the mixture was shaken at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 3 h, the reaction mix was diluted with MilliQ water and dialyzed extensively for 24 h against MilliQ water using regenerated cellulose membrane tubes (ZelluTrans, Carl Roth) with a MWCO of 12-14 kDa. After dialysis, the Cy3-PAMAM-SO1861-DBCO solution was concentrated using centrifugal filtration at 4,000 rpm (15 °C) via Amicon Ultra 15 filters with a MWCO of 3 kDa. The conjugate was stored as solution in the fridge and aliquots were taken for analysis. Yield: 0.1 mg (22%).

[0328] MALDI-TOF-MS (Figure 42 D) (LP mode): m/z 92.5 kDa ($[M+H]^+$, Cy3-PAMAM-SO1861-DBCO), 53.0 kDa ($[M+H]^{2+}$, Cy3-PAMAM-SO1861-DBCO).

[0329] The synthesis of Cy3-PAMAM-$(SO1861)_5$-$(DBCO)_{38}$, and Cy3-PAMAM-$(SO1861)_{27}$-$(DBCO)_{10}$, have been performed via the above described methodology. The respective feed equivalents of the starting material and the respective mass of the conjugates are highlighted in Table 15.

Table 15. Reaction parameter for Cy3-PAMAM-SO1861-DBCO synthesis.

Used Cy3-PAMAM-saponin batch	**DBCO-PEG13-NHS feed equivalents**	**Mass via MALDI-TOF-MS**	**DBCO molecules attached per PAMAM**	**Resulting conjugate**
Cy3-PAMAM-$(SO1861)_5$	40	76.3 kDa	~38	Cy3-PAMAM-$(SO1861)_5$-$(DBCO)_{38}$, Figure 42 C
Cy3-PAMAM-$(SO1861)_{27}$	40	92.5 kDa	~10	Cy3-PAMAM-$(SO1861)_{27}$-$(DBCO)_{10}$, Figure 42 D

Used Cy3-PAMAM-saponin batch	DBCO-PEG13-NHS feed equivalents	Mass via MALDI-TOF-MS	DBCO molecules attached per PAMAM	Resulting conjugate

Cy3-PAMAM-NC-SO1861-DBCO synthesis

[0330] Cy3-PAMAM-NC-$(SO1861)_{17}$ (0.3 mg, 4.8 nmol) was freeze-fried and dissolved in 100 µL DMSO. DBCO-PEG_{13}-NHS ester (0.202 mg, 194 nmol) dissolved in DMSO was added to the Cy3-PAMAM-NC-SO1861 solution and the mixture was shaken at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 3 h, the reaction mix was diluted with MilliQ water and dialyzed extensively for 24 h against MilliQ water using regenerated cellulose membrane tubes (ZelluTrans, Carl Roth) with a MWCO of 12-14 kDa. After dialysis, the Cy3-PAMAM-SO1861-DBCO solution was concentrated using centrifugal filtration at 4,000 rpm (15 °C) via Amicon Ultra 15 filters with a MWCO of 3 kDa. The conjugate was stored as solution in the fridge and aliquots were taken for analysis. Yield: 0.1 mg (22%). Mass spectrometry indicates the conjugation of 30 DBCO moieties per PAMAM molecule.

[0331] MALDI-TOF-MS (Figure 42 B) (LP mode): m/z 93.2 kDa ($[M+H]^+$, Cy3-PAMAM-NC-SO1861-DBCO), 49.6 kDa ($[M+H]^{2+}$, Cy3-PAMAM-NC-SO1861-DBCO).

EGFDianthin and dianthin expression

[0332] Plasmid-DNA (His-dianthin-EGF-pET11d or His-dianthin-pET11d) [20] was transformed into chemically competent Escherichia coli NiCo21 (DE3) (New England Biolabs®, Inc.) and grown in 3 mL lysogeny broth supplemented with 50 µg/mL ampicillin at 37 °C for 5 h at 200 rpm. These bacteria were used to inoculate 500 mL lysogeny broth supplemented with 50 µg/mL ampicillin for overnight culture at 37 °C. Subsequently, the culture volume was scaled up to 2 L and bacteria were grown until an optical density (A600) of 0.9. Protein expression was induced by the addition of isopropyl β-D-1-thiogalactopyranoside (IPTG) at a final concentration of 1 mM. Cells were further grown for 3 h at 37 °C and 200 rpm. After centrifugation (5 min, 5,000 g, 4 °C) cell pellets were resuspended in 20 mL phosphate buffered saline (Dulbecco's phosphate-buffered saline (PBS) with Ca^{2+} and Mg^{2+}, pH 7.4) and stored at -20 °C. After thawing, proteins were released by ultrasound device (Branson Sonifier 250, G. Heinemann). The solution was centrifuged (15,800 × g, 30 min, 4 °C) and adjusted to 20 mM imidazole concentration. The construct contained an N-terminal His-tag and was purified by nickel nitrilotriacetic acid chromatography (Ni-NTA Agarose, Qiagen, Hilden, Germany). After elution with imidazole (20-250 mM) the eluates were analyzed by sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) (12%). Fractions containing dianthin-EGF or dianthin were dialyzed against 2 L chitin binding domain buffer (20 mM tris(hydroxymethyl)-aminomethane/HCl, 500 mM NaCl, 1 mM EDTA, 0.1% Tween-20, pH 8.0) at 4 °C. Further purification by chitin column affinity chromatography served to remove bacterial proteins with binding activity for Ni-NTA agarose. After elution with chitin binding domain buffer, the fractions were analyzed by SDS-PAGE (12%). Fractions containing dianthin-EGF or dianthin were dialyzed against 5 L PBS at 4 °C. Purified proteins were concentrated by Amicon centrifugal filter devices (10 kDa, Millipore, Eschborn, Germany). The protein concentration was determined by a bicinchoninic acid assay (Pierce, Rockford, USA).

Dianthin-EGF-Alexa488 synthesis

[0333] Dianthin-EGF (240 µg, 6.7 nmol) solution in PBS was placed into an Amicon Ultra 15 filter with a MWCO of 3 kDa and centrifuged at 4,000 rpm and 4 °C for 30 min three times. After each cycle, the Amicon filter was refilled with 0.1 M sodium carbonate buffer at pH 9. After the third centrifugation cycle, the volume was reduced to 0.5 mL via centrifugation. The dianthin-EGF sodium carbonate solution was placed into a 2 mL reaction tube and Alexa Fluor™ 488 5-TFP (50 µg, 56 nmol) dissolved in 10 µL DMSO was added to the protein solution. The mix was shaken at 800 rpm and room temperature on a ThermoMixer C (Eppendorf) for 80 min. After shaking, the mix was run through a Sephadex G25 M size exclusion column (GE Healthcare, PD10 column). The dianthin-EGF-Alexa488 conjugate was stored in solution in 0.1 M sodium carbonate buffer at pH 9 in the fridge and aliquots were taken for analysis. Yield: 210 µg (85%).

[0334] MALDI-TOF-MS (Figure 43D) (LP mode): *m/z* 36.8 kDa ($[M+H]^+$, dianthin-EGF-Alexa488), *m/z* 33.6 kDa ($[M+H]^+$, dianthin-EGF-Alexa488), 18.8 kDa ($[M+H]^{2+}$, dianthin-EGF-Alexa488), 16.6 kDa ($[M+H]^{2+}$, dianthin-EGF-Alexa488).

Dianthin-Alexa488 synthesis

[0335] Dianthin (184 µg, 6.2 nmol) solution in PBS was placed into an Amicon Ultra 15 filter with a MWCO of 3 kDa and centrifuged at 4,000 rpm and 4 °C for 30 min three times. After each cycle, the Amicon filter was refilled with 0.1 M sodium carbonate buffer at pH 9. After the third centrifugation cycle, the volume was reduced to 0.5 mL via centrifugation. The dianthin sodium carbonate solution was placed into a 2 mL reaction tube and Alexa Fluor™ 488 5-TFP (16.7 µg, 19 nmol) dissolved in 3.5 µL DMSO was added to the protein solution. The mix was shaken at 800 rpm and room temperature on a ThermoMixer C (Eppendorf) for 80 min. After shaking, the mixwas run through a Sephadex G25 M size exclusion column (GE Healthcare, PD 10 column). The dianthin-Alexa488 conjugate was stored in solution in 0.1 M sodium carbonate buffer at pH 9 in the fridge and aliquots were taken for analysis. Yield: not determined MALDI-TOF-MS (Figure 44 D) (LP mode): *m/z* 30.7 kDa ($[M+H]^+$, dianthin-Alexa488).

Dianthin-EGF-Alexa488-S-S-PEG-N_3, and Dianthin-EGF-Alexa488-PEG_{12}-N_3 synthesis

[0336] Procedure is described exemplary for dianthin-EGF-Alexa488-S-S-PEG-N_3. Dianthin-EGF-Alexa488 (70 µg, 1.9 nmol) sodium carbonate solution was placed into a 2 mL reaction tube and azido-PEG_3-S-S-NHS (120 µg, 272 nmol) dissolved in 9 µL DMSO was added to the protein solution. The mix was shaken at 800 rpm and 15 °C on a ThermoMixer C (Eppendorf) for 12 h. After shaking, the reaction mix was diluted with PBS and was washed with PBS via centrifugal filtration at 4,000 rpm and 4 °C using Amicon Ultra 15 filter with a MWCO of 3 kDa.

Yield: 54 µg (70%).

MALDI-TOF-MS (Figure 43 E) (LP mode): *m/z* 40.8 kDa ($[M+H]^+$, dianthin-EGF-Alexa488-S-S-PEG-N_3), *m/z* 37.5 kDa ($[M+H]^+$, dianthin-EGF-Alexa488-S-S-PEG-N_3).

[0337] The synthesis of dianthin-EGF-Alexa488-S-S-PEG-N_3, and dianthin-EGF-Alexa488-PEG_{12}-N_3 have been performed via the above described methodology but differed in the used azido-PEG linker. The respective azido-PEG linker, their feed equivalents, and the respective mass of the conjugates are highlighted

in Table 16.

Table 16. Reaction parameter for dianthin-EGF-Alexa488-PEG-N_3 synthesis

Used toxin batch	Azido-PEG linker used	Azido-PEG linker feed equivalents	Mass of conjugate via MALDI-TOF-MS	Resulting conjugate
Dianthin-EGF-Alexa488	Azido-PEG_3-S-S-NHS	135	40.8 kDa	Dianthin-EGF-Alexa488-S-S-PEG_3-N_3
Dianthin-EGF-Alexa488	Azido-PEG_{12}-NHS	135	43.3 kDa	Dianthin-EGF-Alexa488-PEG_{12}-N_3

Dianthin-Alexa488-S-S-PEG-N_3

[0338] Dianthin-Alexa488 (24.5 µg, 0.8 nmol) sodium carbonate solution was placed into a 2 mL reaction tube and azido-PEG_3-S-S-NHS (34 µg, 78 nmol) dissolved in 9 µL DMSO was added to the protein solution. The mix was shaken at 800 rpm and 15 °C on a ThermoMixer C (Eppendorf) for 12 h. After shaking, the reaction mix was diluted with PBS and was washed with PBS via centrifugal filtration at 4,000 rpm and 4 °C using Amicon Ultra 15 filter with a MWCO of 3 kDa.

Yield: 10.3 µg (39%).

MALDI-TOF-MS (Figure 44 E) (LP mode): m/z 32.9 kDa ($[M+H]^+$, dianthin-Alexa488-S-S-PEG-N_3).

Cy3-PAMAM-Saponin-Toxin conjugate synthesis

[0339] Procedure is described exemplary for Cy3-PAMAM-$(SO1861)_{27}$-DBCO. Cy3-PAMAM-$(SO1861)_{27}$-DBCO (17 µg, 0.184 nmol) solution in MilliQ water was mixed with a dianthin-EGF-Alexa488-SS-PEG3-N_3 (3.6 µg, 0.089 nmol) solution in PBS in a 1.5 mL reaction tube and the reaction mix was shaken at 800 rpm and 15 °C on a ThermoMixer C (Eppendorf) for 2 h. After shaking, small aliquots were taken out for analysis via SDS-PAGE and fluorescence imaging on a Molecular Imager® VersaDoc™ MP 4000 imaging system (Bio-Rad) (Figure 45).

[0340] The synthesis of Cy3-PAMAM-$(SO1861)_5$-S-S-Dianthin-EGF-Alexa488, Cy3-PAMAM-$(SO1861)_{27}$-S-S-Dianthin-EGF-Alexa488, Cy3-PAMAM-NC-$(SO1861)_{17}$-S-S-Dianthin-EGF- Alexa488, Cy3-PAMAM-NC-$(SO1861)_{17}$-S-S-Dianthin-Alexa488, and Cy3-PAMAM-NC-$(SO1861)_{17}$-Dianthin-EGF-Alexa488, have been performed via the above described methodology but differ in the used PAMAM-saponin-DBCO batch, the used azido-toxin batch, and their feed equivalents. The respective feed equivalents of the starting materials are highlighted in Table 17.

Table 17. Reaction parameter for Cy3-PAMAM-saponin-toxin synthesis.

PAMAM-saponin-DBCO batch used	PAMAM-saponin-DBCO feed equivalents	Azido-toxin batch used	Azido-toxin feed equivalents	Resulting conjugate
Cy3-PAMAM-$(SO1861)_5$-$(DBCO)_{38}$	3	Dianthin-EGF-Alexa488-S-S-PEG_3-N_3	1	Cy3-PAMAM-$(SO1861)_5$-S-S-Dianthin-EGF-Alexa488
Cy3-PAMAM-$(SO1861)_{27}$-$(OBCO)_{10}$	2.1	Dianthin-EGF-Alexa488-S-S-PEG_3-N_3	1	Cy3-PAMAM-$(SO1861)_{27}$-S-S-Dianthin-EGF-Alexa488
Cy3-PAMAM-NC-$(SO1861)_{17}$-$(DBCO)_{30}$	2.3	Dianthin-EGF-Alexa488-S-S-PEG_3-N_3	1	Cy3-PAMAM-NC-$(SO1861)_{17}$-S-S-Dianthin-EGF-Alexa488
Cy3-PAMAM-NC-$(SO1861)_{17}$-$(DBCO)_{30}$	7.1	Dianthin-Alexa488-S-S-PEG_3-N_3	1	Cy3-PAMAM-NC-$(SO1861)_{17}$-S-S-Dianthin-Alexa488
Cy3-PAMAM-NC-$(SO1861)_{17}$-$(DBCO)_{3O}$	2.3	Dianthin-EGF-Alexa488-PEG_{12}-N_3	1	Cy3-PAMAM-NC-$(SO1861)_{17}$-Dianthin-EGF-Alexa488

Cy3-PAMAM-NC-SO1861 synthesis via reductive amination

[0341] Cy3-PAMAM (0.19 mg, 13 nmol) and SO1861 (0.73 mg, 0.39 µmol) were dissolved separately in 200 µL 0.1 M acetate buffer at pH 5. SO1861 and Cy3-PAMAM solutions were mixed and shaken for 20 min at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 20 min, $NaCNBH_3$ (5 mg, 81 µmol) was added to the shaking reaction solution and the reaction mixture was allowed to shake for 12 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking for 12 h, the reaction mix was diluted with MilliQ water and dialyzed extensively for 24 h against MilliQ water using regenerated cellulose membrane tubes (ZelluTrans, Carl Roth) with a MWCO of 12-14 kDa. After dialysis, the Cy3-PAMAM-NC-SO1861 solution was concentrated using centrifugal filtration at 4,000 rpm (15 °C) via Amicon Ultra 15 filters with a MWCO of 3 kDa. The conjugate was stored as solution in the fridge and aliquots were taken for analysis. Yield: not determined

[0342] MALDI-TOF-MS (Figure 46 B, C) (LP mode): m/z 88.7 kDa ($[M+H]^+$, Cy3-PAMAM-NC-SO1861), 49.2 kDa ($[M+H]^{2+}$, Cy3-PAMAM-NC-SO1861).

[0343] The synthesis of Cy3-PAMAM-NC-$(SO1861)_{30}$, and Cy3-PAMAM-NC-$(SO1861)_{10}$, have been performed via the above described methodology but differed in the time after which the reducing agent $NaCNBH_3$ was added to the reaction batch. The respective time of the $NaCNBH_3$ addition and the respective mass of the conjugates are highlighted in Table 18.

Table 18. Reaction parameter **Cy3-PAMAM-NC-SO1861** synthesis via reductive amination.

Time of shaking reaction mix before $NaCNBH_3$ addition	Mass via MALDI-TOF-MS	SO1861molecules attached per PAMAM	Resulting conjugate
20 min	88.8 kDa	~30	Cy3-PAMAM-NC-$(SO1861)_{30}$, Figure 46 C
12 h	48.0 kDa	~10	Cy3-PAMAM-NC-$(SO1861)_{10}$, Figure 46 B

Poly(SO1861) synthesis

[0344] SO1861-EMCH (0.13 mg, 63 nmol) was dissolved in 30 µL degased MilliQ water. APS (0.2 µg, 0.8 nmol) dissolved in 4 µL degased MilliQ water was added to the SO1861-EMCH solution and the solution was placed into a ThermoMixer C (Eppendorf) at 60 °C. Then, TMEDA (cat., 0.5 µL) was added to the mix and the mix was shaken at 800 rpm and 60 °C on a ThermoMixer C (Eppendorf) for 2 h. After 2 h, a small aliquot was taken out for analysis via mass spectrometry.

[0345] MALDI-TOF-MS (Figure 48 C) (LP mode): *m/z* 18.2 kDa ($[M+H]^+$, poly$(SO1861)_9$), 16.0 kDa ($[M+H]^+$, poly$(SO1861)_8$), 14.2 kDa ($[M+H]^+$, poly$(SO1861)_7$), 12.2 kDa ($[M+H]^+$, poly$(SO1861)_6$), 10.2 kDa ($[M+H]^+$, poly$(SO1861)_5$), 8.2 kDa ($[M+H]^+$, poly$(SO1861)_4$), 6.2 kDa ($[M+H]^+$, poly$(SO1861)_3$).

SO1861-EMCH peptide coupling

[0346] Customized peptide with the sequence SESDDAMFCDAMDESDSK (0.6 mg, 0.3 µmol) and SO1861-EMCH (0.8 mg, 0.39 µmol) were dissolved separately in 200 µL PBS. SO1861-EMCH and peptide solutions were mixed and shaken for 12 h at 800 rpm and room temperature on a ThermoMixer C (Eppendorf). After shaking small aliquots were taken out for analysis. Yield: not determined MALDI-TOF-MS (Figure 51 B) (RN mode): *m/z* 4.05 kDa ($[M+H]^-$, peptide-SO1861), 3.92 kDa ($[M+H]^-$, peptide-SO1730), 1.98 kDa ([M+H]-, peptide), 1.86 kDa ([M+H]-, SO1861).

Cell viability assay

[0347] HeLa cells were seeded in DMEM (PAN-Biotech GmbH) supplemented with 10% fetal bovine serum (PAN-Biotech GmbH) and 1% penicillin/streptomycin (PAN-Biotech GmbH), in a 96 well plate at 5,000 c/w in 100 µL/well and incubated overnight at 37 °C and 5% CO_2. The next day 20x concentrated stocks of the PAMAM, PAMAM-conjugates, G4-dendron (prepared in Example 3) or chloroquine (Sigma Aldrich) samples were prepared in DMEM. The media was removed from the cell culture plate and replaced by 160 µL culture media, followed by the addition of 10 µL sample/well (from the 20x concentrated stocks) and a 45 min incubation at 37 °C. During this incubation the SO1861 concentration curve was prepared. The SO1861 master stock was heated for 10 min at 50 °C, while shaking at 1,250 rpm. Followed by 15 sec sonication and a brief re-heating at 50 °C for 1 min, while shaking at 1,250 rpm. Subsequently a serial dilution of SO1861 was prepared in PBS. The SO1861 concentration curve was prepared as 10x concentrated stock, from which 20 µL was added/well. After a 15 min incubation at 37 °C, 10 µL dianthin-EGF (prepared in Example 2) diluted in

DMEM to 30 pM) or DMEM containing an equal amount of PBS was added/well, to obtain a final dianthin-EGF concentration of 1.5 pM as well as the indicated SO1861 and the different polymeric structures or chloroquine concentrations in a final volume of 200 µL/well.

[0348] After treatment the cells were incubated for 72 hr at 37 °C before the cell viability was determined by a MTS-assay, performed according to the manufacturer's instruction (CellTiter 96® AQueous One Solution Cell Proliferation Assay, Promega). Briefly, the MTS solution was diluted 20× in DMEM without phenol red (PAN-Biotech GmbH) supplemented with 10% FBS. The cells were washed once with 200 µL/PBS well, after which 100 µL diluted MTS solution was added/well. The plate was incubated for approximately 20-30 minutes at 37 °C. Subsequently, the OD at 492 nm was measured on a Thermo Scientific Multiskan FC plate reader (Thermo Scientific). For quantification the background signal of 'medium only' wells was subtracted from all other wells, before the cell viability percentage of treated/untreated cells was calculated, by dividing the background corrected signal of treated wells over the background corrected signal of the untreated wells (x 100).

FACS analysis

[0349] HeLa cells were seeded in DMEM (PAN-Biotech GmbH) supplemented with 10% fetal calf serum (PAN-Biotech GmbH) and 1% penicillin/streptomycin (PAN-Biotech GmbH), at 500,000 c/plate in 10 cm dishes and incubated for 48 hrs (5% CO_2, 37 °C), until a confluency of 90% was reached. Next, the cells were trypsinized (TryplE Express, Gibco Thermo Scientific) to single cells. 0.75 × 10^6 Cells were transferred to a 15 mL falcon tube and centrifuged (1,400 rpm, 3 min). The supernatant was discarded while leaving the cell pellet submerged. The pellet was dissociated by gentle tapping the falcon tube on a vortex shaker and the cells were washed with 4 mL cold PBS (Mg^{2+} and Ca^{2+} free, 2% FBS). After washing the cells were resuspended in 3 mL cold PBS (Mg^{2+} and Ca^{2+} free, 2% FBS) and divided equally over 3 round bottom FACS tubes (1 mL/tube). The cells were centrifuged again and resuspended in 200 µL cold PBS (Mg^{2+} and Ca^{2+} free, 2% FBS) or 200 µL antibody solution; containing 5 µL antibody in 195 µL cold PBS (Mg^{2+} and Ca^{2+} free, 2% FBS). APC Mouse IgG1, κ Isotype Ctrl FC (#400122, Biolegend) was used as isotype control, and APC anti-human EGFR (#352906, Biolegend) was used to stain the EGFR receptor. Samples were incubated for 30 min at 4 °C on a tube roller mixer. Afterwards, the cells were washed 3x with cold PBS (Mg^{2+} and Ca^{2+} free, 2% FBS) and fixated for 20 min at room temperature using a 2% PFA solution in PBS. Cells were washed 2x with cold PBS, and resuspended in 250-350 µL cold PBS for FACS analysis. Samples were analyzed with a BD FACSCanto II flow cytometry system (BD Biosciences) and FlowJo software. Table 19 shows the eExpression levels of EGFR, HER2 and CD71 of various cells.

Table 19. *Expression levels of EGFR, HER2 and CD71 of various cells*

Cell line		EGFR expression level (MFI)		HER2 expression level (MFI)		CD71 expression level (MFI)	
	MDA-MB-468		1656		1		186
	A431		1593		10		322
	CaSki		481		12		189
	SK-BR-3		28		1162		331
	JIMT-1		58		74		107
	HeLa		91		7		312
	A2058		1		5		59

Results

[0350] Considering available chemical groups for conjugation reactions to the SO1861 molecule, four chemical groups have been identified. The alcohols and diols of the sugar residues, the aldehyde group on the triterpenoid backbone, the carboxylic acid on one of the sugar residues (glucuronic acid), and the alkene group on the triterpenoid backbone as highlighted in Figure 30.

[0351] In view of the pros and cons of each identified chemical group (Table 20), the aldehyde and alcohol groups are best suitable for reversible conjugation reactions, while the alkene and the carboxylic acid (glucuronic acid) are the groups best suitable for irreversible / stable conjugation reactions. The aldehyde group within the molecule structure of SO1861, however, is the most suitable for reversible conjugation reactions over the alcohols. On the one hand, because there is only one aldehyde present in the structure that allows chemoselective reactions. On the other hand, because the aldehyde can perform reversible conjugation reactions with a variety of chemical groups such as amines, hydrazides, and hydroxylamines forming acid-cleavable moieties like imines, hydrazones, and oximes. This factor enables a freedom of choice over the chemical group for the desired reversible conjugation reaction. Contrary, the alcohols are good candidates for reversible conjugation reaction via the formation of acetals and ketals as well, but lack in chemoselectivity since they are present in a large quantity on the glycosidic structure.

[0352] For the formation of an irreversible and stable bond the carboxylic acid is the most suitable since it can form amides and esters with the common tools used in peptide chemistry (e.g. reaction with amines via carbodiimide mediated amide formation).

Table 20. Functional groups that are available for saponin conjugation reactions

Functional Group	Pros	Cons
Alcohol (Diols)	- Suitable for reversible acetal/ketal formation	- Acetal/ketal formation without chemoselectivity
	- Suitable for ester formations with activated carboxylic acids	- Ester formation without chemoselectivity
Aldehyde	- Suitable for chemoselective reversible hydrazone formation with hydrazides	- Not suitable for acetal formation in the presence of unprotected saponin sugar diols
	- Suitable for chemoselective reversible imine formation with amines	
	- Suitable for chemoselective reversible oxime formation with hydroxylamines	
Alkene	- Suitable for chemoselective irreversible radical reactions	- Not suitable for reversible conjugation reactions
		- Not suitable for reactions involving a hydrogenation step
Carboxylic acid	- Suitable for chemoselective amide / ester formation with amines and alcohols after activation	- Not suitable for reversible conjugation reactions under mild conditions

[0353] Thus, for the development of an endosomal escape enhancing saponin (such as SO1861) a methodology has been established that allows the generation of a non-cleavable and cleavable 'ready to conjugate' saponins (Figure 31) using the most suitable chemical groups present on SO1861.

[0354] For producing non-cleavable 'ready to conjugate' saponins the carboxylic group of SO1861 is activated via a reagent used in peptide coupling chemistry to generate an active ester (e.g. 1-

[Bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxide hexafluorophosphate, HATU). The resulting active ester of SO1861 is able to react with amines forming stable amide bonded conjugates (Figure 31 A).

[0355] For producing cleavable 'ready to conjugate' saponins the aldehyde group of SO1861 is reacted with an EMCH (ε-maleimidocaproic acid hydrazide) linker. The hydrazide group of EMCH forms an acid cleavable hydrazone bond with the aldehyde of SO1861. At the same time the EMCH linker presents a maleimide group that is thiol (sulfhydryl group) reactive and thus can be conjugated to thiols (Figure 31 B).

[0356] The maleimide group of SO1861-EMCH performs a rapid and specific Michael addition reaction with thiols and thiol bearing polymeric structures when carried out in a pH range of 6.5-7.5 (Figure 31 B). In addition, the acid sensitive hydrazone linkage between the SO1861 and EMCH can be utilized to perform saponin release from a scaffold in acidic environment (Figure 32). Thus, the EMCH linker fulfills both the need for a pH cleavable strategy and a conjugation strategy to a polymeric structure.

[0357] Regarding an ideal EMCH spacer length for conjugation to a polymeric structure, computer simulation (PerkinElmer, ChemBio3D, Ver. 13.0.0.3015) shows that the maleimide group on SO1861-EMCH is located at the periphery of the molecule and thus should be accessible for thiol bearing polymeric structures (Figure 33).

[0358] To synthesize the SO1861-EMCH, a strategy has been developed that allows the conversion of the aldehyde group on the SO1861 to EMCH (Figure 34 A). The SO1861-EMCH conjugate was isolated and successfully characterized via nuclear magnetic resonance spectroscopy (see materials and methods section, Figure 28) and matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF-MS) as shown on Figure 34 B and 34 C, and Figure 28 A.

[0359] For testing the pH dependent hydrolysis of the hydrazone bond, SO1861-EMCH was dissolved in an HCl solution at pH 3 and MALDI-TOF-MS spectra were recorded at two different points in time (Figure 35). As shown on Figure 35 A and 35 B, a clear decreasing tendency of the peak at m/z 2070 Da that corresponds to SO1861-EMCH is visible in Figure 35 B. Since SO1861 is generated during hydrolysis, an increase of the peak at m/z 1861 Da was recorded that accompanied the decreasing tendency at *m/z* 2070 Da. These results show that the hydrazone bond is responsive towards hydrolysis and gets cleaved even if it is attached on SO1861.

[0360] In order to conjugate SO1861-EMCH to a polymeric structure, accessible amines of the polymeric structure are converted into thiols with the aid of the agent 2-iminothiolane. The generated free thiols on the polymeric structure act then as the nucleophile for the thiol-ene Michael-type addition to the maleimide group of SO1861-EMCH (Figure 36). This developed methodology is suitable for the conjugation of SO1861-EMCH to any available polymeric structure that obtains accessible amine groups and allows furthermore a control over the number of conjugated SO1861 molecules depending on the polymeric structure, respectively.

[0361] First proof of concept for conjugation of 'ready-to conjugate saponins' to a polymeric structure was obtained by use of the amine of a protein (poly amino acid scaffold example), bovine serum albumin (BSA). After conjugation, mass spectrometry obtained the corresponding peaks of BSA-SO1861 at m/z ~ 70, ~72, and ~ 74 kDa (Figure 37 A). In comparison with the detected mass of BSA with *m/z* 66 kDa (Figure 37 B), the obtained masses of BSA-SO1861 correspond to a mixture of BSA-SO1861 conjugates consisting of 2, 3, and 4 SO1861 molecules per BSA.

[0362] Next proof of concept for conjugation of 'ready-to conjugate saponins' to a polymeric structure was obtained by the use of the amine bearing generation 5 (G5) dendrimer poly(amidoamine) (PAMAM with covalently coupled red-fluorescent dye (Cy3)). PAMAM-Cy3 was utilized as the polymeric structure for the conjugation to both SO1861-EMCH and SO1861-HATU and served as a model for conjugation of SO1861 to a polymeric structure (Figure 38).

[0363] All accessible amine groups of Cy3-PAMAM were converted into thiols using a 3 fold excess of 2-iminothiolane per Cy3-PAMAM amines followed by the reaction with SO1861-EMCH. Three different feed equivalents (5, 20 and 57) of SO1861-EMCH were used for the three reaction batches. After reaction, the recorded masses of the Cy3-PAMAM-SO1861 conjugates at MALDI-TOF-MS show an increment of the corresponding masses with increasing the SO1861-EMCH feed (Figure 39). The three different feeds corresponded to an obtained mass of *m/z* 38.4 kDa, *m/z* 53.9 kDa and *m/z* 133.8 kDa for the Cy3-PAMAM-SO1861 conjugates that correspond to 6, 13 and 51 SO1861 molecules attached per PAMAM dendrimer as shown on Figure 39 B-D.

[0364] In another reaction, only a certain number of PAMAM amines were converted into thiols prior to reaction with SO1861-EMCH. Here, two different feed equivalents of 2-Iminothiolane (8 and 32) and two different feed equivalents of SO1861-EMCH (5 and 30) were used, respectively. After reaction, the respective spectra of the Cy3-PAMAM-SO1861 conjugates at MALDI-TOF-MS show peaks at *m/z* 37.7 kDa (5 feed equivalents of SO1861-EMCH) and at *m/z* 87.0 kDa (30 feed equivalents of SO1861-EMCH) as shown on Figure 40. The obtained masses at *m/z* 37.7 kDa and *m/z* 87.0 kDa correspond to Cy3-PAMAM-SO1861 conjugates with 5 and 30 SO1861 molecules attached and demonstrate that with this method almost all feed of SO1861-EMCH were conjugated.

[0365] For the generation of a non-pH-cleavable saponin conjugate the carboxylic acid of SO1861 was activated with HATU and then reacted with the amines of Cy3-PAMAM forming a pH stable amide bound between Cy3-PAMAM and SO1861. The resulting mass of the conjugate was detected via MALDI-TOF-MS with a mass of *m/z* 62.3 kDa that corresponds to Cy3-PAMAM-NC-SO1861 (NC = non-cleavable) conjugate with 17.5 SO1861 molecules attached per PAMAM (Figure 38 B, Figure 41).

[0366] Next, the saponin conjugated scaffolds were conjugated to linking points for a possible conjugation to targeted therapeutics (e.g. targeted toxins) via the so-called strain-promoted alkyne-azide cycloaddition (SPAAC, click chemistry) to obtain a functionalized scaffold. For this reaction, Cy3-PAMAM-SO1861 (Figure 42 C, D) and Cy3-PAMAM-NC-SO1861 (Figure 42 B) were conjugated to a heterobifunctional NHS-PEG_{13}-DBCO linker that generated a strained alkyne on the conjugates' surface (Figure 42 A). The NHS (N Hydroxysuccinimide) moiety of the linker reacted with remaining amines of the PAMAM-saponin conjugates forming an amide bond between the scaffold and the linker. The resulting DBCO (dibenzocyclooctyne) moiety on the conjugates is able to perform SPAAC with corresponding azides on the targeted therapeutics.

[0367] Dianthin-EGF served as a model targeted toxin and dianthin served as a non-targeted toxin. Both toxins were labeled with Alexa Fluor™ 488 using the tetrafluorophenyl ester (TFP) derivative of the dye. The dye labeled proteins were then conjugated to a heterobifunctional NHS-SS-PEG_3-azide linker to obtain the corresponding chemical moiety for the SPAAC to the PAMAM-saponin conjugates. Maldi-TOF-MS measurements showed that one Alexa Fluor™ 488 dye and 9 NHS-SS-PEG3-azide molecules were attached per dianthin-EGF molecule (Figure 43, Figure 44). Furthermore, Alexa Fluor™ 488 labeled dianthin-EGF was conjugated to a heterobifunctional NHS-PEG_{12}-azide linker that lacked the disulfide bond and would thus generate a non-toxin-cleavable construct.

[0368] The Cy3-PAMAM-NC-SO1 861 -DBCO and Cy3-PAMAM-SO1861-DBCO conjugates were reacted with Alexa Fluor™ 488 labeled azido-toxins to perform a strain-promoted alkyne-azide cycloaddition. The conjugation between the reacting agents was indicated via gel electrophoresis and the co-localization of the fluorescent signals of Cy3 that is only attached on the PAMAM polymer and Alexa Fluor™ 488 that is only attached on the toxins on a polyacrylamide gel after gel electrophoresis (Figure 45).

[0369] As an alternative polymeric structure to the PAMAM dendrimer, a G4-dendron (PFd-G4-Azide-NH-BOC,

Polymer Factory) with 16 functional amino end groups and an azido group at the focal point was utilized for the conjugation to SO1861 (Figure 52). The advantage of using a dendron over a dendrimer is the focal point that the dendron structure is exhibiting. This focal point allows the direct conjugation to a targeted toxin without the need of its post-modification with orthogonal click functions (Figure 53). As shown in Figure 53, the dendron underwent the same methodology as described for the PAMAM dendrimer. Briefly, after partial dye labeling and deprotection (Figure 54), the amino groups of the dendron were converted into thiols using the thiolating reagent 2-iminothiolane followed by conjugation to SO1861-EMCH. For the conjugation to SO1861-EMCH three different feed equivalents of SO1861-EMCH were used. The dendron-SO1861 conjugates were analyzed via MALDI-TOF-MS. As expected, the conjugate species of 1 and 2 SO1861 molecules per dendron molecule were obtained when low SO1861-EMCH feed equivalents of 3 and 10 were used (Figure 55 B, C). Higher dendron-SO1861 conjugate species of up to 9 SO1861 molecules per dendron were obtained (Figure 55 A) when using a feed equivalent of 22 SO1861-EMCH molecules per dendron molecule. In further experiments, the saponin functionalized dendron will be conjugated to targeted toxins over its focal point to yield a functionalized scaffold and will be evaluated biologically.

[0370] The previous examples demonstrate that a methodology has been developed that allows the conjugation of a determined amount of SO1861 molecules or other endosomal escape enhancer molecules to a polymeric structure for enhanced cytoplasmic delivery of therapeutic substances such as targeted toxins.

[0371] To further test other conjugation methodologies of SO1861 to a polymeric structure, the reductive amination pathway was used. For this, the aldehyde group of SO1861 was directly conjugated to PAMAM amines forming an imine bound. The imine bond formation was followed a reductive amination step through the addition of the reductive agent sodium cyanoborohydride forming a pH-stable amine bond between SO1861 and PAMAM (Figure 46 A). Similar to the EMCH and HATU approach, this methodology allows a control over the number of conjugated saponins per polymer as shown on Figure 46 B, C. Here, PAMAM-saponin conjugates were produced which obtained a number of 10 (Figure 46B) and 30 (Figure 46 C) SO1861 molecules per PAMAM.

[0372] Another approach for the development of a SO1861 scaffold among the discussed polymer, and protein approach is the poly(SG1861) approach. The idea of this approach is to generate a polymerthat consists of SO1861 molecules only, with pH sensitive cleavable bonds that release the SO1861. In addition, the poly(SO1861) should be able to perform conjugation reactions to toxins and biopolymers. The main goal with this approach is to keep it as simple and cost effective as possible. Since a protocol for the generation of acid cleavable SO1861 has been developed already (SO1861-EMCH approach) it would be interesting to see if it is possible to polymerize the SO1861-EMCH through simple addition of a polymerization initiator without further modifying the SO1861 or identifying other conjugation sites on the SO1861 molecule. In the past, several papers have discussed the polymerization of maleimide groups by using radical initiators which attack the double bond of the maleimide group and thus initiate a radical polymerization along the double bonds of the maleimides (29-31). Since SO1861-EMCH reveals a maleimide group in its structure this group could potentially be explored for radical polymerization reactions to yield a poly(SO1861) with acid cleavable function. If the polymerization reaction has a reasonable reaction time the generated SO1861 polymers could be quenched with a radical quencher that not only quenches the reaction but also generates a functional group for toxin or biopolymer conjugation. Such a reaction scheme is illustrated in Figure 47. Here, the system of ammonium persulfate (APS) and tetramethylethylenediamine (TMEDA) is shown in an exemplary way as radical generator and aminopropanethiol serves as a model radical quencher. Using aminopropanethiol as a quencher exemplary, the generated amine group could be specifically further modified to a click-able group or being used to directly conjugate the poly(SO1861) to a toxin.

[0373] In free radical polymerization the reaction conditions have a huge influence on the polymer properties and the reaction outcome. For instance, temperature, monomer concentration, and initiator concentration play a major role for forming the polymer and have to be fine-tuned according to the desired polymer properties. As

radical polymerizations are usually carried out at temperatures above 50 °C, the first reactions have been performed at a temperature of 60 °C. It was interesting to see if the SO1861-EMCH polymerization can be initiated spontaneously and if APS and TMEDA would have an influence on the polymerization degree. Thus, three reactions have been carried out, using the same SO1861-EMCH concentration, but differ in their APS / TMEDA composition. In the first reaction only the SO1861-EMCH was heated up to 60 °C for 3 h, while the second reaction contained SO1861-EMCH and APS, and the third reaction contained SO1861-EMCH, APS, and TMEDA. (For these experiments the same amount and concentration of starting materials have been used which are mentioned in the Materials and Methods section "Poly(SO1861) synthesis"). The batches have been analyzed via MALDI-TOF-MS as shown on Figure 48 A-C. Interestingly it has been shown that SO1861-EMCH started to form oligomers consisting of 2, 3, 4, 5, and 6 SO1861 molecules spontaneously when heated up to 60 °C (Figure 48 A). The addition of 11^{-3} equivalents APS at the same temperature had no influence on this trend (Figure 48 B). When using the initiator system of APS / TMEDA, however, SO1861 oligomers of up to 9 SO1861 molecules with a molecular weight of 18.2 kDa could be detected (Figure 48 C). In addition, the obtained peaks for the oligomers seemed much bigger in comparison with the peaks in Figure 48 A and 48 B, indicating a higher consumption of SO1861-EMCH for this reaction.

[0374] To further fine-tune the reaction conditions, other initiators such as azo-initiators like 2,2'-azobis[2-methyl-N-(2-hydroxyethyl)propionamide] and azobisisobutyronitrile will be tested, as well as other polymerization techniques such as controlled radical polymerization (atom-transfer radical-polymerization, reversible addition-fragmentation chain transfer, etc). Moreover, another hydrazide linker as a substitute for EMCH could be considered which obtains a functional group (such as an acryl or acrolyol residue) that is more suitable for radical polymerization than the maleimide group.

[0375] Another approach for the development of a SO1861 scaffold is the DNA approach. The idea of this approach is to utilize the concept of the so-called DNA-origami (32, 33). DNA-origami as the polymeric or assembled polymeric structure to conjugate saponins to it, can offer several inherent advantages including stability, scalability, and precise control of the final size and shape of the resulting DNA-saponin scaffold. Since these DNA nanocarriers are comprised of natural DNA, they are biocompatible and do not show toxicity to living cells, and can ease the release of cargo from internal cellular compartments. The multivalency of such a structure can further allow fine-tuning targeting capabilities and high capacity for a variety of payloads such as fluorophores and toxins. Thus, in this approach DNA strands are identified that offer chemical functional groups on the 3' and 5' endings respectively, and that are able to hybridize only in certain wanted areas of the sequence that allow a control over the final shape of the construct. The chemical groups should be utilized to couple saponins, for instance though a thiol-ene reaction between the already developed SO1861-EMCH and a thiol group on one of the 3' and 5' DNA strands. The complementary DNA strand can offer a click function group that can be used for coupling to a targeted toxin. The concept is illustrated in Figure 49.

[0376] A similar approach is imaginable by using a specific peptide sequence instead of DNA strands that is able to bind and release saponins and that can be polymerized forming a large poly(peptide)-like structure. In this approach, a peptide sequence has been identified and purchased that has a length fitting the calculated size of a SO1861-EMCH molecule, that offers a cysteine residue in the middle of the sequence, and that obtains an amine group at both the N-terminus and C-terminus. The cysteine residue can be utilized to conjugate SO1861-EMCH via a thiol-ene reaction of the maleimide group of SO1861-EMCH and the thiol group of the cysteine residue. The two amine groups can be utilized to polymerize the peptide-SO1861 conjugate with a suitable crosslinker as shown on Figure 50.

