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(54) Title: COMPOUNDS WITH A BRANCHED LINKER

Ac-EUEUEU

Bi-UKKUEUEK

Ac-EUEUEU

(57) Abstract: The present invention concerns new compounds comprising a branched linker and their use for producing conjugates for applications in diagnostic or therapeutic methods.

Compounds with a branched linker

The present invention concerns new linkers and their use for producing conjugates for applications in diagnostic or therapeutic methods.

Conjugates comprising several binding groups or/and effector groups e.g. labeling or solid phase binding groups or toxins are often used in diagnostic or therapeutic methods. Such

5 conjugates can be prepared by direct coupling or by using the bridge or linker structures known in the prior art. Interfering intramolecular and intermolecular interactions between the conjugate partners or other components are often disadvantageous for the properties of such conjugates.

10 In diagnostic tests these undesired intramolecular and intermolecular interactions often lead to an impairment of important assay parameters such as the dynamic range of the signal, signal-to-noise ratio, breadth of the measuring range, blank value, lower limit of detection and thus to a considerable impairment of the assay. In therapeutic procedures the interactions in turn result in a reduction of efficacy or of target specificity.

15 The use of linkers that are known in the prior art for the conjugation of luminescent metal complexes (EP-A-0 178 450, EP-A-0 580 979, WO 87/06706) for example leads to a worsening of the dynamic range of an assay. Other disadvantages of such conjugates are a high unspecific binding to proteins and high blank values. However, similar problems also occur with other labeling and solid phase binding groups.

20 WO 96/03409 and WO 96/03410 disclose that the introduction of free positive or/and negative charge carriers in the linkers that link the reactive coupling group of the metal complex to one of the ligands or the introduction of hydrophilic groups into these luminescent metal complexes reduces the unspecific adsorption of conjugates of these complexes and thus improves the test sensitivity as well as the stability and recovery of the conjugates in immunoassays. Moreover in some cases it is possible to achieve an increase in 25 quantum yield.

Bredehorst, R., et al., Anal Biochem 193 (1991) 272-9 describe a trifunctional hapten-fluorophore conjugate which contains the 21 amino acid residues of the insulin A chain molecule as a backbone. The insulin A chain thus working as a linker between the fluorescent and hapten groups is a linear linker and not a branched linker.

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In recent investigations it was found that the use of hydrophilic or charged linkers according to WO 96/03409 or WO 96/03410 results in considerable advantages in test performance but even when using such complexes the blank value is considerably higher than the blank value of the system. Hence a further reduction of the blank value by 5 reducing unspecific binding would be desirable. In addition unspecific intramolecular and intermolecular interactions between the labeling group and other test components should be reduced without adversely affecting the signal yield and the accessibility of the labeling group.

It was surprisingly found that the said disadvantages can be eliminated by using branched 10 linkers with charged carriers or/and hydrophilic groups especially in the side chains. These branched linkers also result in improvements in other types of conjugates used in diagnostic or therapeutic methods or for screening purposes.

Hence a subject matter of the present invention is the use of a polyfunctional compound of the general formula (I):



in which Z denotes at least one reactive functional group or a binding group, X is a reactive functional group which is bound covalently to Z via a linker Y where the linker is a branched linker which has a molecular weight of ≥ 1000 Da and contains at least one charge carrier or/and at least one hydrophilic group, n is an integer from 1 to 10 and 20 preferably from 1 to 4 and m is 1 or 2 and preferably 1, for the production of conjugates.

The group Z can occur once or several times and can in each case be independently a reactive functional group or a binding group. Examples of binding groups are labeling groups or effector groups. Effector groups are for example partners of a bioaffine binding pair which can specifically interact with the other partner of the bioaffine binding pair.

25 The labeling groups can be selected from any detectable known groups such as dyes, luminescent labeling groups such as chemiluminescent groups e.g. acridinium esters or dioxetanes or fluorescent dyes e.g. fluorescein, coumarin, rhodamine, oxazine, resorufin, cyanine and derivatives thereof. Other examples of labeling groups are luminescent metal complexes such as ruthenium or europium complexes, enzymes as used for CEDIA 30 (Cloned Enzyme Donor Immunoassay, e.g. EP 0 061 888), microparticles or nanoparticles e.g. latex particles or metal sols, and radioisotopes.

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In a preferred embodiment the labeling group is a luminescent metal complex and the compound has a structure of the general formula (II):



in which M is a divalent or trivalent metal cation selected from rare earth or transition metal ions, L₁, L₂ and L₃ are the same or different and denote ligands with at least two nitrogen-containing heterocycles in which L₁, L₂ and L₃ are bound to the metal cation via nitrogen atoms, X is a reactive functional group which is covalently bound to at least one of the ligands L₁, L₂ and L₃ via a linker Y, n is an integer from 1 to 10, preferably 1 to 4, m is 1 or 2 and preferably 1 and A denotes the counterion which may be required to equalize the charge.

The metal complex is preferably a luminescent metal complex i.e. a metal complex which undergoes a detectable luminescence reaction after appropriate excitation. The luminescence reaction can for example be detected by fluorescence or by electrochemiluminescence measurement. The metal cation in this complex is for example a transition metal or a rare earth metal. The metal is preferably ruthenium, osmium, rhenium, iridium, rhodium, platinum, indium, palladium, molybdenum, technetium, copper, chromium or tungsten. Ruthenium, iridium, rhenium, chromium and osmium are particularly preferred. Ruthenium is most preferred.

The ligands L₁, L₂ and L₃ are ligands with at least two nitrogen-containing heterocycles. Aromatic heterocycles such as bipyridyl, bipyrazyl, terpyridyl and phenanthrolyl are preferred. The ligands L₁, L₂ and L₃ are particularly preferably selected from bipyridine and phenanthroline ring systems.

The complex can additionally contain one or several counterions A to equalize the charge. Examples of suitable negatively charged counterions are halogenides, OH⁻, carbonate, alkylcarboxylate, e.g. trifluoroacetate, sulphate, hexafluorophosphate and tetrafluoroborate groups. Hexafluorophosphate, trifluoroacetate and tetrafluoroborate groups are particularly preferred. Examples of suitable positively charged counterions are monovalent cations such as alkaline metal and ammonium ions.

On the other hand the group Z can be an effector group which interacts specifically and preferably non-covalently with a binding partner. Examples of suitable binding partners are hapten or antigen/antibody, biotin or biotin analogues such as aminobiotin, iminobiotin or

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desthiobiotin/avidin or streptavidin, sugar/lectin, nucleic acid or nucleic acid analogue/complementary nucleic acid, receptor/ligand e.g. steroid hormone receptor/steroid hormone in which one partner of the binding pair is the effector group and thus a component of compound (I).

- 5 In a further preferred embodiment Z can also be a therapeutically active substance e.g. a toxin or protoxin e.g. an anti-tumour substance.

The compounds (I) are used as linker molecules to prepare conjugates. In this process a binding partner and in particular a binding partner as stated above is coupled covalently to the at least one free functional group of the compound (I).

- 10 The resulting coupling product contains at least two, preferably different, binding groups which are linked together via the branched linker Y.

The reactive functional group X or Z of the compound (I) or of the complex (II) is a reactive group which can be coupled covalently to a biological substance. The group X is preferably an activated carboxylic acid group such as a carboxylic acid halogenide, a carboxylic acid anhydride, a carboxylic acid hydrazide, a carboxylic acid azide or an active ester e.g. an N-hydroxy-succinimide, a p-nitrophenyl, pentafluorophenyl, imidazolyl or N-hydroxybenzotriazolyl ester, an amine, a maleimide, a thiol or a photoactivatable group e.g. an azide. The compound can contain one or several functional groups X or Z which can be the same or different. X and Z are preferably different. If Z is a functional group, it preferably only occurs once. If Z is a binding group, it can be present several times e.g. up to 10 times. The functional groups or binding groups Z, respectively can be the same or different and be optionally blocked by protective groups. However, the total number of groups X plus Z is at least 2 i.e. the compound is at least a bifunctional compound, preferably at least a hetero-bifunctional compound. Appropriate active groups for hetero-bifunctional linkers are described in Aslam M., Dent A., Bioconjugation (1998) Mcmillan Reference Ltd., London, p 216-363.

The molecular weight of the linker is at least 1000 Da, because then the advantages of the linker become particularly apparent. The molecular weight of the linker is preferably in the range of 1000 to 50,000 Da, particularly in the range of 1000 to 20,000 Da and most preferably in the range of 1000 to 10,000 Da.

The compound (I) and the metal complex (II) differ from the prior art in that the linker Y between X and Z is a branched linker with at least one charge carrier or/and at least one hydrophilic group. In the sense of the present invention the term "charge carrier" means a group which is present mainly in an ionic form at a pH value in the range 6 to 8. The linker 5 preferably contains up to 70, particularly preferably 1 to 40 and most preferably 2 to 20 such charge carriers.