[0377] Conjugation studies have shown that the conjugation of SO1861-EMCH to the customized peptide (sequence: SESDDAMFCDAMDESDSK) was successful. The peptide that bears a maleimide reactive cysteine in the middle of the sequence and its conjugation to SO1861-EMCH was analyzed via MALDI-TOF-MS (Figure 51). The MALDI-TOF-MS spectra shows the expected peak for the peptide-SO1861 conjugate at *m/z* 4053 Da and an additional peak at *m/z* 3920 Da which is the peptide-SO1861 conjugate of the corresponding saponin-

EMCH of SO1730. As SO1861-EMCH has been used in slight excess (1.3 equivalents) and no SO1861-EMCH peak was detected after reaction, it can be assumed that the conjugation was quantitative. For starting first polymerization reactions of the peptide-SO1861, disuccinimidyl tartrate will be utilized as the amine reactive cross-linker.

REFERENCES

[0378]

Weng, A.; Thakur, M.; Beceren-Braun, F.; Bachran, D.; Bachran, C.; Riese, S.B.; Jenett-Siems, K.; Gilabert-Oriol, R.; Melzig, M.F.; Fuchs, H. The toxin component of targeted anti-tumor toxins determines their efficacy increase by saponins. Molecular oncology 2012, 6, 323-332.

saponinum album in a synergistic way. Journal of immunotherapy 2009, 32, 713-725.

Sama, S.; Jerz, G.; Schmieder, P.; Joseph, J.F.; Melzig, M.F.; Weng, A. Plant derived triterpenes from Gypsophila elegans M.Bieb. enable non-toxic delivery of gene loaded nanoplexes. Journal of Biotechnology 2018, 284, 131-139

Kolb, H.C.; Finn, M.G.; Sharpless, K.B. Click Chemistry: Diverse Chemical Function from a Few Good Reactions. Angewandte Chemie 2001, 40, 2004-2021.

Bird, R.E.; Hardmann, K.D.; Jacobson, J.W.; Johnson, S.; Kaufman, B.M.; Lee, S.M.; Lee, T.; Pope, S.H.; Riordan, G.S.; Whitlow, M. Single-chain antigen-binding proteins. Science 1988, 242, 423-426.

Y Zhang, Z Qu, S Kim, V Shi, B Liao1, P Kraft, R Bandaru, Y Wu, LM Greenberger and ID Horak, Down-modulation of cancer targets using locked nucleic acid (LNA)-based antisense oligonucleotides without transfection, Gene Therapy (2011) 18, 326-333

REFERENCES CITED IN THE DESCRIPTION

Cited references

This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

Patent documents cited in the description

- WO9305789A [0012]

Non-patent literature cited in the description

- **B. YUD. LIU**Antibody-drug conjugates in clinical trials for lymphoid malignancies and multiple myelomaJournal of Hematology & Oncology, 2019, vol. 12, 94- [0006]
- **LUIS M. ALVAREZ-SALAS**(Nucleic Acids as Therapeutic AgentsCurrent Topics in Medicinal Chemistry, 2008, vol. 8, 1379-1404 [0007]
- **BHARGAVA et al.**Targeted dianthin is a powerful toxin to treat pancreatic carcinoma when applied in combination with the glycosylated triterpene SO1861Molecular Oncology, 2017, vol. 11, 111527-154 [0012]
- **GILABERT-ORIOL et al.**Combinatorial approach to increase efficacy of Cetuximab, Panitumumab and Trastuzumab by dianthin conjugation and co-application of SO1861Biochemical Pharmacology, 2015, vol. 97, 3247-255 [0012]
- **ALEXANDER WENG et al.**Saponins modulate the intracellular trafficking of protein toxinsJournal of Controlled Release, 2015, vol. 164, 174-86 [0012]
- **FROKIER H et al.**Optimization of hapten-protein conjugation by high-performance capillary electrophoresisJournal of Chromatography, 1995, vol. 717, 175-81 [0012]
- **DIRK C. VAN SETTENGERRIT VAN DE WERKENGIJSBERT ZOMERGIDEON F. A. KERSTEN,**Glycosyl Compositions and Structural Characteristics of the Potential Immuno-adjuvant Active Saponins in the Quillaja saponaria Molina Extract Quil ARAPID COMMUNICATIONS IN MASS SPECTROMETRY, 1995, vol. 9, 660-666 [0063]
- **JULIANE DEISE FLECKANDRESA HEEMANN BETTIFRANCINI PEREIRA DA SILVAEDUARDO ARTUR TROIANCRISTINA OLIVAROFERNANDO FERREIRASIMONE GASPARIN VERZA**Saponins from Quillaja saponaria and Quillaja brasiliensis: Particular Chemical Characteristics and Biological ActivitiesMolecules, 2019, vol. 24, 171- [0064]
- **Y ZHANG,Z QUS KIMV SHIB LIAO1P KRAFTR BANDARU,Y WU,LM GREENBERGERID HORAK**Down-modulation of cancer targets using locked nucleic acid (LNA)-based antisense oligonucleotides without transfection,Gene Therapy, 2011, vol. 18, 326-333 [0091]
- **SCHROT JWENG AMELZIG MF et al.**Ribosome-inactivating and related proteins. Toxins (Basel), 2015, vol. 7, 51556-615 [0154]
- **CONRADO PEDEBOSLAÉRCIO POL-FACHINRAMON PONSCILAINE V. TEIXEIRA HUGO VERLI**Atomic Model and Micelle Dynamics of QS-21 SaponinMolecules, 2014, vol. 19, 3744-3760 [0187]
- **WENG, A.THAKUR, MBECEREN-BRAUN, FBACHRAN, D.BACHRAN, CRIESE, S.BJENETT-SIEMS, KGILABERT-ORIOL, RMELZIG, M.FFUCHS, H**The toxin component of targeted anti-tumor toxins determines their efficacy increase by saponinsMolecular oncology, 2012, vol. 6, 323-332 [0378]
- saponinum album in a synergistic wayJournal of immunotherapy, 2009, vol. 32, 713-725 [0378]
- **SAMA, SJERZ, GSCHMIEDER, PJOSEPH, J.F.MELZIG, M.FWENG, A**Plant derived triterpenes from Gypsophila elegans M.Bieb. enable non-toxic delivery of gene loaded nanoplexesJournal of Biotechnology, 2018, vol. 284, 131-139 [0378]
- **KOLB, H.CFINN, M.GSHARPLESS, K.B**Click Chemistry: Diverse Chemical Function from a Few Good ReactionsAngewandte Chemie, 2001, vol. 40, 2004-2021 [0378]
- **BIRD, R.E.HARDMANN, K.DJACOBSON, J.WJOHNSON, SKAUFMAN, B.MLEE, S.MLEE, TPOPE, S.HRIORDAN, G.S.WHITLOW, M**Single-chain antigen-binding proteinsScience, 1988, vol. 242, 423-426 [0378]
- **Y ZHANGZ QUS KIMV SHIB LIAO1P KRAFTR BANDARUY WULM GREENBERGERID HORAK**Down-modulation of cancer targets using locked nucleic acid (LNA)-based antisense oligonucleotides without transfectionGene Therapy, 2011, vol. 18, 326-333 [0378]

Patentkrav

1. Terapeutisk kombination til anvendelse som et medikament, hvor den terapeutiske kombination omfatter:

(a) en første farmaceutisk sammensætning med et første proteinholdigt molekyle, der omfatter et bindingssted til binding til en epitop på et celleoverflademolekyle og mindst én saponin, der er kovalent bundet til det første proteinholdige molekyle, fortrinsvis til en aminosyrerest af det første proteinholdige molekyle, hvor den første farmaceutiske sammensætning eventuelt yderligere omfatter en farmaceutisk acceptabel excipiens; og

(b) en anden farmaceutisk sammensætning med et andet proteinholdigt molekyle, hvor det andet proteinholdige molekyle omfatter et bindingssted til binding til epitopen på celleoverflademolekylet af (a) og en effektorenhed, hvor den anden farmaceutiske sammensætning eventuelt yderligere omfatter en farmaceutisk acceptabelt excipiens,

hvor bindingsstedet af det første proteinholdige molekyle og bindingsstedet for det andet proteinholdige molekyle er det samme, og hvor celleoverflademolekylet og epitopen på celleoverflademolekylet, som det første proteinholdige molekyle kan binde til, og celleoverflademolekylet og epitopen på celleoverflademolekylet, som det andet proteinholdige molekyle kan binde til, er de samme;

og hvor bindingsstedet for det første proteinholdige molekyle og det andet proteinholdige molekyle omfatter eller består af et immunglobulin eller et bindingsfragment eller bindingsdomæne af immunglobulinet, såsom et hvilket som helst eller flere af et antistof, et IgG, et molekyle omfattende eller bestående af af et Vhh-domæne eller Vh-domæne, et Fab, et scFv, et Fv, et dAb, et F(ab)2, Fcab-fragment og/eller omfatter eller består af mindst én ligand, hvor liganden er epidermal vækstfaktor (EGF);

og hvor det mindst ene saponin er en triterpenoid saponin og/eller en bisdesmosidisk triterpensaponin, der tilhører typen af en 12,13-dehydrooleanan med en aldehydfunktion i position C-23;

og hvor effektorenheden omfatter eller består af et hvilket som helst eller flere af et oligonukleotid, en nukleinsyre og en xenonukleinsyre; eller omfatter eller består af mindst et proteinholdigt molekyle, valgt blandt et eller flere af et viralt toksin, et bakterielt toksin, et svampetoksin, et plantetoksin inklusive ribosominaktiverende proteiner og A-kæden af type 2 ribosominaktiverende proteiner, et animalsk toksin eller et humant toksin.

2. Terapeutisk kombination til anvendelse i behandlingen eller forebyggelsen af en cancer hos et menneske, hvor den terapeutiske kombination omfatter:
(a) den første farmaceutiske sammensætning ifølge krav 1; og
(b) den anden farmaceutiske sammensætning ifølge krav 1,
hvor celleoverflademolekylet er udtrykt på en tumorcelleoverflade, og celleoverflademolekylet fortrinsvis er et tumorcellespecifikt overflademolekyle, og hvor epitopen fortrinsvis er en tumorcellespecifik epitop,
fortrinsvis hvor det andet farmaceutiske præparat ifølge krav 1 og det første farmaceutiske præparat ifølge krav 1 indgives til en patient med behov derfor.

3. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-2, hvor bindingsstedet af det første proteinholdige molekyle og det andet proteinholdige molekyle omfatter eller består af et immunoglobulin eller et bindende fragment eller bindingsdomæne af immunglobulinet, såsom et hvilket som helst eller flere af et antistof, et IgG, et molekyle omfattende eller bestående af et Vhh-domæne eller Vh-domæne, et Fab, et scFv, et Fv, et dAb, et F(ab)2, Fcab-fragment, fortrinsvis hvor tumorcelleoverflademolekylet er en celleoverfladereceptor, der er specifikt til stede ved en tumorcelle, og hvor epitopen fortrinsvis er en tumorcellespecifik epitop.

4. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-3, hvor det mindst ene saponin omfatter en glucuronsyrefunktion i en kulhy-

dratsubstituent ved C-3beta-OH-gruppen af saponinen, fortrinsvis hvor saponinen er SO1861 og/eller QS-21, mest foretrukket hvor saponinen er SO1861.

5. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-4, hvor den mindst ene saponin er en bisdesmosidisk triterpensaponin, der tilhører typen af en 12,13-dehydrooleanan med en aldehydfunktion i position C-23, hvor saponinen er kovalent koblet til det første proteinholdige molekyle, fortrinsvis kovalent koblet til en aminosyrerest af det første proteinholdige molekyle, via en aldehydfunktion i saponinen, fortrinsvis aldehydfunktionen i position C-23, fortrinsvis via mindst én linker, og/eller via mindst én spaltelig linker, hvor aminosyreresten fortrinsvis er valgt blandt cystein og lysin, fortrinsvis hvor aldehydfunktionen i position C-23 af den mindst ene saponin er kovalent koblet til linker N-ε-maleimidocaproinsyre-hydrazid, hvilken linker er kovalent koblet via en thioetherbinding til en sulfhydrylgruppe i det første proteinholdige molekyle, såsom en sulfhydrylgruppe i et cystein.

6. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-5, hvor den mindst ene saponin er et bisdesmosidisk triterpen-saponin, der tilhører typen af en 12,13-dehydrooleanan med en aldehydfunktion i position C-23 og omfatter en glucuronsyrefunktion i en kulhydratsubstituent ved C-3beta-OH-gruppen af saponinen, hvor saponinen er kovalent koblet til en aminosyrerest i det første proteinholdige molekyle via glucuronsyrefunktionen i kulhydratsubstituenten ved C-3beta-OH-gruppen af saponinen, fortrinsvis via mindst én linker, hvor aminosyreresten fortrinsvis er valgt blandt cystein og lysin, fortrinsvis hvor glucuronsyrefunktionen i kulhydratsubstituenten ved C-3beta-OH-gruppen af den mindst ene saponin er kovalent koblet til linker 1-[Bis(dimethylamino)methylen]-1H-1,2,3-triazolo[4,5-b]pyridinium-3-oxid-hexafluorphosphat, hvilken linker er kovalent koblet via en amidbinding til en amingruppe i det første proteinholdige molekyle, såsom en amingruppe af en lysin eller en N-terminal af det første proteinholdige molekyle.

7. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-6, hvor epitopen er en epitop på en tumorcellereceptor, fortrinsvis en tumorcellespecifik epitop, og hvor receptoren fortrinsvis er en tumorcellespecifik receptor, mere fortrinsvis en receptor valgt blandt CD71, CA125, EpCAM(17-1A), CD52, CEA, CD44v6, FAP, EGF-IR, integrin, syndecan-1, vaskulært integrin-alfa-V-beta-3, HER2, EGFR, CD20, CD22 , Folatreceptor 1, CD146, CD56, CD19, CD138, CD27L-receptor, PSMA, CanAg, integrin-alphaV, CA6, CD33, mesothelin, Cripto, CD3, CD30, CD239, CD70, CD123, CD352, DLL3, CD25, ephrinA4, MUC1, Trop2, CEACAM5, CEACAM6, HER3, CD74, PTK7, Notch3, FGF2, C4.4A, FLT3, CD38, FGFR3, CD7, PD-L1, CTLA4, CD52, PDGFRA, VEGFR1, VEGFR2, mest fortrinsvis valgt blandt CD71 HER2 og EGFR, fortrinsvis hvor tumorcellereceptoren, fortrinsvis den tumorcellespecifikke receptor, internaliseres af tumorcellen efter binding til det første proteinholdige molekyle og/eller efter binding til det andet proteinholdige molekyle, og hvor fortrinsvis binding af det første proteinholdige molekyle og/eller det andet proteinholdige molekyle til tumorcellereceptoren, såsom den tumorcellespecifikke receptor, efterfølges af tumorcellereceptormedieret internalisering, f.eks. via endocytose, af et kompleks af det første proteinholdige molekyle og tumorcellereceptoren og af et kompleks af det andet proteinholdige molekyle og tumorcellereceptoren.

8. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-7, hvor bindingsstedet af det første proteinholdige molekyle og det andet proteinholdige molekyle omfatter eller består af et hvilket som helst af 3B7, ADCT-502, Affilutin, AGS-16M18, AGS-67E, alemtuzumab, AMG-172, AMG-595, ASG-15ME, ASG-5ME, ASP-6183, AVE-9633, Avicidin, BAY-794620, B-B4, Bismab-A, bivatuzumab, BIWI-1, brentuximab, cantuzumab-ravtansin, CDX-014, cetuximab, Cixutumumab, clivatuzumab-tetraxetan, CNTO 95, coltuximab-ravtansin, daratumumab, DCDS-0780A, denintuzumab-mafodotin, depatuxizumab-mafodotin, DS6, edrecolomab, epratuzumab, etaracizumab, gemtuzumab, glembatumumab-vedotin, Herceptin, HTI-1511, huB4, huC242,

HuMax-CD74-ADC, huMy9-6, huN901, ibritumomab, ibritumomab-tiuxetan, IMGN-289, IMGN-388, IMGN-779, ImmuRAIT-LL2, indusatumab-vedotin, Inotuzumab, inotuzumab-ozogamicin, labetuzumab, lifastuzumab-vedotin, lorvotuzumab-mertansin, MLN-2704, MM-302, MM-310, Moxetumomab, Neuradiab, nimotuzumab, Obinutuzumab, ofatumumab, OKT-10-anti-CD38-monoklonalt antistof, OKT-9-anti-CD71-monoklonalt antistof af IgG-typen, Oncolysin B, Oncolysin S, oregovomab, panitumumab, pertuzumab, PF-06263507, PF-06647263, PF-06650808, Pinatuzumab, pinatuzumab-vedotin, Polatuzumab, RG-7598, RG-7600, RG-7636, RG-7841, RG-7882, rituximab, rovalpituzumab-tesirin, sacituzumab-govitecan, SAR-428926, SAR-566658, SC-004, SC-006, SGN-15, SGNCD-123A, SGNCD-19B, SGNCD-352A, SGN-CD70A, sibrotuzumab, sofituzumab-vedotin, tositumomab, Trastuzumab, trastuzumab-emtansin, vadastuximab-talirin, vandortuzumab-vedotin, vorsetuzumab-mafodotin, XMT-1522 eller ZW-33, fortrinsvis cetuximab eller trastuzumab eller OKT-9, eller mindst ét celleoverflademolekyle-bindende fragment og/eller -domæne deraf, forudsat, at det første bindingssted er det samme som det andet bindingssted.

9. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-8, hvor effektorenheden, der er omfattet af det andet proteinholdige molekyle, omfatter eller består af et hvilket som helst eller flere af et oligonukleotid, en nukleinsyre og en xenonukleinsyre, valgt blandt en hvilken som helst eller flere af en vektor, et gen, et cellesuicid-inducerende transgen, deoxyribonukleinsyre (DNA), ribonukleinsyre (RNA), antisense-oligonukleotid (ASO, AON), kort interfererende RNA (siRNA), mikroRNA (miRNA), DNA-aptamer, RNA-aptamer, mRNA, minicirkel-DNA, peptidnukleinsyre (PNA), phosphoramidatmorpholinoligomer (PMO), låst nukleinsyre (LNA), broforbundet nukleinsyre (BNA), 2'-deoxy-2'-fluorarabino-nukleinsyre (FANA), 2'-O-methoxyethyl-RNA (MOE), 2'-O,4'-aminoethylenbroforbundet nukleinsyre, 3'-fluorhexitolnukleinsyre (FHNA), et plasmid, glycolnukleinsyre (GNA) og threose-nukleinsyre

(TNA), mere fortrinsvis en BNA, for eksempel en BNA til at dæmpe HSP27-proteinekspression, og/eller

hvor effektorenheden, der er omfattet af det andet proteinholdige molekyle, omfatter eller består af mindst ét proteinholdigt molekyle, valgt blandt et eller flere af apoptin, Shiga-toksin, Shiga-lignende toksin, Pseudomonas aeruginosa-exotoxin (PE) eller exotoxin A af PE, fuld-længde eller trunkeret difteritoksin (DT), koleratoksin, dianthin f.eks. dianthin-30 eller dianthin-32, saporin, f.eks. saporin-S3 eller saporin-S6, bouganin eller afimmuniseret derivat debouganin af bouganin, shiga-lignende toksin A, antiviralt protein af kermesbær, ricin, ricin-A-kæde, modeccin, modeccin-A-kæde, abrin, abrin A-kæde, volkensin, volkensin-A-kæde, viscumin, viscumin-A-kæde, frø-RNase, granzym B eller angiogenin fra mennesker; fortrinsvis er proteintoksinet dianthin og/eller saporin,

fortrinsvis hvor den effektorenhed, der er omfattet af det andet proteinholdige molekyle, er en BNA til at dæmpe HSP27-proteinekspression, dianthin eller saporin.

10. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-9, hvor det andet proteinholdige molekyle omfatter eller består af et hvilket som helst af ADCT-502, Affilutin, AGS-16M18, AGS-67E, AMG-172, AMG-595, ASG-15ME, ASG-5ME, ASP-6183, AVE-9633, Avicidin, BAY-794620, Bismab-A, BIWI-1, Brentuximab-vedotin, cantuzumab-ravtansin, CDX-014, clivatuzumab-tetraxetan, coltuximab-ravtansin, DCDS-0780A, denintuzumab-mafodotin, depatuxizumab-mafodotin, Gemtuzumab-ozogamicin, glembatumumab-vedotin, HTI-1511, HuMax-CD74-ADC, ibritumomab-tiuxetan, IMGN-289, IMGN-388, IMGN-779, ImmuRAIT-LL2, indusatumab-vedotin, inotuzumab-ozogamicin, lifastuzumab-vedotin, lorvotuzumab-mertansin, MLN-2704, MM-302, MM-310, Moxetumomab-pasudotox, Neuradiab, Oncolysin B, Oncolysin S, PF-06263507, PF-06647263, PF-06650808, pinatuzumab-vedotin, Polatuzumab-vedotin, RG-7598, RG-7600, RG-7636, RG-7841, RG-7882, rovalpituzumab-tesirin, sacituzumab-govitecan, SAR-428926, SAR-

566658, SC-004, SC-006, SGN-15, SGNCD-123A, SGNCD-19B, SGNCD-352A, SGN-CD70A, sofituzumab-vedotin, trastuzumab-emtansin, vadastuximab-talirin, vandortuzumab-vedotin, vorsetuzumab-mafodotin, XMT-1522 og ZW-33, eller mindst ét tumorcellereceptorbindingsdomæne deraf og/eller mindst ét tumorcellereceptorbindingsfragment deraf, hvor domænet/domænerne eller fragmentet/fragmenterne omfatter effektorenheden og fortrinsvis er (a) tumorcellespecifikke receptorbindingsdomæne(r) og/eller (a) tumorcellespecifikke receptorbindingsfragment(er).

11. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-10, hvor det første proteinholdige molekyle omfatter mere end én kovalent bundet saponin, fortrinsvis 2, 3, 4, 5, 6, 8, 10, 16, 32, 64, 128 eller 1-100 saponiner eller et hvilket som helst antal saponiner derimellem, såsom 7, 9, 12 saponiner.

12. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 1-11, hvor den mindst ene saponin er kovalent bundet direkte til en aminosyrerest af det første proteinholdige molekyle, fortrinsvis til et cystein og/eller til et lysin, og/eller er kovalent bundet via mindst én linker og/eller via mindst én spaltelig linker og/eller via mindst ét oligomer eller polymer stillads, fortrinsvis 1-8 af sådanne stilladser eller 2-4 af sådanne stilladser, hvor det mindst ene stillads er eventuelt baseret på en dendron, hvor 1-32 saponiner, såsom 2, 3, 4, 5, 6, 8, 10, 16, 32 saponiner, eller et hvilket som helst antal saponiner derimellem, såsom 7, 9, 12 saponiner, er kovalent bundet til det mindst ene stillads,
fortrinsvis hvor den mindst ene saponin er kovalent bundet via mindst en spaltelig linker, og hvor den spaltelige linker er genstand for spaltning under sure betingelser, reduktive betingelser, enzymatiske betingelser eller lys-inducerede betingelser, og fortrinsvis omfatter den spaltelige linker en spaltelig binding udvalgt fra en hydrazonbinding og en hydrazidbinding, der er udsat for

spaltning under sure forhold, og/eller en binding, der er modtagelig for proteolyse, for eksempel proteolyse af Cathepsin B, og/eller en binding, der er modtagelig for spaltning under reduktive betingelser, såsom en disulfidbinding, mere foretrukket, hvor den spaltelige linker udsættes for spaltning *in vivo* under sure betingelser som til stede i endosomer og/eller lysosomer af pattedyrceller, fortrinsvis humane celler, fortrinsvis ved pH 4,0 - 6,5, og mere foretrukket ved pH < 5,5.

13. Terapeutisk kombination til anvendelse ifølge krav 12, hvor det oligomere eller polymere stillads omfatter en polymer eller oligomer struktur og omfatter mindst én kemisk gruppe, hvor den mindst ene kemiske gruppe til kovalent kobling af stilladset til aminosyreresten af det første proteinholdige molekyle, fortrinsvis hvor den mindst ene saponin er kovalent bundet til den polymere eller oligomere struktur af stilladset via den spaltelige linker som defineret i krav 12, og/eller hvor den mindst ene saponin er kovalent bundet til den polymere eller oligomere struktur af stilladset via en eller flere af en iminbinding, en hydrazonbinding, en hydrazidbinding, en oximbinding, en 1,3-dioxolanbinding, en disulfidbinding, en thioetherbinding, en amidbinding, en peptidbinding eller en esterbinding, fortrinsvis via mindst én linker.

14. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 12-13, hvor den mindst ene saponin omfatter en aldehydfunktion i position C-23 og eventuelt en glucuronsyrefunktion i kulhydratsubstituenten ved C-3beta-OH-gruppen af den mindst ene saponin, hvilken aldehydfunktion er involveret i den kovalente binding til den polymere eller oligomere struktur af stilladset, og/eller, hvis den er til stede, glucuronsyrefunktionen er involveret i den kovalente binding til den polymere eller oligomere struktur af stilladset enten via direkte binding eller via mindst én linker, fortrinsvis hvor aldehydfunktionen i position C-23 af den mindst ene saponin er kovalent koblet til linker N-ε-maleimidocaproinsyrehydrazid, hvilken linker er kovalent koblet via en thioetherbinding til en sulfhydrylgruppe i den polymere eller oligomere struktur af

stilladset, såsom en sulfhydrylgruppe i en cystein, og/eller hvor glucuronsyren fungerer i kulhydratsubstituenten ved C-3beta-OH-gruppen i den mindst ene saponin er kovalent koblet til linker 1-[Bis(dimethylamino)methylen]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxid-hexafluorphosphat, hvilken linker er kovalent koblet via en amidbinding til en amingruppe i den polymere eller oligomere struktur af stilladset, såsom en amingruppe af en lysin eller en N-terminal af et proteinholdigt molekyle.

15. Terapeutisk kombination til anvendelse ifølge et hvilket som helst af kravene 12-14, hvor den mindst ene kemiske gruppe af stilladset, til kovalent kobling af det oligomere eller polymere stillads til aminosyreresten af det første proteinholdige molekyle, er en klik-kemigruppe, fortrinsvis valgt blandt en tetrazin, et azid, en alken eller en alkyn, eller et cyklisk derivat af disse grupper, mere fortrinsvis er klik-kemigruppen et azid, og/eller hvor den polymere eller oligomere struktur af stilladset omfatter en lineær, forgrenet og/eller cyklisk polymer, oligomer, dendrimer, dendron, dendroniseret polymer, dendroniseret oligomer, polylysin, en polyethylenglycol eller en samling af disse polymere eller oligomere strukturer, hvilken samling fortrinsvis er bygget op af kovalent krydsbinder.

DRAWINGS

Figure 1

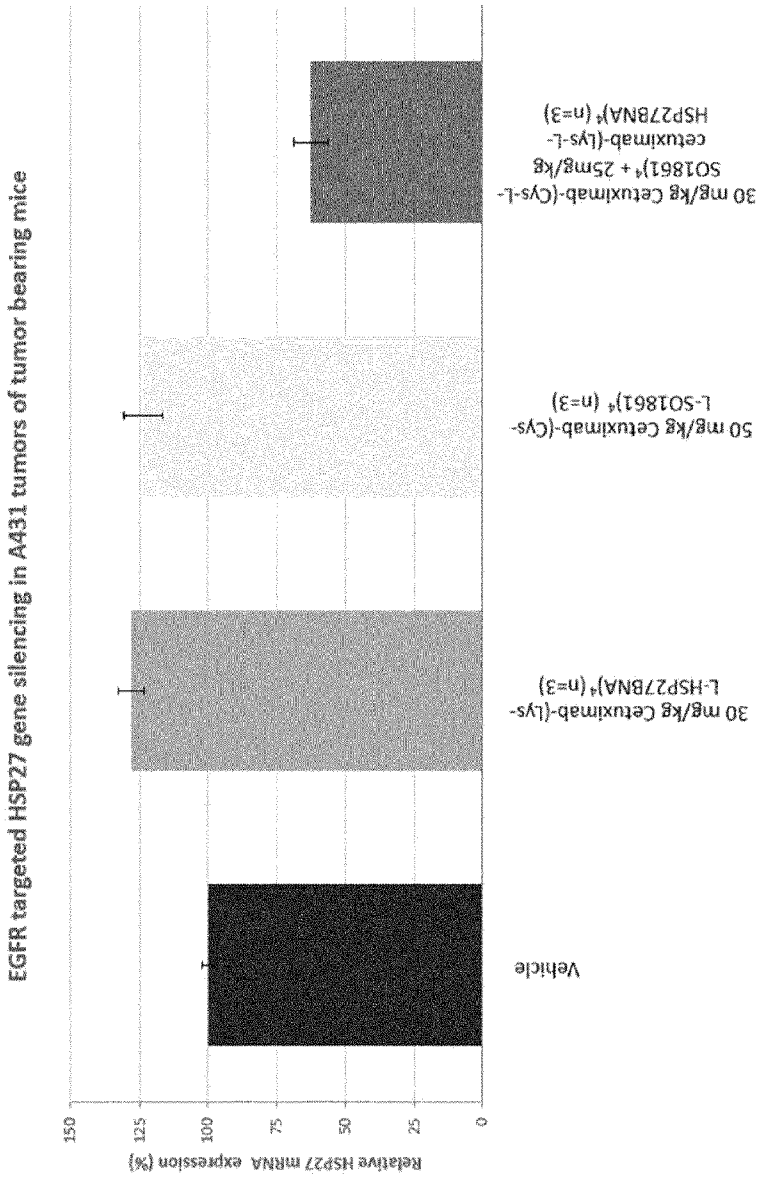


Figure 2

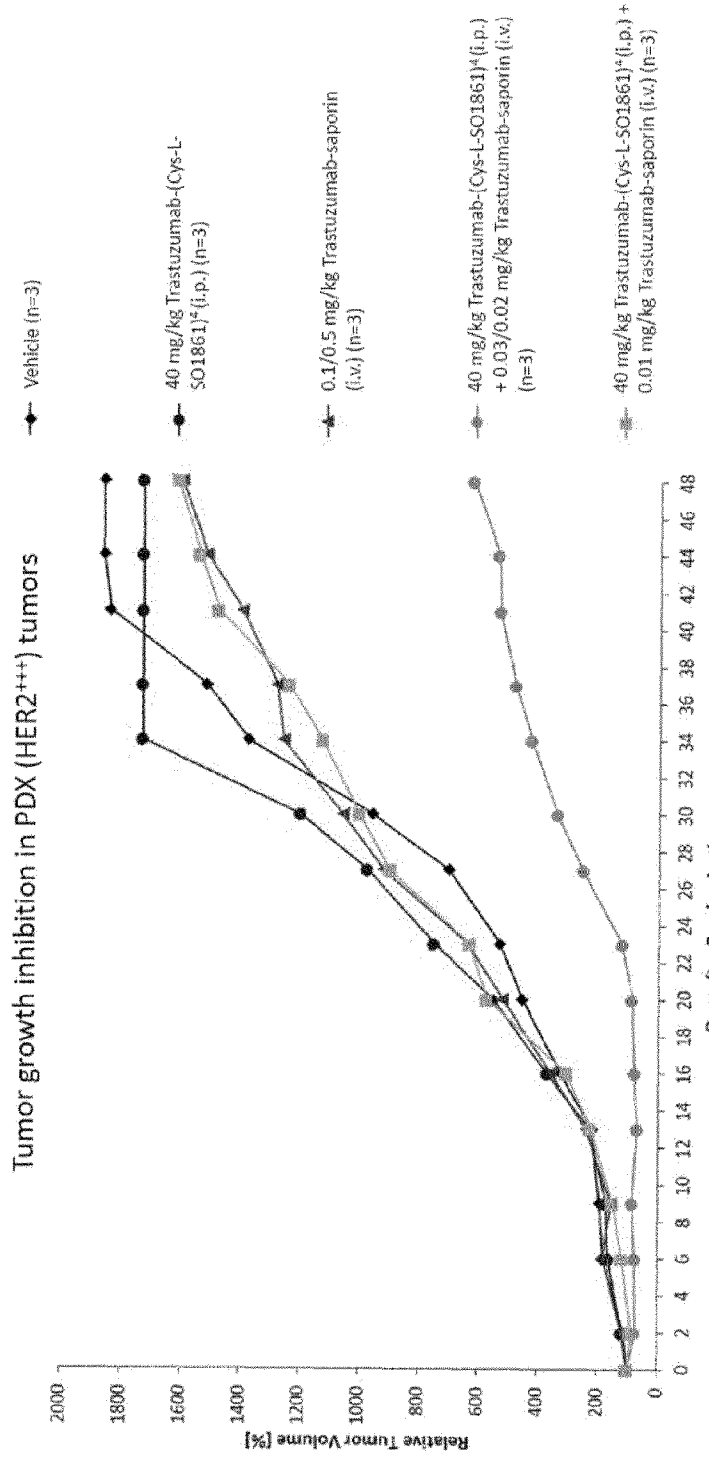

Tumor growth inhibition in PDX (HER2+++) tumors
Vehicle (n=3)
40 mg/kg Trastuzumab-(Cys-L-SO1861)4 (i.p.) (n=3)
0.1/0.5 mg/kg Trastuzumab-saporin (i.v.) (n=3)
40 mg/kg Trastuzumab-(Cys-L-SO1861)4 (i.p.) + 0.03/0.02 mg/kg Trastuzumab-saporin (i.v.) (n=3)
40 mg/kg Trastuzumab-(Cys-L-SO1861)4 (i.p.) + 0.01 mg/kg Trastuzumab-saporin (i.v.) (n=3)
Relative Tumor Volume [%]
Days after Randomization

Figure 3

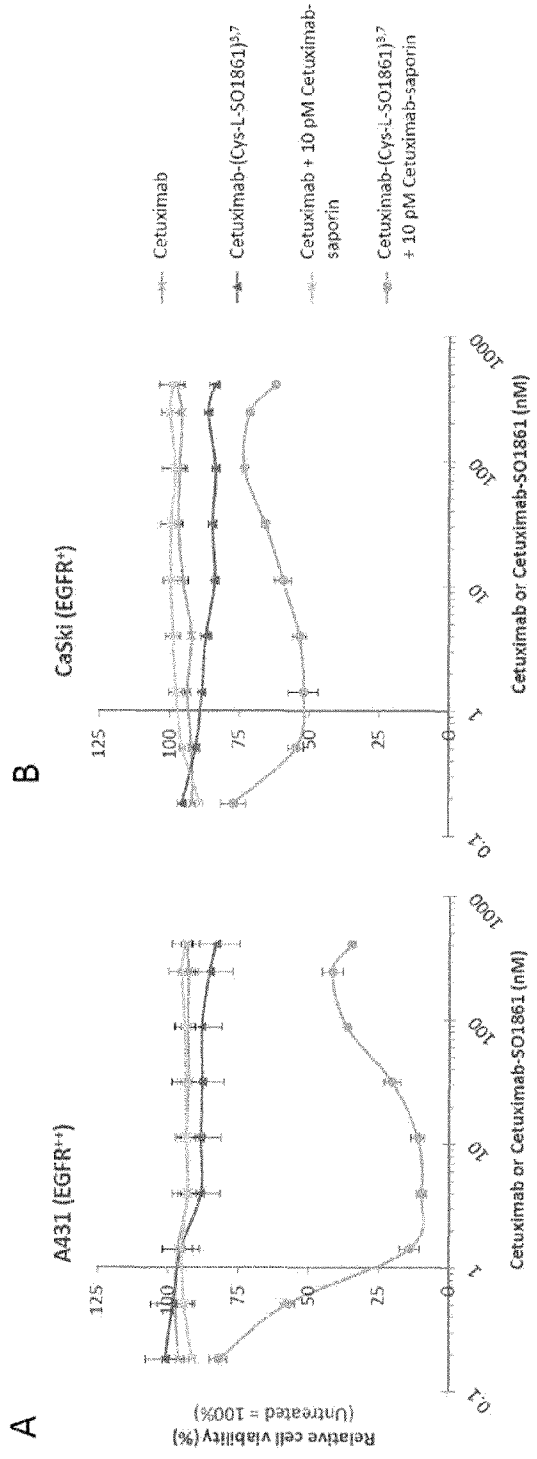

A
A431 (EGFR++)
B
CaSki (EGFR+)
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0.1
1
10
100
1000
Cetuximab or Cetuximab-SO1861 (nM)
Cetuximab
Cetuximab-(Cys-L-SO1861)3.7
Cetuximab + 10 pM Cetuximab-saporin
Cetuximab-(Cys-L-SO1861)3.7 + 10 pM Cetuximab-saporin

Figure 3

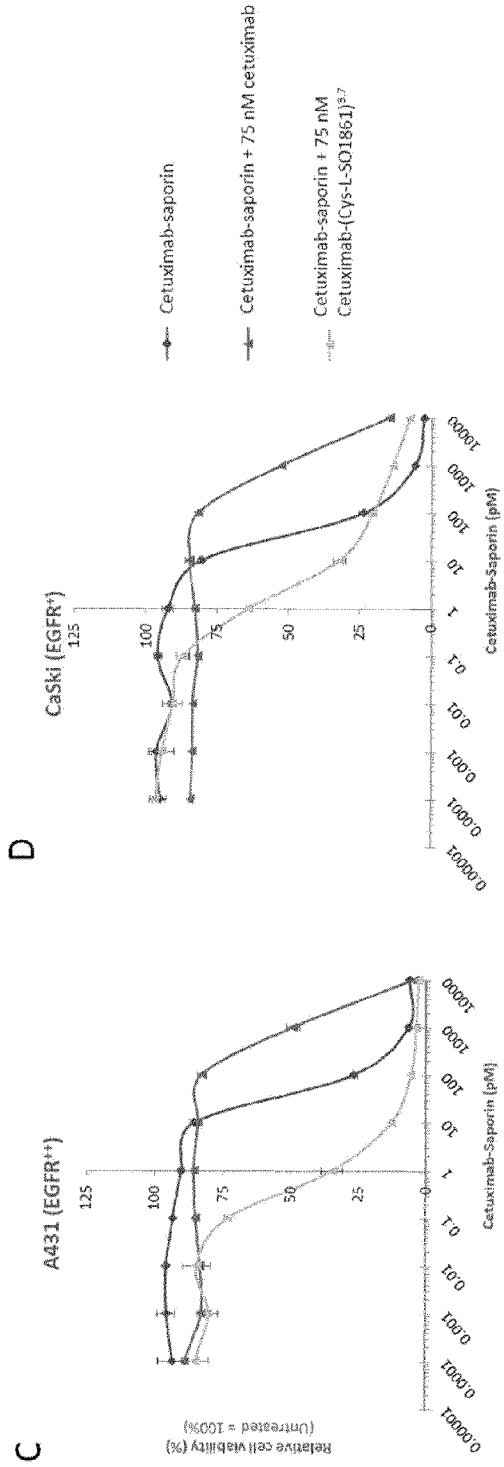

C
A431 (EGFR++)
D
CaSki (EGFR+)
Relative cell viability (%)
(Untreated = 100%)
Cetuximab-Saporin (pM)
Cetuximab-saporin
Cetuximab-saporin + 75 nM cetuximab
Cetuximab-saporin + 75 nM
Cetuximab-(Cys-L-SO1861)3,7

Figure 4

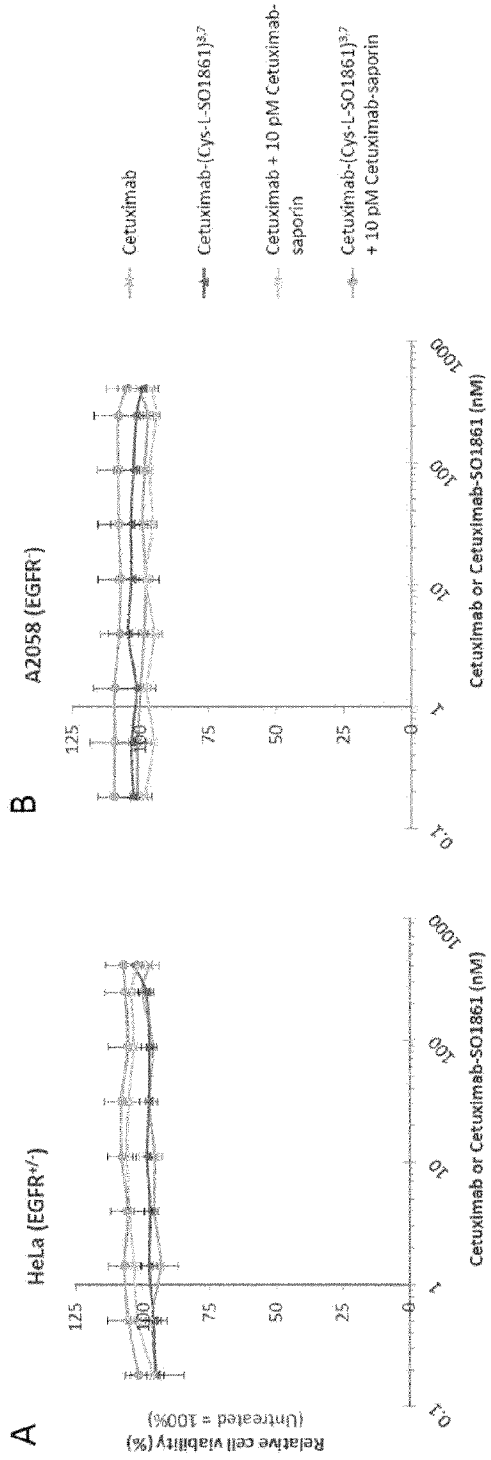

A
HeLa (EGFR+/-)
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0.1
1
10
100
1000
Cetuximab or Cetuximab-SO1861 (nM)
B
A2058 (EGFR-)
Cetuximab
Cetuximab-(Cys-L-SO1861)3,7
Cetuximab + 10 pM Cetuximab-saporin
Cetuximab-(Cys-L-SO1861)3,7 + 10 pM Cetuximab-saporin