The linker particularly preferably contains at least one negative charge carrier. Examples of suitable negative charge carriers are phosphate, phosphonate, sulphinate, sulphonate, sulphate and carboxylate groups, carboxylate groups and phosphate groups being most 10 preferred.

Examples of positive charge carriers are amino and mono-substituted or polysubstituted amino groups such as mono-, di- or trialkyl amino groups, in which alkyl denotes a straight-chained or branched alkyl residue with 1 to 6 C atoms or a cyclic alkyl residue with 3 to 6 C atoms. The positive charge carriers are particularly preferably selected from basic 15 amino acids such as lysine or substituted amino acids such as diethyllysine, or dipropyllysine. Amines and substituted amines can also be used as electron donors for the detection of metal complexes by electrochemiluminescence.

The linkers can also contain uncharged hydrophilic groups as an alternative to or in addition to the charge carriers. Preferred examples of uncharged hydrophilic groups are 20 ethylene oxide or polyethylene oxide groups with preferably at least three ethylene oxide units, sulphoxide, sulphone, carboxylic acid amide, carboxylic acid ester, phosphonic acid amide, phosphonic acid ester, phosphoric acid amide, phosphoric acid ester, sulphonic acid amide, sulphonic acid ester, sulphuric acid amide and sulphuric acid ester groups. The amide groups are preferably primary amide groups, particularly preferably carboxylic acid 25 amide residues in amino acid side groups e.g. the amino acids asparagine and glutamine. The esters are preferably derived from hydrophilic alcohols, in particular C₁-C₃ alcohols or diols or triols.

In the sense of the present invention the term "branched" means that the linker contains a main chain between the groups Z and X and in addition one or several side chains starting 30 from the main chain. The charge carriers and hydrophilic groups can be located in the main chain or/and in a side chain. If the linker according to the invention contains several groups Z and X, the main chain itself can already be branched. However, the linker in any case additionally contains one or several side chains which contain none of the groups Z

and X. The number of side chains is preferably 1 to 10, particularly preferably 2 to 6 and most preferably 2 to 4.

In a preferred embodiment of the invention the linker contains a main chain which contains one or several uncharged hydrophilic groups as mentioned above in particular 5 carboxylic acid amide groups or/and polyethylene glycol groups while there is at least one charge carrier in one or several of the side chains. In this case 1 to 10 charge carriers and in particular 1 to 5 charge carriers can for example be present per side chain. Alternatively the linker can also contain charge carriers in the main chain and uncharged hydrophilic groups in one or several side chains. Furthermore embodiments are also conceivable in which the 10 main chain and the side chains contain uncharged hydrophilic groups as well as charge carriers.

In case the branched linker comprises groups other than X and Z, which might interfere with the intended coupling chemistry, e.g., like $-COO^-$ groups or $-NH_2$ groups, appropriate protective groups, which are known to the skilled artisan are used during synthesis and/or 15 coupling. Terminal $-NH_2$ groups in peptidic side chains preferably are inactivated, e.g., by acetylation or succinylation.

The length of the main chain of the linker is preferably 7 to 200 atoms, particularly preferably 7 to 100 atoms. The main chain is an alkylene chain modified by the incorporation of heteroatoms e.g. O atoms or amide groups and contains at least one 20 branch site. The side chains formed at the branching site preferably having a length of 4 to 100 atoms.

The charge carriers are preferably located in the linker in such a manner that a H atom of an alkylene unit of the main chain or/and in a side chain is replaced by a group containing a charge carrier e.g. NH_3^+ or CO_2^- .

25 The branched linker which contains the free charge carriers or/and hydrophilic groups is preferably at least partially composed of aminocarboxylic acid units that are linked together by peptide bonds. In such a linker the branching points can be derived from polyfunctional aminocarboxylic acids which contain at least three functional groups e.g. amino or carboxylate groups such that one functional group is still present after incorporation into 30 the main chain which can be used as the starting point for the synthesis of the side chain. The branches are particularly preferably generated with diaminocarboxylic acids such as lysine, ornithine, hydroxylysine etc.

The charge carriers of the branched linker can be preferably derived from free positively or/and negatively charged groups of polyfunctional amino-carboxylic acids which contain a total of at least three charged groups e.g. amino, carboxylate or phosphate groups such that after incorporation into the linker and the concomitant reaction of two of the charged 5 groups, at least one free charge carrier is still present. For example the charge carriers can be derived from trifunctional aminocarboxylic acids which contain (a) an amino group and two carboxylate groups or (b) two amino groups and one carboxylate group. Examples of such trifunctional aminocarboxylic acids are lysine, ornithine, hydroxylysine, aspartic acid and glutamic acid, symmetric trifunctional carboxylic acids like those described in EP 0 618 10 192 or US 5,519,142. Alternatively one of the carboxylate groups in the trifunctional aminocarboxylic acids (a) can be replaced by a phosphate, sulphonate or sulphate group. An example of such a trifunctional amino acid is phosphoserine.

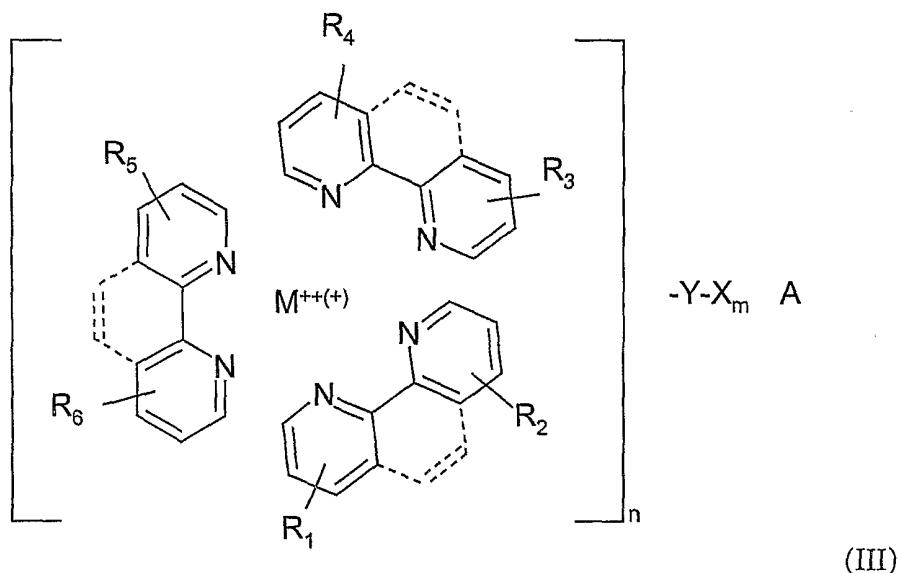
Alternatively the branched linker can also be composed at least partially of phosphate-sugar units e.g. a DNA backbone without nucleobases or composed of glyco-peptidic structures. 15 Furthermore the linker can also be composed at least partially of saccharide units. In any case the side chain of the linker is preferably situated at a branch site of the main chain which is formed by a trifunctional unit and the length of a side chain is at least two of the building blocks used for the synthesis e.g. natural or synthetic amino acids or other components such as ethylene glycol.

20 Preferably, the branched hydrophilic linker is used in immunological procedures to reduce problems caused by non-specific binding.

It has been found advantageous to use non-naturally occurring amino acids and non-naturally occurring sequence motives, like e.g., the di-peptides UE or UQ to construct a linker molecule according to the present invention. This has proven especially 25 advantageous in serological assays, i.e. in assays set up to detect antibodies in patient sera. Linkers with non-naturally occurring β -amino acids have proven rather stable against proteases, e.g., against proteases in serum or plasma, and therefore represent a further preferred embodiment of the present invention.

In one embodiment of the present invention the branched linker comprising a metal 30 complex has the general formula (III):

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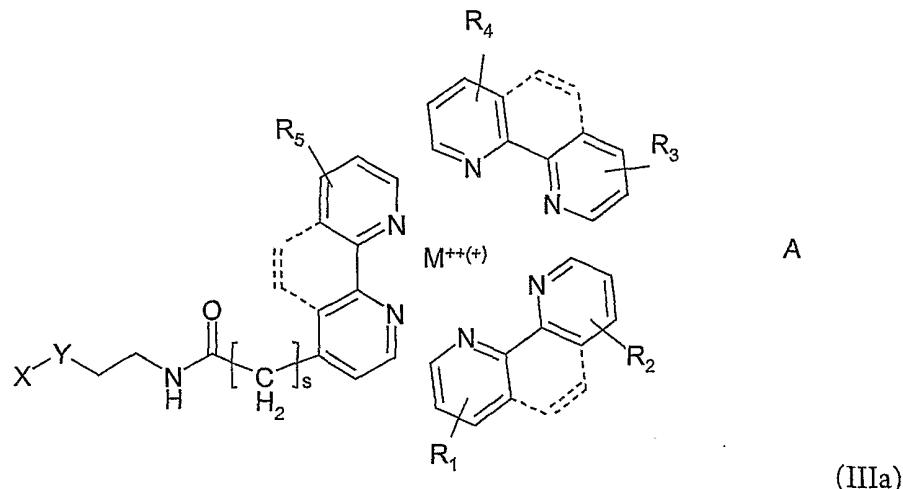


in which M, X, A, n and m are defined as above, R₁, R₂, R₃, R₄, R₅ and R₆ are the same or different and each denotes one or several substituents provided that X is bound to one of the ligands via one of the substituents R₁, R₂, R₃, R₄, R₅ or R₆ and the linker Y.