Figure 4

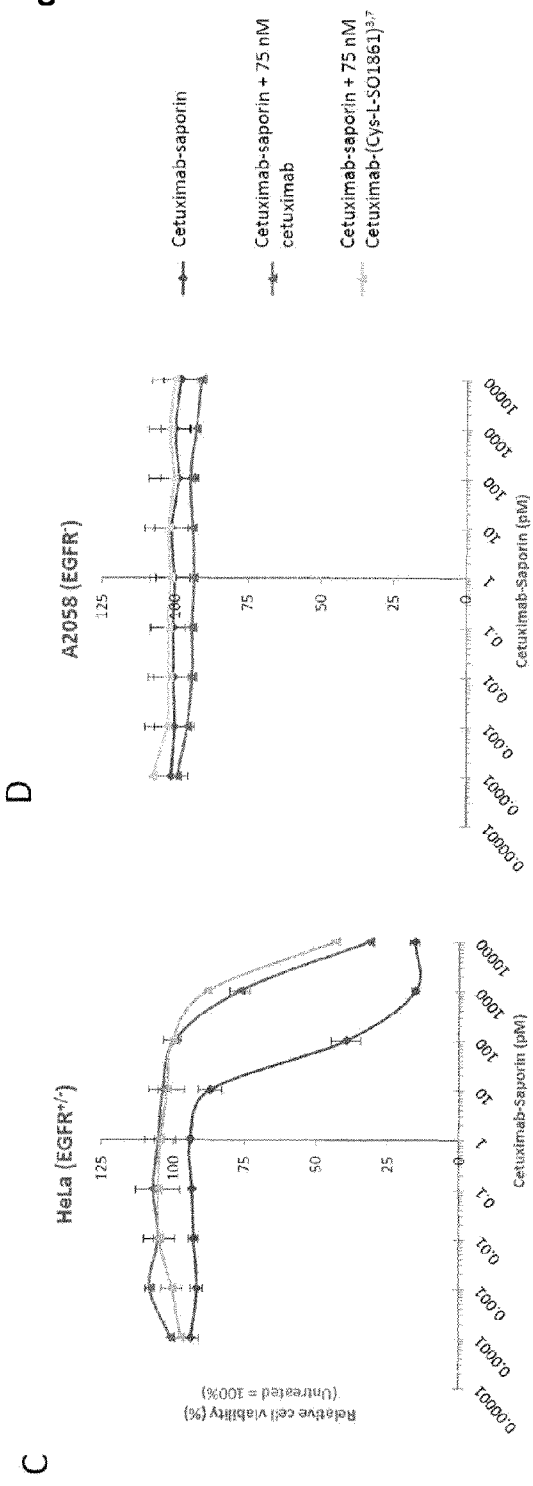

C
HeLa (EGFR+/-)
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0,00001
0,0001
0,001
0,01
0,1
1
10
100
1000
10000
Cetuximab-Saporin (pM)
D
A2058 (EGFR-)
Cetuximab-saporin
Cetuximab-saporin + 75 nM cetuximab
Cetuximab-saporin + 75 nM Cetuximab-(Cys-LSO1861)3,7

Figure 5

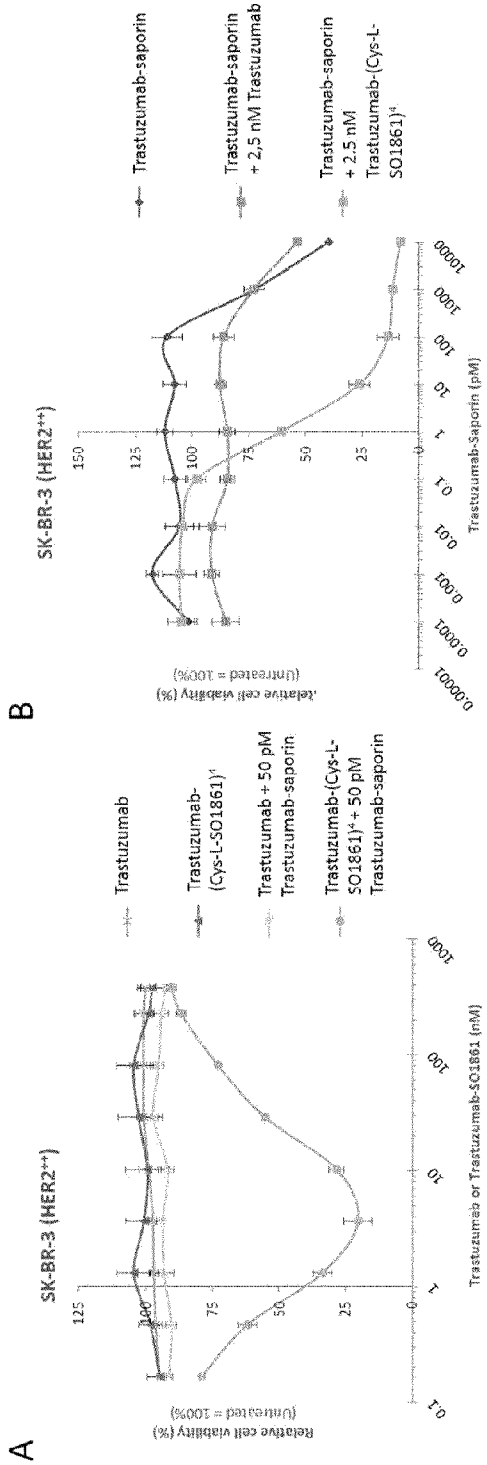

A
SK-BR-3 (HER2+++)
Relative cell viability (%)
(Untreated = 100%)
Trastuzumab or Trastuzumab-SO1861 (nM)
Trastuzumab
Trastuzumab-(Cys-L-SO1861)1
Trastuzumab + 50 pM Trastuzumab-saporin
Trastuzumab-(Cys-L-SO1861)4 + 50 pM Trastuzumab-saporin
B
SK-BR-3 (HER2+++)
Relative cell viability (%)
(Untreated = 100%)
Trastuzumab-Saporin (pM)
Trastuzumab-saporin
Trastuzumab-saporin + 2,5 nM Trastuzumab
Trastuzumab-saporin + 2.5 nM Trastuzumab-(Cys-L-SO1861)4

Figure 6

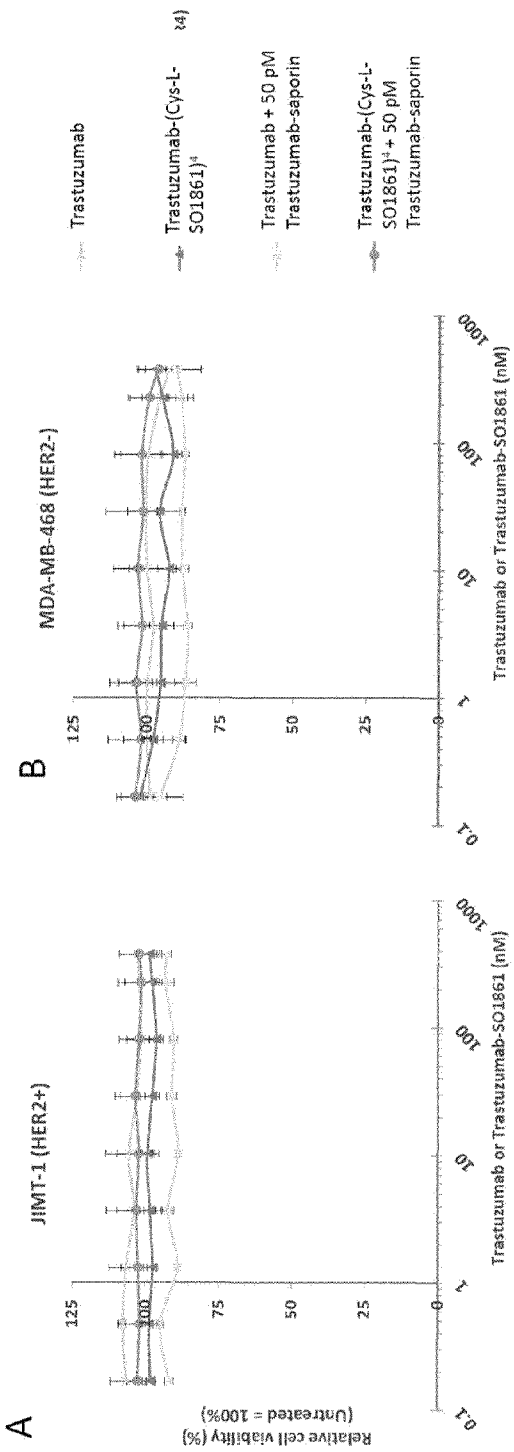

A
JIMT-1 (HER2+)
B
MDA-MB-468 (HER2-)
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0.1
1
10
100
1000
Trastuzumab or Trastuzumab-SO1861 (nM)
Trastuzumab
Trastuzumab-(Cys-L-SO1861)4
Trastuzumab + 50 pM Trastuzumab-saporin
Trastuzumab-(Cys-L-SO1861)4 + 50 pM Trastuzumab-saporin

Figure 6

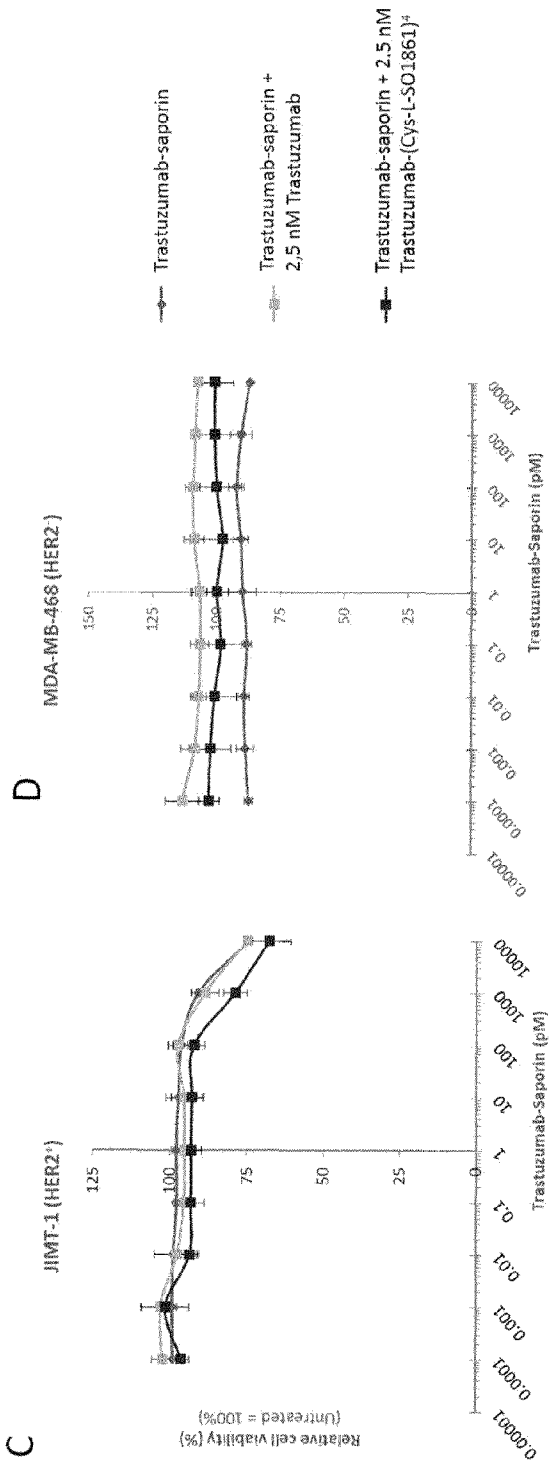

C
JIMT-1 (HER2+)
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0,00001
0,0001
0,001
0,01
0,1
1
10
100
1000
10000
Trastuzumab-Saporin (pM)
D
MDA-MB-468 (HER2+)
150
125
100
75
50
25
0
0,00001
0,0001
0,001
0,01
0,1
1
10
100
1000
10000
Trastuzumab-Saporin (pM)
Trastuzumab-saporin
Trastuzumab-saporin + 2,5 nM Trastuzumab
Trastuzumab-saporin + 2.5 nM Trastuzumab-(Cys-L-SO1861)4

Figure 8

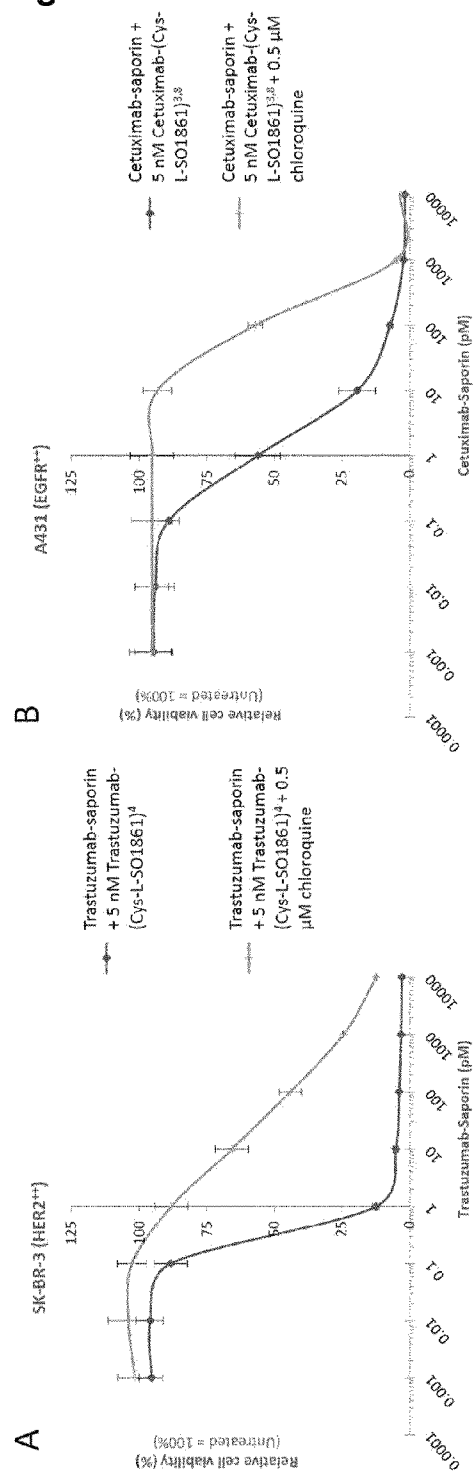

A
SK-BR-3 (HER2++)
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0.0001
0.001
0.01
0.1
1
10
100
1000
10000
Trastuzumab-Saporin (pM)
Trastuzumab-saporin + 5 nM Trastuzumab-(Cys-L-SO1861)4
Trastuzumab-saporin + 5 nM Trastuzumab-(Cys-L-SO1861)4 + 0.5 µM chloroquine
B
A431 (EGFR++)
Relative cell viability (%)
(Untreated = 100%)
Cetuximab-Saporin (pM)
Cetuximab-saporin + 5 nM Cetuximab-(Cys-L-SO1861)3,8
Cetuximab-saporin + 5 nM Cetuximab-(Cys-L-SO1861)3,8 + 0.5 µM chloroquine

Figure 9

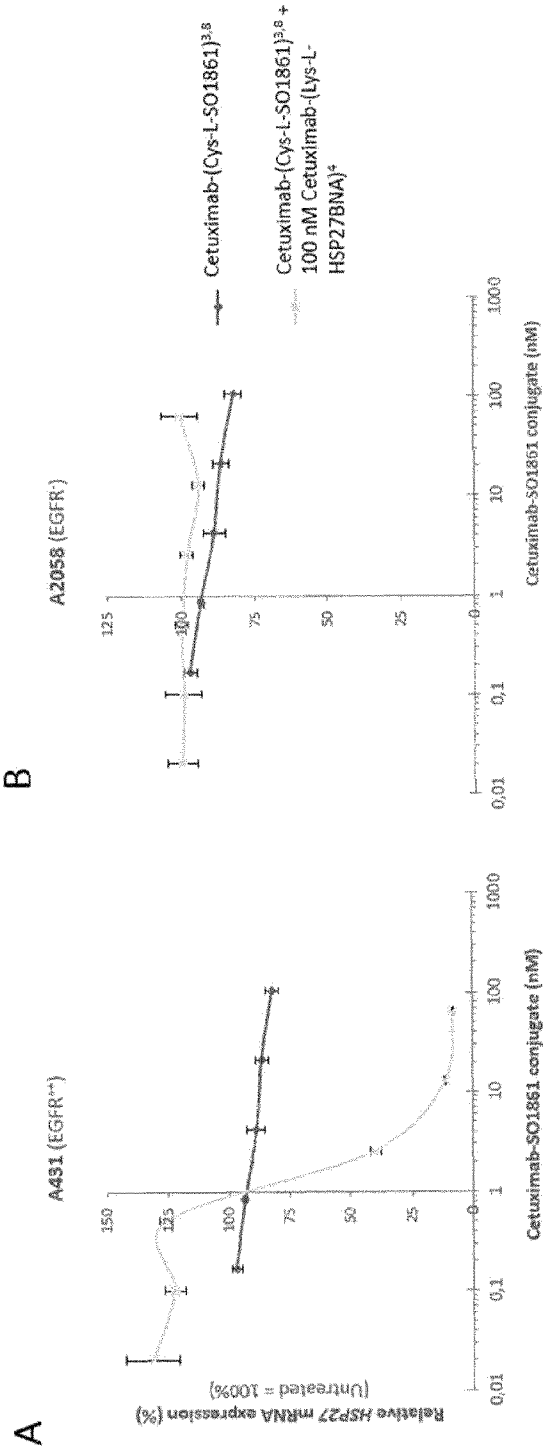

A
A431 (EGFR++)
Relative HSP27 mRNA expression (%)
(Untreated = 100%)
Cetuximab-SO1861 conjugate (nM)
B
A2058 (EGFR-)
Cetuximab-(Cys-L-SO1861)3.8
Cetuximab-(Cys-L-SO1861)3.8 + 100 nM Cetuximab-(Lys-L-HSP27BNA)4
Cetuximab-SO1861 conjugate (nM)

Figure 9

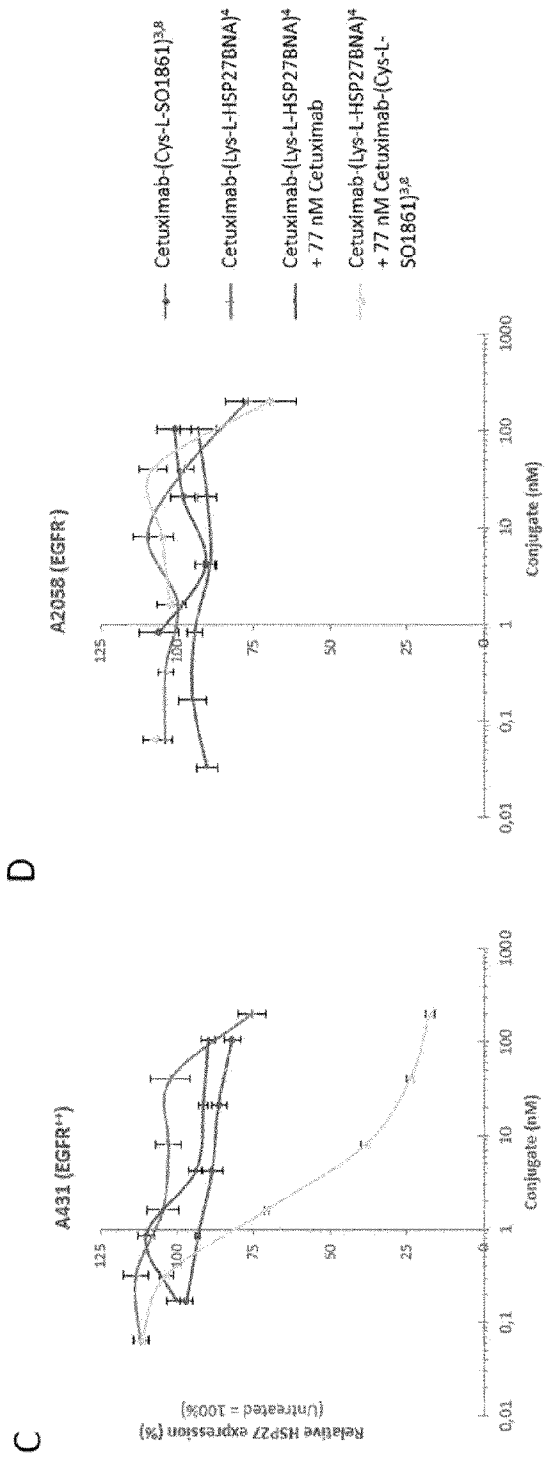

C
A431 (EGFR++)
Relative HSP27 expression (%)
(Untreated = 100%)
Conjugate (nM)
D
A2058 (EGFR+)
Conjugate (nM)
Cetuximab-(Cys-L-SO1861)3,8
Cetuximab-(Lys-L-HSP27BNA)4
Cetuximab-(Lys-L-HSP27BNA)4 + 77 nM Cetuximab
Cetuximab-(Lys-L-HSP27BNA)4 + 77 nM Cetuximab-(Cys-L-SO1861)3,8

Figure 10

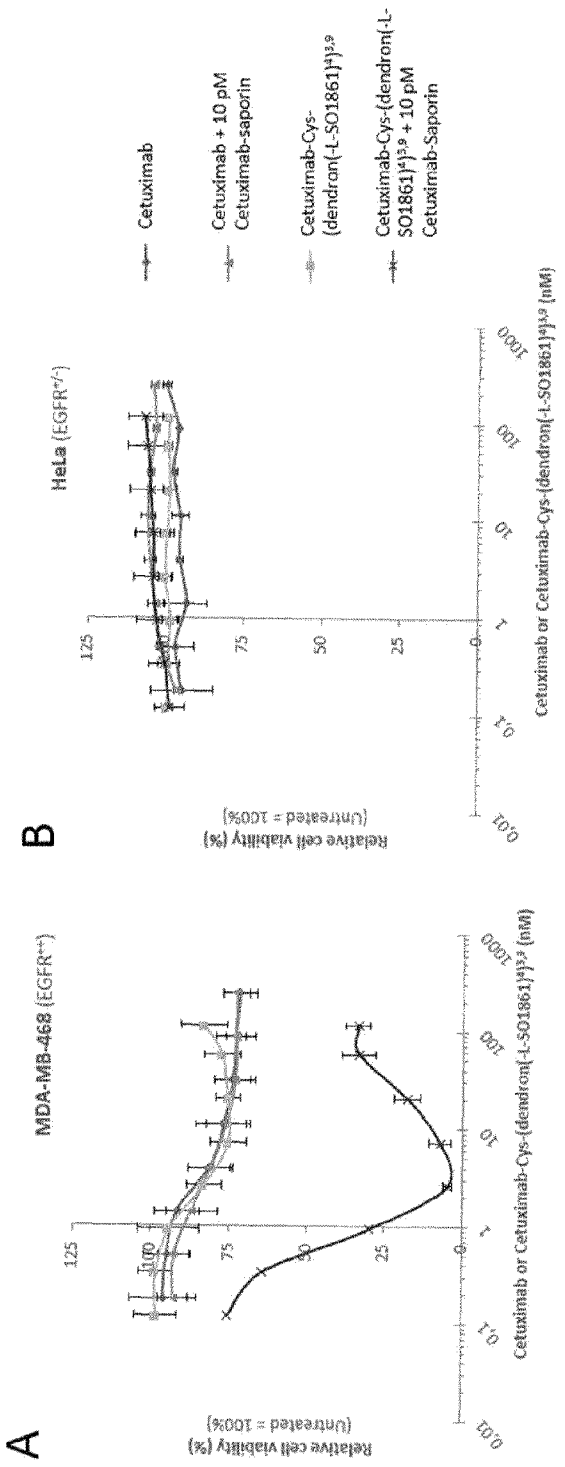

A
MDA-MB-468 (EGFR+++)
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0.01
0.1
1
10
100
1000
Cetuximab or Cetuximab-Cys-(dendron[-L-SO1861]4)3,9 (nM)
B
HeLa (EGFR+/-)
Cetuximab
Cetuximab + 10 pM Cetuximab-saporin
Cetuximab-Cys-(dendron[-L-SO1861]4)3,9
Cetuximab-Cys-(dendron[-L-SO1861]4)3,9 + 10 pM Cetuximab-Saporin

Figure 10

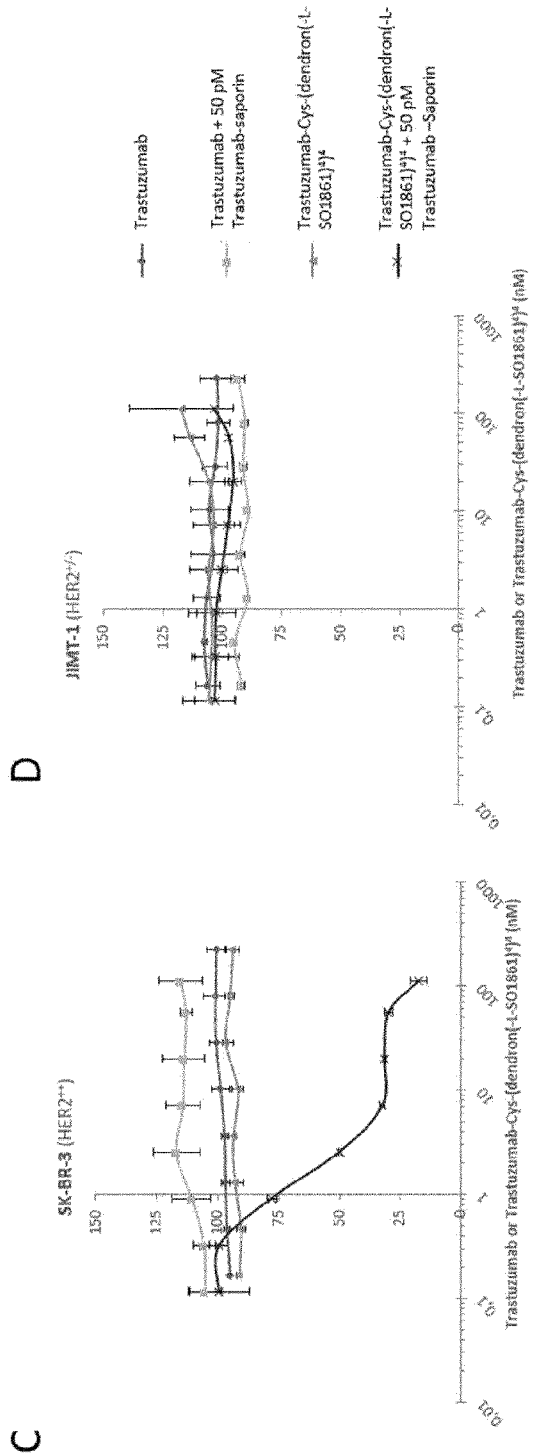

D
JIMT-1 (HER2+/-)
Trastuzumab
Trastuzumab + 50 pM Trastuzumab-saporin
Trastuzumab-Cys-(dendron[-L-SO1861])4
Trastuzumab-Cys-(dendron[-L-SO1861])4 + 50 pM Trastuzumab –Saporin
Trastuzumab or Trastuzumab-Cys-(dendron[-L-SO1861])4 (nM)
C
SK-BR-3 (HER2++)
Trastuzumab or Trastuzumab-Cys-(dendron[-L-SO1861])4 (nM)

Figure 11

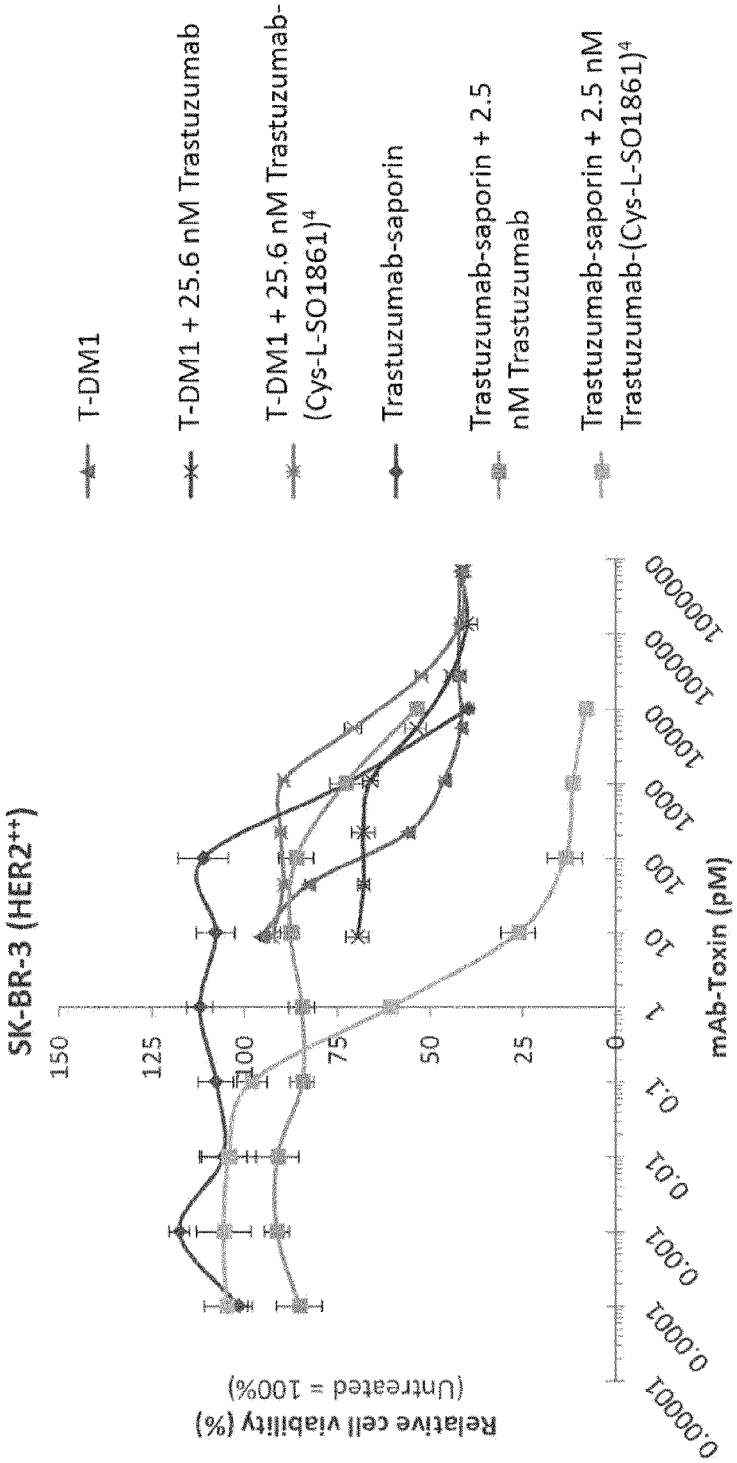

T-DM1
T-DM1 + 25.6 nM Trastuzumab
T-DM1 + 25.6 nM Trastuzumab-(Cys-L-SO1861)4
Trastuzumab-saporin
Trastuzumab-saporin + 2.5 nM Trastuzumab
Trastuzumab-saporin + 2.5 nM Trastuzumab-(Cys-L-SO1861)4
SK-BR-3 (HER2++)
Relative cell viability (%)
(Untreated = 100%)
150
125
100
75
50
25
0
mAb-Toxin (pM)
0.00001
0.0001
0.001
0.01
0.1
1
10
100
1000
10000
100000
1000000

Figure 12

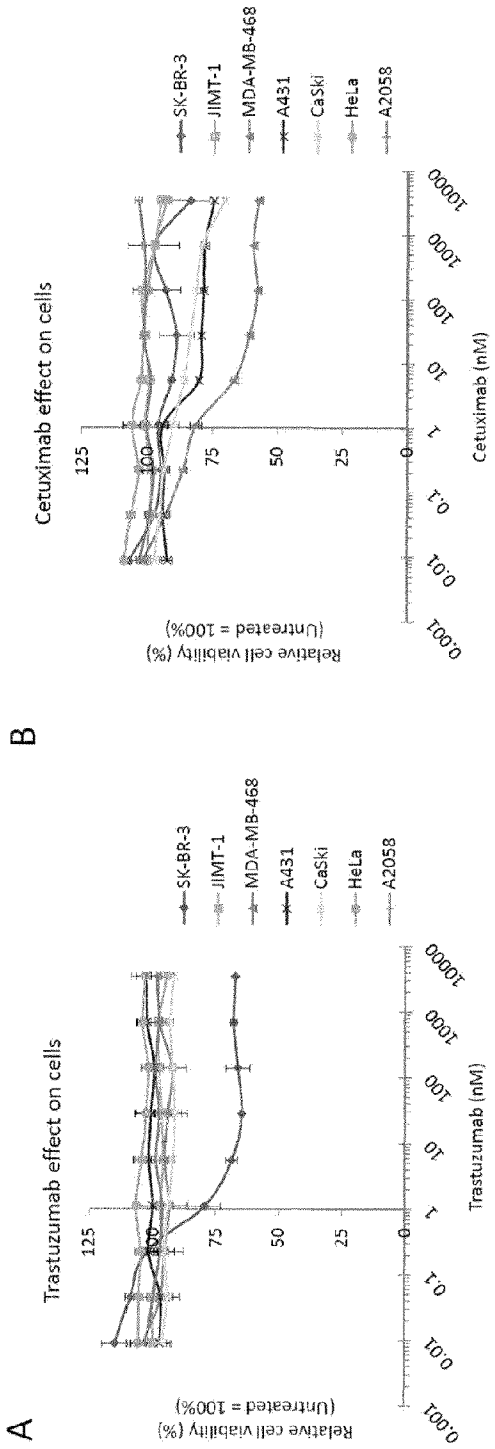

A
Trastuzumab effect on cells
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0.001
0.01
0.1
1
10
100
1000
10000
Trastuzumab (nM)
SK-BR-3
JIMT-1
MDA-MB-468
A431
CaSki
HeLa
A2058
B
Cetuximab effect on cells
Relative cell viability (%)
(Untreated = 100%)
Cetuximab (nM)

Figure 12

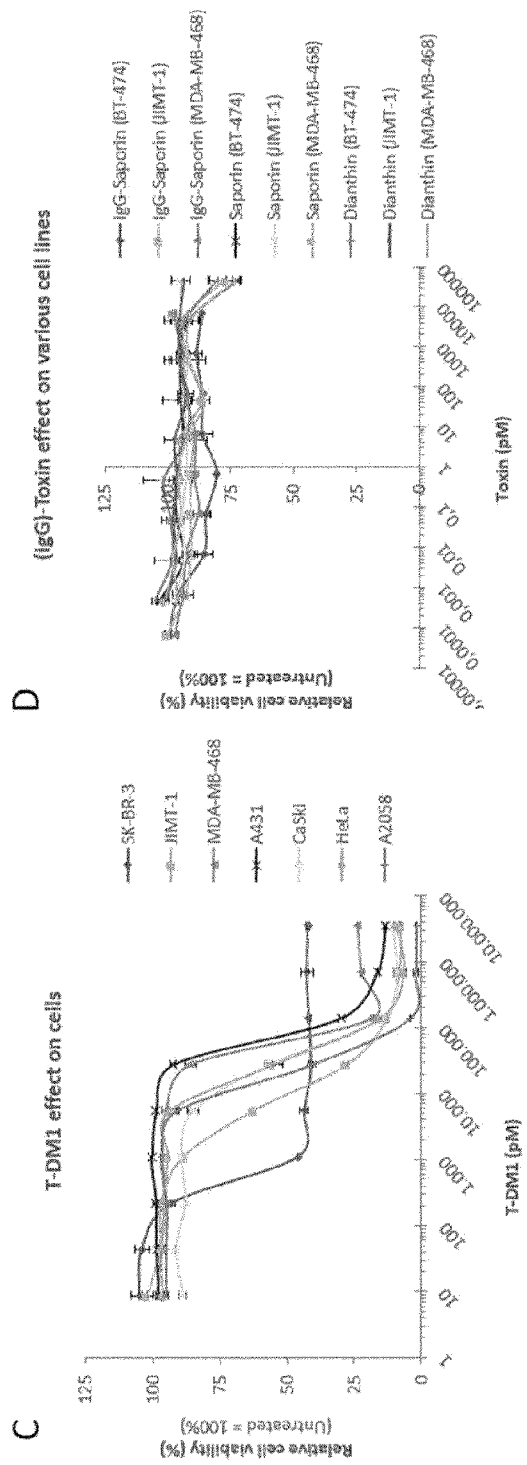

C
T-DM1 effect on cells
Relative cell viability (%)
(Untreated = 100%)
T-DM1 (pM)
SK-BR-3
JIMT-1
MDA-MB-468
A431
CaSki
HeLa
A2058
D
(IgG)-Toxin effect on various cell lines
Relative cell viability (%)
(Untreated = 100%)
Toxin (pM)
IgG-Saporin (BT-474)
IgG-Saporin (JIMT-1)
IgG-Saporin (MDA-MB-468)
Saporin (BT-474)
Saporin (JIMT-1)
Saporin (MDA-MB-468)
Dianthin (BT-474)
Dianthin (JIMT-1)
Dianthin (MDA-MB-468)

Figure 13

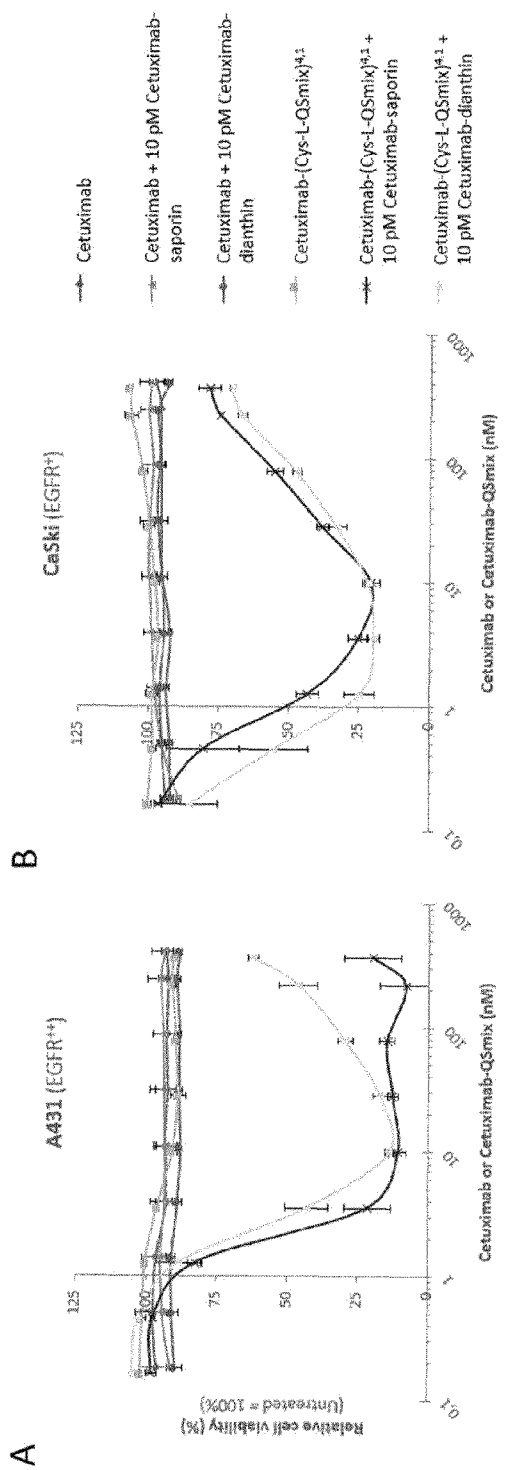

A
A431 (EGFR++)
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0.1
1
10
100
1000
Cetuximab or Cetuximab-QSmix (nM)
B
CaSki (EGFR+)
Cetuximab
Cetuximab + 10 pM Cetuximab-saporin
Cetuximab + 10 pM Cetuximab-dianthin
Cetuximab-(Cys-L-QSmix)4:1
Cetuximab-(Cys-L-QSmix)4:1 + 10 pM Cetuximab-saporin
Cetuximab-(Cys-L-QSmix)4:1 + 10 pM Cetuximab-dianthin

Figure 13

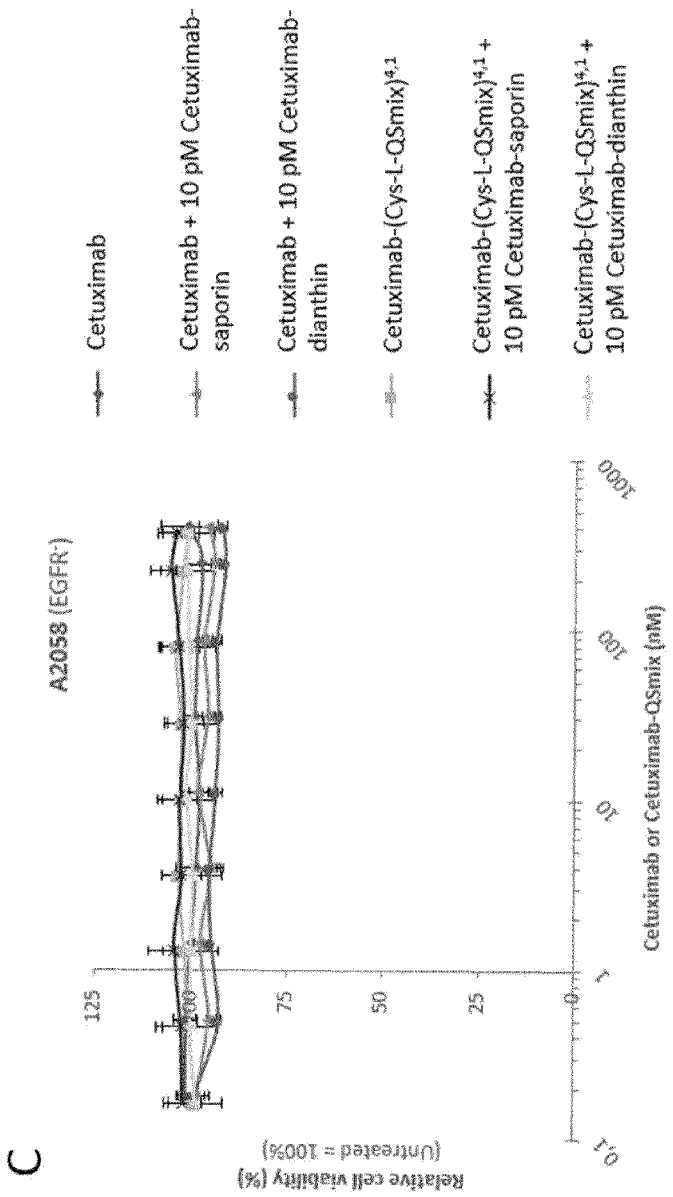

C
A2058 (EGFR+)
Cetuximab
Cetuximab + 10 pM Cetuximab-saporin
Cetuximab + 10 pM Cetuximab-dianthin
Cetuximab-(Cys-L-QSmix)4,1
Cetuximab-(Cys-L-QSmix)4,1 + 10 pM Cetuximab-saporin
Cetuximab-(Cys-L-QSmix)4,1 + 10 pM Cetuximab-dianthin
Relative cell viability (%)
(Untreated = 100%)
125
100
75
50
25
0
0,1
1
10
100
1000
Cetuximab or Cetuximab-QSmix (nM)

Figure 14

L1-(S)n + L1-E concept: mAb1-SO1861 + mAb1-toxin

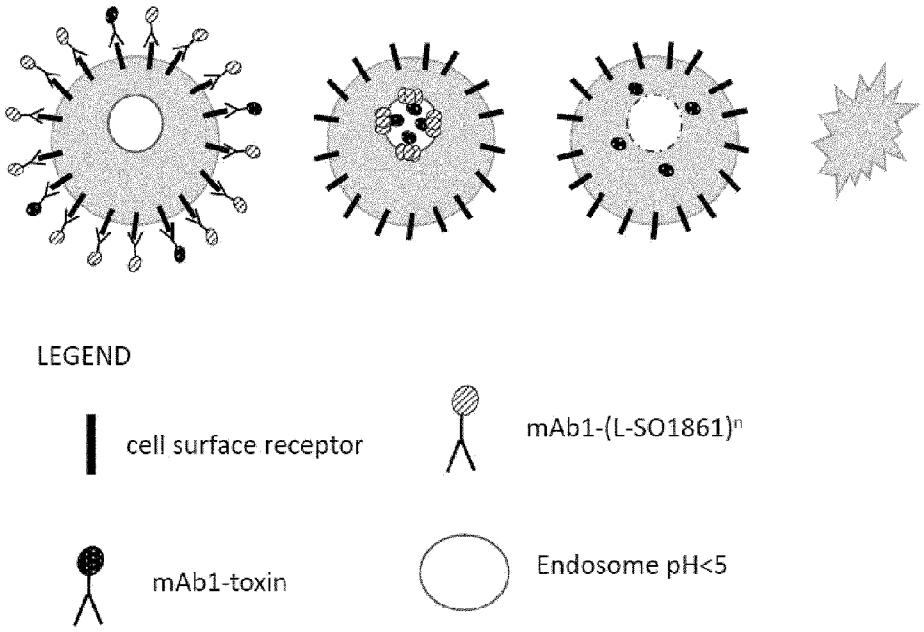


Figure 15

L1-(S)n + L1-E concept: mAb1-SO1861 + mAb1-BNA oligo

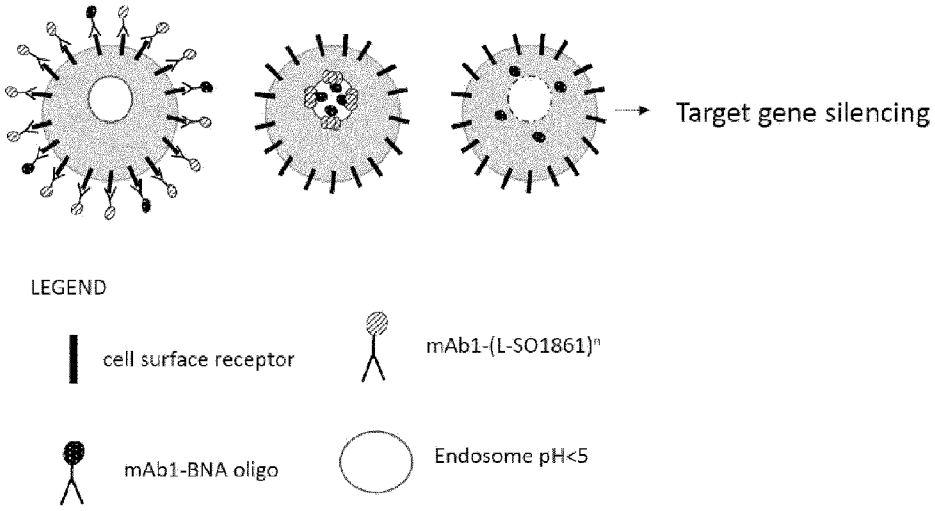


Figure 16

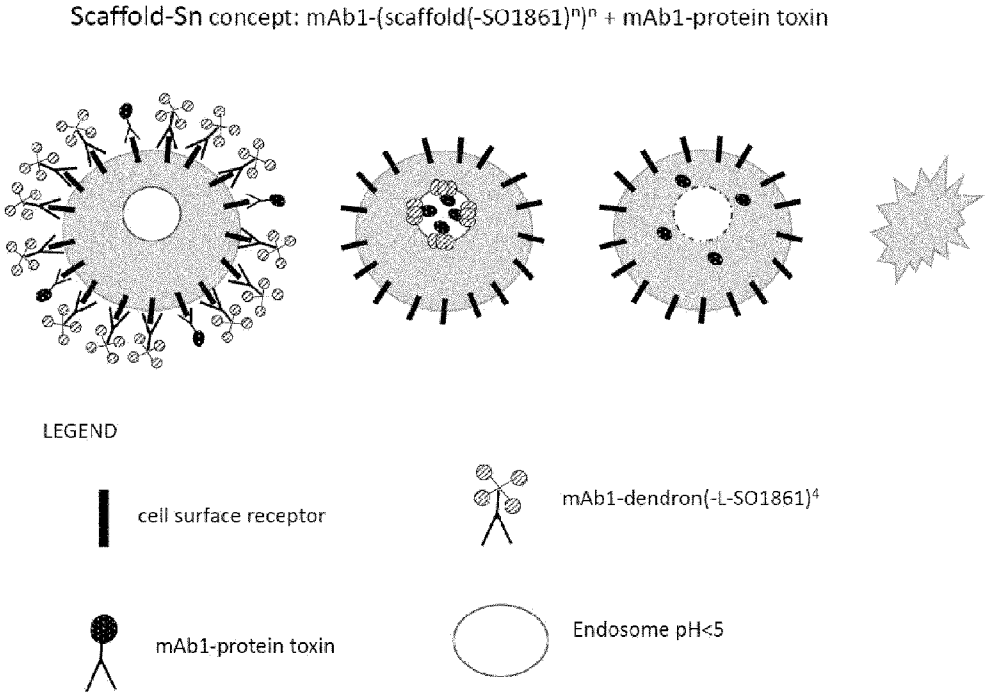


Figure 17

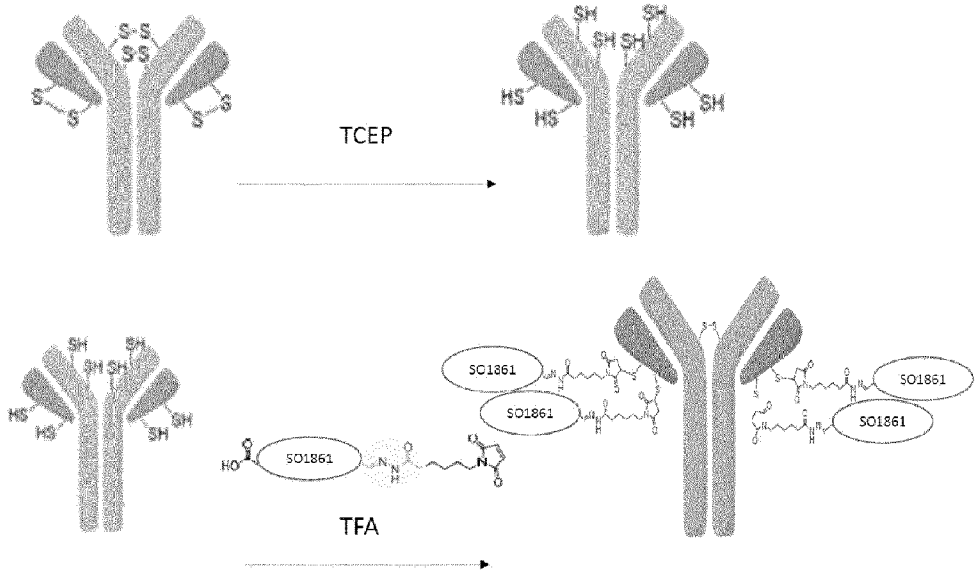


Figure 18

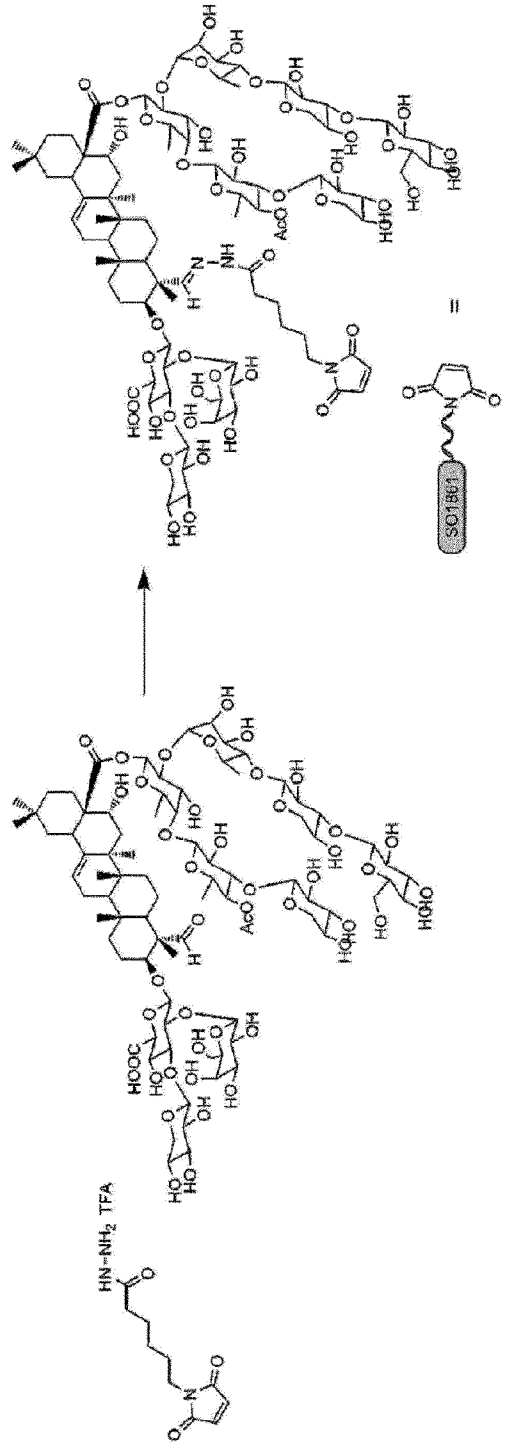


Figure 19

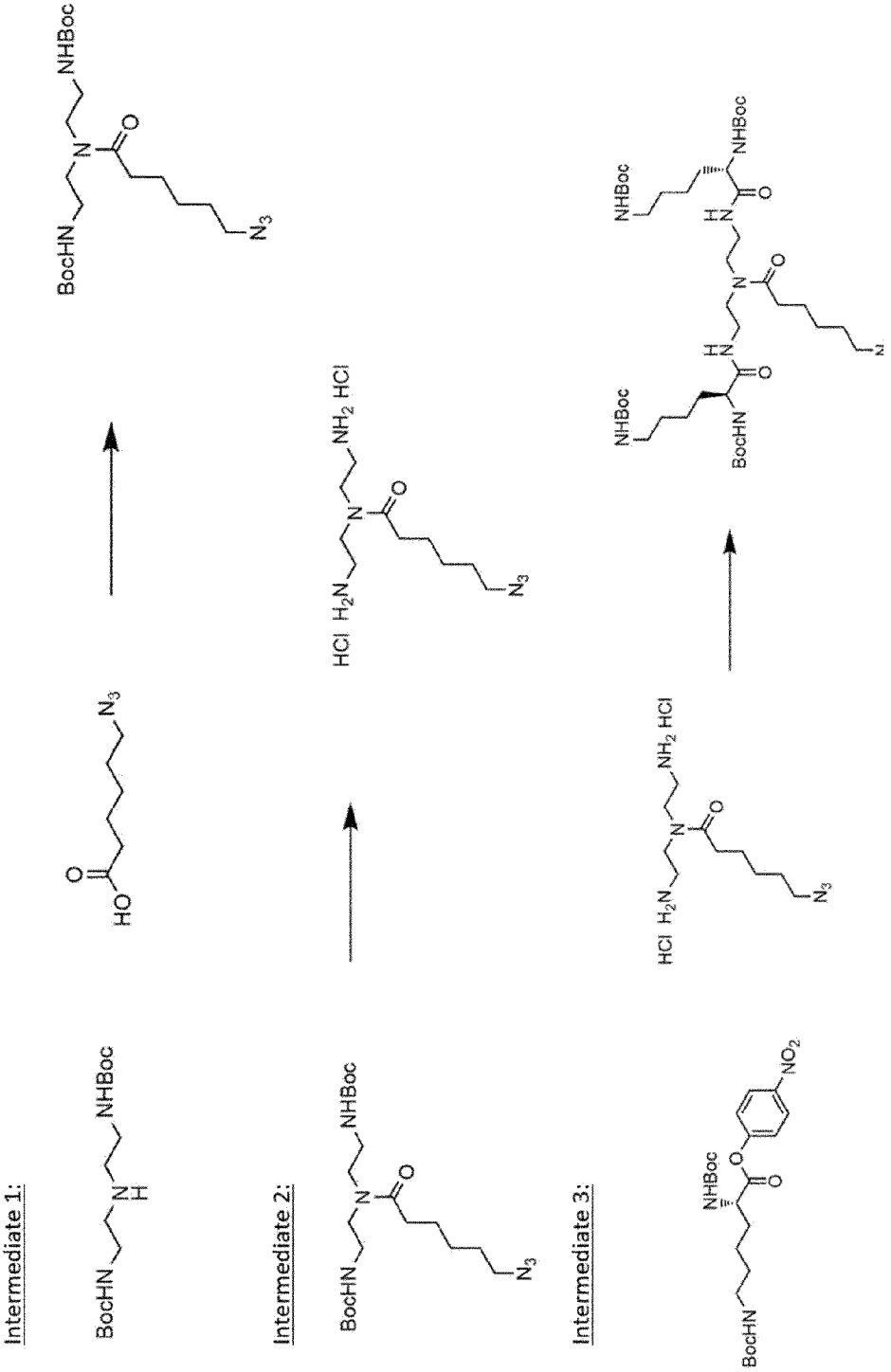

Intermediate 1:
Intermediate 2:
Intermediate 3:
NHBoc
BocHN
HO
N_3
NH_2 HCl
HCl H_2N
NO_2

Figure 19

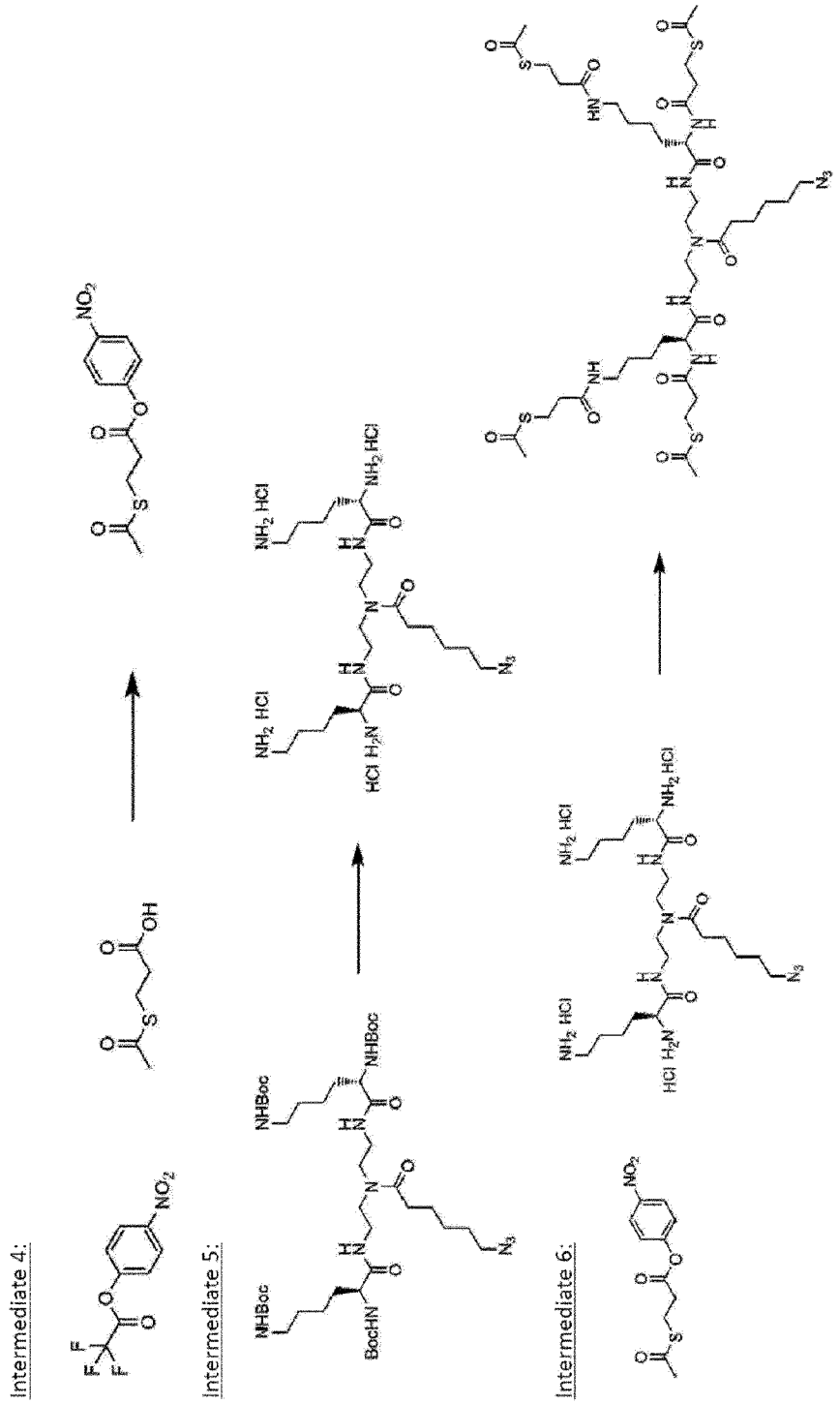


Figure 19

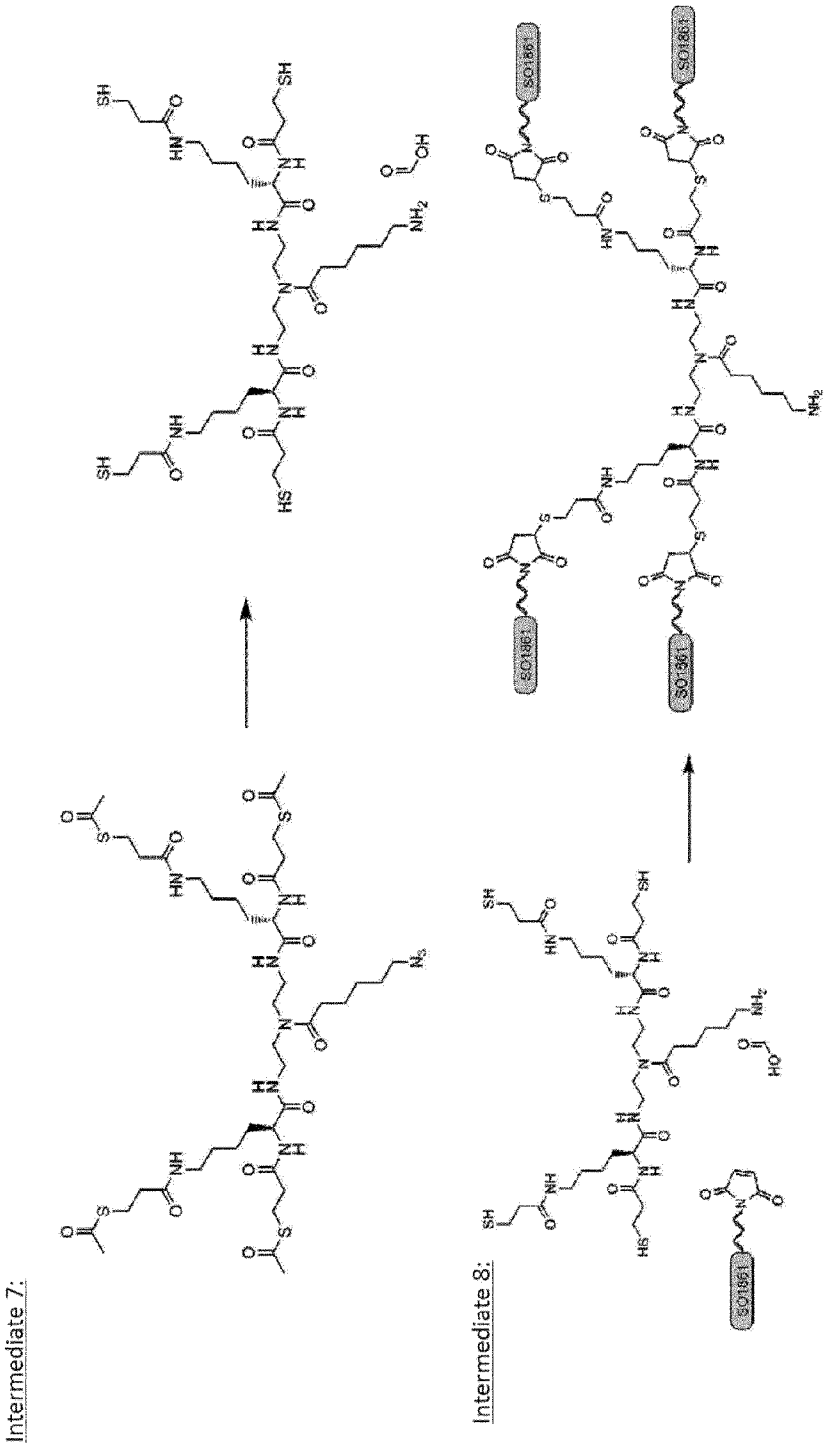

Intermediate 7:
Intermediate 8:
SO1861
SO1861
SO1861
SO1861
SO1861

Figure 19

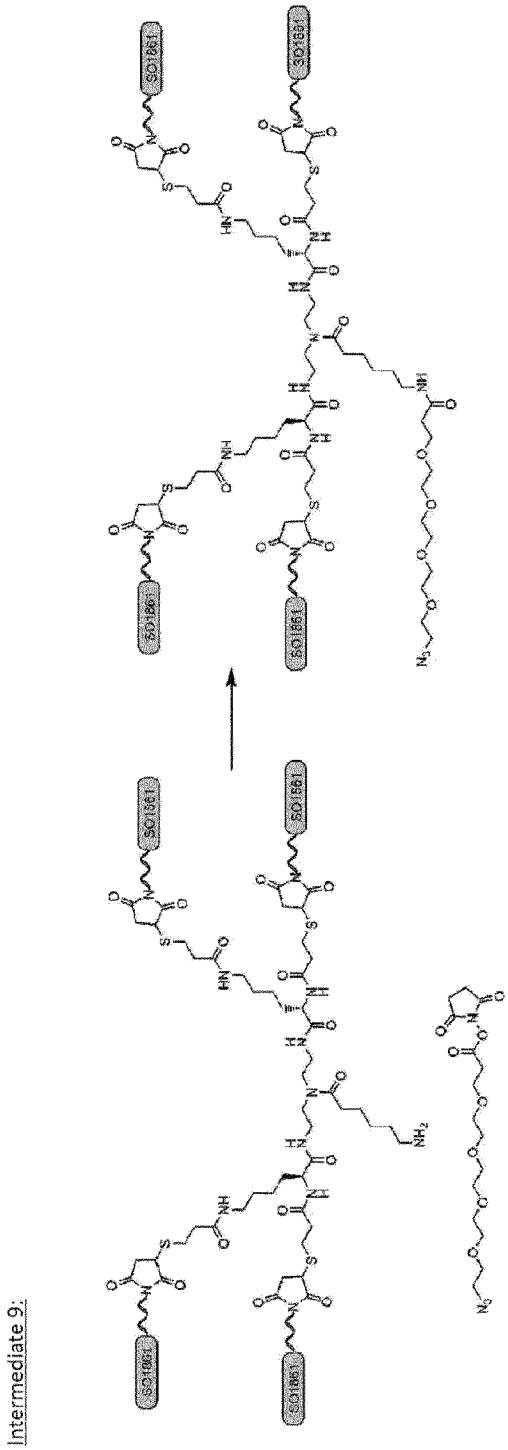

SO1861
SO1861
SO1861
SO1861
SO1861
SO1861
SO1861
SO1861
Intermediate 9:

Figure 19

Intermediate 10:

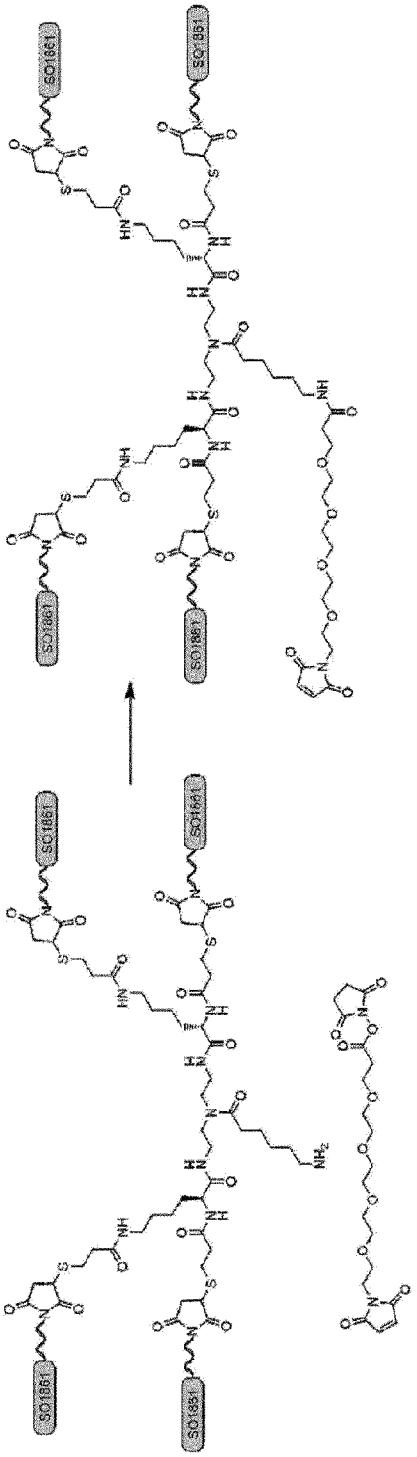


Figure 19

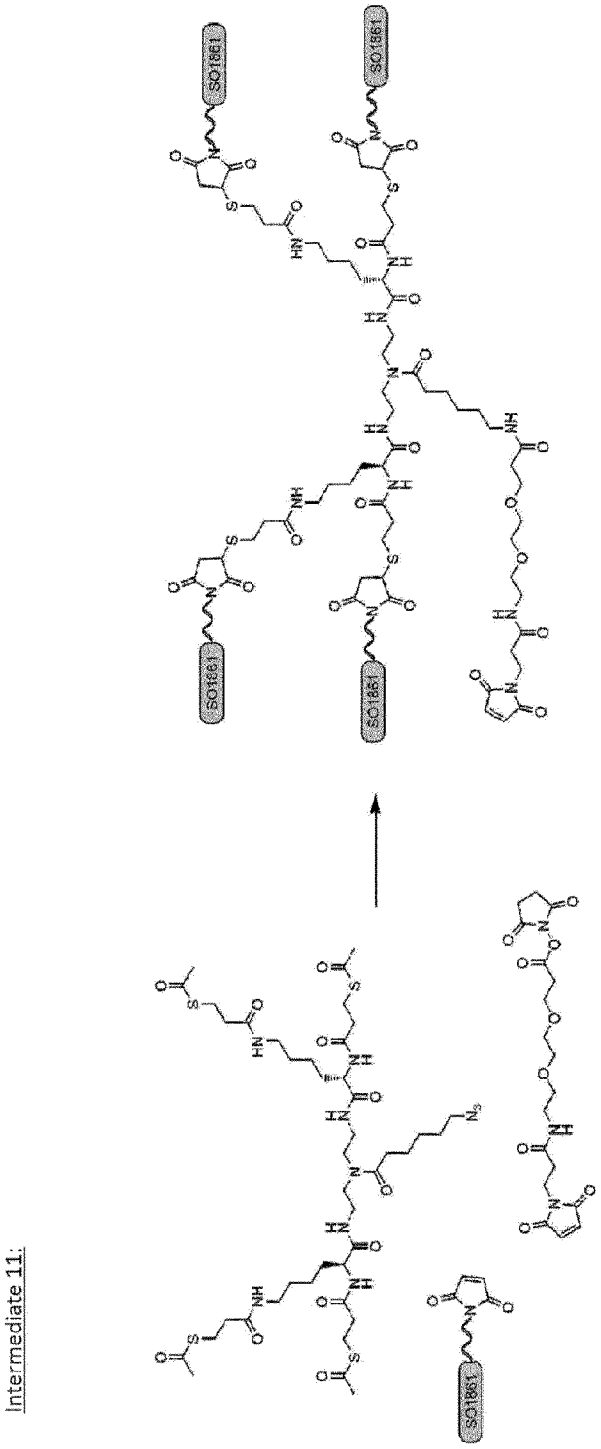


Figure 20

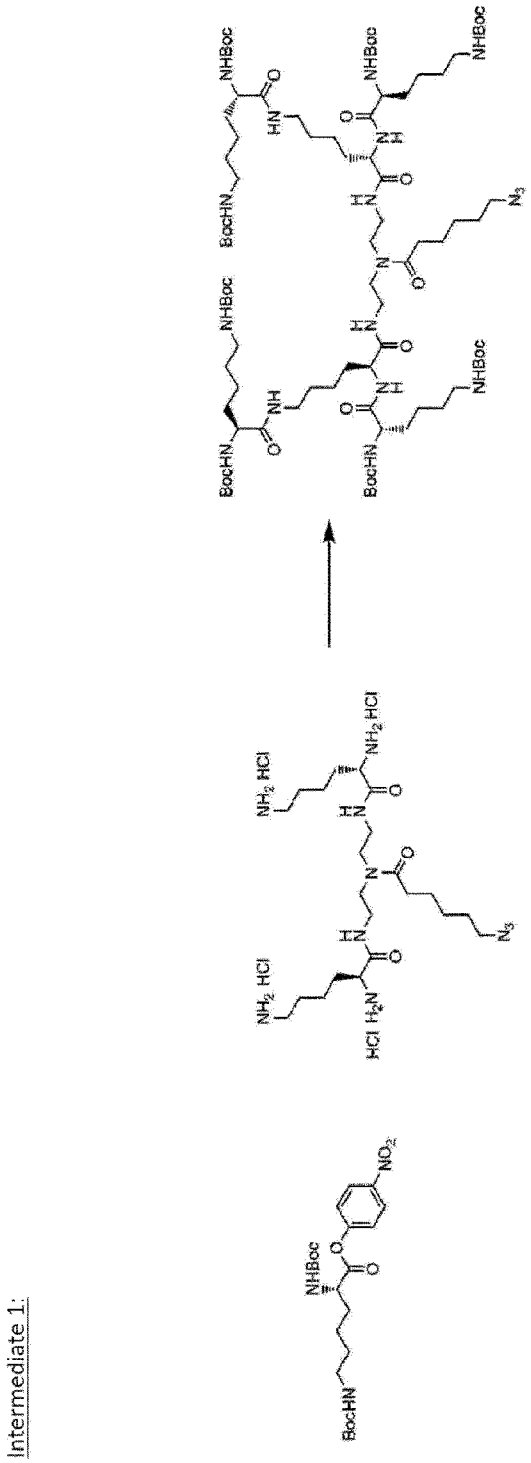


Figure 20

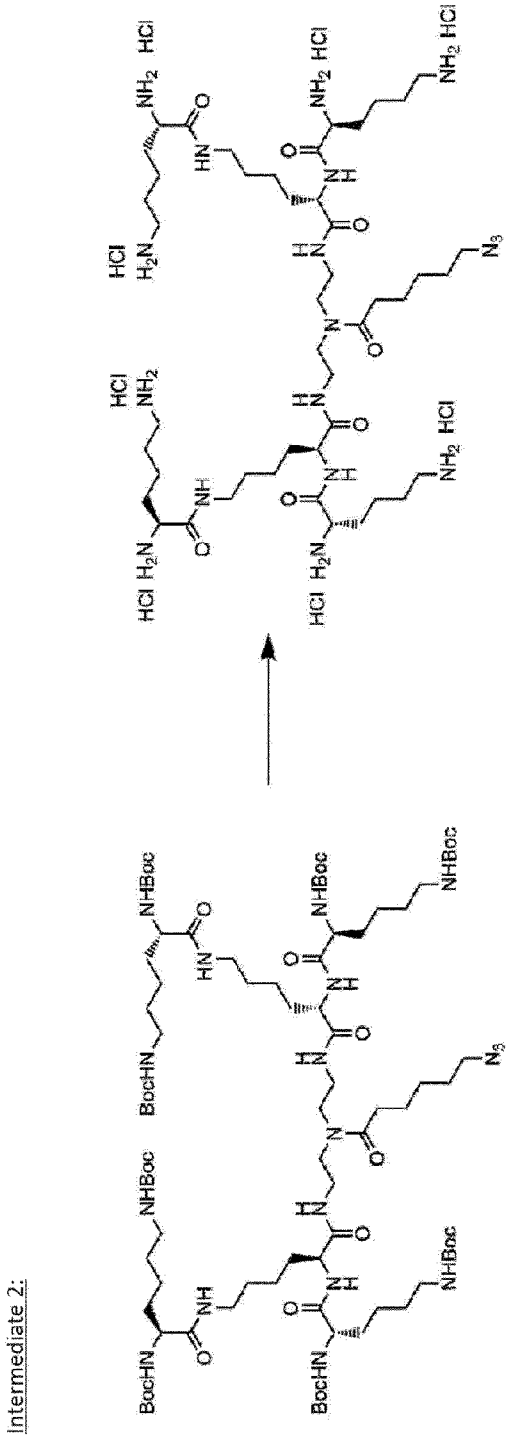


Figure 20

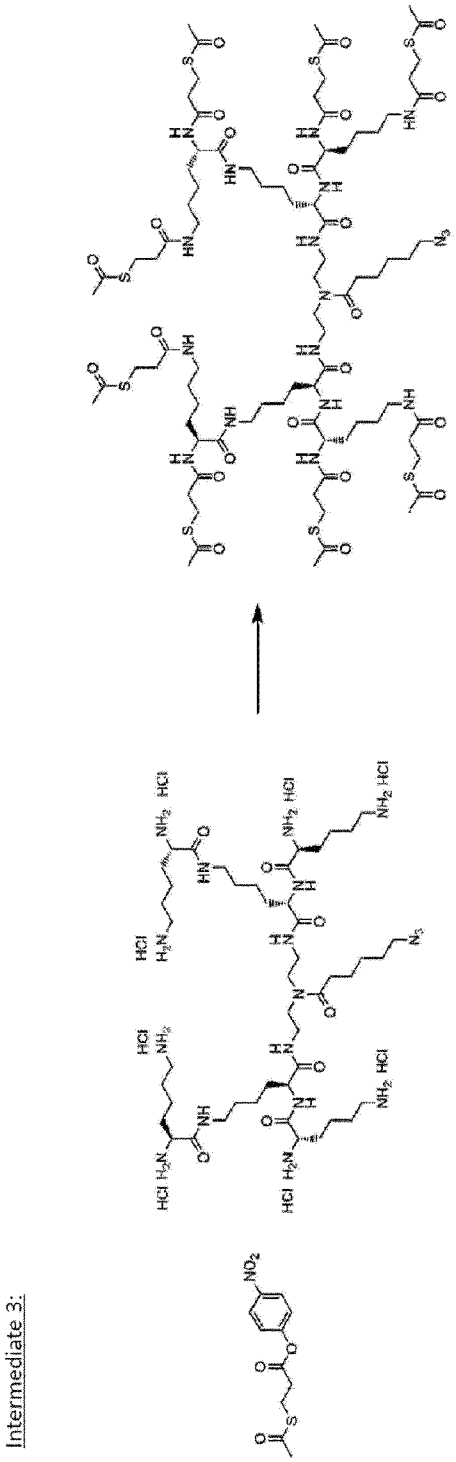
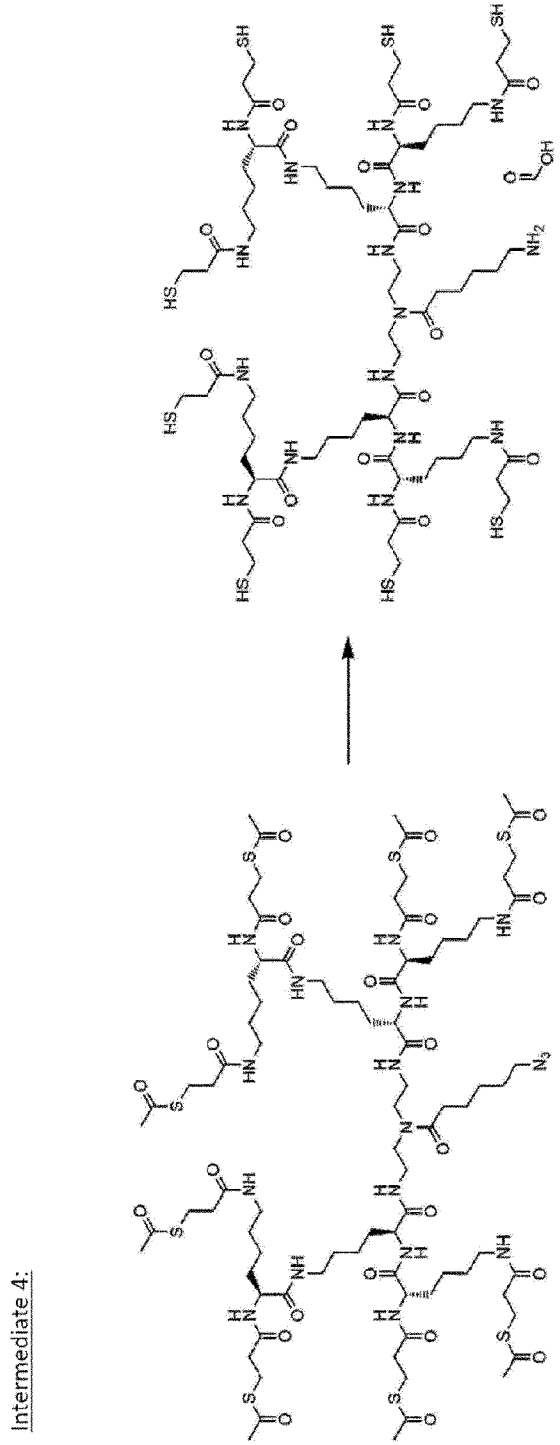


Figure 20



Intermediate 4:

Figure 20

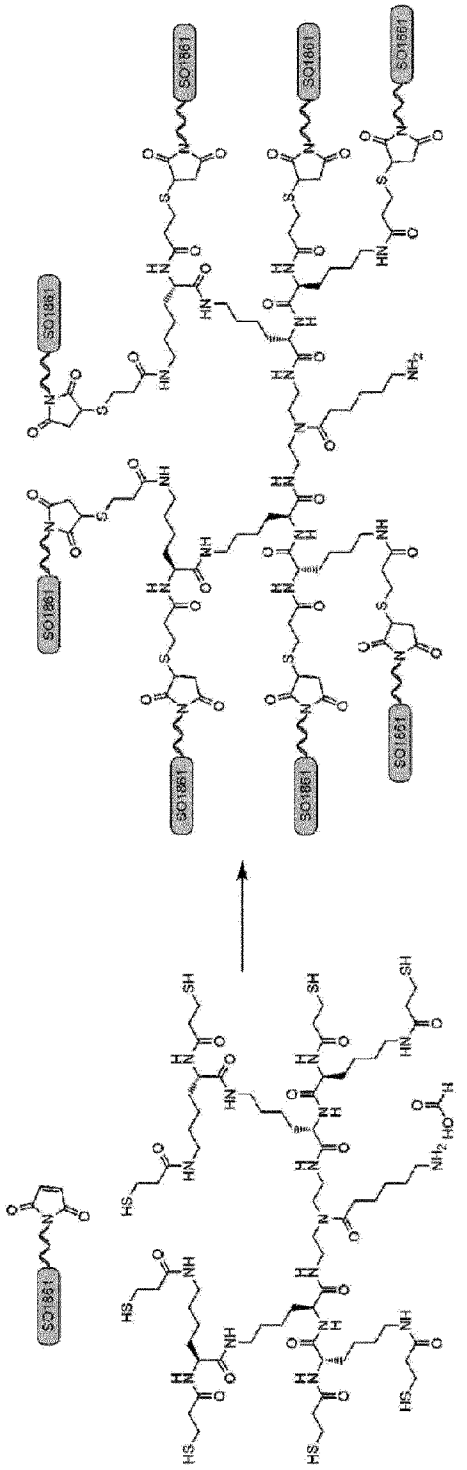


Figure 21

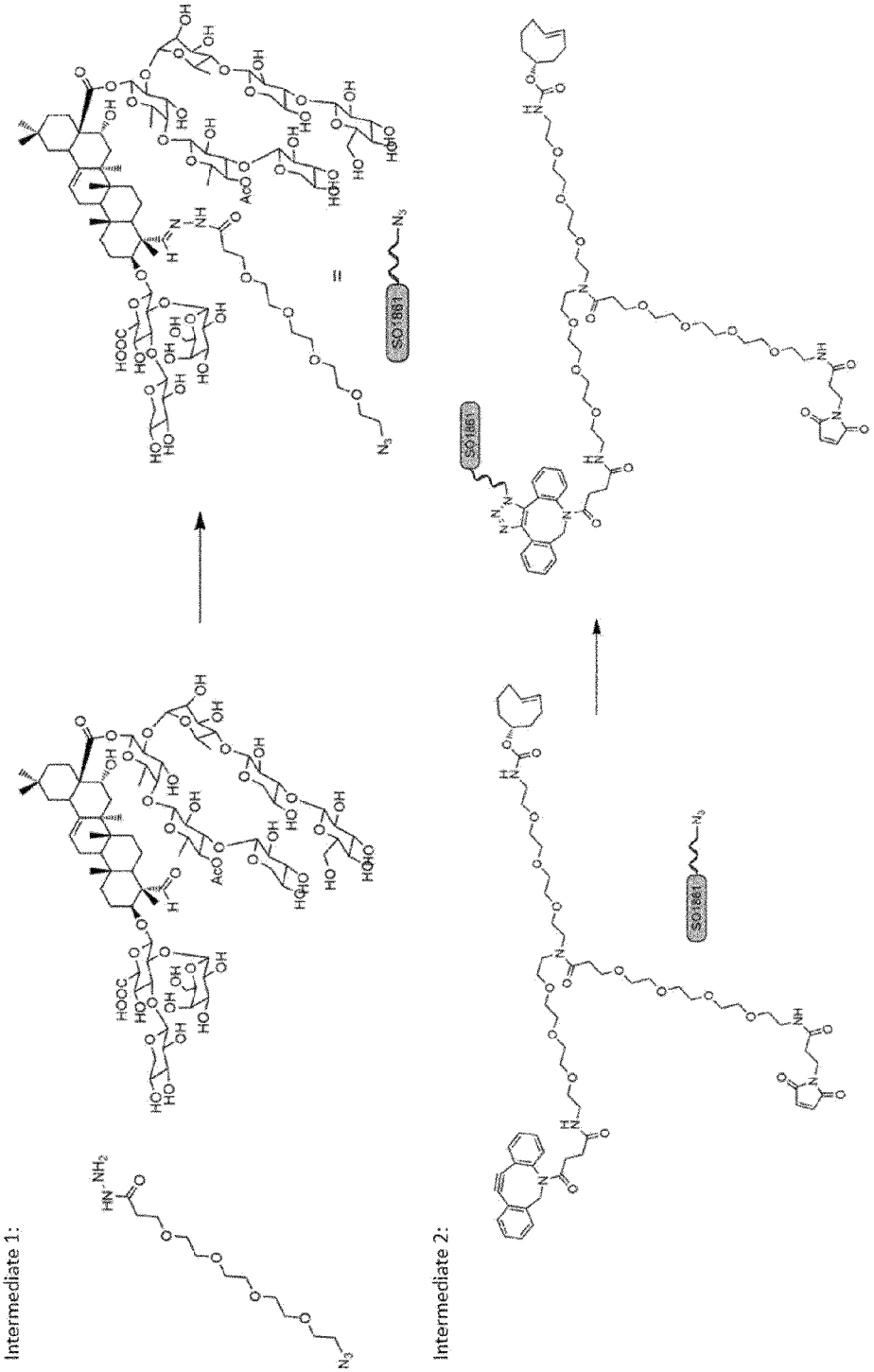


Figure 21

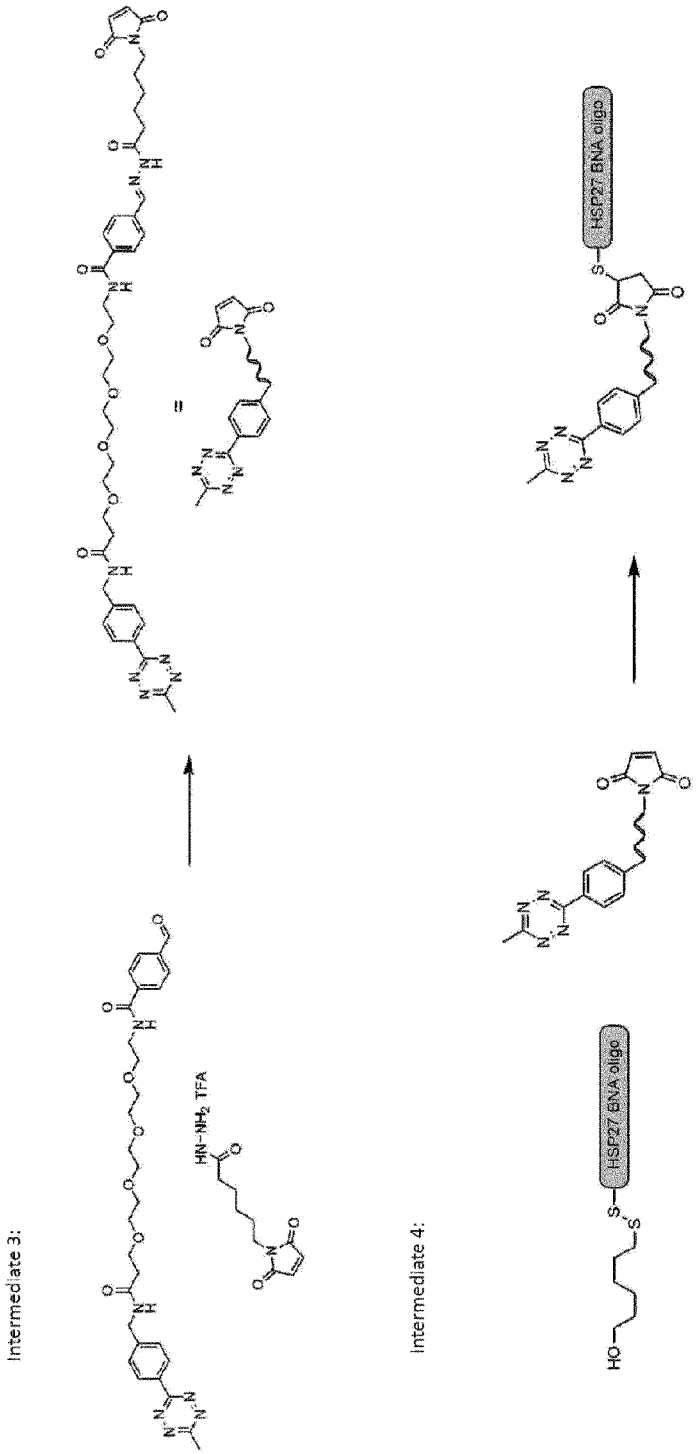
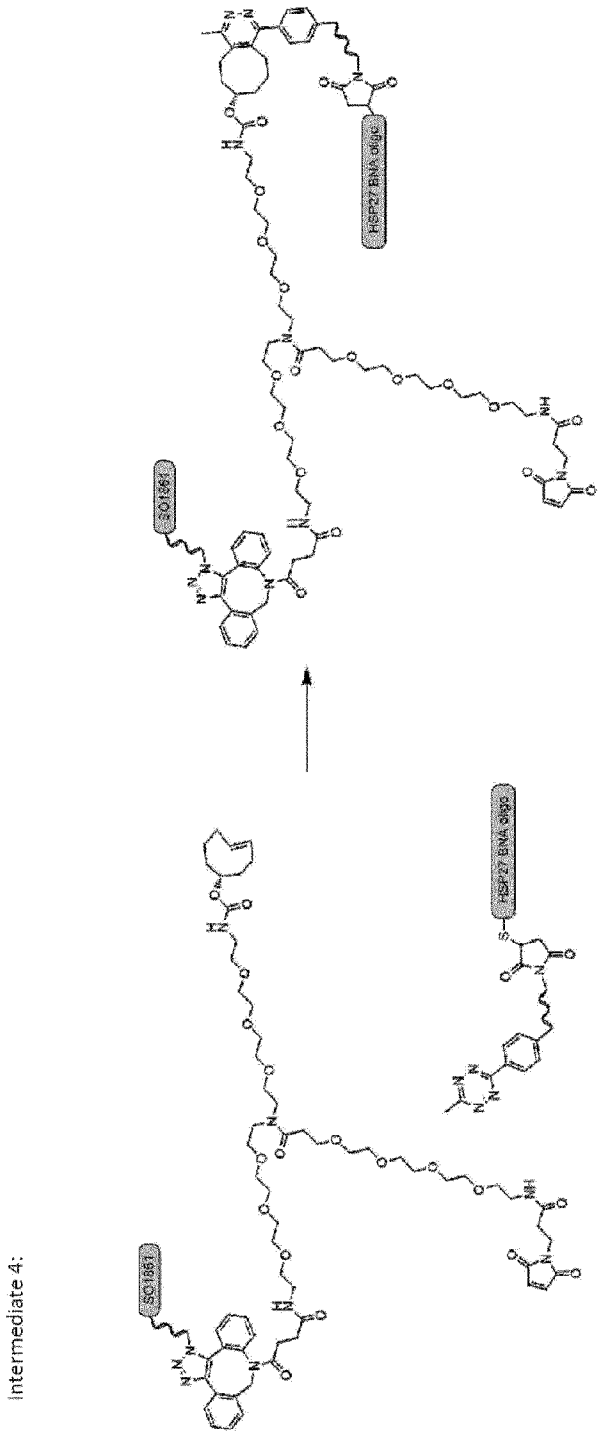

Intermediate 3:
HN-NH2, TFA
Intermediate 4:
HSP27 BNA oligo
HSP27 BNA oligo
HO

Figure 21



Intermediate 4:

Figure 22

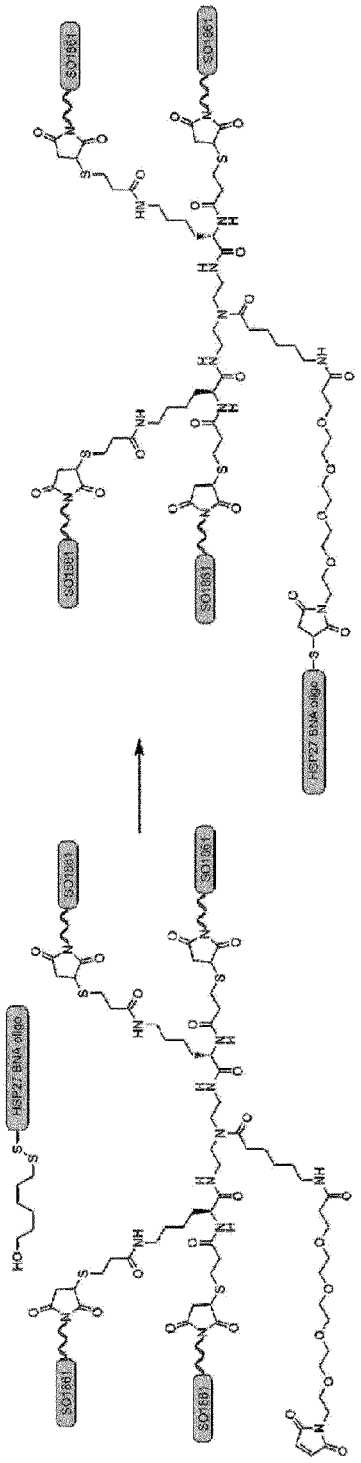

SG1681
SG1681
SG1681
SG1681
HSP27 Ab Hinge
SG1681
SG1681
SG1681
SG1681
HSP27 Ab Hinge

Figure 23

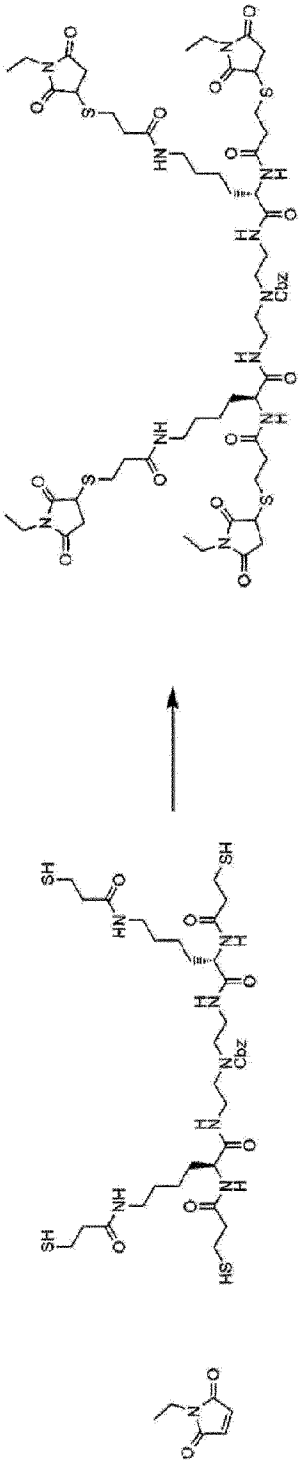


Figure 24

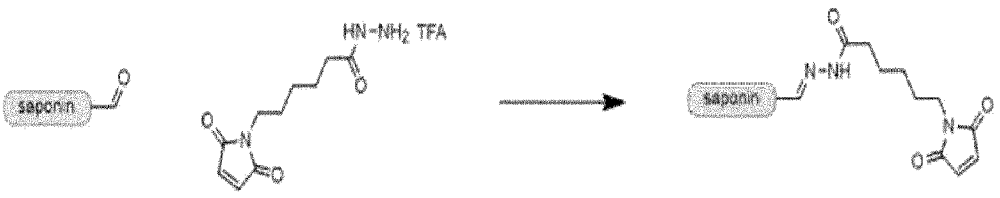


Figure 25

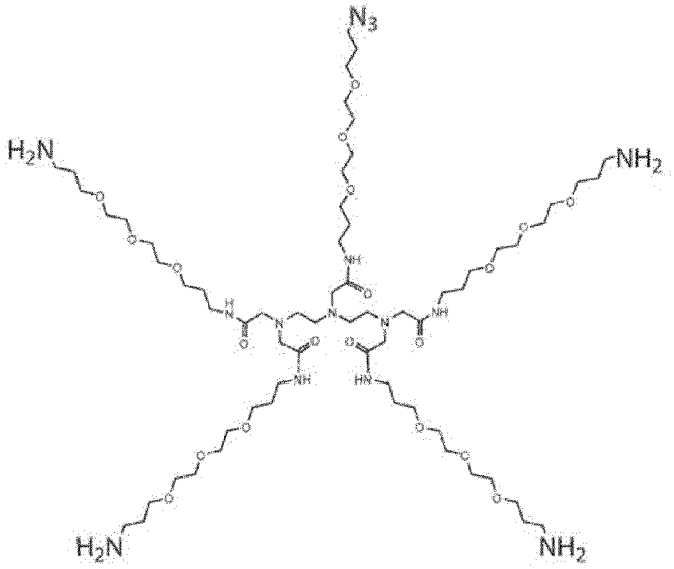


Figure 26

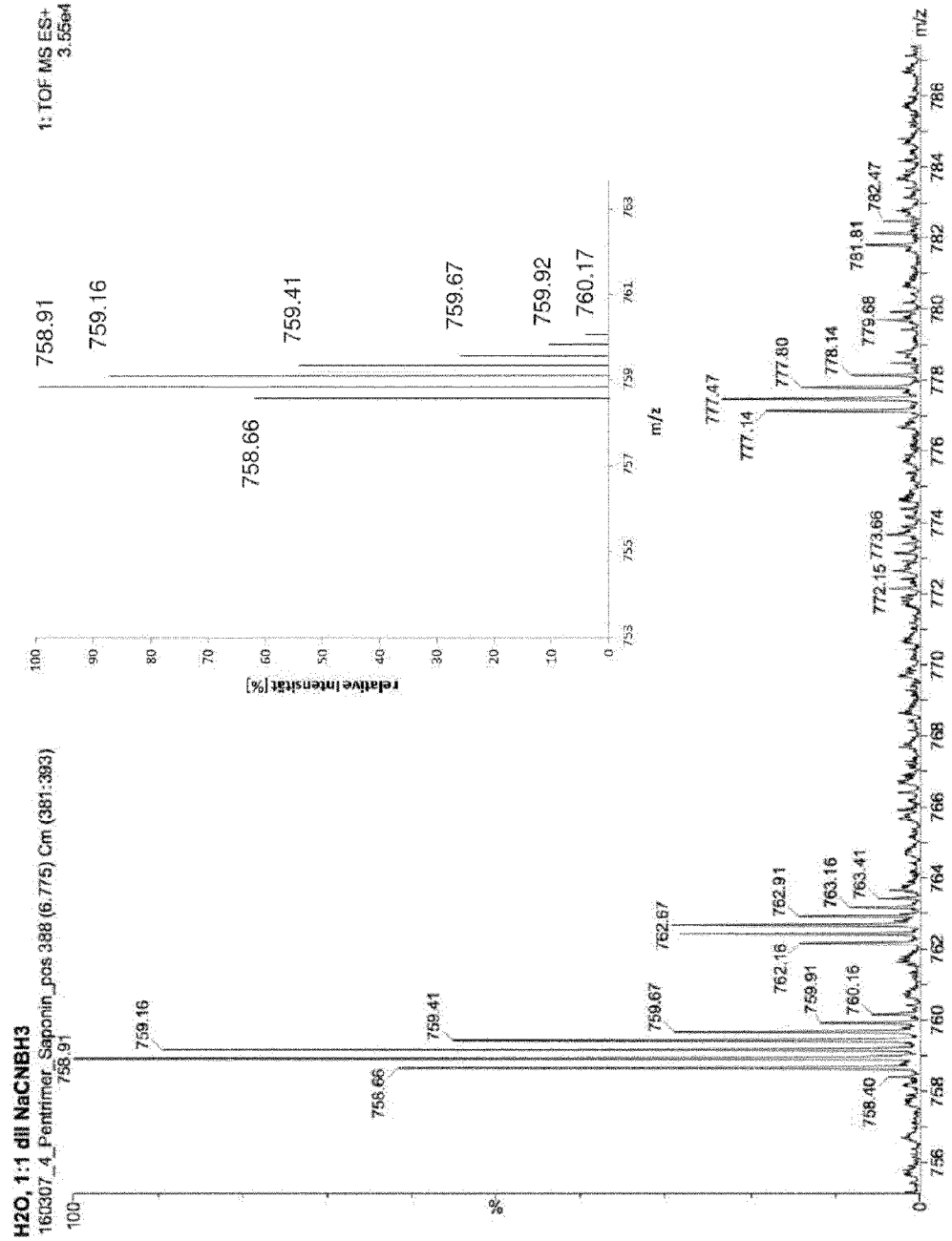

H2O, 1:1 dil NaCNBH3
160307_4_Pentrimer_Saponin_pos 388 (6.775) Cm (381:393)
1: TOF MS ES+
3.55e4
758.91
759.16
759.41
759.67
759.92
760.17
758.66
relative Intensität [%]
m/z
%
m/z

Figure 27

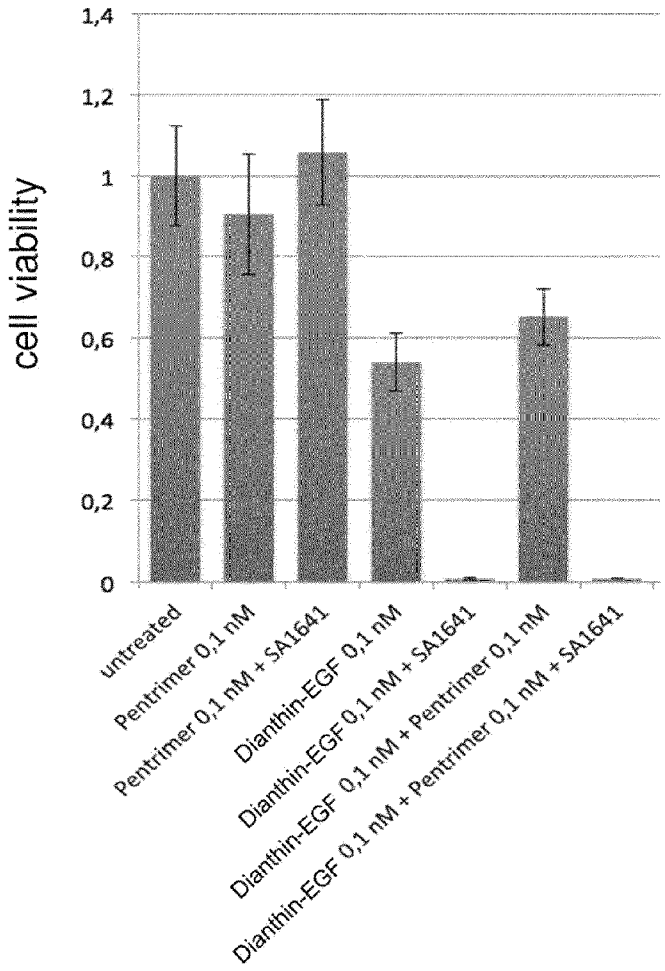

cell viability
1,4
1,2
1
0,8
0,6
0,4
0,2
0
untreated
Pentrimer 0,1 nM
Pentrimer 0,1 nM + SA1641
Dianthin-EGF 0,1 nM
Dianthin-EGF 0,1 nM + SA1641
Dianthin-EGF 0,1 nM + Pentrimer 0,1 nM
Dianthin-EGF 0,1 nM + Pentrimer 0,1 nM + SA1641

Figure 28

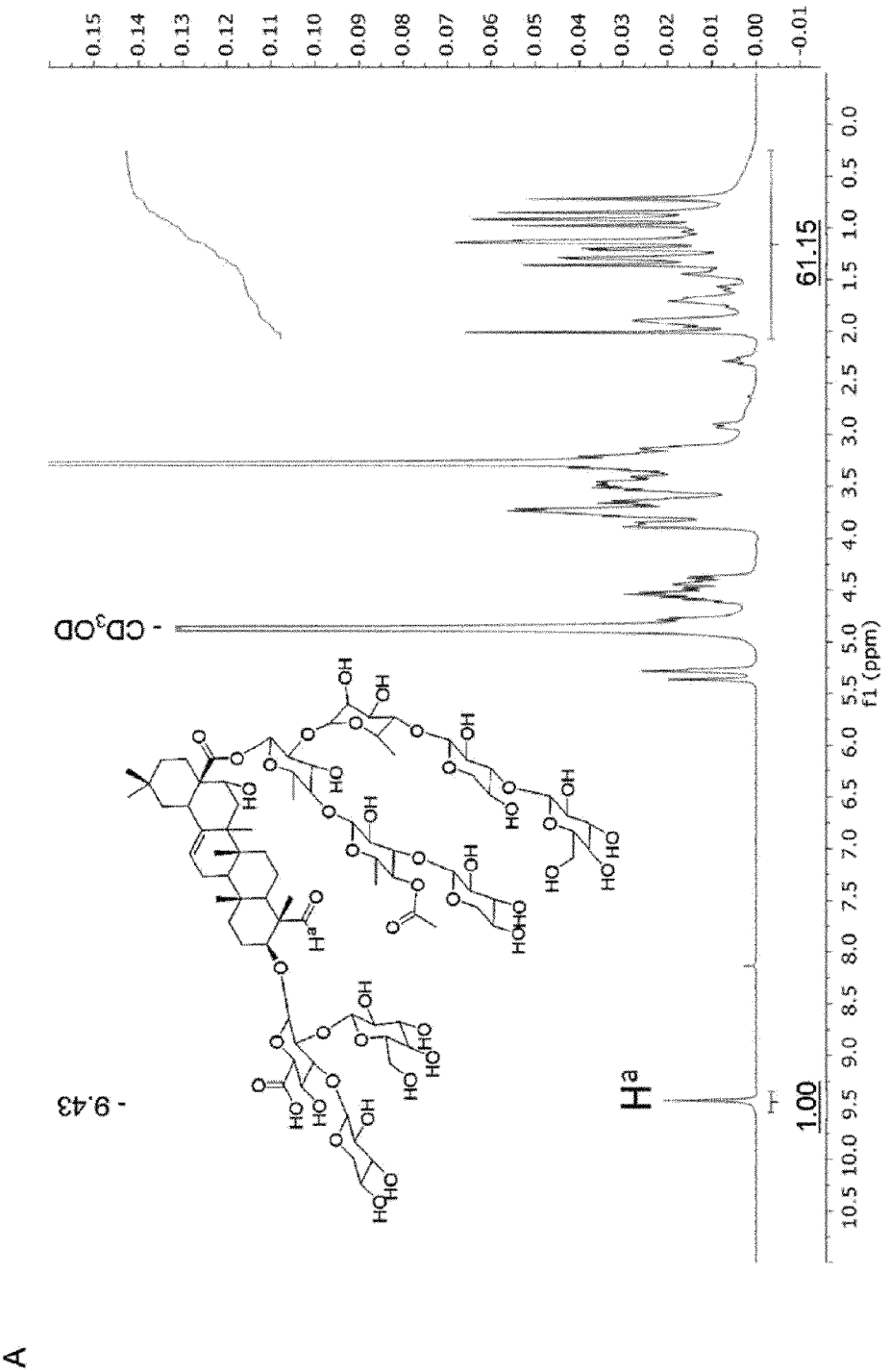
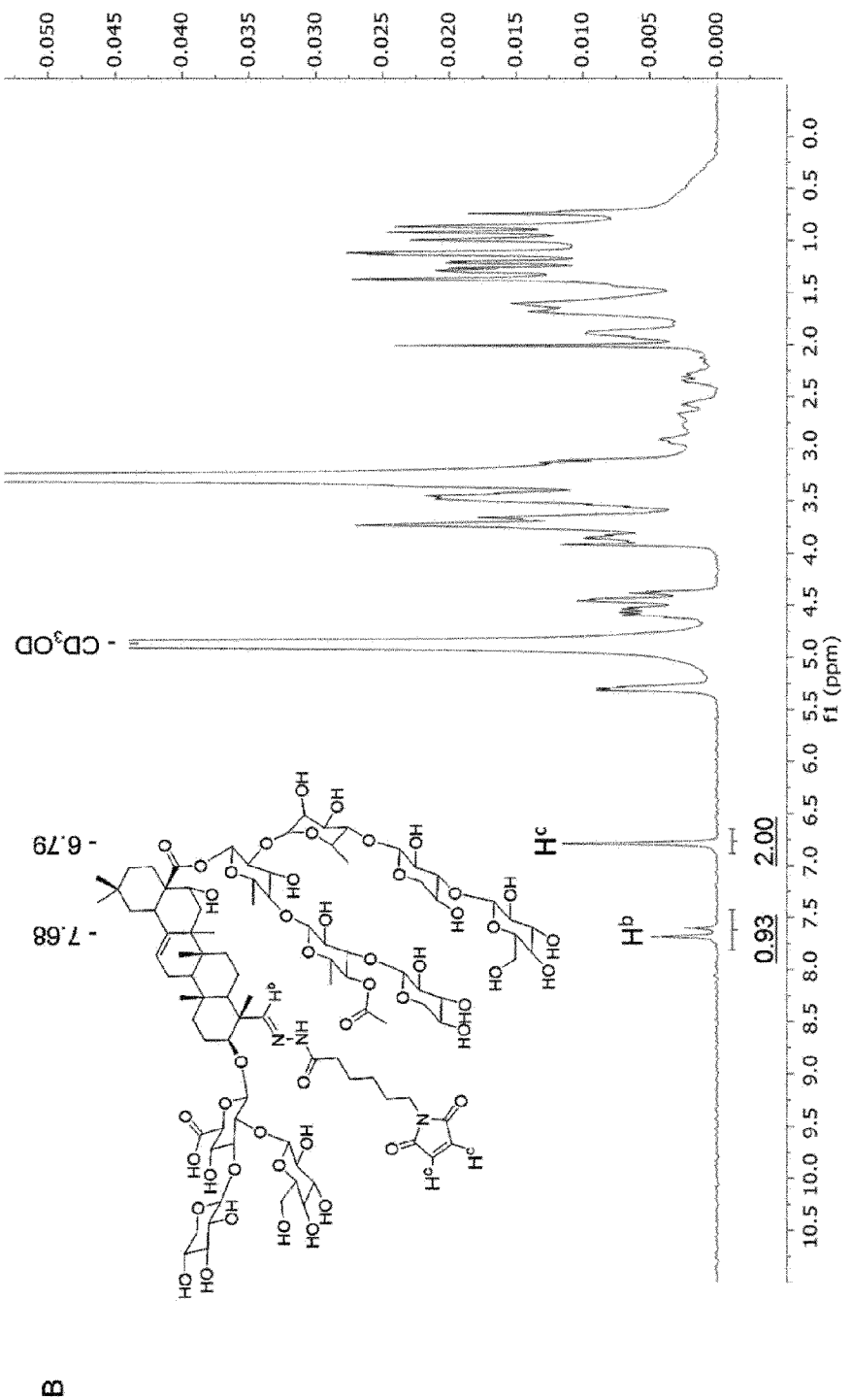


Figure 28


0.050
0.045
0.040
0.035
0.030
0.025
0.020
0.015
0.010
0.005
0.000
CD3OD
7.68
6.79
Hc
Hb
0.93
2.00
10.5 10.0 9.5 9.0 8.5 8.0 7.5 7.0 6.5 6.0 5.5 5.0 4.5 4.0 3.5 3.0 2.5 2.0 1.5 1.0 0.5 0.0
f1 (ppm)

B

Figure 29

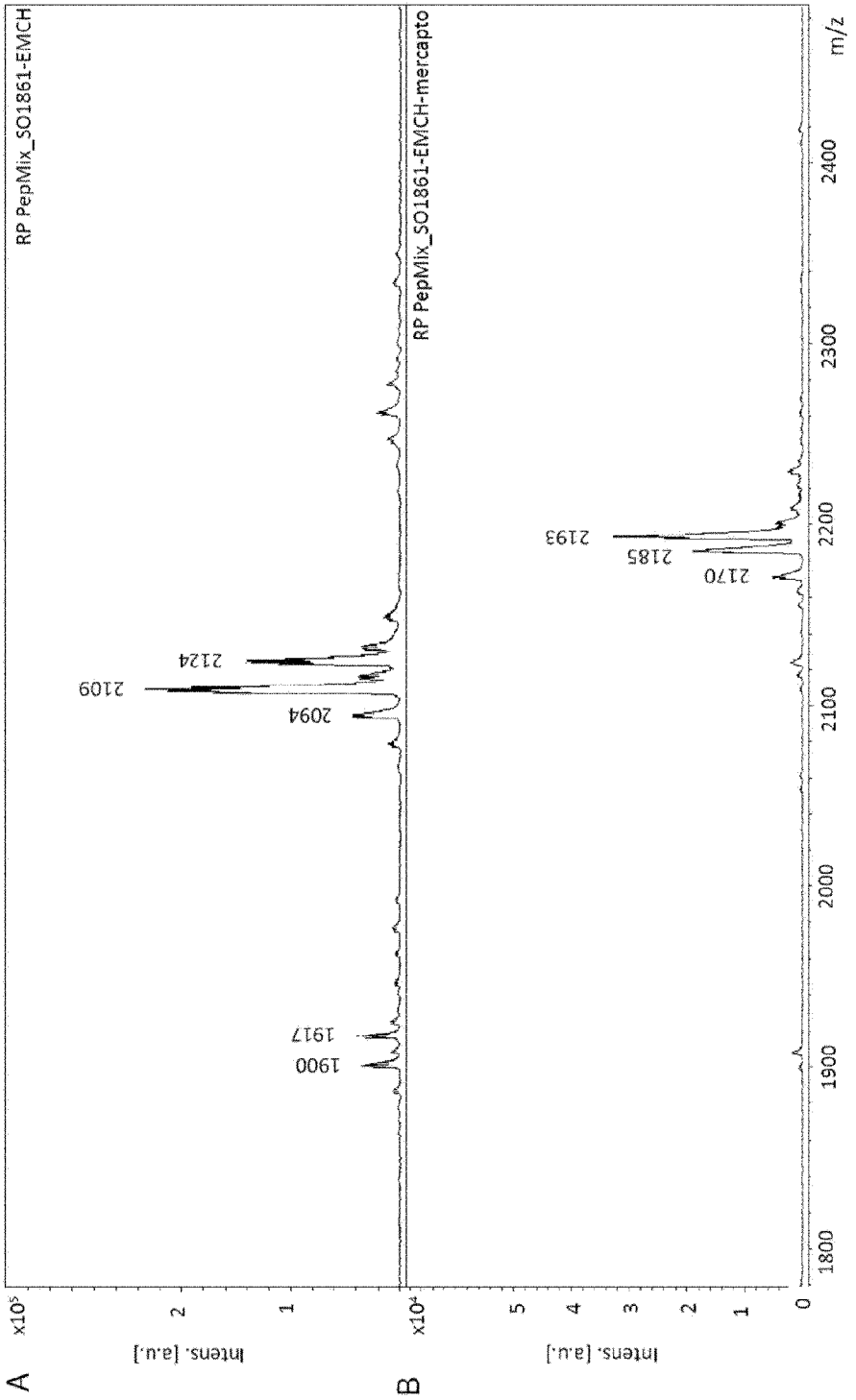

RP PepMix_SO1861-EMCH
RP PepMix_SO1861-EMCH-mercapto
A
B
Intens. [a.u.]
Intens. [a.u.]
x10⁵
x10⁴
m/z
1800
1900
2000
2100
2200
2300
2400
1900
1917
2094
2109
2124
2170
2185
2193

Figure 30

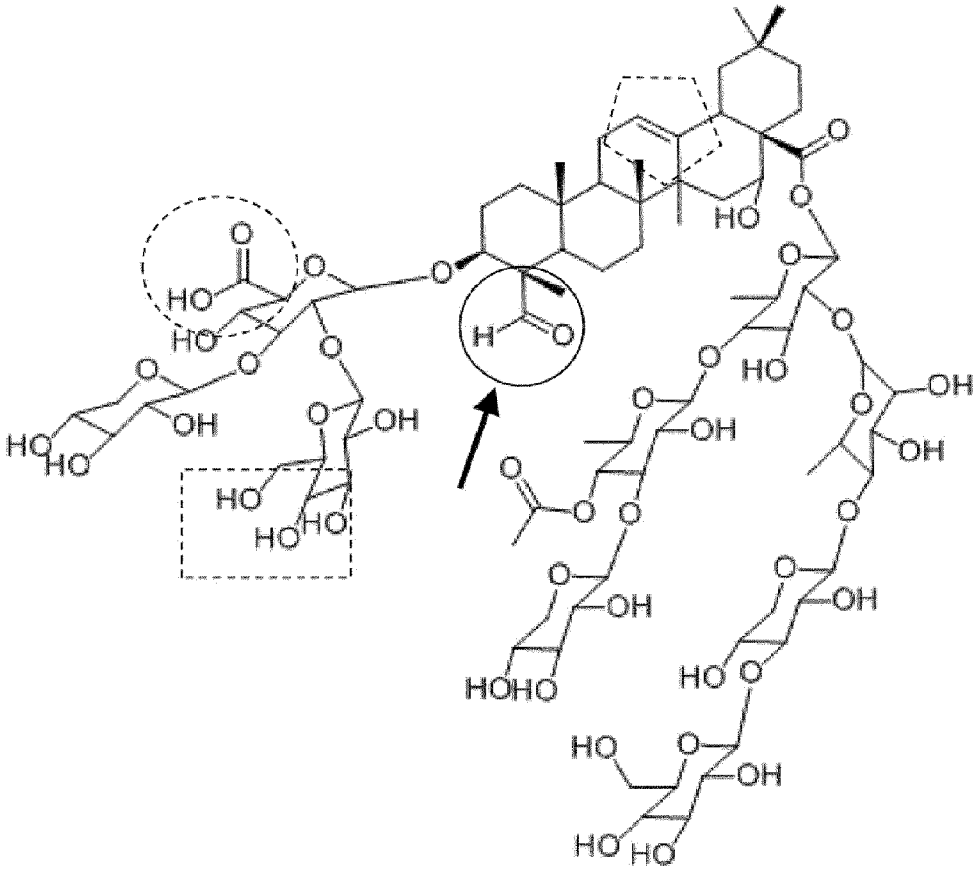


Figure 31

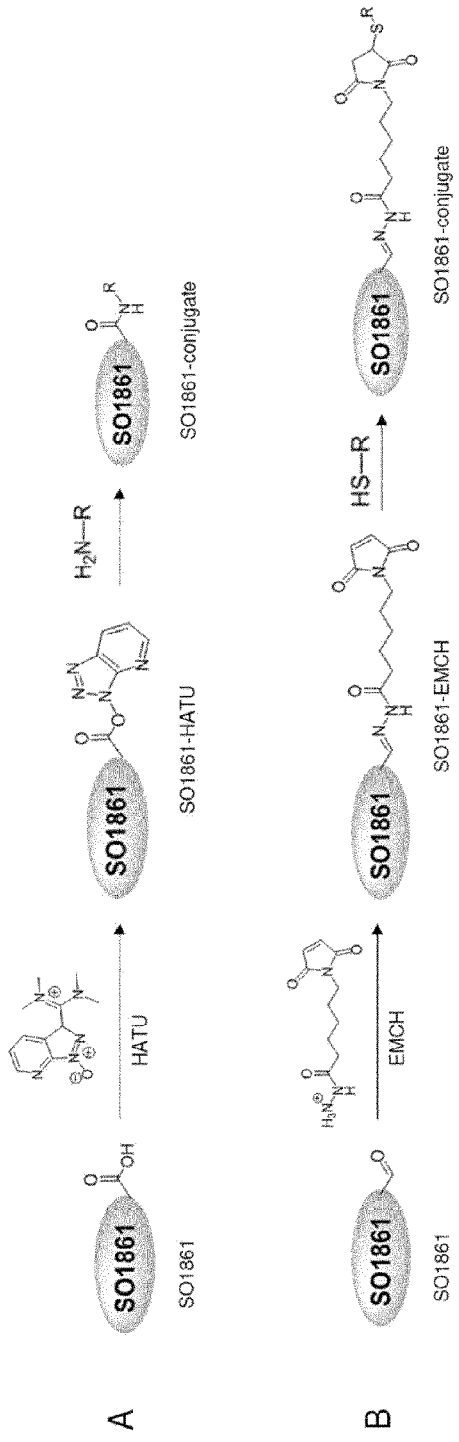

A
SO1861
SO1361
HATU
SO1861-HATU
H_2N—R
SO1861-conjugate
B
SO1861
SO1861
EMCH
SO1861-EMCH
HS—R
SO1861-conjugate

Figure 32

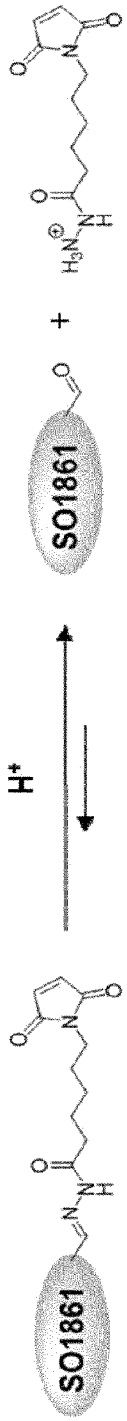
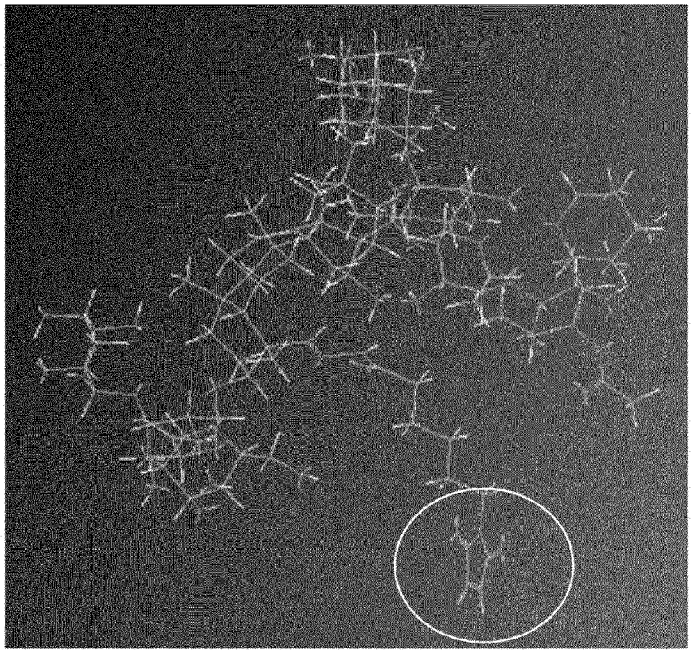
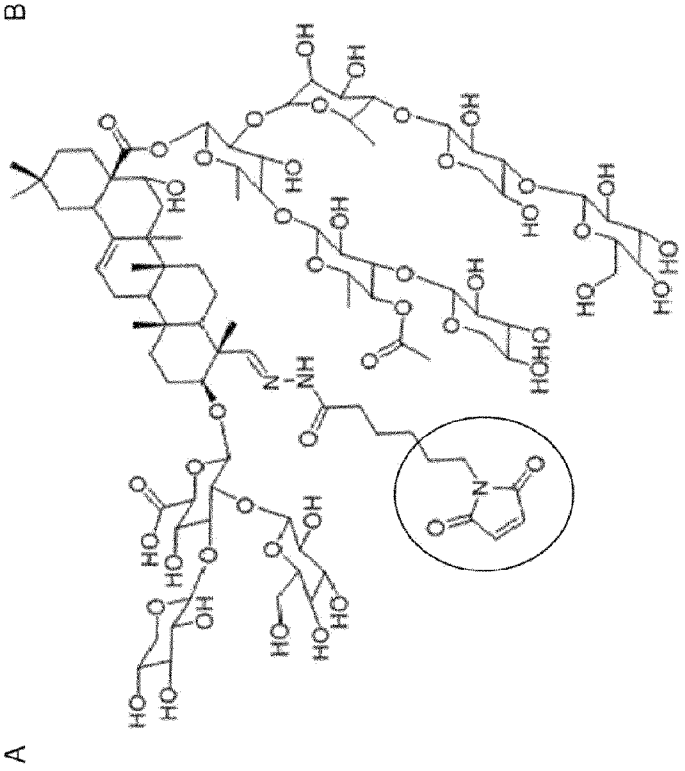