The ligands of the complex are optionally substituted phenanthroline or bipyridene systems depending on whether the groups denoted by the broken lines are present or not.

The substituents R₁, R₂, R₃, R₄, R₅ and R₆ on the ligands are preferably hydrogen, C₁-C₅ alkyl, in particular C₁-C₃ alkyl, phenyl or a hydrophilic group as defined above provided they do not contain the linker Y.

In a particularly preferred embodiment the branched linker comprising a metal complex has the general formula (IIIa):



in which M, X and A are defined as above, R₁, R₂, R₃, R₄, R₅ are as defined above, s is an integer from 0 to 6 preferably from 1 to 4 and Y denotes the branched linker with free charge carriers or/and hydrophilic groups.

Examples of compounds of formula (I) with metal complexes as labeling groups or biotin

5 as an effector group and the amino side group of lysine as the reactive functional group are shown in figs. 1-7. The branching points of the linker are formed by the trifunctional amino acid lysine which has two amino groups and one carboxylate group. One amino group and one carboxylate group are used to form peptide bonds in the main chain of the linker whereas the second amino group is used as the starting point for the synthesis of the

10 side chain. The free charge carriers are formed from glutamic acid side chains. In Fig. 8 the lysine amino group is blocked by phenylacetyl (Phac). The blocking of one or several reactive groups in the linker by protective groups e.g. phenylacetyl or/and other protective groups compatible with the overall structure also enables labile binding groups to be introduced into the compound (I).

15 The reactive carboxylate group can for example be converted into an active ester by reaction with N-hydroxysuccinimide or disuccinimidyl suberate (DSS) (see Fig. 12). Alternatively the primary amino group can also be converted into a maleimide group by reaction with maleimidohexanoyl-N-hydroxysuccinimide ester (MHS) (see figs. 9 - 11).

20 The preparation of the compounds according to the invention is described in detail in the following using metal complexes as an example. Other compounds that for example contain biotin or peptide antigens as the effector group can be prepared in an analogous manner.

25 The synthesis of a charged and branched linker on a labeling group or solid phase binding group such as the N-heterocyclic ligand of a metal complex can be carried out as a coupling reaction in solution by coupling an optionally partially protected aminocarboxylic acid to a reactive group of the ligand e.g. a carboxylic acid. This coupling stage can optionally be repeated until a branched linker of the desired length has been synthesized. In this process at least one polyfunctional aminocarboxylic acid is introduced which contains one or several charged side groups.

30 Subsequently the reactive group X is introduced and protective groups that may be present on the side groups of the aminocarboxylic acids are cleaved off. This synthesis of the ligand by successively coupling amino acids in solution can take place on a single ligand and also

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on a ligand that has already been bound to a metal complex as the starting material. A suitable starting material is for example a luminescent metal complex which contains a free carboxylate group. Such complexes are known from the above-mentioned documents and are also offered commercially for example by the Igen Inc. Company, Rockville, MD, USA.

5 On the other hand the branched linker can also be prepared by solid phase peptide synthesis. In a first embodiment of the solid phase synthesis, an amino acid is coupled via its carboxylate group to the solid phase support and then the desired linker is synthesized by successively coupling further amino acids. In this process at least one amino acid which contains a charged group as a side group e.g. an amino group or a carboxylate group and at 10 least one amino acid which serves as the branching site and is optionally in a protected form are used to prepare a linker according to the invention. After completion of the desired linker sequence, an activated metal complex, e.g. an active ester, can be coupled to the free N-terminal amino group of the peptide bound to the solid phase. After cleavage from the solid phase the reactive group X can be coupled to the carboxy terminus of the 15 peptide linker and protective groups that may be present are cleaved off.

In another mode of the solid phase synthesis an amino acid-metal complex conjugate which contains a protected amino group and a carboxylate group e.g. Fmoc-Lys(-Ru(bipyridyl)₃-OH) can be anchored to a solid phase by means of the free carboxylate group and a peptide linker can be synthesized after release of the blocked amino group. 20 After completion of the desired linker sequence, the complex is cleaved from the solid phase to obtain a linker which contains at least the original carboxylate anchor group as the free charge carrier. The reactive group X can be coupled to the amino terminus of the resulting peptide linker.

25 In a further procedure of solid phase synthesis the branched linker sequence with charge carriers can also be synthesized directly on a selected peptide epitope.

A combination of the above-mentioned synthesis variants can also be used to prepare the compounds according to the invention. Amino acid-metal complex conjugates that are suitable for the solid phase synthesis of the complexes according to the invention with a charged linker are described in DE 44 30 998.8. Reference is herewith made to this 30 disclosure.

A further subject matter of the present invention is a compound of the general formula (I) as defined above.

Yet a further subject matter of the present invention is a conjugate comprising at least one biological substance to which at least one compound (I) according to the invention is coupled. Examples of suitable biological substances are cells, viruses, subcellular particles, proteins, lipoproteins, glycoproteins, peptides, polypeptides, nucleic acids, peptidic nucleic acids (PNA), oligosaccharides, polysaccharides, lipopoly-saccharides, cellular metabolites, haptens, hormones, pharmacological substances, alkaloids, steroids, vitamins, amino acids and sugars.

The compound is coupled to the biological substance by means of a reactive functional group of the compound that can covalently couple to a functional group of the biological substance. If the functional group is an active ester, it can for example be coupled to free amino groups of the biological substance. If the functional group is a maleimide residue, it can for example be coupled to free SH groups of the biological substance.

In a particularly preferred embodiment of the present invention the compounds are coupled to peptides which preferably have a maximum length of 50 amino acids and particularly preferably a maximum of 30 amino acids. These peptides are preferably prepared by synthesizing a peptide of the desired amino acid sequence on a solid phase during which a) a solid phase binding group or/and a labeling group e.g. an activated metal complex, preferably a metal complex-active ester derivative is coupled to the N-terminal amino group of the peptide or/and b) an amino acid derivative which is coupled covalently to an effector or/and labeling group e.g. a hapten or metal complex is introduced during the synthesis in at least one position of the peptide. The coupling of the effector group or/and labeling group e.g. to the N-terminal amino acid of the peptide is preferably carried out before cleaving the peptide from the solid phase and before cleaving protective groups on reactive side groups of the amino acid derivatives used for the peptide synthesis.

The peptides preferably contain one or several immunologically reactive epitope regions. These epitope regions are preferably derived from pathogenic organisms, e.g. bacteria, viruses and protozoa or from autoimmune antigens. The epitope region is particularly preferably derived from viral antigens and corresponds to the amino acid sequences of HIVI, HIVII, HIVO or hepatitis C virus (HCV).

Further preferred examples of biological substances are biotin, toxins, protoxins, nucleic acids, antibodies or antibody fragments, polypeptide antigens i.e. immunologically reactive polypeptides or haptens i.e. organic molecules having a molecular weight of 150 to 2000, in particular molecules with a steroid backbone such as cardenolides, cardenolide glycosides

(e.g. digoxin, digoxigenin), steroid alkaloids, sex hormones (e.g. progesterone), glucocorticoids etc. Other examples of haptens are prostaglandins, leucotreines, leuco-endoines, thromboxanes etc.

Yet a further subject matter of the present invention is the use of the compounds according
5 to the invention or the conjugates according to the invention in a detection method e.g. in
an immunological detection method or an nucleic acid hybridization method, in particular
in a luminescence assay.

If a metal complex is used as the labeling group, it is preferably detected by
electrochemiluminescence in which luminescent species are generated electrochemically on
10 the surface of an electrode. Examples for carrying out luminescence assays with metal
complexes of the prior art can be found in EP 0 580 979, WO 90/05301, WO 90/11511 and
WO 92/14138. Reference is herewith made to the methods and devices for luminescence
assays disclosed therein. The electrochemiluminescence assays are carried out in the
presence of a solid phase which is preferably composed of microparticles, in particular of
15 magnetic microparticles which are provided with a reactive coating e.g. with streptavidin.
In this manner it is possible to detect immune or hybridization complexes containing a
metal complex as the labeling group that are bound to the solid phase.

The electrochemiluminescence measurement is preferably carried out in the presence of a
reducing agent for the metal complex e.g. an amine. Aliphatic amines are preferred and in
20 particular primary, secondary and tertiary alkylamines whose alkyl groups each have 1 to 3
carbon atoms. Tripropylamine is particularly preferred. The amine can, however, also be an
aromatic amine such as aniline or a heterocyclic amine. The reducing agent can already be
integrated into the ligand sphere of the complex.

In addition a surface active agent e.g. a non-ionic agent such as an ethoxylated phenol may
25 be present as an amplifier. Such substances are for example commercially available under
the names Triton X100 or Triton N401.

On the other hand the luminescent metal complex can also be detected by measuring the
fluorescence or time-resolved fluorescence in which the metal chelate is excited by
irradiation with light of a suitable wavelength and the resulting fluorescence radiation is
30 measured. Examples for carrying out fluorescence assays are to be found in EP 0 178 450
and EP 0 255 534. Reference is herewith made to this disclosure.

The principle described in detail above of using metal complexes with branched and charged or hydrophilic linkers can be applied in an analogous manner to other labeling groups or/and effector groups. Other preferred test formats in which the branched linkers can be used are homogeneous assays. Such assays are based for example on the 5 measurement procedures known as CEDIA or FRET (fluorescence resonance energy transfer, cf. e.g. Pope, A. J., et al., Drug Discov Today 4 (1999) 350-362) such as time-resolved FRET.

Considerable advantages compared to known test formats can be achieved by using the branched linkers according to the invention. Thus, e.g., positively charged luminescent 10 metal complexes can be better handled within a conjugate comprising a negatively charged branched linker. An improved solubility and thus a lower unspecific binding is generally found when the branched linkers are used in combination with hydrophobic labeling groups or/and biological substances. In many cases this can be used to increase the number 15 of labeling groups and thus to increase signal yields. Furthermore the sterically demanding branched linkers prevent interactions between hydrophobic labeling groups and hydrophobic biological substances which ensures an improved accessibility of the labeling group.

The branched linkers according to the invention can be very advantageous in diagnostic methods by for example reducing the blank value, improving the dynamic range of the test, 20 lowering the lower limit of detection, broadening the test range or/and improving the signal-to-noise ratio. A reduction of the dose of active substance or/and a reduction of side effects can be achieved in therapeutic applications.

Yet a further aspect of the present invention is that linkers that carry one or several charge carriers or/and one or several hydrophilic groups as defined above cause a large shift of the 25 apparent molecular weight in chromatographic methods such as gel electrophoresis e.g. agarose gel electrophoresis, SDS gel electrophoresis, gel filtration, hydrophobic interaction chromatography and ion exchange chromatography. This effect occurs with the branched linkers described in the present application as well as with the linear linkers described in WO 96/03409. As a result of this shift in the apparent molecular weight, i.e. the linkers 30 appear to have a higher molecular weight than is actually the case, they can be used to prepare conjugates with a defined stoichiometry and homogeneous composition. After the linker which for example carries a defined number of labeling groups or effector groups, has been coupled to a binding group e.g. a biomolecule, the reaction products of the preparation can be simply obtained by chromatographic methods according to their

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stoichiometry (e.g. one molecule of linker per binding group, two molecules of linker per binding group, three molecules of linker per binding group etc.) in the form of separate fractions.

5 The linker used to prepare a particular conjugate should in this case have an apparent molecular weight of preferably $\geq 20\%$, particularly preferably of $\geq 30\%$ and most preferably of $\geq 40\%$ of the apparent molecular weight of the binding group in the same chromatographic separation system.

10 Furthermore reagent kits (linker plus labeling group(s) or effector group(s) plus binding group e.g. biomolecule to be labeled), a system (including a measuring device to detect the respective labeling group) and a composition containing a reagent of defined stoichiometry and functionality are provided.

A preferred example of such conjugates with a defined stoichiometry are mono-digoxigenylated Fab' antibody fragment conjugates.

The present invention is further elucidated by the following examples and figures.

15 The following examples, references, sequence listing and figures are provided to aid the understanding of the present invention, the true scope of which is set forth in the appended claims. It is understood that modifications can be made in the procedures set forth without departing from the spirit of the invention.

Figures:

20 Fig. 1 to 12 and 16 to 20 as well as 23 to 25 show compounds according to the invention Fig. 13 to 15 show amino acid sequences of reference antigens, and Figures 21 and 22 show the solid-phase-bound branched linker (with and without the Mtt-protective group and with protective groups on the amino acid side chains) used to produce the conjugates of Fig. 23 to 25.

25 Example 1: Preparation of branched linkers by means of solid phase peptide synthesis

The branched linkers were synthesized by means of fluorenylmethyloxycarbonyl-(Fmoc)-solid phase peptide synthesis on a batch peptide synthesizer e.g. from Applied Biosystems A433. In each case 4.0 equivalents of the amino acid derivative shown in table 1 were used for this.

Table 1:

A	Fmoc-Ala-OH
C	Fmoc-Cys(Trt)-OH
D	Fmoc-Asp(OtBu)-OH
E	Fmoc-Glu(OtBu)-OH
gE	Fmoc-Glu-OtBu
F	Fmoc-Phe-OH
G	Fmoc-Gly-OH
H	Fmoc-His(Trt)-OH
I	Fmoc-Ile-OH
K1	Fmoc-Lys(Boc)-OH
K2	Fmoc-Lys(Fmoc)-OH
K3	Fmoc-Lys(Dde)-OH
K4	Fmoc-Lys(Alloc)-OH
K5	Fmoc-Lys(PhAc)-OH
K6	Fmoc-Lys-(label)-OH
K7	Boc-Lys(Fmoc)-OH
L	Fmoc-Leu-OH
M	Fmoc-Met-OH
N	Fmoc-Asn(Trt)-OH
P	Fmoc-Pro-OH
Q	Fmoc-Gln(Trt)-OH
R	Fmoc-Arg(Pmc)-OH
S	Fmoc-Ser(tBu)-OH
T	Fmoc-Thr(tBU)-OH
U	Fmoc- β -alanine-OH
V	Fmoc-Val-OH
W	Fmoc-Trp-OH
Y	Fmoc-Tyr(tBU)-OH
Z	Fmoc- ϵ -amino caproic acid
Ps	Fmoc-Ser(PO(OBzl)OH)-OH
Cs	Fmoc-Cys(SO ₃ H)-OH

M _{tt} -U	Mtt-β-alanine-OH
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The amino acids and amino acid derivatives were dissolved in N-methyl-pyrrolion. The peptide is synthesized on Wang resin (Wang, S. S., J Am Chem Soc 95 (1973) 1328-33). The resin is loaded with 0.2 to 0.4 mMol/g. The coupling reactions were carried out for 20 minutes using 4 equivalents dicyclohexylcarbodiimide and 4 equivalents N-hydroxybenzotriazole in dimethyl-formamide relative to the Fmoc-amino acid derivative in dimethylformamide as the reaction medium. The Fmoc group was cleaved after each step of the synthesis with 20 % piperidine in dimethylformamide for 20 min. The amount of resin was selected such that after the last branch, 4 equivalents Fmoc-amino acid relative to the amino groups are used. Fmoc-Lys(Fmoc)-OH is used for the branch and subsequent synthesis of two identical arms. The unsymmetric branches are achieved by amino acid derivatives with orthogonal side chain protective groups such as Fmoc-Lys(Dde) or Fmoc-Lys(Alloc). These orthogonal protective groups are cleaved on the resin by methods known in the literature (Bycroft, B. W., et al., J. Chem. Soc., Chem. Commun. 9 (1993) 778-9; Merzouk, A., et al., Tetrahedron Letters 33 (1992) 477-80). Terminal amino groups on the solid phase are optionally acetylated or succinylated with acetic anhydride or succinic anhydride.

The hapten, label or functional group in those cases where the corresponding amino acid derivative is stable during the solid phase synthesis was already introduced on the resin e.g. on the N-terminal amino acid of the peptide.

The introduction of e.g. a metal chelate label was carried out via appropriate active ester derivatives at the free N-terminal amino group of the carrier-bound peptide. For this four equivalents ruthenium(bipyridyl)₃ complex (BPRu) per free primary amino function were activated with N-hydroxybenzotriazole/dicyclohexyl-carbodiimide and dissolved in a small amount of DMSO and this was added dropwise and stirred for 2 h at room temperature.

A hapten or a label can also be introduced at the C-terminus already during the solid phase synthesis by the direct incorporation of for example a metal chelate or biotin-coupled amino acid derivatives (described in WO 96/03409).