Figure 33



B



A

Figure 34

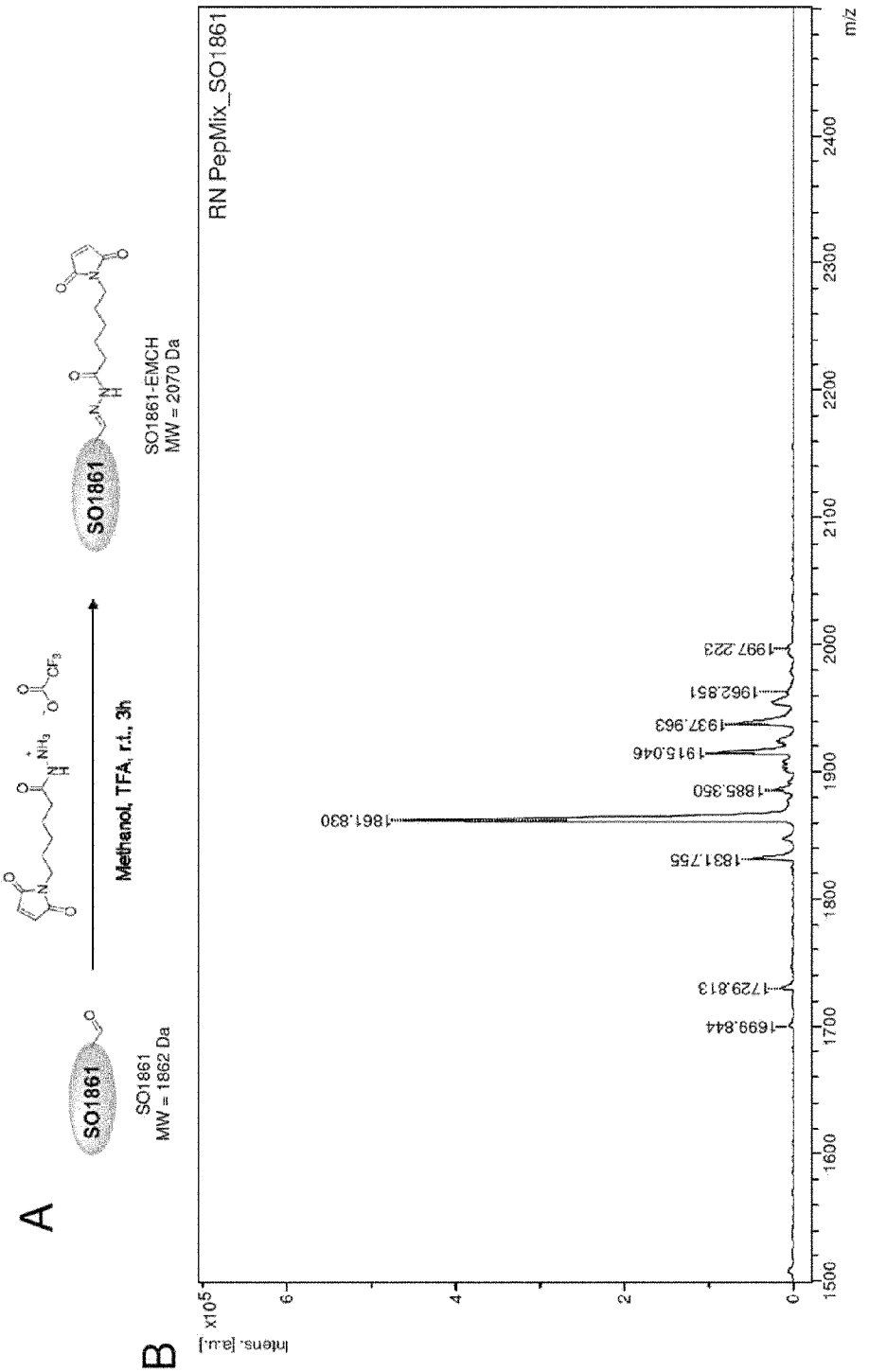
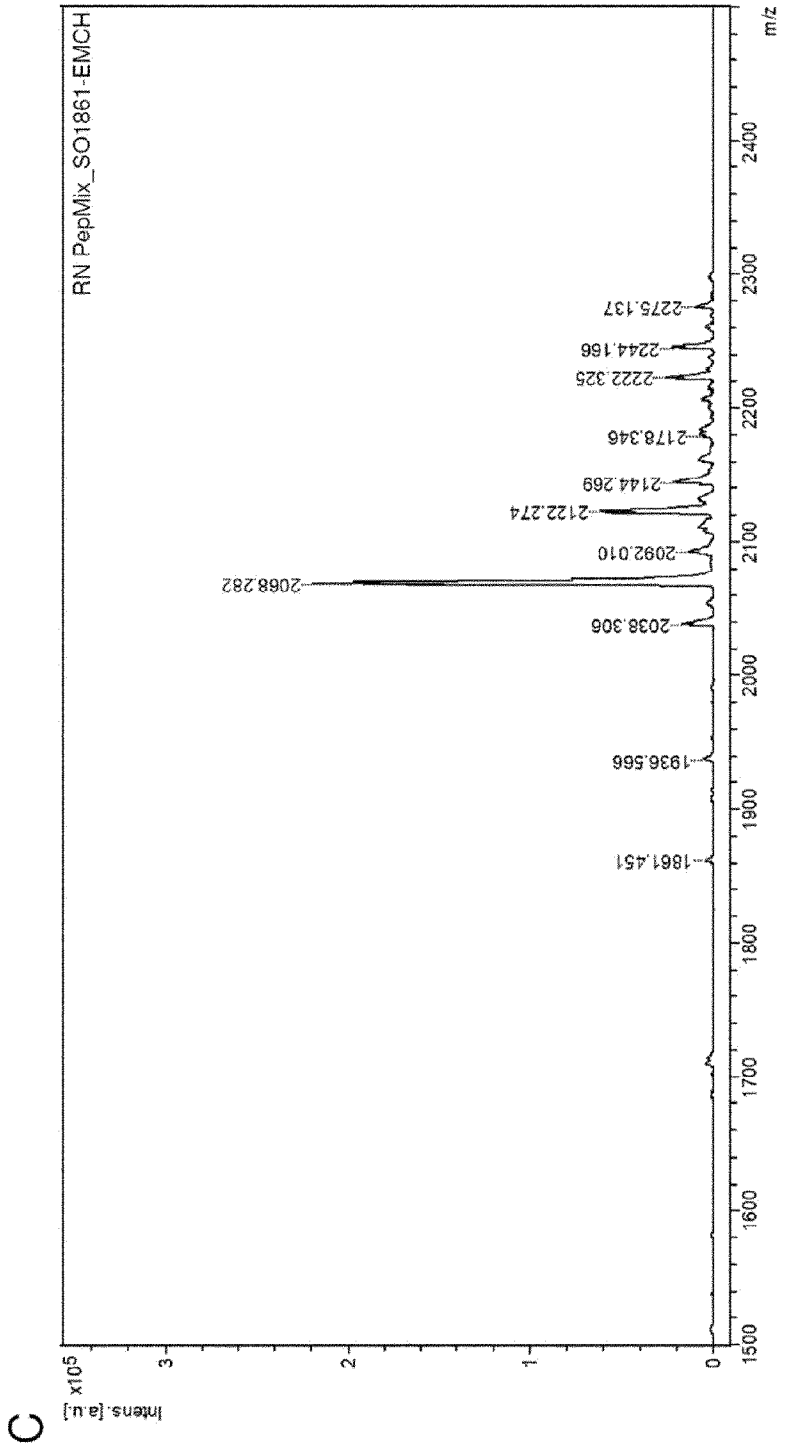

A
SO1861
SO1861
MW = 1862 Da
Methanol, TFA, r.t., 3h
SO1861
SO1861-EMCH
MW = 2070 Da
B
RN PepMix_SO1861
Intens. [a.u.]
x10^5
m/z
1699.844
1729.813
1831.755
1861.830
1885.350
1915.046
1937.963
1962.851
1997.223

Figure 34


RN PepMix_SO1861-EMCH
Intens. [a.u.]
x10^5
3
2
1
0
1500
1600
1700
1800
1900
2000
2100
2200
2300
2400
m/z
1861.451
1936.566
2038.306
2068.282
2092.010
2122.274
2144.269
2178.346
2222.325
2244.166
2275.137

C

Figure 35

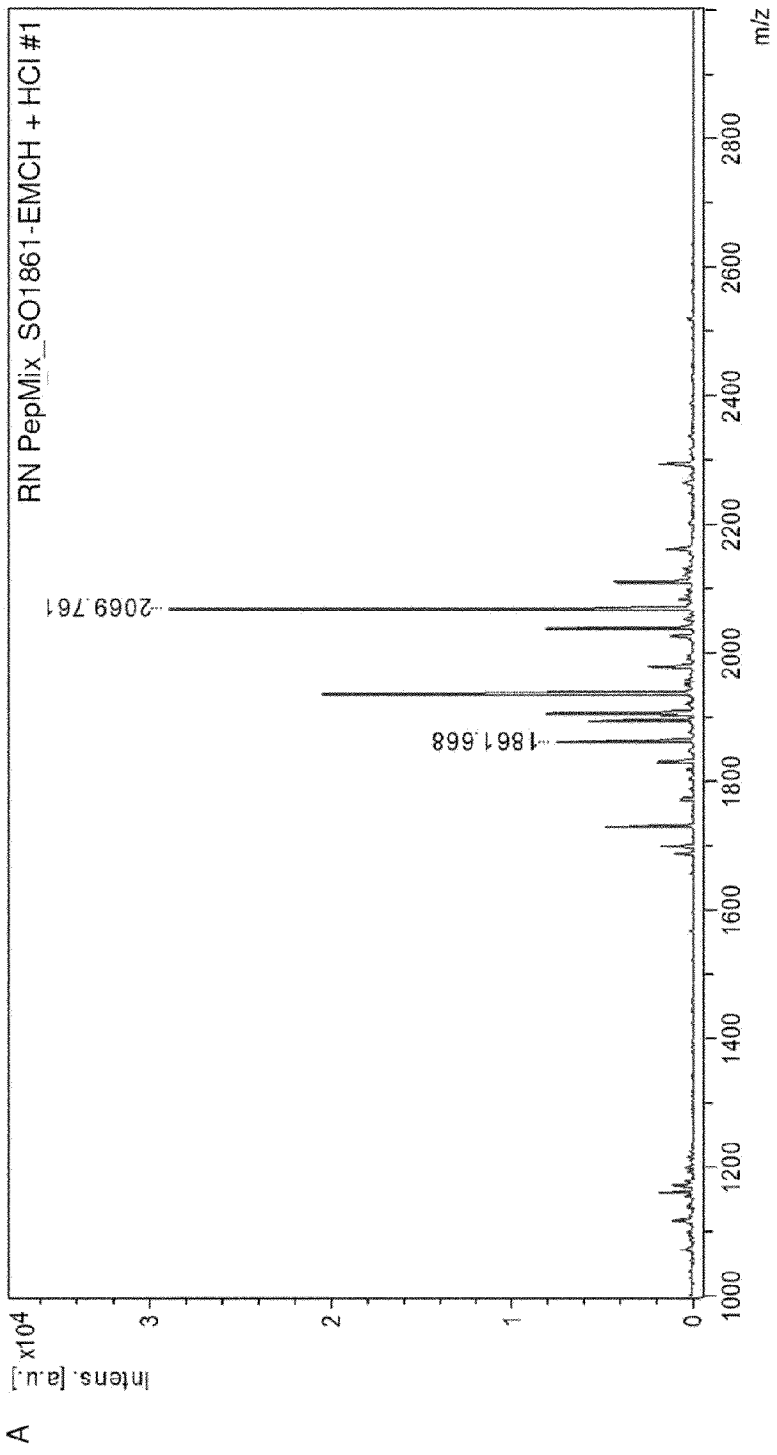

A
Intens. [a.u.]
x10^4
3
2
1
0
1000
1200
1400
1600
1800
2000
2200
2400
2600
2800
m/z
RN PepMix_SO1861-EMCH + HCl #1
2069.761
1861.668

Figure 35

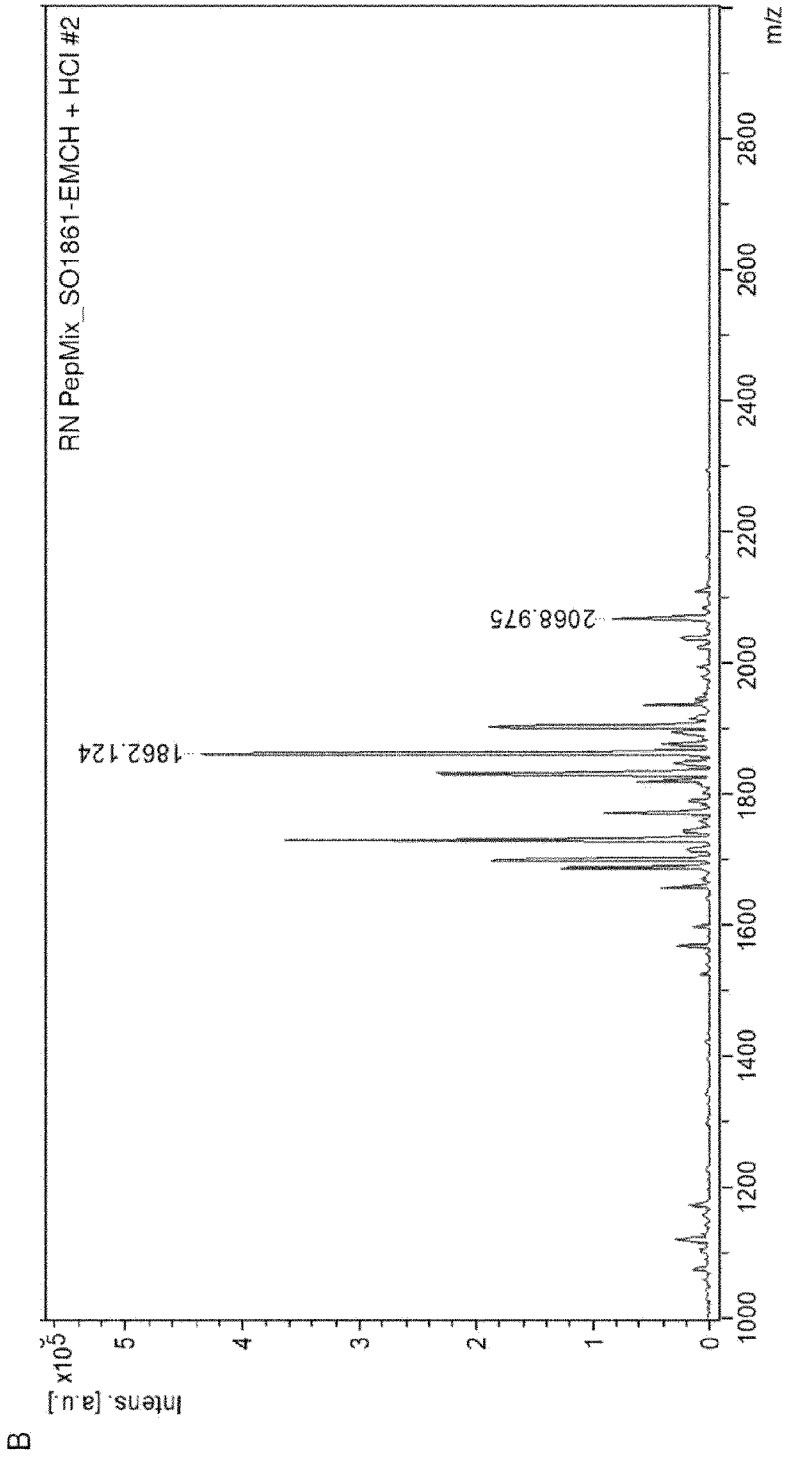

B
RN PepMix_SO1861-EMCH + HCl #2
Intens. [a.u.]
x10^5
0
1
2
3
4
5
1000
1200
1400
1600
1800
2000
2200
2400
2600
2800
m/z
1862.124
2068.975

Figure 36

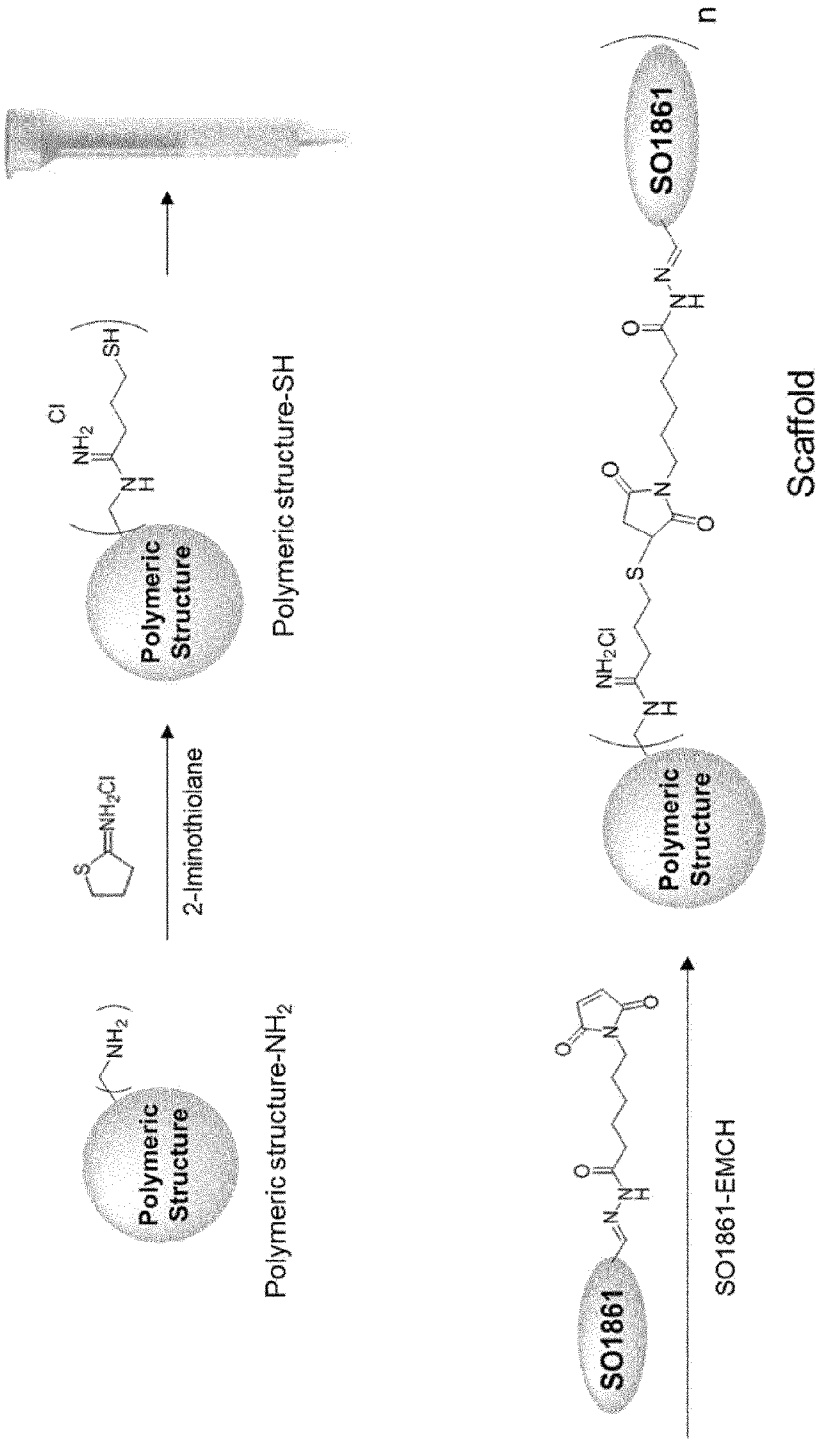

Polymeric Structure
Polymeric structure-NH2
2-Iminothiolane
Polymeric structure-SH
SO1861-EMCH
SO1861
Scaffold
n

Figure 37

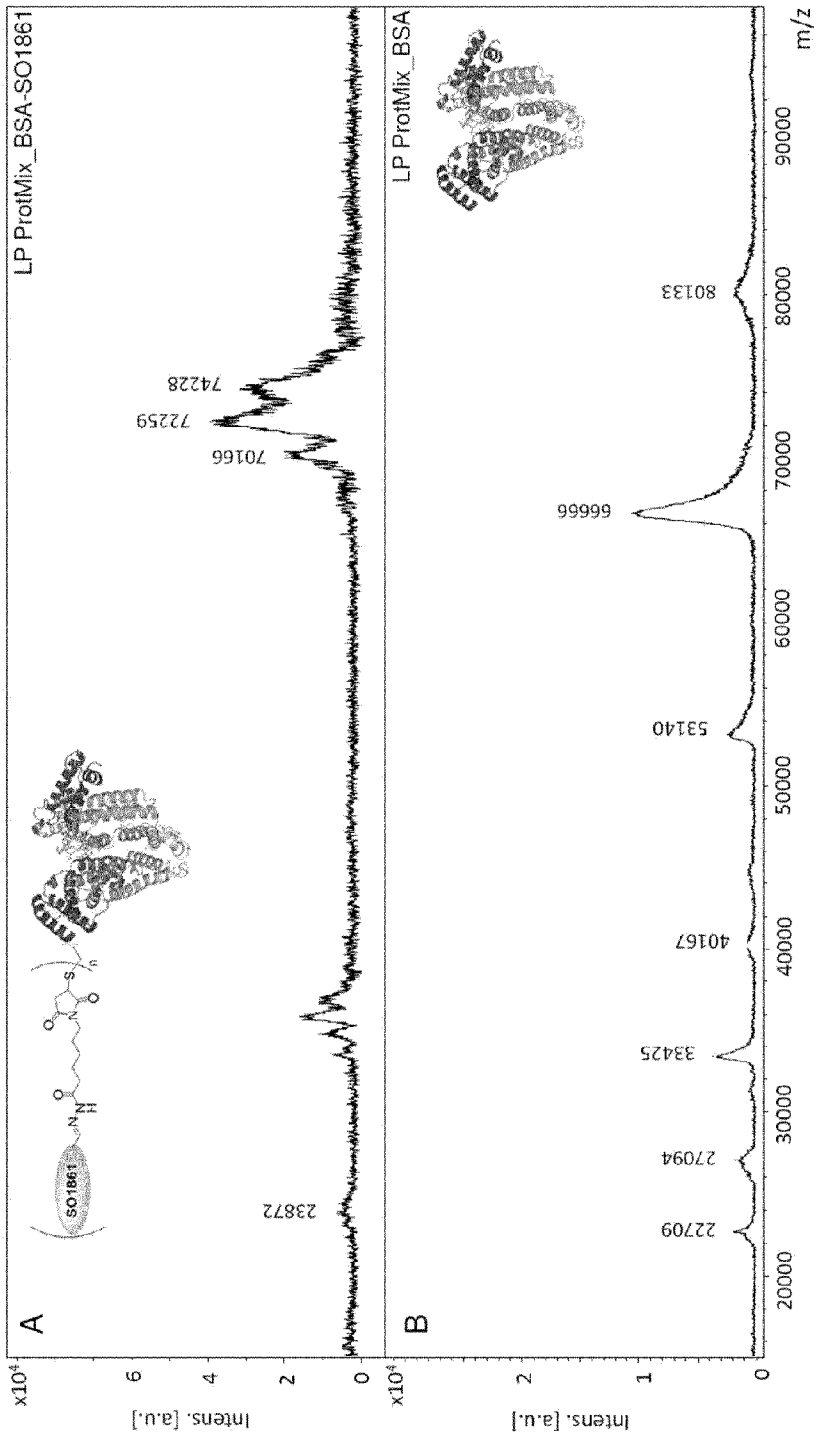

A
LP ProtMix_BSA-SO1861
SO1861
23872
70166
72259
74228
Intens. [a.u.]
x10⁴
B
LP ProtMix_BSA
22709
27094
33425
40167
53140
66666
80133
m/z
Intens. [a.u.]
x10⁴

Figure 38

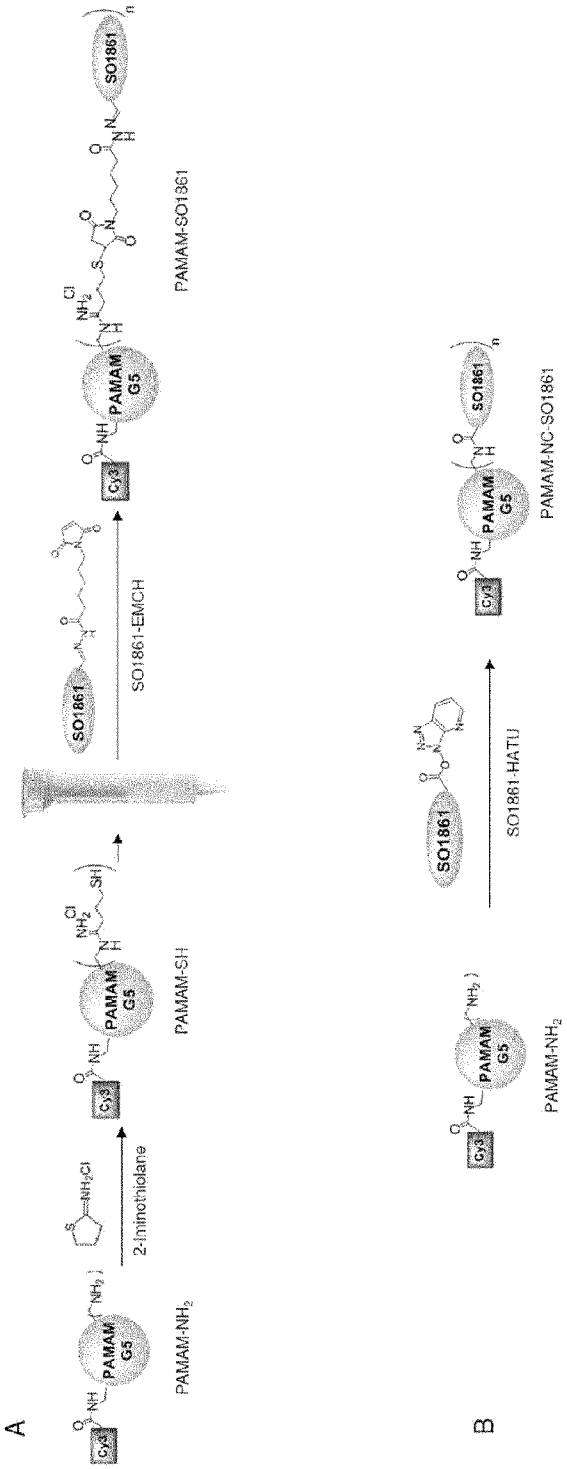

A
PAMAM-NH2
2-Iminothiolane
PAMAM-SH
SO1861-EMCH
PAMAM-SO1361
B
SO1861-HATU
PAMAM-NH2
PAMAM-NC-SO1861

Figure 39

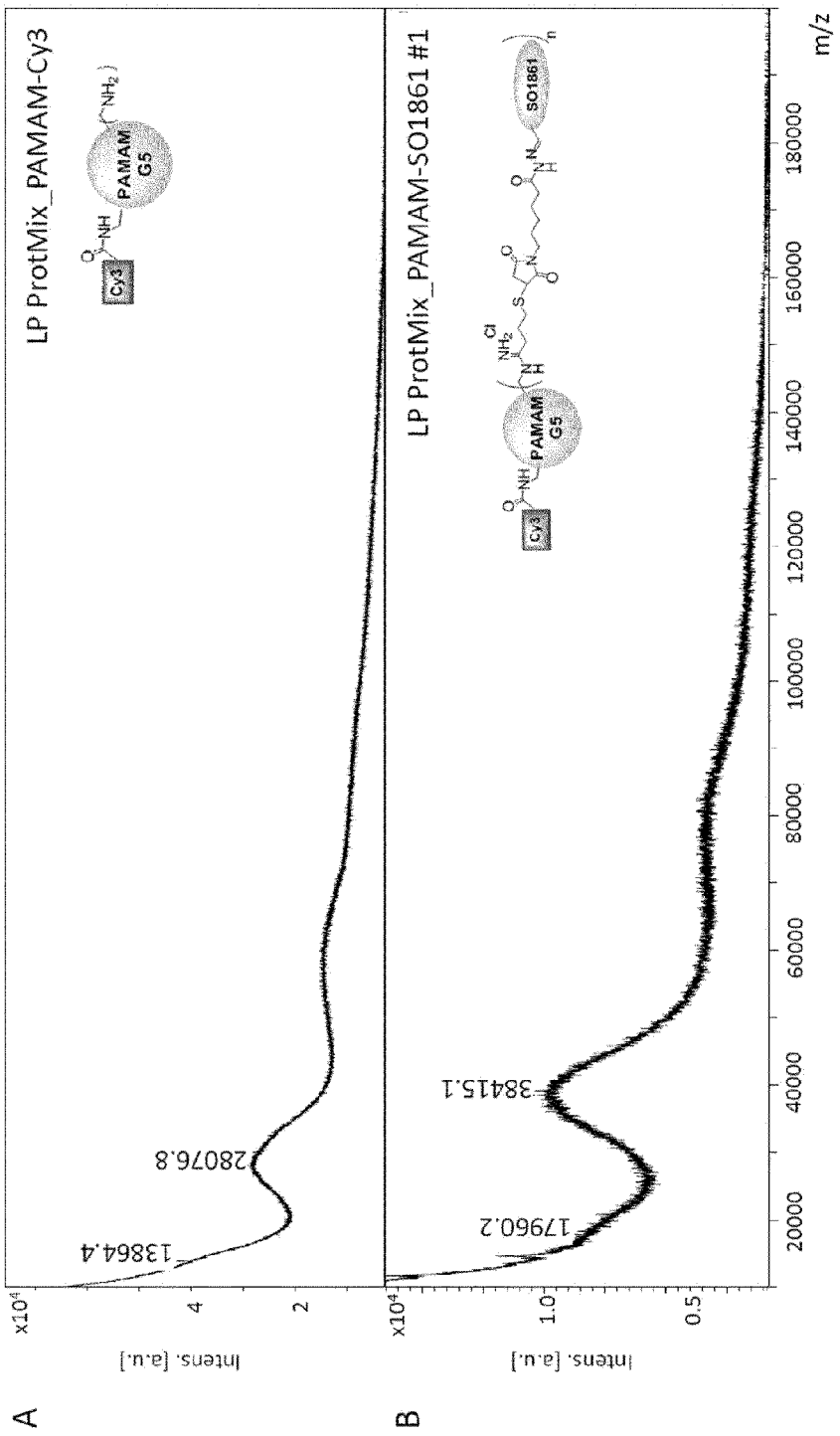

A
LP ProtMix_PAMAM-Cy3
PAMAM G5
Cy3
13864.4
28076.8
Intens. [a.u.]
x10^4
4
2
B
LP ProtMix_PAMAM-SO1861 #1
PAMAM G5
Cy3
SO1861
38415.1
17960.2
Intens. [a.u.]
x10^4
1.0
0.5
20000
40000
60000
80000
100000
120000
140000
160000
180000
m/z

Figure 39

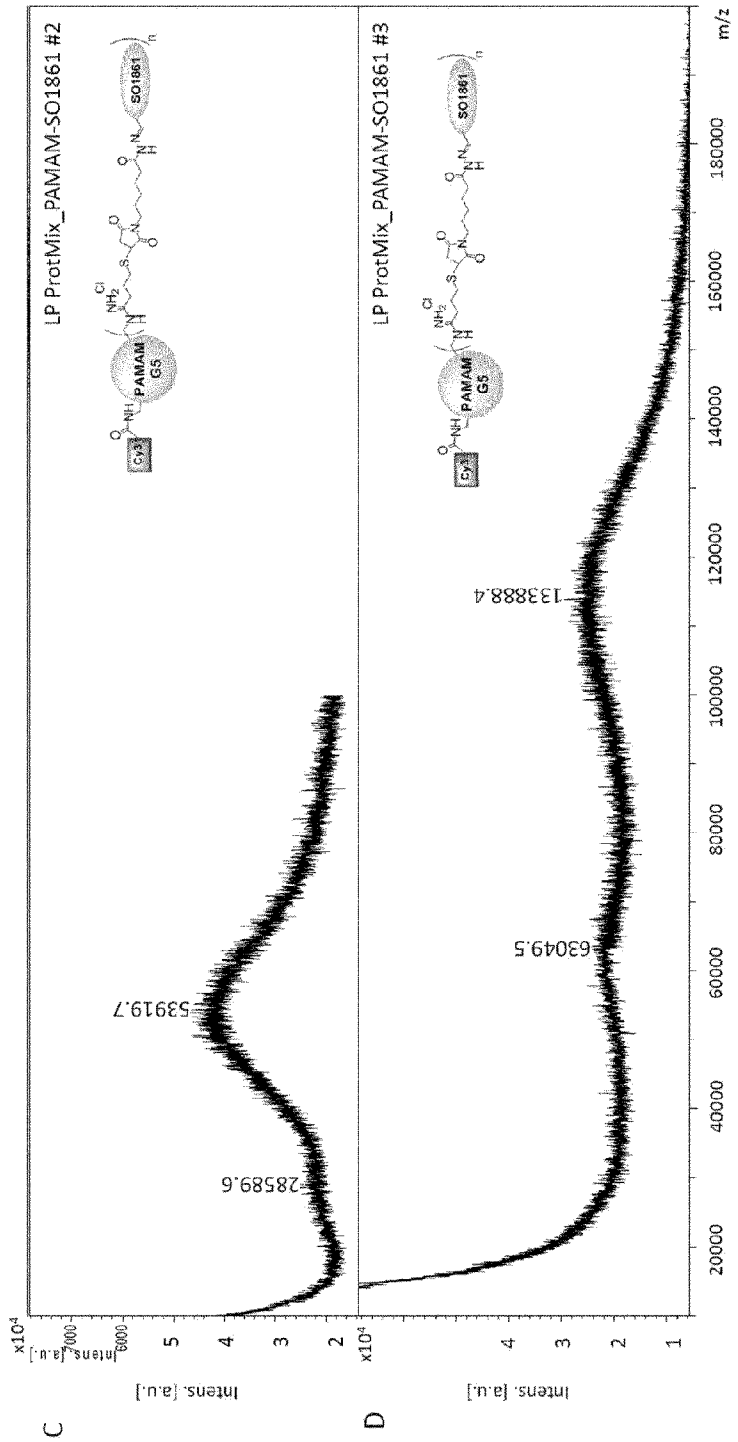

C
LP ProtMix_PAMAM-SO1861 #2
PAMAM G5
Cy3
SO1861
28589.6
53919.7
Intens. [a.u.]
x10^4
D
LP ProtMix_PAMAM-SO1861 #3
PAMAM G5
Cy3
SO1861
63049.5
133888.4
Intens. [a.u.]
x10^4
m/z
20000
40000
60000
80000
100000
120000
140000
160000
180000

Figure 40

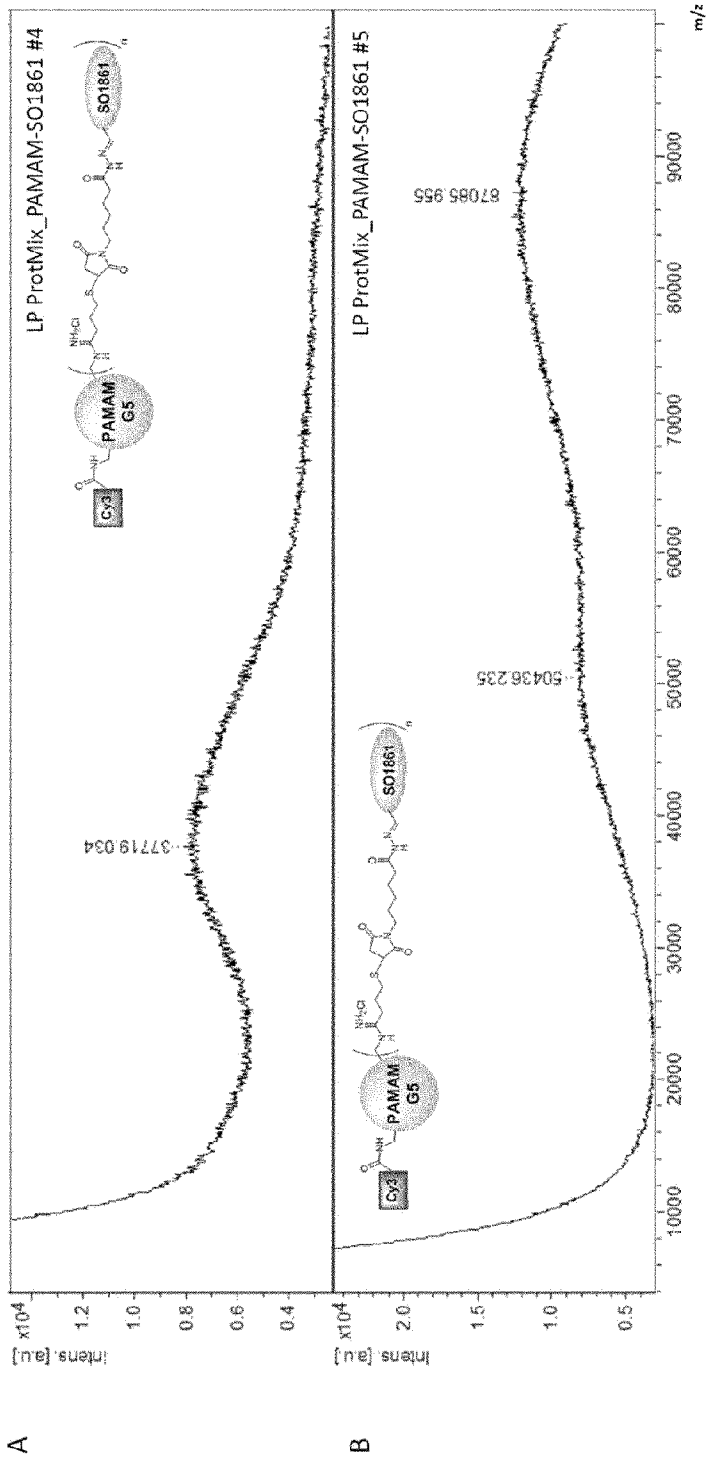

A
B
LP ProtMix_PAMAM-SO1861 #4
LP ProtMix_PAMAM-SO1861 #5
PAMAM G5
Cy3
SO1861
377190.34
870659.55
504362.35
Intens. [a.u.]
x10^4
m/z

Figure 41

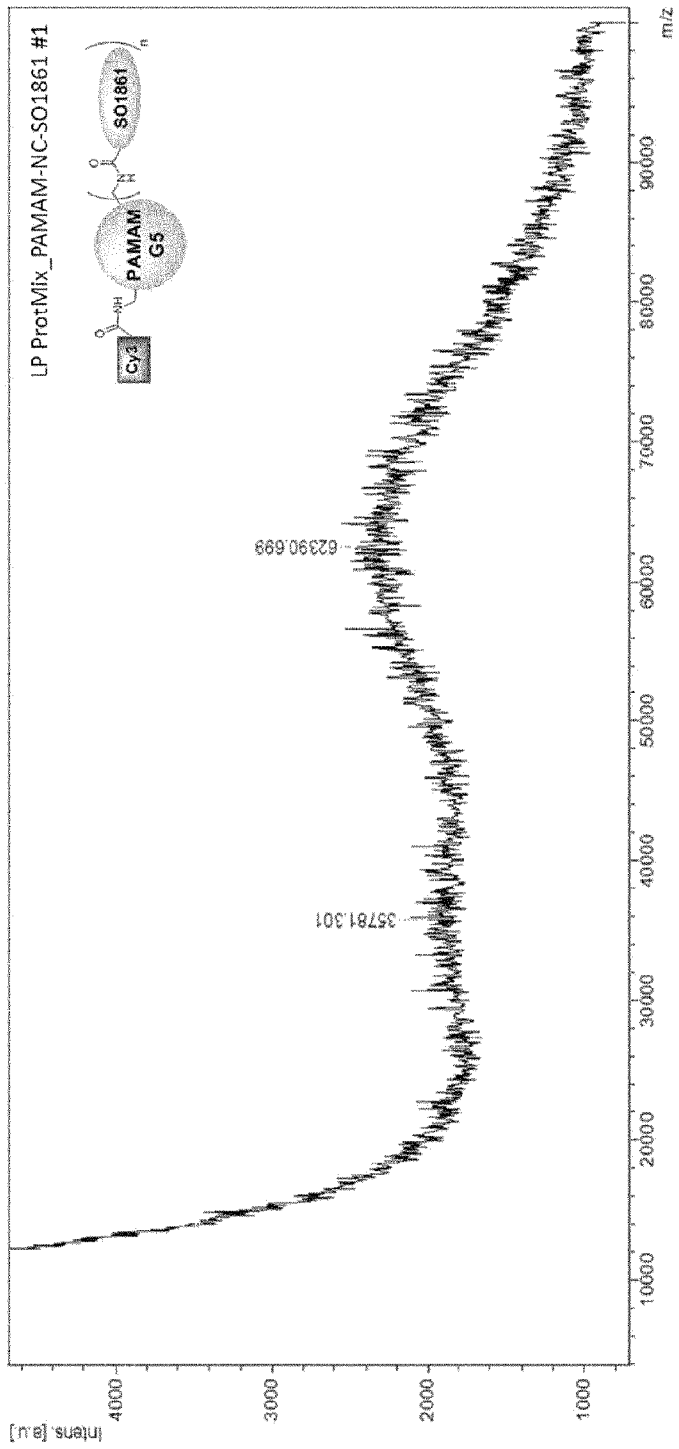

LP ProtMix_PAMAM-NC-SO1861 #1
SO1861
PAMAM G5
Cy3
62390.699
35781.301
Intens. [a.u.]
4000
3000
2000
1000
m/z
10000
20000
30000
40000
50000
60000
70000
80000
90000

Figure 42

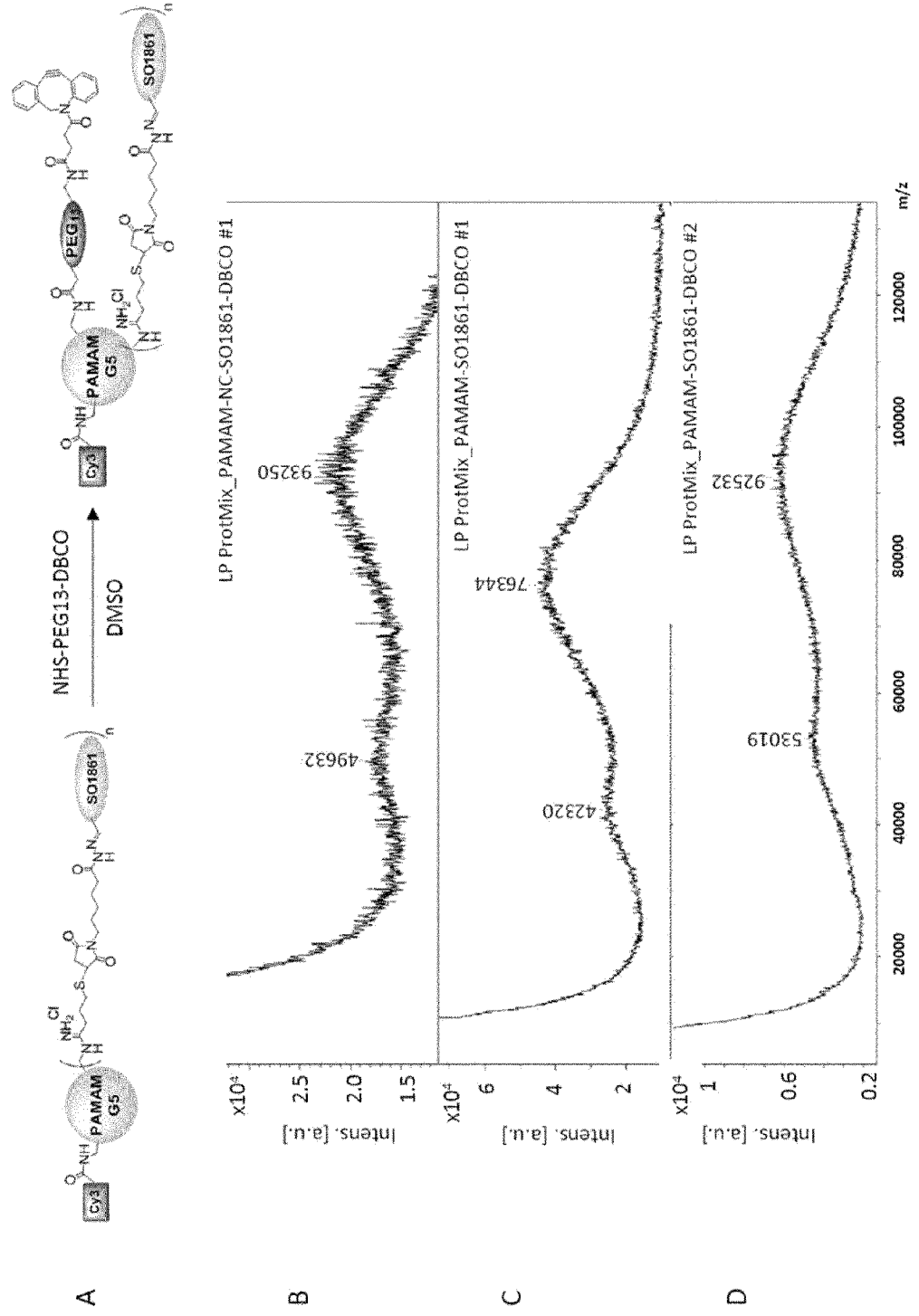

A
Cy3
PAMAM G5
NHS-PEG13-DBCO
DMSO
PEG 13
SO1861
B
LP ProtMix_PAMAM-NC-SO1861-DBCO #1
93250
49632
C
LP ProtMix_PAMAM-SO1861-DBCO #1
76344
42320
D
LP ProtMix_PAMAM-SO1861-DBCO #2
92532
53019
Intens. [a.u.]
x10⁴
m/z
20000
40000
60000
80000
100000
120000