The peptide is released from the support and the acid-labile protective groups are cleaved with 20 ml tri-fluoroacetic acid, 0.5 ml ethanediol, 1 ml thioanisole, 1.5 g phenol and 1 ml water within 40 min at room temperature. Depending on the amino acid derivatives that

are used, it is also possible to use cocktails containing fewer radical traps. 300 ml cooled diisopropyl ether was subsequently added to the reaction solution and was kept for 40 min at 0°C in order to completely precipitate the peptide. The precipitate was filtered, washed with diisopropyl ether and dissolved in a small amount of 50 % acetic acid and lyophilized.

5 The crude material obtained was purified by means of preparative HPLC on Delta-PAK RP C18 (column 50 x 300 mm, 100 Å; 15 µ) over an appropriate gradient (eluant A: water, 0.1 % trifluoroacetic acid, eluant B: acetonitrile, 0.1 % trifluoroacetic acid) within ca. 120 min. The eluted material was identified by mass spectrometry.

10 Examples of such compounds prepared by solid phase synthesis are shown in figs. 1 to 7 and 16.

Alternatively the labeling group (label), the effector group (hapten) or the functional group can also be introduced after cleavage from the resin. For this it may be necessary to block other groups that should not be derivatized with a protective group which is stable during the solid phase peptide synthesis as well as during the cleavage (e.g. phenylacetyl (Phac)).

15 The protective group can be removed enzymatically with PenG amidase (described in PCT/EP95/02921).

An example of a compound protected with Phac is shown in Fig. 8.

Example 2: Introduction of a maleinimide function into a branched peptidic linker

In order to introduce the maleinimide function, a peptide according to example 1 was 20 dissolved in 0.1 M potassium phosphate buffer pH 7.0, admixed with one equivalent maleinimidohexanoic acid N-hydroxysuccinimide ester (MH = maleinimidohexanoyl) in DMSO and stirred for 16 h at 25°C. The preparation was purified by preparative HPLC (see above). The identity of the eluted material was checked by means of mass spectrometry.

The compounds shown in Fig. 9 to 11 and 17 were prepared.

25 **Example 3: Introduction of N-hydroxysuccinimide ester groups into branched peptidic linkers**

The experiment was carried out analogously to WO 96/03409, example 6.

The compound shown in Fig. 12 was prepared.

Example 4: Use of metal complex-antigen conjugates with branched and charged linkers in immunological tests

A double antigen bridge test was carried out to detect specific antibodies to HIV. In this method the sample liquid was incubated with a ruthenium-labeled antigen and a

5 biotinylated antigen for the antibody to be determined in the presence of a streptavidin-coated solid phase. The presence of anti-HIV antibodies in the sample liquid is determined by measuring the label on the solid phase by means of electrochemiluminescence using the Elecsys® system. A partial sequence (SEQ ID NO: 1) of the gp36 protein of HIV2 is known to comprise antigenic epitopes as recognized by important antibodies in patient sera.

10 A peptide comprising SEQ ID NO: 1 extended at the N-terminus with an SH functional linker (SEQ ID NO: 2 and Fig. 13) was prepared as described in WO 96/03652. The full line between the two cysteines (C) of Fig. 13 shall indicate a -SS- cystine bridge.

The peptide of SEQ ID NO: 2 (Fig. 13) was conjugated within 2 h at room temperature to the respective maleimidooctyl-activated ruthenium linker in 0.1 mol/l potassium phosphate

15 buffer pH 7 in order to derivatize the HIV peptide with maleimidooctyl-activated ruthenium complexes. Non-reacted components were either separated by means of preparative HPLC or gel chromatography. The purified products were lyophilized.

compound A: BPRu linker from Fig. 9 with gp36 antigen

compound B: BPRu linker from Fig. 10 with gp36 antigen

20 compound C: BPRu linker from Fig. 11 with gp36 antigen

compound D: BPRu linker from Fig. 17 with gp36 antigen

The conjugates comprising SEQ ID NO: 2 and one of the various linker variants (compounds A-D) were evaluated with the test format described above. All evaluations were carried out with the same biotinylated peptide comprising the identical gp36 epitope of

25 SEQ ID NO: 1 (Fig. 14) and at the same concentration. Labeled detection antigens were used in an equimolar concentration to the biotinylated capture antigen.

The conjugate shown in Fig. 15 and prepared as described in PCT/EP95/02921 was used as a labeled reference antigen according to the prior art (comparison in tables 2 to 4).

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The antigens according to the invention were used in equimolar amounts relative to the antigen of the prior art. The concentration was 0.018 nmol/l.

The result of the experiments with conjugates A and C compared to the compound from Fig. 15 is shown in ECL counts in table 2. It can be seen that considerably lower blank values with constant positive signals are only obtained by using the linker according to the invention which thus results in a better differentiation between positive signals and negative signals. These improved signal-to-noise ratios lead to an improvement of the measuring range.

Table 2:

Experiment	Comparison	A	C
negative sample	6014	2044	1975
positive sample	345247	484681	391007
ratio	57.5	237.1	198
positive/negative			

10

The result of the experiment with conjugate B in comparison to the compound of Fig. 15 is shown in ECL counts in table 3. It can be seen that the advantageous effects of the branched linker also enable the introduction of several labels without significantly increasing the blank value. It is also surprising that the positive signal is not quenched and even an increase in the measured counts is observed.

Table 3:

Experiment	Comparison	B
negative sample	6096	2366
positive sample	393197	765298
ratio	64.5	323.5
positive/negative		

- 20 -

Table 4 shows the result from the experiment with conjugate D compared to the compound of Fig. 15 in ECL counts. Surprisingly uncharged branched, hydrophilic linkers also have a positive effect on the blank value. The signal-to-noise ratio is improved.

Table 4:

Experiment	Comparison	D
negative sample	6718	2015
positive sample	553816	759947
ratio positive/negative	82.4	377.1

5

Example 5: Preparation of antibody fragment conjugates

1. Description of the procedure

Preparation of the Fab' from IgG

A monoclonal anti-Dig antibody was cleaved by pepsin to form $F(ab')_2$ -fragments. After 10 quantitative cleavage the pepsin was inactivated by increasing the pH and adding pepstatin. The $F(ab')_2$ was reduced by means of cysteamine to Fab' without prior purification. Cysteamine cleaves almost selectively the disulfide bridges in the hinge region. It was subsequently dialysed. This removes most of the Fc cleavage products generated by pepsin since they are small enough to pass through the pores of the dialysis tube (> 10,000 15 Dalton).

Fab'-BPRU linker conjugate

The conjugate synthesis was carried out by reacting the Fab' with an excess of BPRU-linker-MH. In this process an SH group in the hinge region was mainly converted. Small amounts of polyruthenylated Fab' were formed as a side reaction most likely as a result of reduced 20 intramolecular disulphide bridges in the light and in the Fd chain.

Purification of the crude conjugate

The crude conjugate was purified by a molecular sieve. In this process the monoruthenylated material was separated from the polyruthenylated material.

2. Procedure

5 Cleavage of the antibody to form F(ab')₂

The lyophilisate of the monoclonal antibody anti-DIG-M19.11 IgG was reconstituted with H₂O to obtain a concentration of 20 mg/ml. 20 µl 1 M citrate pH 3.5 were added per ml solution (final concentration citrate = 20 mM). The pH was adjusted with HCl to 3.60. It was filtered through a 0.45 µm filter. The concentration was determined at OD 280 nm (10 OD_{280nm} = 1.4 mg/ml). It was adjusted to 10 mg/ml with 20 mM citrate pH 3.60. The solution was heated in a water bath to 37°C. 100 µl pepsin solution (3 mg/ml) was added per ml antibody solution and incubated at 37°C in a water bath. After complete cleavage the reaction was stopped by increasing the pH value and adding pepstatin.

Reduction to Fab'

15 52.6 µl 0.1 M dithiothreitol (DTT) was added per ml cleavage mixture and incubated for 30 min at 25°C in a water bath. The Fab' was dialysed against 0.1 M NaH₂PO₄/NaOH pH 6.5, 30 mM NaCl, 2 mM EDTA.

Synthesis of the Fab'-BPRu-linker conjugate

20 The BPRu-linker-MH was dissolved in DMSO. The stoichiometry Fab':BPRu-linker-MH was 1:3 (mole/mole). The final concentration of Fab' in the mixture was 3.9 mg/ml. The maximum concentration of DMSO in the mixture was 10 %. The reaction time was 1 h at room temperature.

Purification

25 The crude conjugate was concentrated 2-3-fold using an AMICON PM 10 and purified by means of Superdex 200 (buffer: 25 mM MOPS/NaOH pH 6.5, 50 mM NaCl, 10 % DMSO; applied amount: max 1.5 % of the gel bed, fractions: 0.5 % of the gel bed). The fractions containing the Fab'-BPRu-linker conjugate were pooled.

Example 6: Use of metal complex-antibody fragment conjugates with branched and charged linkers in immunological tests

A double antigen bridge test was carried out to detect specific antibodies to HIV. In this method the sample liquid was incubated with a biotinylated antigen and a digoxigenin-labeled antigen to the antibody to be determined in the presence of a streptavidin-coated solid phase and anti-Dig-BPRu antibody. The presence of anti-HIV antibodies in the sample liquid was determined by determining the label bound via the double antigen bridge to the solid phase by electrochemiluminescence using the Elecsys® system.

A HIV peptide from the gp41 region of HIV1 (SEQ ID NO: 3), which was labeled at the N-terminus was used as the antigen. The preparation of the biotinylated and the digoxigenylated antigens is described in PCT/EP 95/02921. Two anti-Dig-BPRu conjugates were used for detection of ruthenylated gp41 peptide.

Compound E: anti-Dig-IgG-BPRu without linker

Compound F: anti-Dig-Fab'-BPRu with the linker of Fig. 9

15 Antigens were used in equimolar amounts at a concentration of 20 ng/ml.

The result of the experiments with compound E in comparison to compound F is shown in table 5. Digoxigenylated antigen and antibody conjugate (concentration 180 ng/ml) were pre-incubated. It can be seen that use of the linker according to the invention results in considerably lower blank values which is associated with a better differentiation between 20 positive and negative signals. This is easily seen from the normalized values (table 5/2).

Table 5:

Experiment [counts]	without BPRu conjugate = system blank value	compound E	compound F
negative sample	441	2117	582
positive sample 1	437	1215275	668410
positive sample 2	453	49187	25819

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Table 5/2:

Experiment (normalized to negative sample)	without BPRu conjugate = system blank value	compound E	compound F
negative sample	1.0	1.0	1.0
positive sample 1	0.99	574.1	1148.5
positive sample 2	1.03	23.1	44.4

In table 6 the test procedure was changed such that the magnetic beads with bound immune complexes were washed before adding the antibody conjugate (concentration 5 600 ng/ml). Also in this case the blank values were considerably lower and this was associated with an improved differentiation. The improved signal-to-noise ratio is especially evident from table 6/2, wherein the values have been normalized to the system blank as measured with the negative sample.

Table 6

Experiment [counts]	without BPRu conjugate = system blank value	compound E	compound F
negative sample	464	2771	758
positive sample 1	453	1503690	686607
positive sample 2	480	47040	27133

Table 6/2:

Experiment pos./neg. sera	without BPRu conjugate = system blank value	compound E	compound F
negative sample	1.0	1.0	1.0
positive sample 1	0.98	592.6	905.8
positive sample 2	1.03	17.0	35.8

5 Table 7 shows the unspecific binding of the antibody conjugate to the streptavidin solid phase. In this "test procedure" only buffer was used and neither the biotinylated nor the digoxigenylated antigens. The concentration of the ruthenylated antibody conjugate was 600 ng/ml. The linker according to the invention again exhibits improved blank values.

Table 7:

Experiment [counts]	without BPRu conjugate = system blank value	compound E	compound F
negative sample	333	3188	566
buffer	329	983	436

Example 7: Preparation of further conjugates**1. Synthesis of a testosterone derivative (Fig. 18)**

8.5 mg BPRu-linker-NH₂ (Fig. 1) was dissolved in ca. 2 ml phosphate buffer pH 8.5 and 1.8 mg of the activated testosterone derivative (testosterone-3-dimethyl-carboxyoxime-NHS) dissolved in 2 ml dioxane was added dropwise. It was allowed to stir for 6 h at room temperature with the exclusion of light.

The crude product is purified by means of preparative HPLC. The molecular weight was confirmed by means of mass spectrometry (MALDI) as 3180.

2. Synthesis of a T3 derivative (Fig. 19)

10 The synthesis was carried out analogous to 7.1. MS-MALDI corresponded to the expected molecular weight.

3. Synthesis of a PEG-Lys-MP-gp36 derivative (Fig. 20)

Starting with a lysine derivative of a ruthenium complex the free α -amino group of the lysine was reacted in the first step by conventional methods with maleinimido-propionic acid-(MP)-NHS ester. Then the carboxylic acid was activated by standard methods.

In the next step 3.64 mg of the active ester was reacted with 25.5 mg of an amino-modified polyethylene glycol H₂N-PEG-OCH₃-5000 (Shearwater) in 20 ml acetonitrile at room temperature. The product mixture was rotary evaporated and purified by means of gel chromatography (MALDI corresponded to the expected molecular weight).

20 The further coupling of the maleinimide to the gp36-peptide was carried out analogously to the already described method. The molecular weight determined with mass spectrometry corresponded to the expected molecular weight of 6990.

4. Synthesis of a fluorescence dye-labeled branched linker (Fig. 23)

25 The Mtt-protected branched peptidic linker (Fig. 21) is synthesized according to standard procedures.

The Mtt protecting group is cleaved off specifically according to the procedure described in Aletras A., et al., Int. J. Peptide Res. (1995) 45, 488 to yield the solid phase bound linker of Fig. 22.

50 mg of solid phase bound branched linker molecule (Fig. 22) with a single unprotected
5 N-terminal amino group was suspended in 4 ml DMF. Afterwards 10 μ l triethylamine and the activated fluorescence label (16 mg) was added and conjugated to the solid phase bound linker. (The synthesis of the activated label, the rhodamine-N-hydroxy succinimide ester, is described in DE 4137934.). The reaction mixture was stirred for 12 h at room temperature.

10 After cleavage from the solid phase under standard conditions (trifluoro acetic acid 95%), purification was performed with prep. HPLC.

Reaction of 16 mg of the intermediate with 18.5 mg disuccinimidyl suberate (DSS) in 10 ml DMF with 21 μ l triethylamine for 5 h and standard purification led to 7 mg product (NHS-activated rhodamine-labeled branched linker as shown in Fig. 23). This was analyzed by
15 MALDI-TOF and the molecular weight was found to correspond to the theoretical value.

Example 8: Acridinium-labeled branched linker structures

1. Acridinium ester labeled branched linker

13 mg of the acridinium ester derivative (synthesis according to EP 82636) were added to 50 mg of solid phase bound linker (cf. Fig. 22) dissolved in 4 ml DMF and reacted as
20 described in Example 7.4.

The acridinium-labeled linker was purified by prep. HPLC and lyophilised.

Product yield was found with 7 mg.

The product (cf. Fig. 24) was analyzed by MALDI-TOF and the molecular weight was found to correspond to the theoretical value.

25 2. Synthesis of acridinium sulfonyl labeled branched linker

The acridinium-labeled branched solid phase bound compound is synthesised as described in example 7.4 (synthesis of the acridinium sulfonamide see US 5,543,524). After cleavage

- 27 -

from the solid phase and purification (see above) the free C-terminal amino group of the peptidic linker (4 mg linker) is reacted with 0,7 mg maleimidopropionyl-oxy succinimide ester (MPS) in 1 ml DMF (room temperature, 150 h) and purified by prep. HPLC.

Product yield was found with 2 mg.

5 The product (cf. Fig. 25) was analyzed by MALDI-TOF and the molecular weight was found to correspond to the theoretical value.

The MPS-activated acridinium sulfonamide labeled linker is conjugated to SH groups. In this example TSH (thyroid-stimulating hormone) was used and coupling performed according to a standard protocol (see Greg T. Hermanson, Bioconjugate Techniques, 10 Academic Press, 1996, p. 456 ff) .

In a comparison study using a two step immunoassay (with a washing step after the incubation of the sample with the first antibody) for TSH, the TSH-conjugate of this example was compared to a TSH-conjugate without linker. The TSH-concentrations are given as μ IU/ml in table 8. Cal 1 and Cal 2 are commercial products, Roche Diagnostics 15 order numbers TSH Cal Set: Id. Nr: 1731483, the samples marked Tris (100mM Tris, 1% BSA, 0,1% Thesit, 0,1% Oxaban, pH 7,4) comprise 0 and 50 μ IU TSH per ml, respectively.

Table 8:

	TSH μ IU/ml	Signal-to-noise ratio (without linker)	Signal-to-noise ratio (with linker)
Cal1	0		
Cal2	1.43	2	7
Tris 0	0		
Tris 50	50	17	127

It is obvious from table 8 that the TSH-conjugate comprising the branched linker according to the present invention shows a significant improvement with regard to signal-to-noise ratio as compared to a conjugate without such linker.