Figure 43

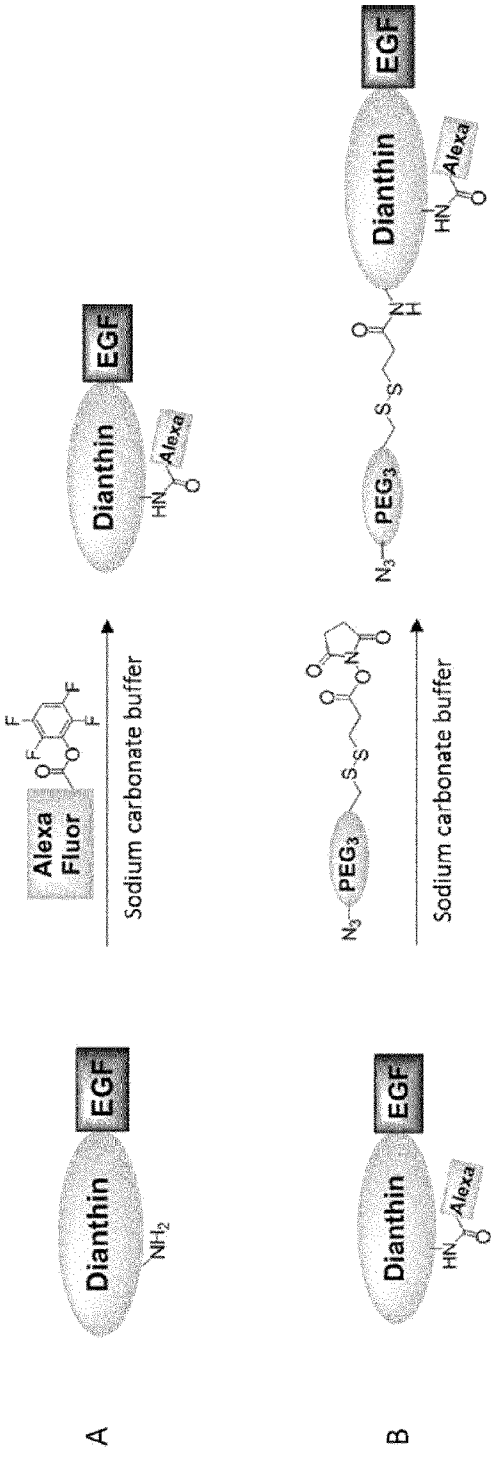


Figure 43

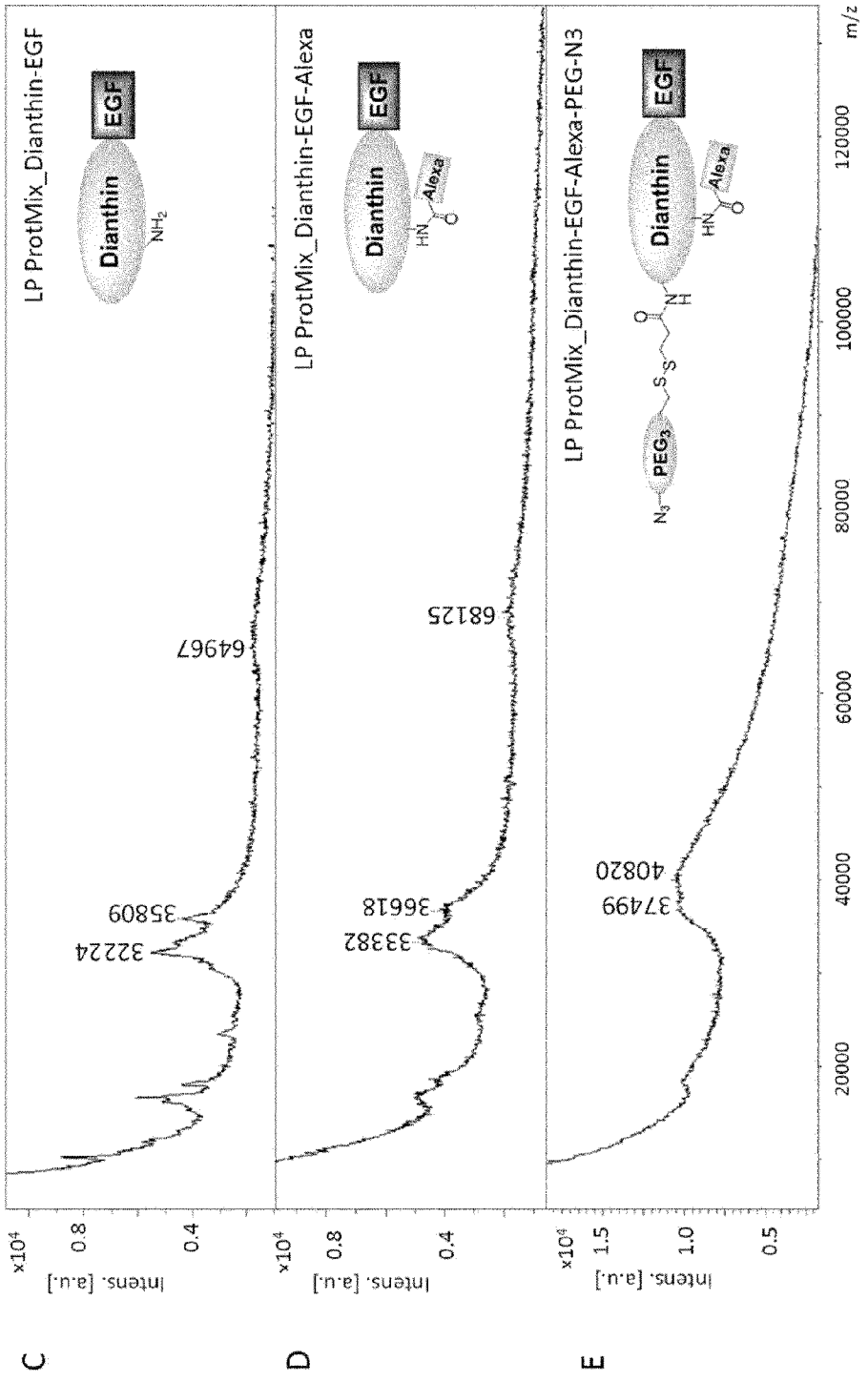

C
LP ProtMix_Dianthin-EGF
Dianthin
EGF
NH2
64967
35809
32224
Intens. [a.u.]
x10^4
0.8
0.4
D
LP ProtMix_Dianthin-EGF-Alexa
Dianthin
EGF
Alexa
68125
36618
33382
Intens. [a.u.]
x10^4
0.8
0.4
E
LP ProtMix_Dianthin-EGF-Alexa-PEG-N3
Dianthin
EGF
Alexa
PEG3
N3
40820
37499
Intens. [a.u.]
x10^4
1.5
1.0
0.5
20000
40000
60000
80000
100000
120000
m/z

Figure 44

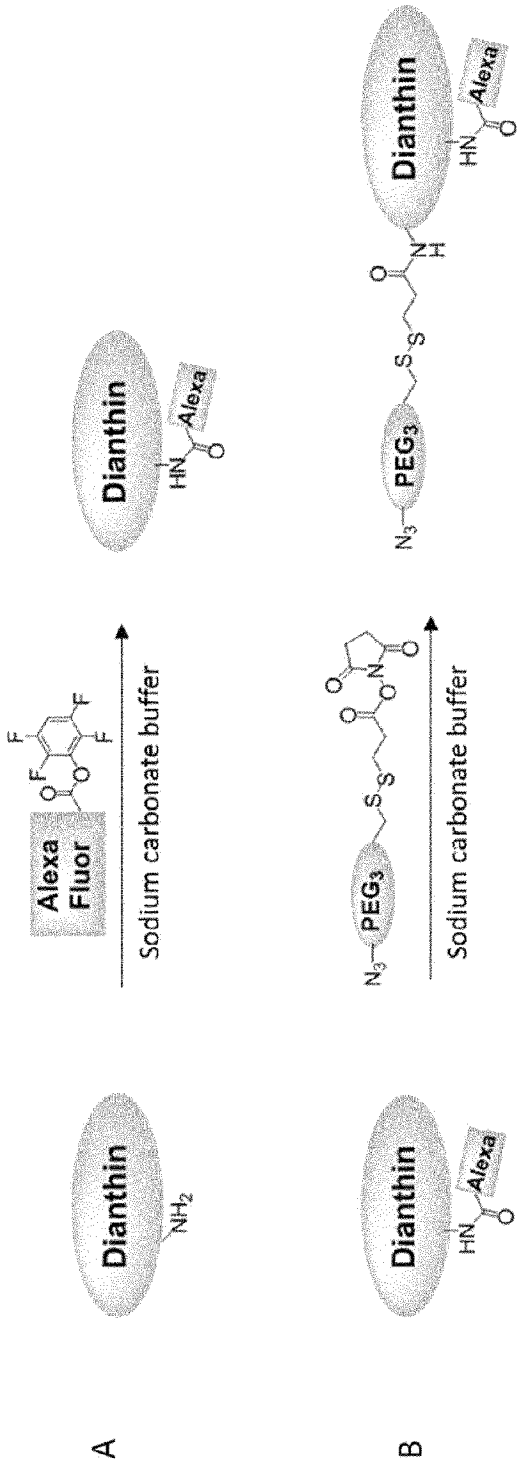

A
Dianthin
NH2
Alexa Fluor
Sodium carbonate buffer
Dianthin
Alexa
B
Dianthin
Alexa
PEG3
Sodium carbonate buffer
Dianthin
Alexa
PEG3

Figure 44

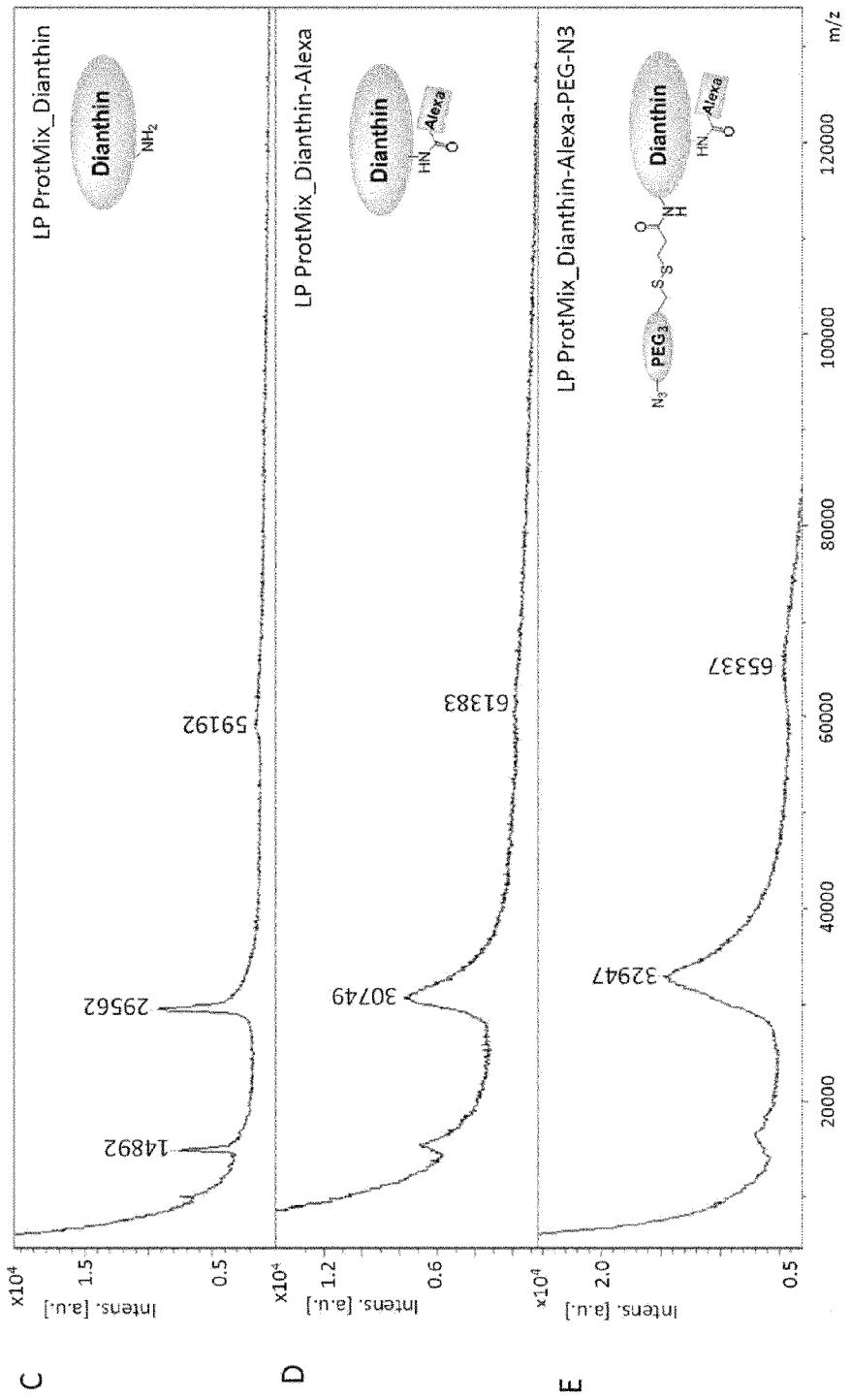

C
LP ProtMix_Dianthin
Dianthin
NH2
14892
29562
59192
Intens. [a.u.]
x10^4
1.5
0.5
D
LP ProtMix_Dianthin-Alexa
Dianthin
Alexa
HN
O
30749
61383
Intens. [a.u.]
x10^4
1.2
0.6
E
LP ProtMix_Dianthin-Alexa-PEG-N3
Dianthin
Alexa
HN
O
PEG3
N3
S
S
32947
65337
Intens. [a.u.]
x10^4
2.0
0.5
20000
40000
60000
80000
100000
120000
m/z

Figure 45

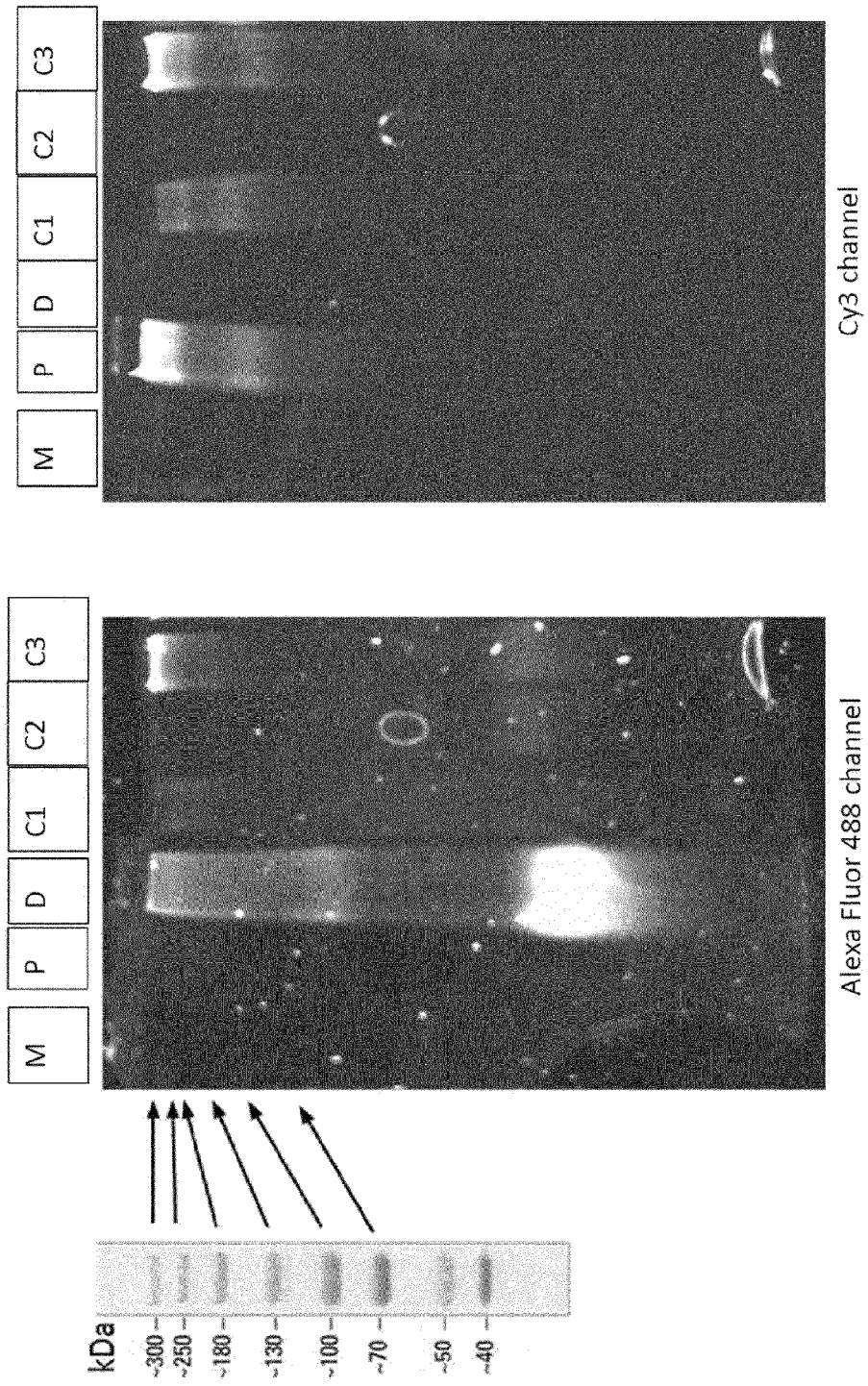

M
P
D
C1
C2
C3
Cy3 channel
M
P
D
C1
C2
C3
Alexa Fluor 488 channel
kDa
~300
~250
~180
~130
~100
~70
~50
~40

Figure 46

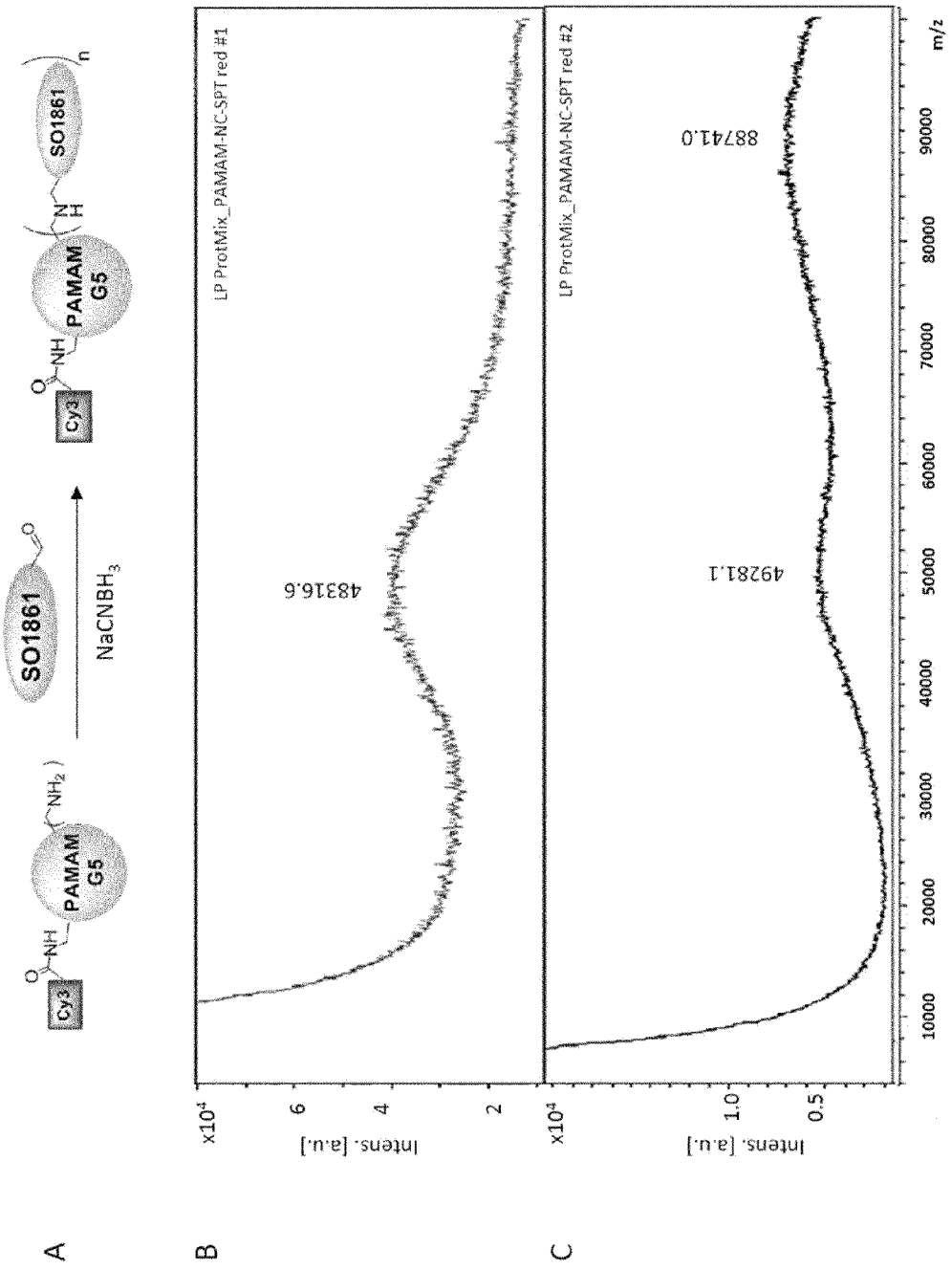

A
B
C
SO1861
NaCNBH3
PAMAM G5
Cy3
LP ProtMix_PAMAM-NC-SPT red #1
LP ProtMix_PAMAM-NC-SPT red #2
48316.6
49281.1
88741.0
Intens. [a.u.]
x10^4
m/z

Figure 47

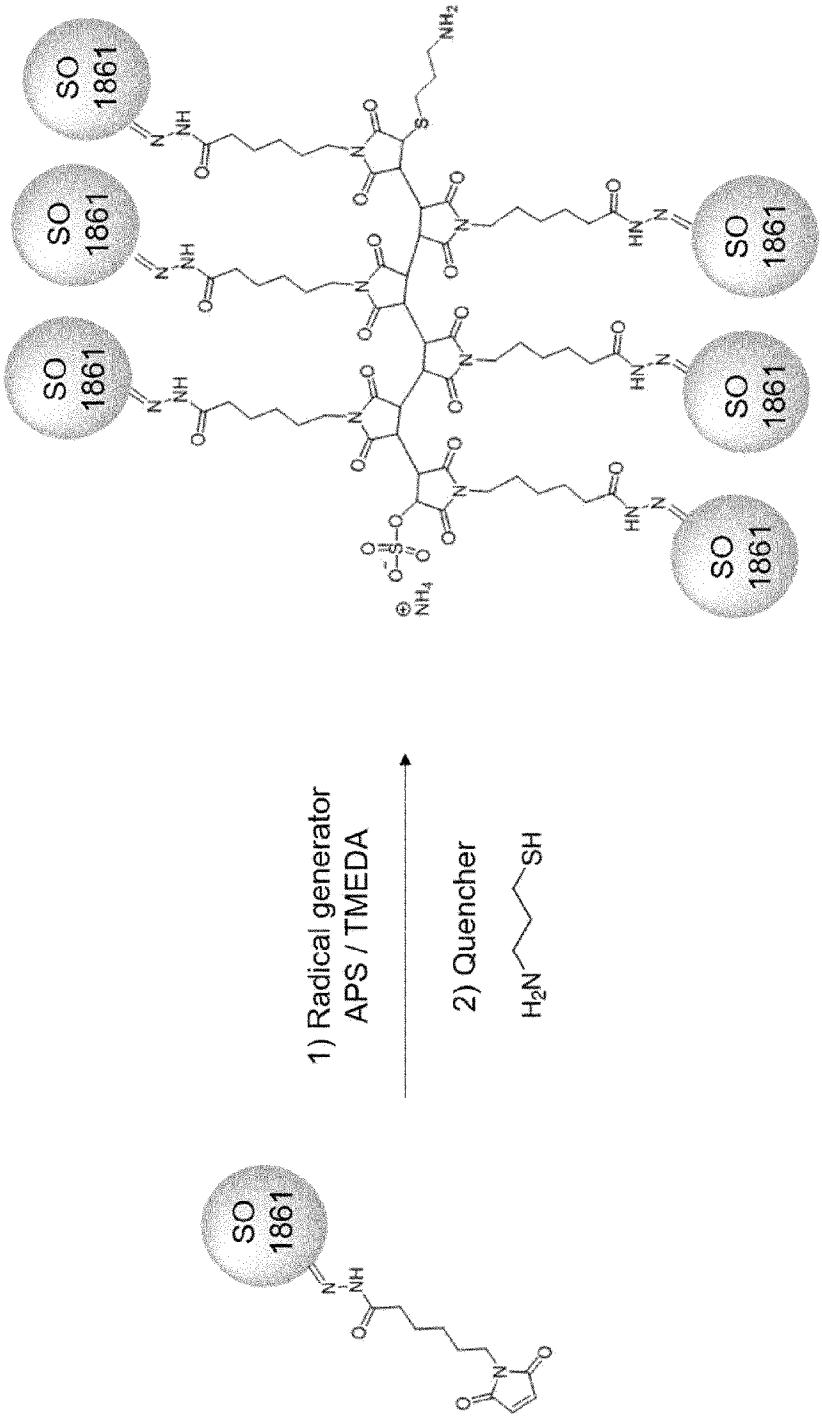


Figure 48

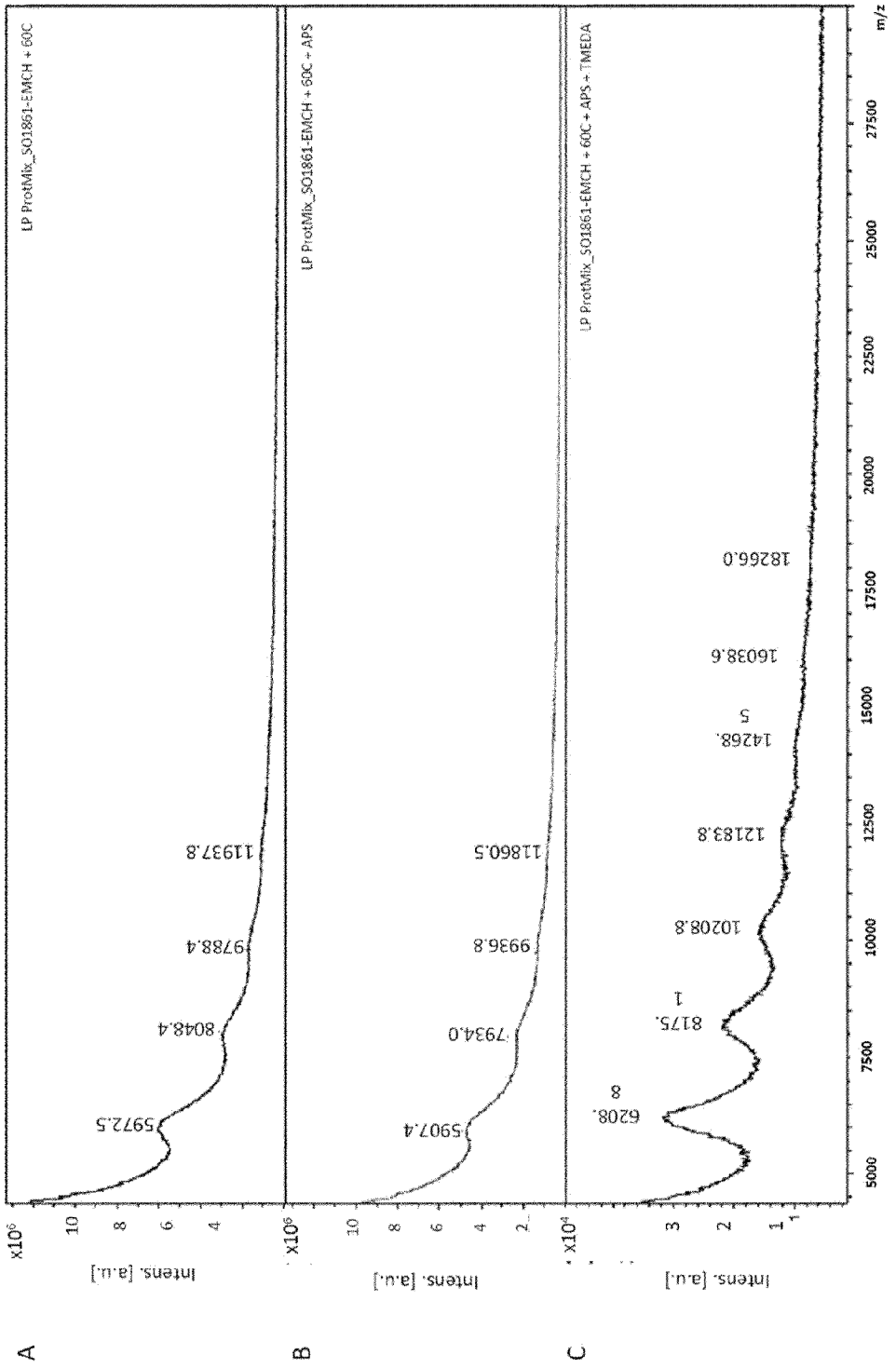

A
B
C
LP ProtMix_SO1861-EMCH + 60C
LP ProtMix_SO1861-EMCH + 60C + APS
LP ProtMix_SO1861-EMCH + 60C + APS + TMEDA
Intens. [a.u.]
x10^6
x10^4
m/z
5000
7500
10000
12500
15000
17500
20000
22500
25000
27500
5972.5
8048.4
9788.4
11937.8
5907.4
7934.0
9936.8
11860.5
6208.8
8175.1
10208.8
12183.8
14268.5
16038.6
18266.0

Figure 49

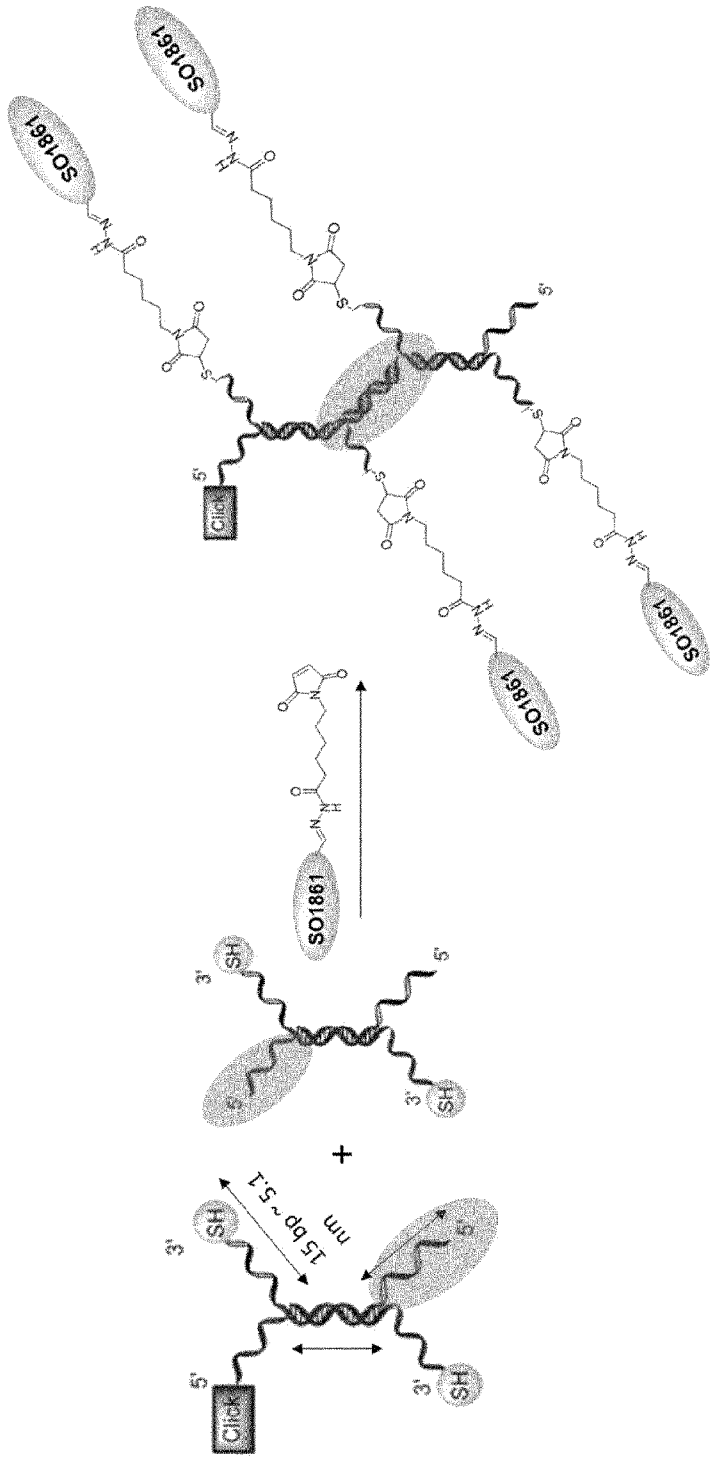

SO1861
SO1861
SO1861
SO1861
SO1861
Click
5'
SO1861
SH
3'
5'
5'
3'
SH
+
SH
3'
15 bp ~ 5.1
nm
5'
5'
Click
3'
SH

Figure 50

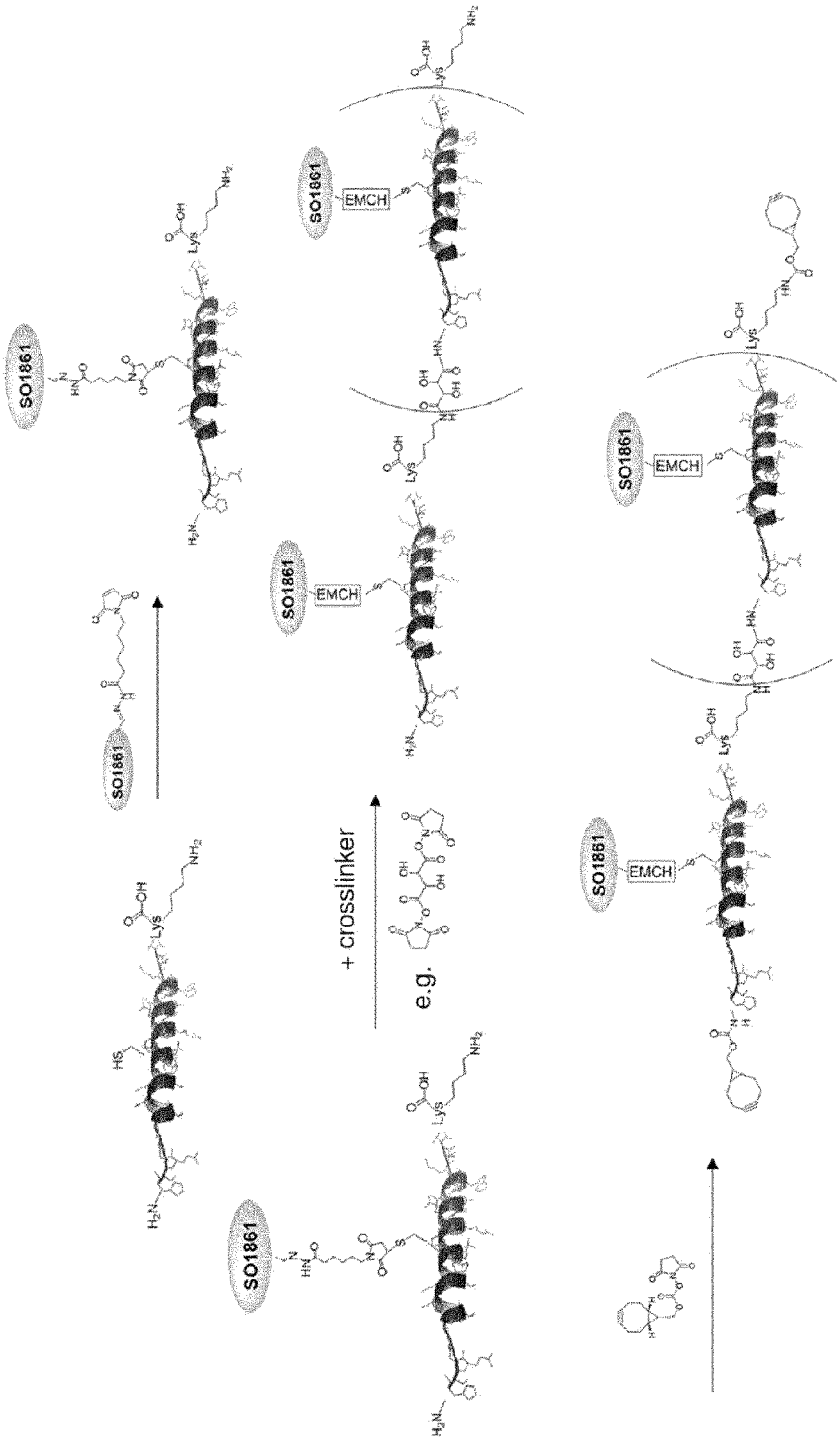


Figure 51

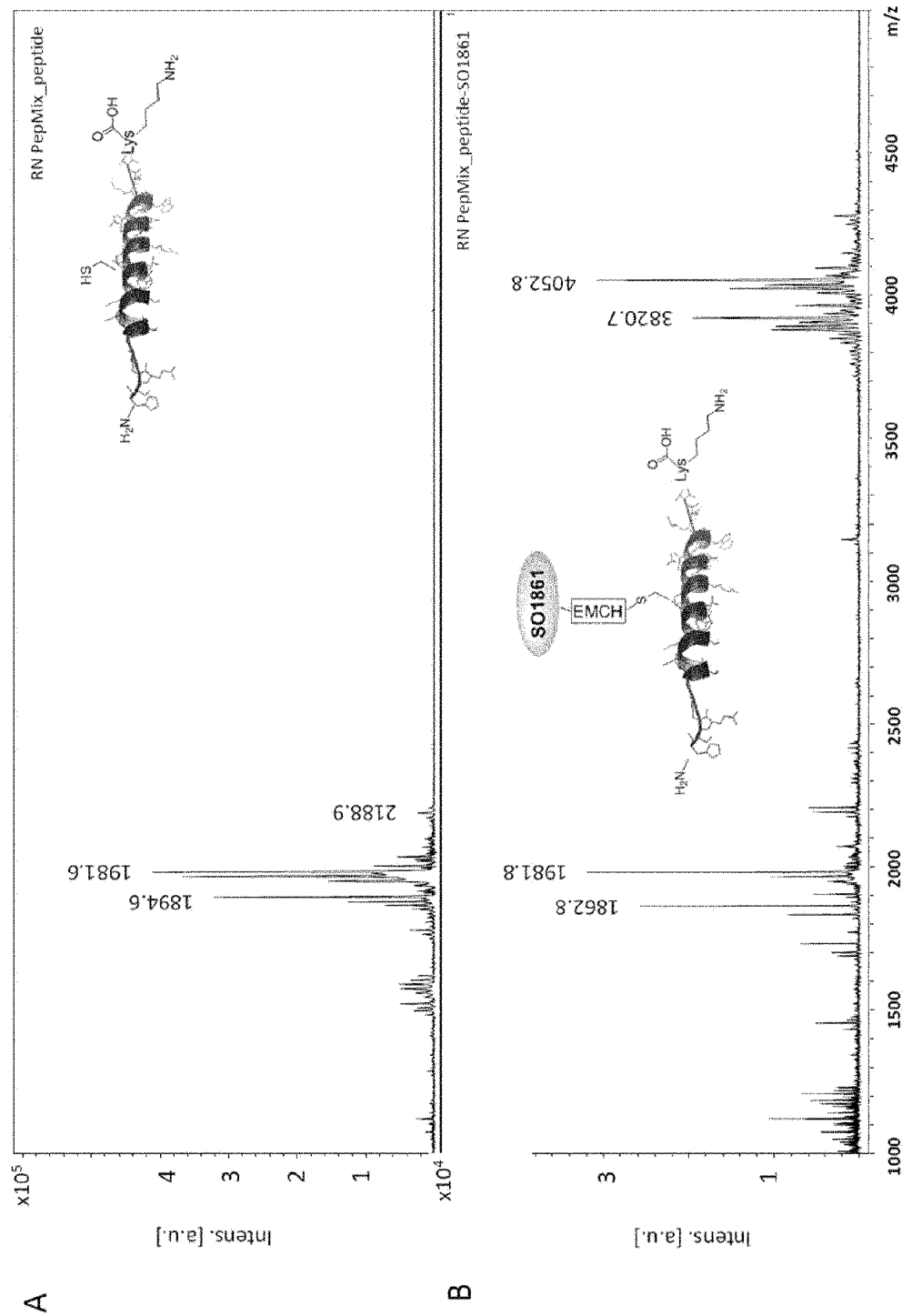

A
RN PepMix_peptide
Intens. [a.u.]
x10^5
x10^4
1000
1500
2000
2500
3000
3500
4000
4500
m/z
1894.6
1981.6
2188.9
B
RN PepMix_peptide-SO1861
1862.8
1981.8
3820.7
4052.8
SO1861
EMCH
Lys
HS