List of References

Aletras A., et al., Int. J. Peptide Res. (1995) 45, 488

Aslam M., Dent A., Bioconjugation (1998) Mcmillan Reference Ltd., London, p 95 ff

Bredehorst, R., et al., Anal Biochem 193 (1991) 272-9

5 Bycroft, B. W., et al., J. Chem. Soc., Chem. Commun. 9 (1993) 778-9

Hermanson, G. T., Bioconjugate Techniques, Academic Press, 1996, p. 456 ff

Merzouk, A., et al., Tetrahedron Letters 33 (1992) 477-80

Pope, A. J., et al., Drug Discov Today 4 (1999) 350-362

Wang, S. S., J Am Chem Soc 95 (1973) 1328-33

10 PCT/EP95/02921

DE 4137934

DE 44 30 998.8

EP 0 061 888

15 EP 0 178 450

EP 0 255 534

EP 0 580 979

EP 0 618 192

US 5,519,142

20 WO 87/06706

WO 90/05301

WO 90/11511

WO 92/14138

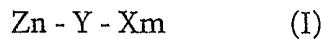
WO 96/03409

25 WO 96/03410

WO 96/03652

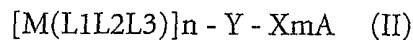
Patent Claims

1. Use of a compound of the general formula (I):



in which Z denotes at least one reactive functional group or a binding group and X is
5 a reactive functional group which is bound covalently to Z via a linker Y wherein the
linker is a branched linker which has a molecular weight of > 1000 Da and contains at
least one charge carrier or/and at least one hydrophilic group, n is an integer from 1
to 10 and m is 1 or 2, for the production of conjugates.

2. Use as claimed in claim 1, wherein the binding group is a labeling group or an
10 effector group.
3. Use as claimed in claim 1 or 2, wherein the binding group is a labeling group selected
from luminescent and fluorescent detection groups, enzymes, microparticles or
nanoparticles and radio-isotopes.
4. Use as claimed in claim 3, wherein the labeling group is metal chelate complex and
15 the compound is of the general formula (II):



in which

- M is a divalent or trivalent metal cation selected from rare earth or transition metal ions,
- 20 - L1, L2 and L3 are the same or different and denote ligands containing at least two nitrogen-containing heterocycles, L1, L2 and L3 being bound to the metal caption via nitrogen atoms,
- X is a reactive functional group which is covalently bound to at least one of the ligands L1, L2 and L3 via the branched linker Y,
- 25 - n is an integer from 1 to 10,

- m is 1 or 2 and
- A denotes the counterion which may be required to equalize the charge.

5. Use as claimed in one of the claims 1 to 4, wherein the binding group is an effector group selected from partners of specific bioaffine binding pairs.

5 6. Use as claimed in claim 5, wherein the effector group is selected from biotin and analogues thereof, streptavidin, avidin, antigens, haptens, antibodies, nucleic acids, nucleic acid analogues, sugars, lectins, receptors and receptor ligands.

7. Use as claimed in one of the claims 1 to 6, wherein the reactive functional group is a carboxylic acid, a carboxylic acid halogenide, a carboxylic acid anhydride, a carboxylic acid hydrazide, a carboxylic acid azide, an amine, an active ester, a maleimide, a thiol or a photoactivatable group.

10 8. Use as claimed in one of the claims 1 to 7, wherein the branched linker contains at least one negative charge carrier selected from the group consisting of phosphate, phosphonate, sulphinate, sulphonate, sulphate and carboxylate groups.

15 9. Use as claimed in claim 8, wherein the branched linker contains at least one carboxylate group or/and phosphate group.

10. Use as claimed in one of the claims 1 to 9, wherein the branched linker contains at least one positive charge carrier selected from amino groups and substituted amino groups.

20 11. Use as claimed in one of the claims 1 to 10, wherein the branched linker contains up to 70 charge carriers.

12. Use as claimed in claim 11, wherein the branched linker contains 1 to 40 charge carriers.

25 13. Use as claimed in one of the previous claims, wherein the branched linker contains at least one uncharged hydrophilic group selected from ethylene oxide, polyethylene oxide, sulphoxide, sulphone, carboxylic acid amide, carboxylic acid ester, phosphonic acid amide, phosphonic acid ester, phosphoric acid amide, phosphoric acid ester,

sulphonic acid amide, sulphonic acid ester, sulphuric acid amide and sulphuric acid ester groups.

14. Use as claimed in claim 13, wherein at least one uncharged hydrophilic group is a primary carboxylic acid amide group.
- 5 15. Use as claimed in one of the previous claims, wherein the molecular weight of the linker is in the range from 1000 to 50,000 Da.
16. Use as claimed in one of the claims 1 to 15, wherein the branched linker is at least partially composed of aminocarboxylic acid units which are linked together by peptide bonds.
- 10 17. Use as claimed in claim 16, wherein the charge carriers are derived from polyfunctional aminocarboxylic acids which still contain at least one free charge carrier after incorporation into the linker.
- 15 18. Use as claimed in claim 16, wherein the hydrophilic groups are derived from polyfunctional aminocarboxylic acids which still contain at least one hydrophilic group after incorporation into the linker.
19. Use as claimed in claim 16, wherein the branching positions are derived from polyfunctional aminocarboxylic acids.
- 20 20. Use as claimed in claim 17, 18 or 19, wherein the polyfunctional aminocarboxylic acids are selected from lysine, ornithine, hydroxylsine, aspartic acid, glutamic acid, asparagine, glutamine, phosphoserine and synthetic trifunctional aminocarboxylic acids.
21. Compound of the general formula (I)

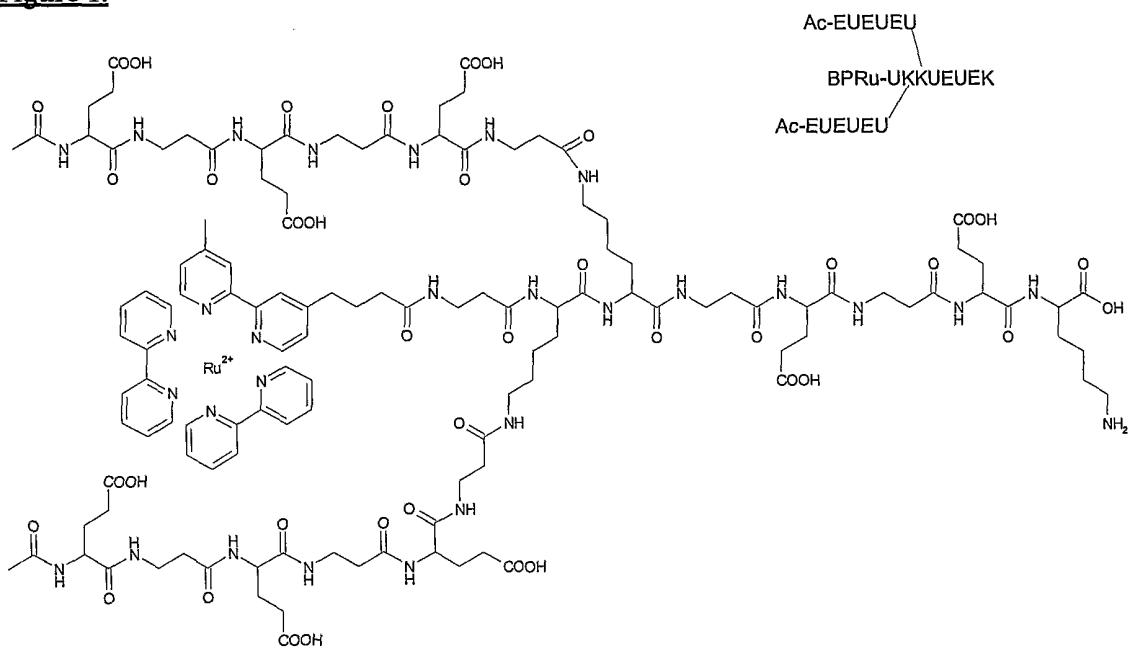
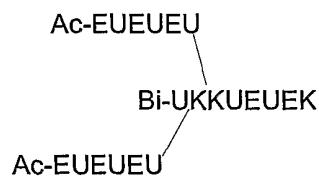
Zn - Y - Xm

25 wherein Z denotes at least one reactive functional group or a binding group and X is at least one reactive functional group which is covalently bound to Z via a linker Y where the linker is a branched linker that has a molecular weight of > 1000 Da and

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contains at least one charge carrier or/and at least one hydrophilic group, n is an integer from 1 to 10 and m is 1 or 2.

22. Compound as claimed in claim 21, wherein it has one or several of the features claimed in one of the claims 2 to 20.
- 5 23. Conjugate comprising at least one biological substance and at least one compound of the general formula (I) as claimed in claim 21 or 22.
24. Conjugate as claimed in claim 23, wherein the biological substance is an antibody or antibody fragment, a nucleic acid, a polypeptide antigen, an immunologically reactive peptide or a hapten.
- 10 25. Use of compounds as claimed in claim 21 or 22 or the conjugates as claimed in claim 23 or 24 in an immunological detection method or in a nucleic acid hybridization method.
26. Use as claimed in claim 25 in a luminescence method.
27. Use as claimed in claim 25 or 26 in an electrochemiluminescence method.
- 15 28. Use as claimed in claim 27 to improve the solubility of labeling groups or effector groups or of their conjugates.