Figure 52

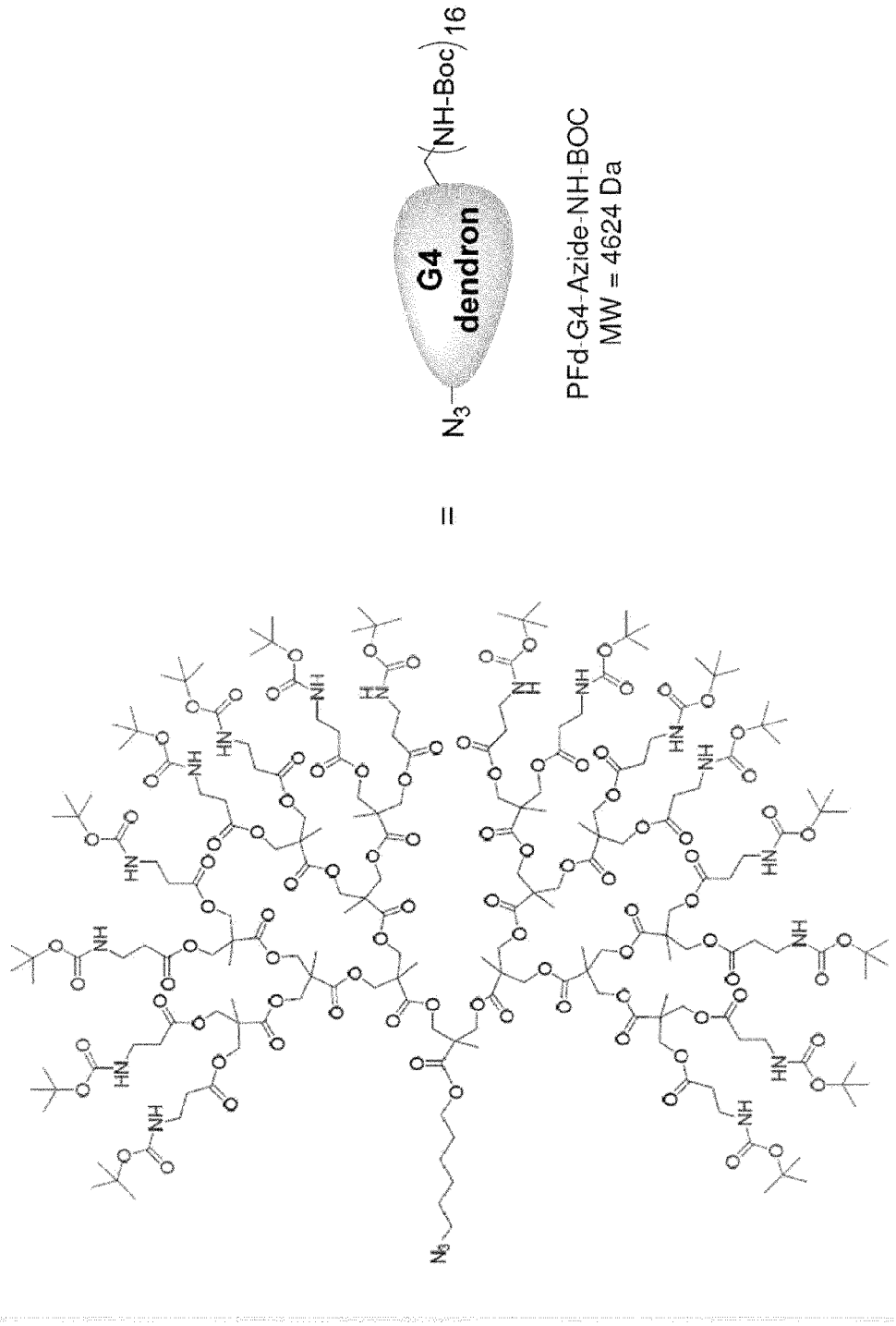


Figure 53

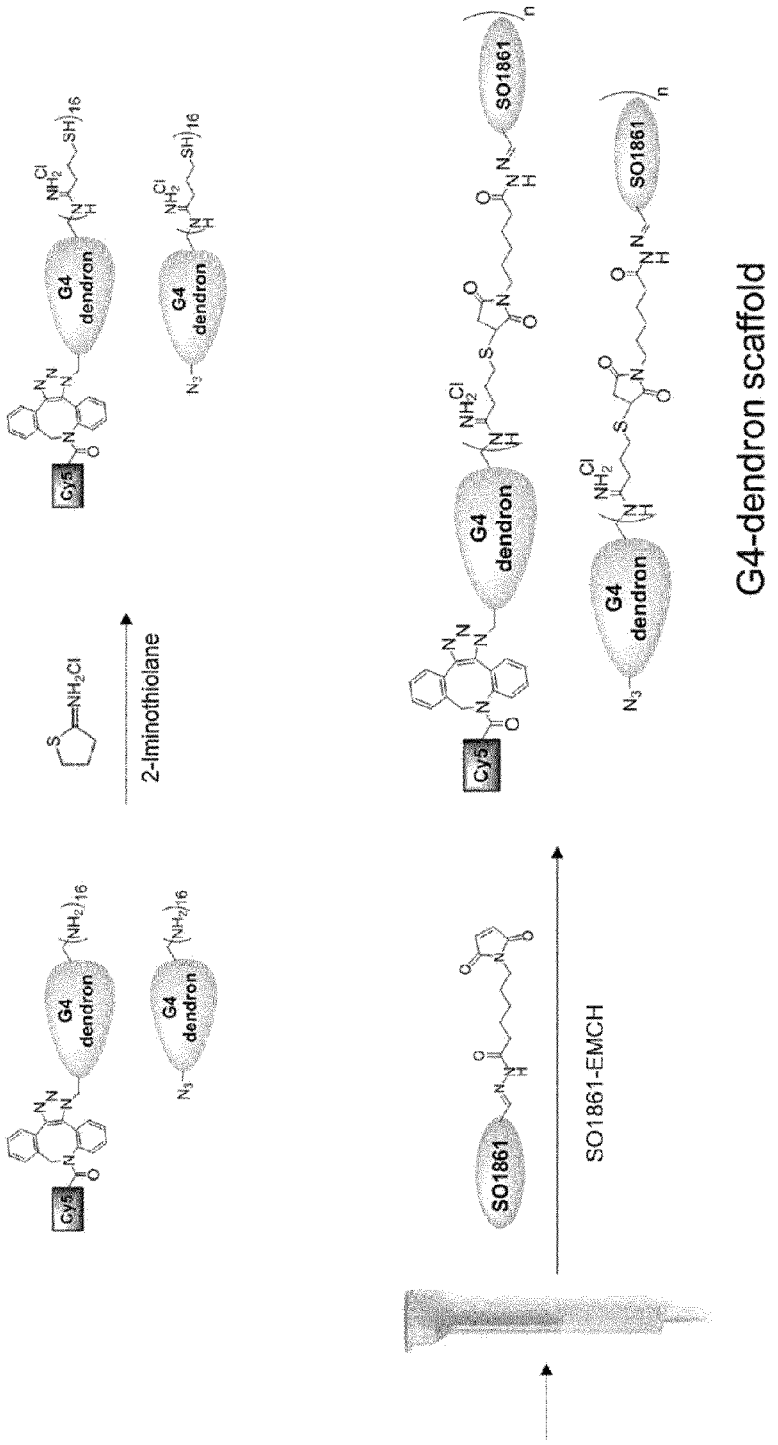

G4 dendron
Cy5
2-Iminothiolane
SO1861
SO1861-EMCH
G4-dendron scaffold

Figure 54

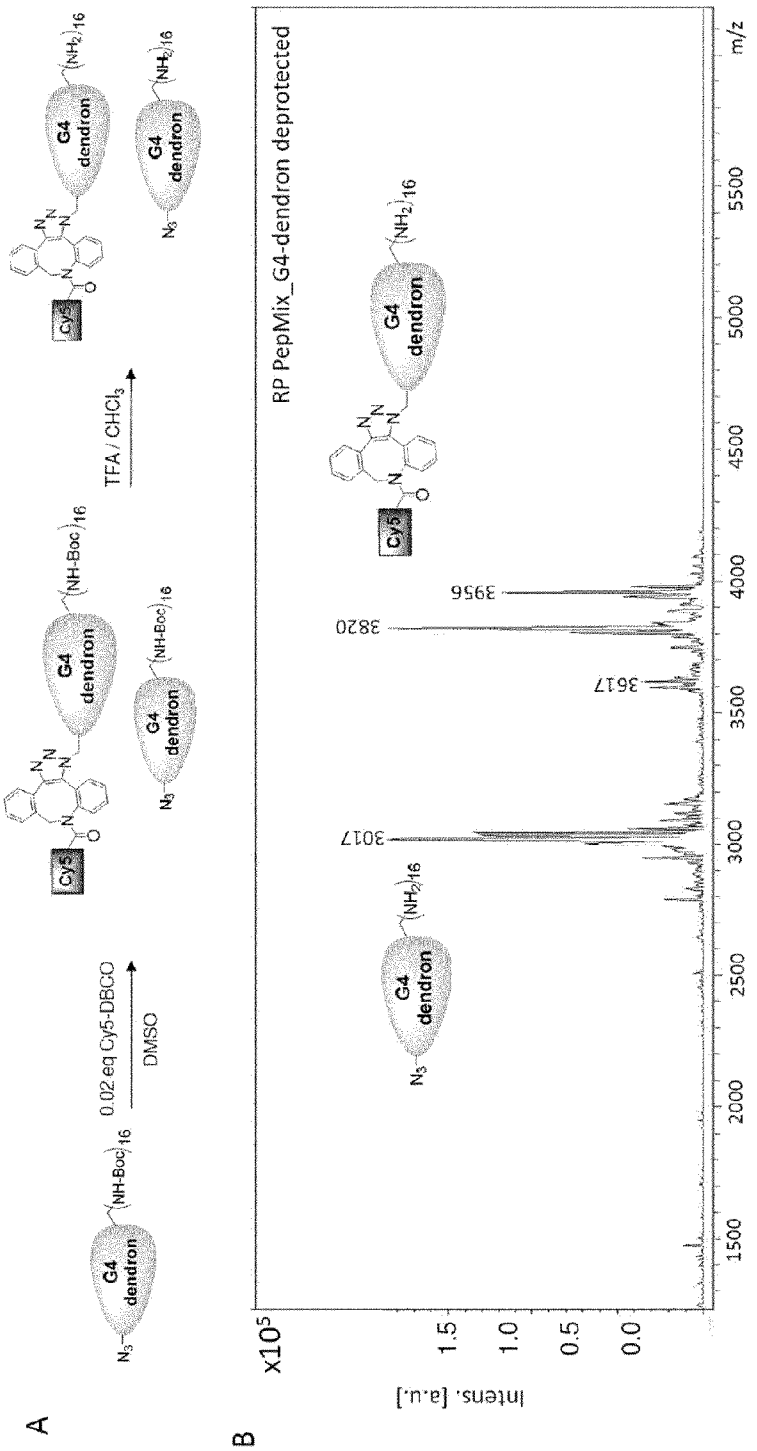

A
G4 dendron
(NH-Boc)16
N3
0.02 eq Cy5-DBCO
DMSO
Cy5
TFA / CHCl3
(NH2)16
B
RP PepMix_G4-dendron deprotected
x10^5
Intens. [a.u.]
1.5
1.0
0.5
0.0
1500
2000
2500
3000
3500
4000
4500
5000
5500
m/z
3017
3617
3820
3956

Figure 55

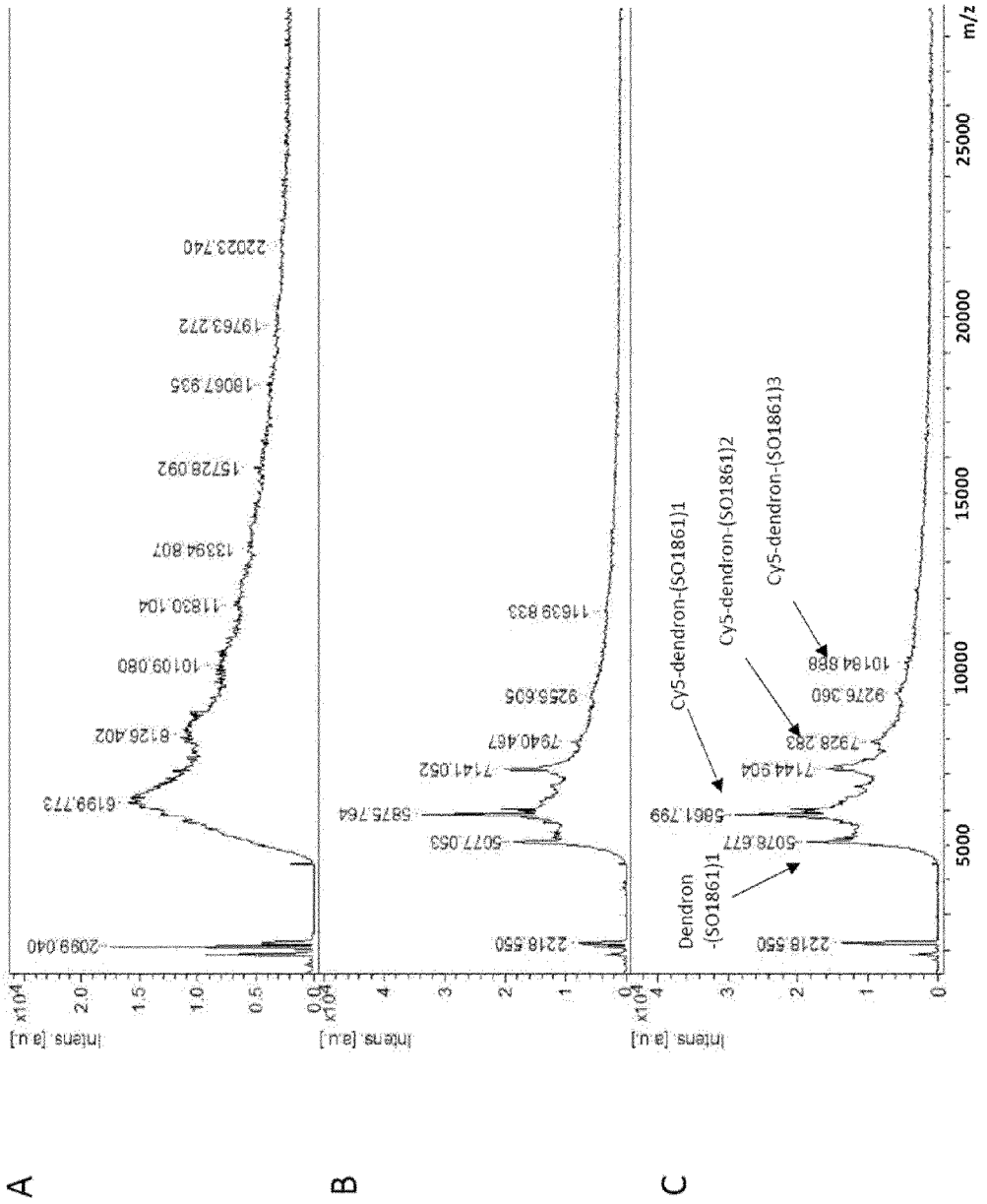

A
B
C
Intens. [a.u.]
m/z
2099.040
6199.773
8126.402
10109.080
11830.104
13394.807
15728.092
18067.935
19763.272
22023.740
2218.550
5077.053
5875.764
7141.052
7940.467
9255.635
11639.833
Dendron -(SO1861)1
Cy5-dendron-(SO1861)1
Cy5-dendron-(SO1861)2
Cy5-dendron-(SO1861)3
2218.550
5078.677
5861.799
7144.504
7928.283
9276.360
10184.868

Figure 56

A

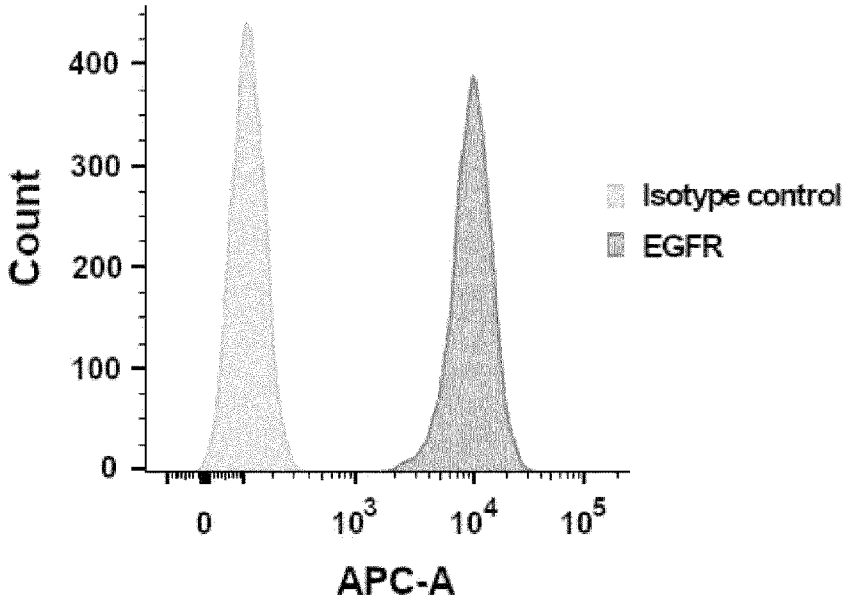


Figure 56

B

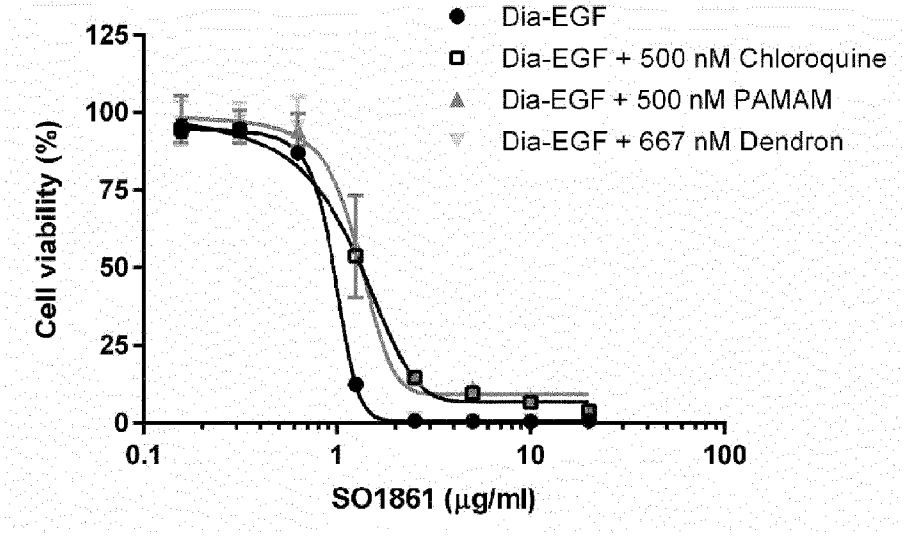


Figure 56

C

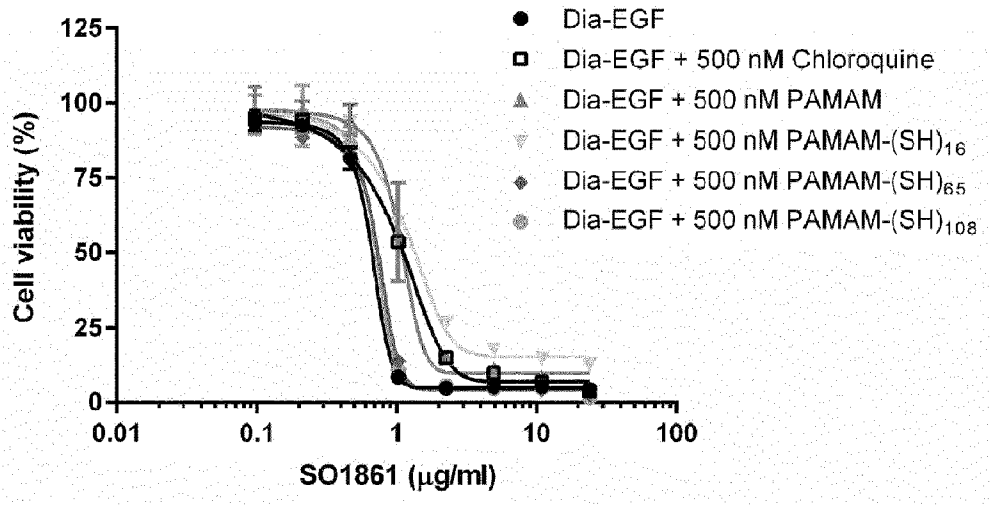


Figure 56

D

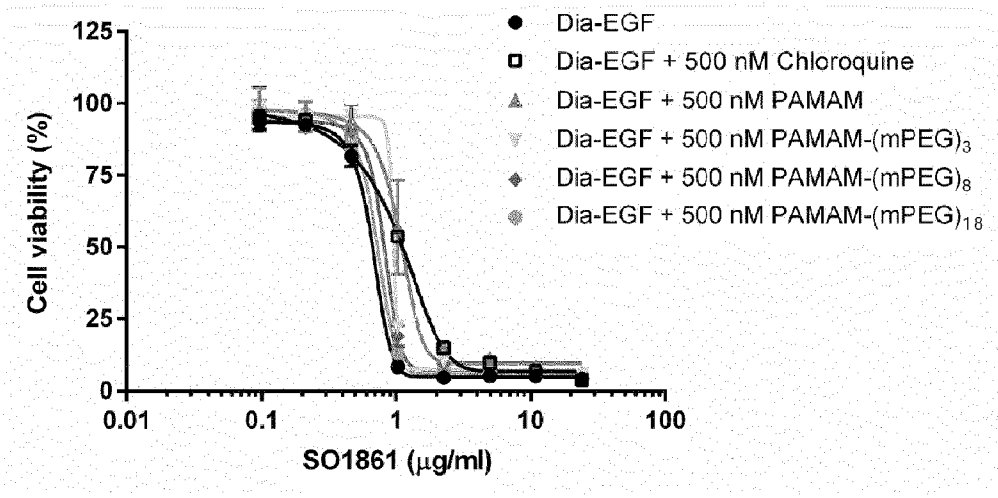


Figure 57

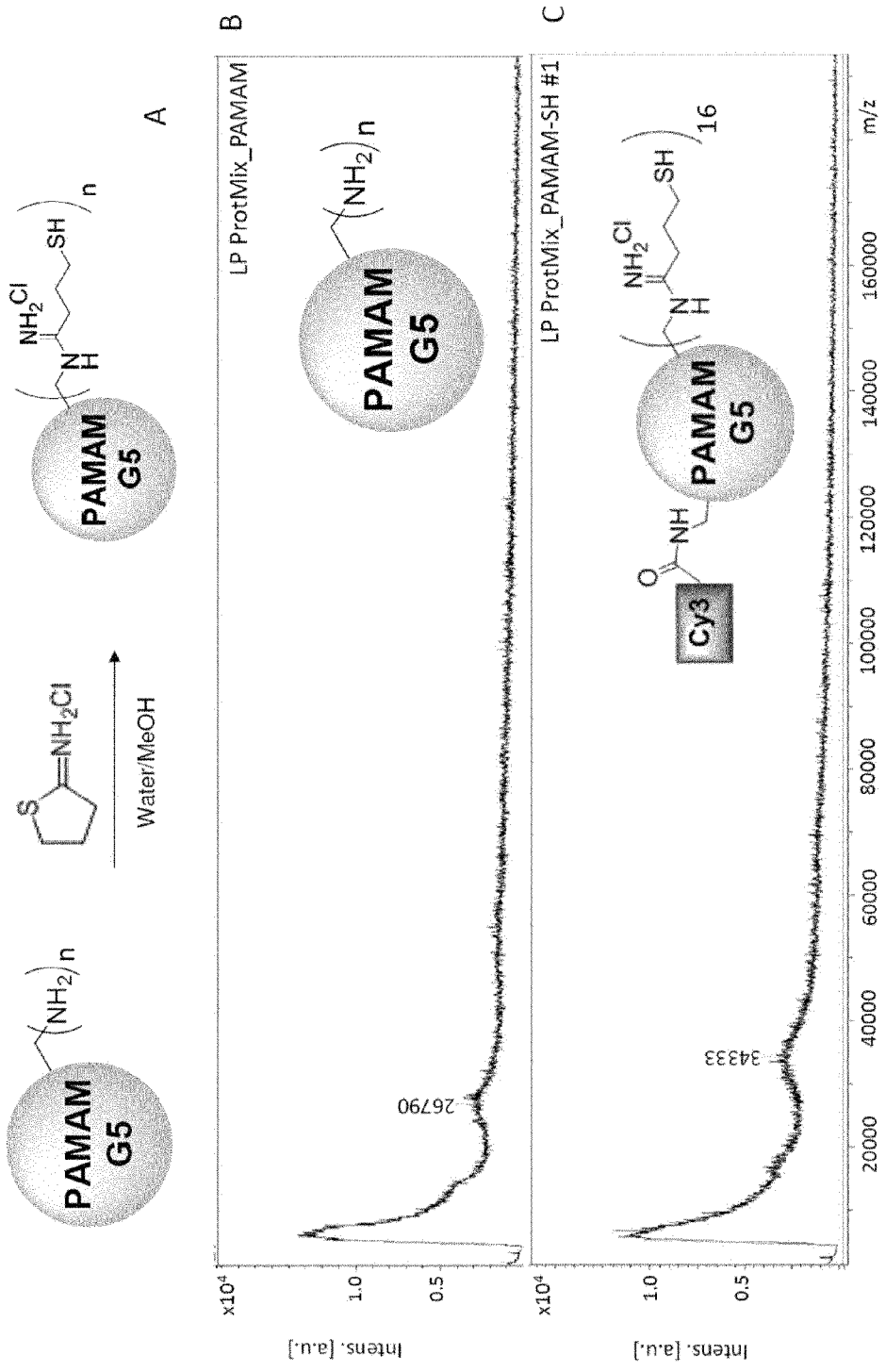

A
B
C
PAMAM G5
Water/MeOH
LP ProtMix_PAMAM
LP ProtMix_PAMAM-SH #1
26790
34333
Cy3
Intens. [a.u.]
m/z
20000
40000
60000
80000
100000
120000
140000
160000
0.5
1.0
x10⁴

Figure 57

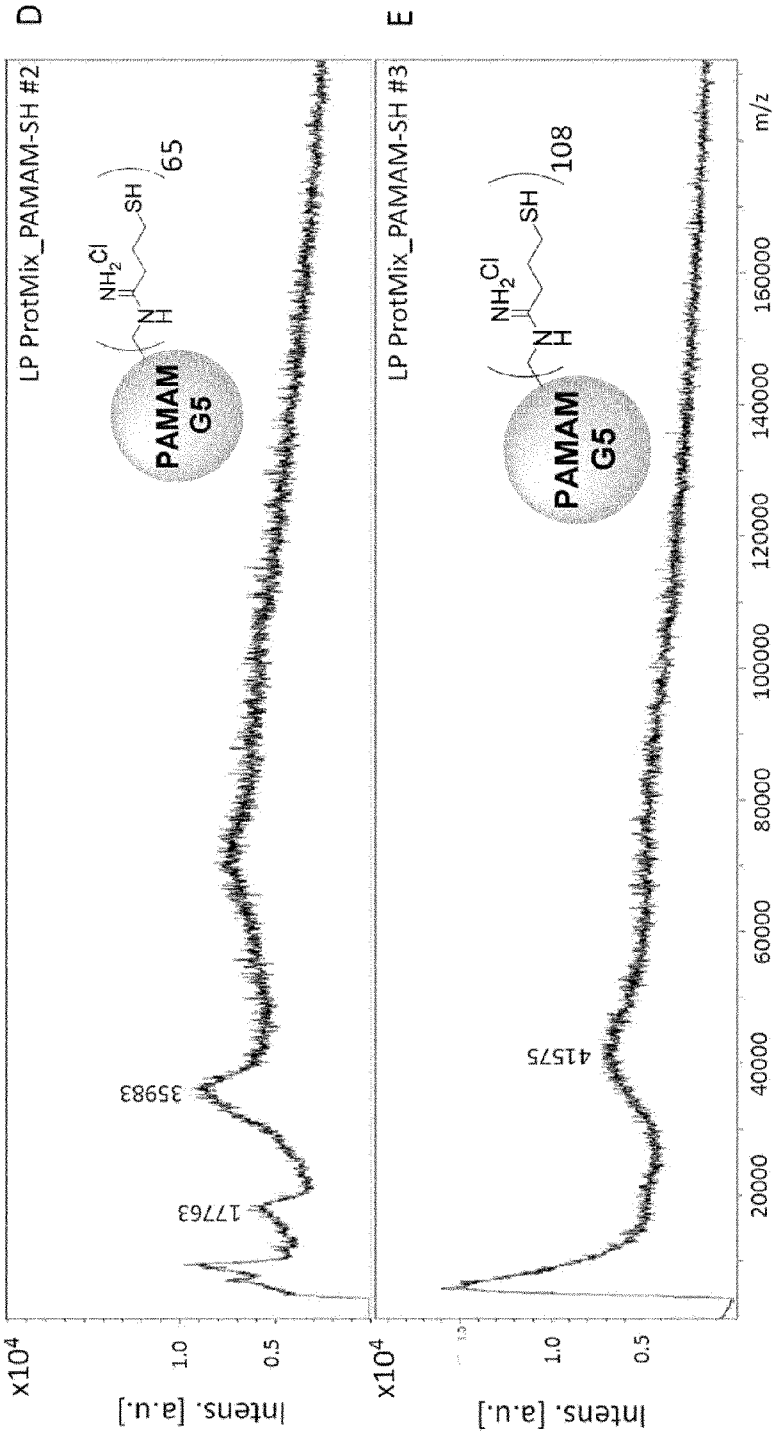

D
LP ProtMix_PAMAM-SH #2
PAMAM G5
65
35983
17763
x10^4
Intens. [a.u.]
E
LP ProtMix_PAMAM-SH #3
PAMAM G5
108
41575
x10^4
Intens. [a.u.]
m/z
20000
40000
60000
80000
100000
120000
140000
160000

Figure 58

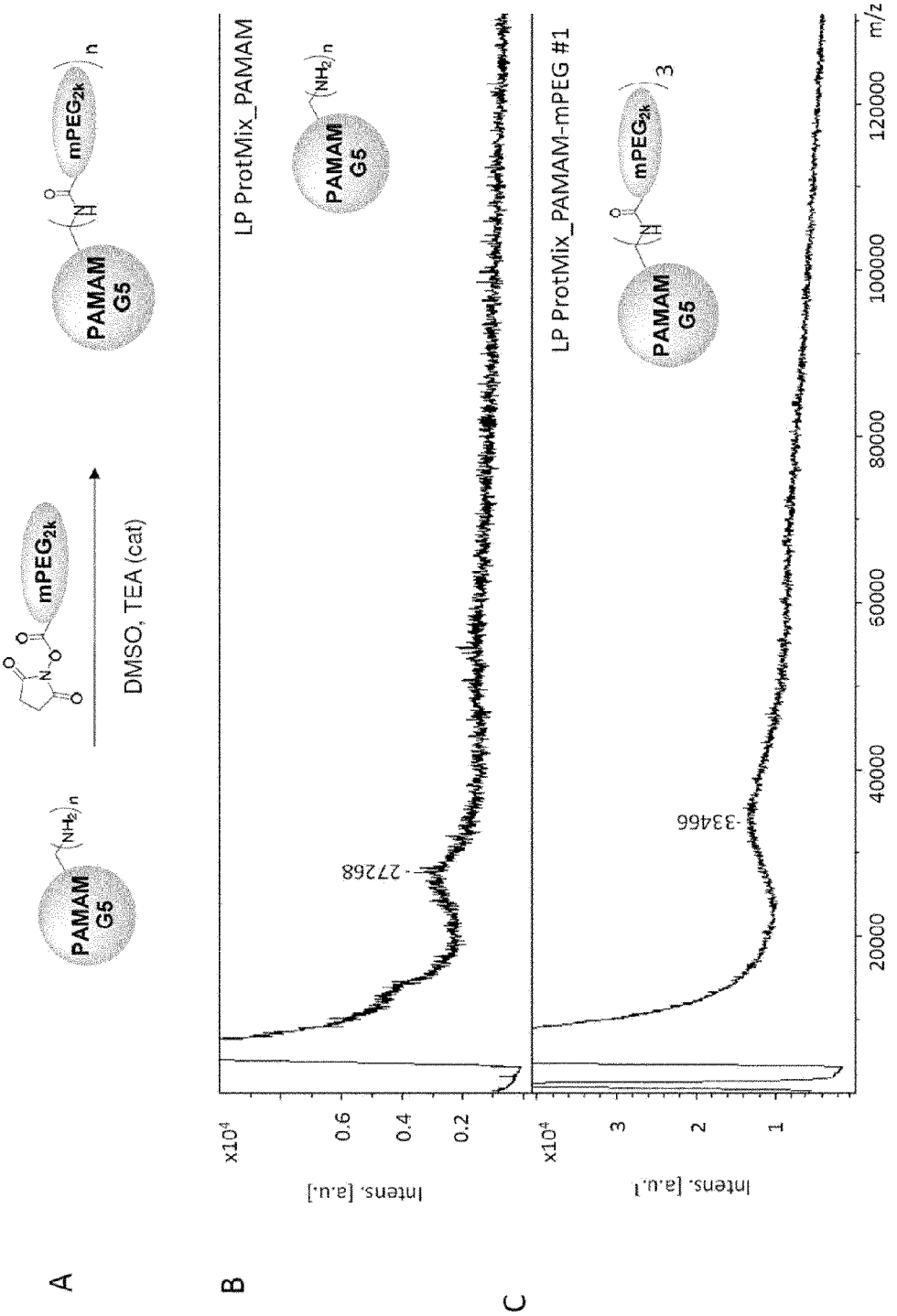

A
PAMAM G5
$(NH_2)_n$
DMSO, TEA (cat)
mPEG2k
PAMAM G5
mPEG2k
n
B
LP ProtMix_PAMAM
PAMAM G5
$(NH_2)_n$
27268
x10^4
0.6
0.4
0.2
Intens. [a.u.]
C
LP ProtMix_PAMAM-mPEG #1
PAMAM G5
mPEG2k
3
33466
x10^4
3
2
1
Intens. [a.u.]
20000
40000
60000
80000
100000
120000
m/z

Figure 58

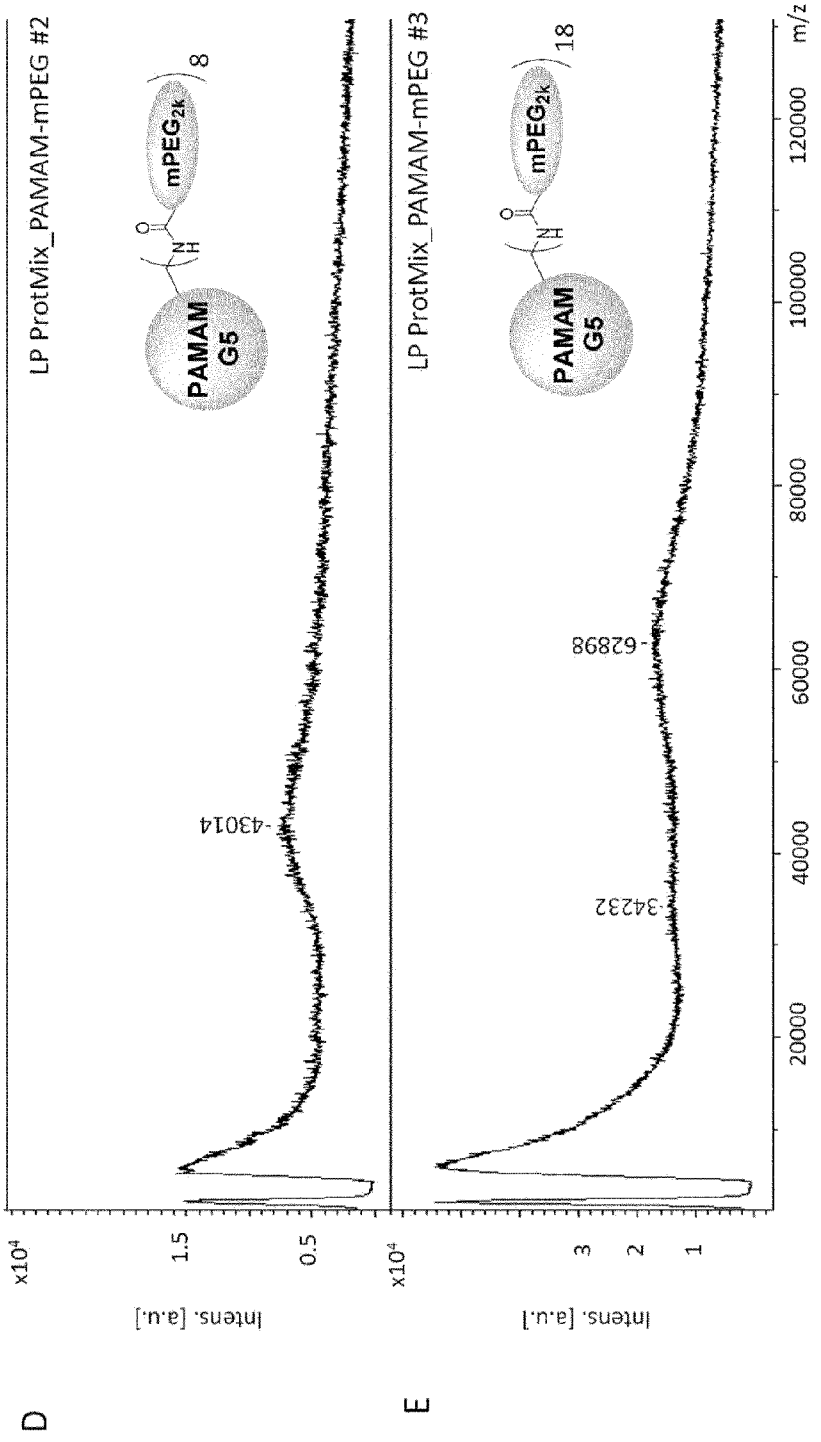

D
LP ProtMix_PAMAM-mPEG #2
PAMAM G5
mPEG2k
8
43014
Intens. [a.u.]
x10^4
1.5
0.5
E
LP ProtMix_PAMAM-mPEG #3
PAMAM G5
mPEG2k
18
62898
34232
Intens. [a.u.]
x10^4
3
2
1
20000
40000
60000
80000
100000
120000
m/z

FIGURE 59

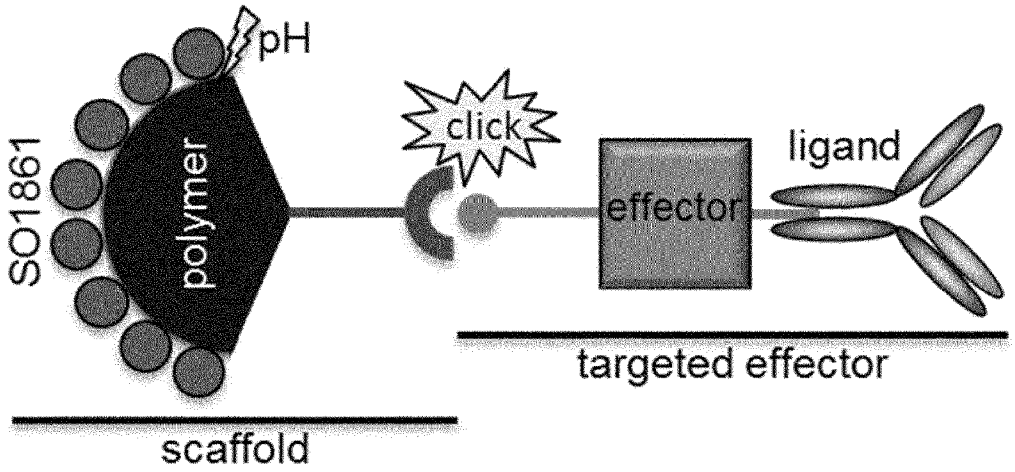
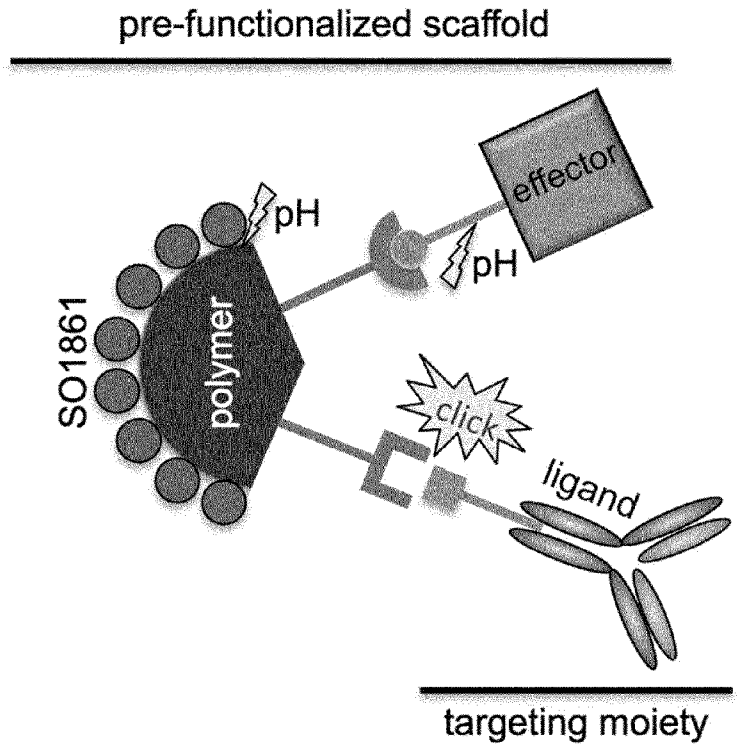


FIGURE 60



SEKVENSLISTE

Sekvenslisten er udeladt af skriftet og kan hentes fra det Europæiske Patent Register.

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