Figure 1:Figure 2:

5

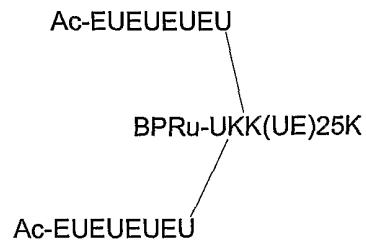
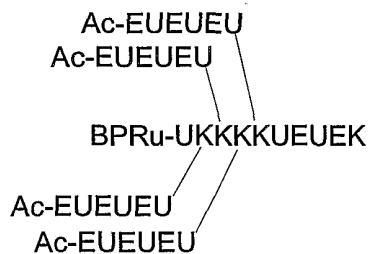
Figure 3:

Figure 4:Figure 5:

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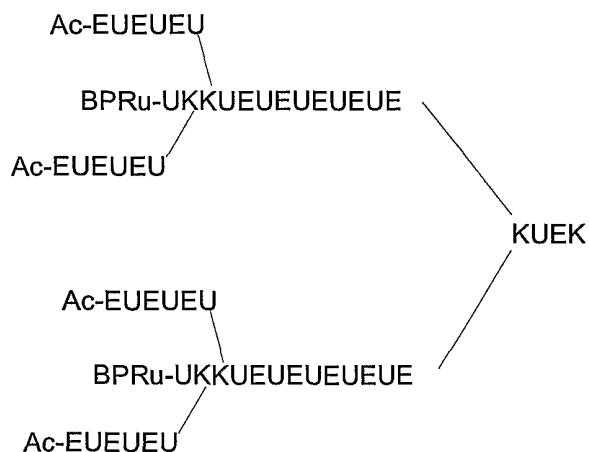
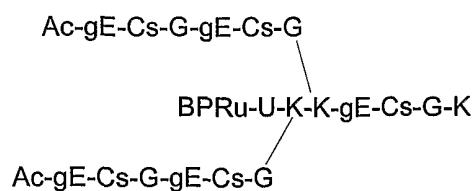
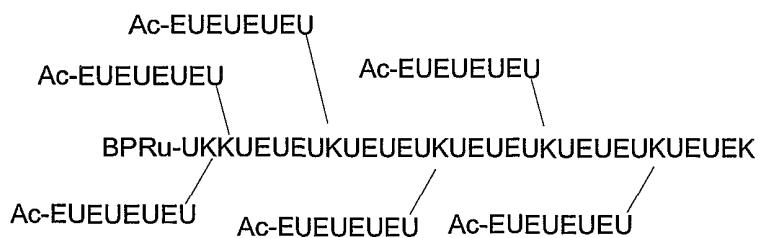
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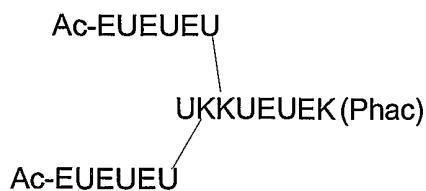
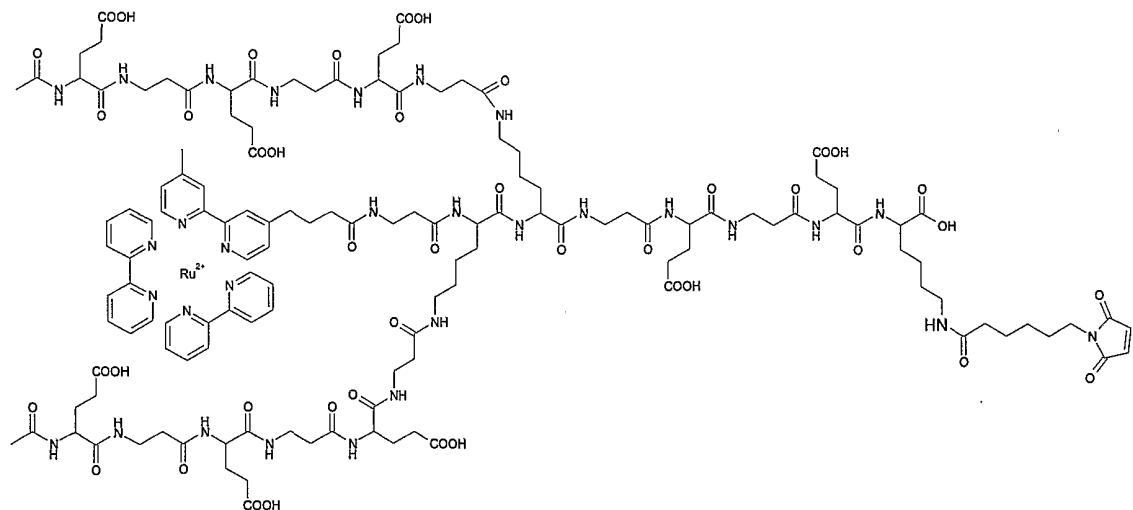
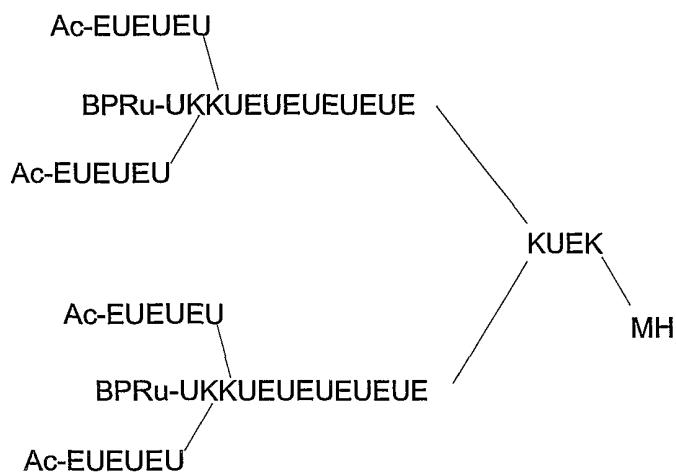
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Figure 10:Figure 11:

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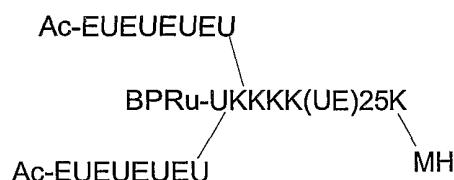
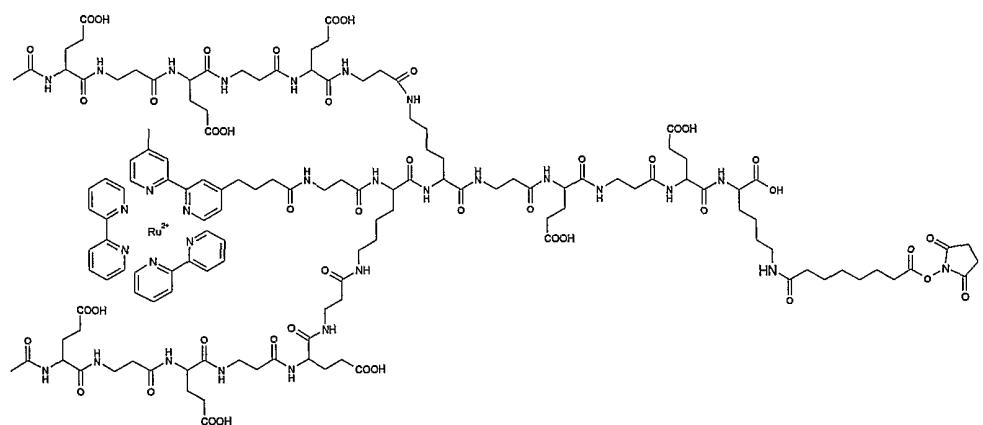
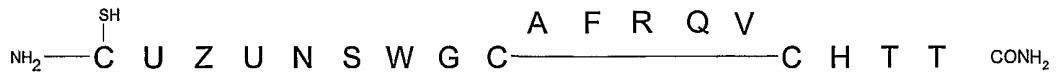
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Figure 13:Figure 14:Figure 15:Figure 16:

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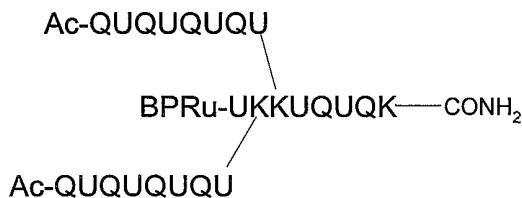
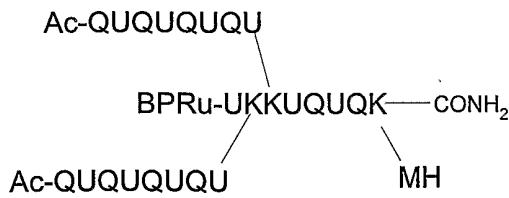
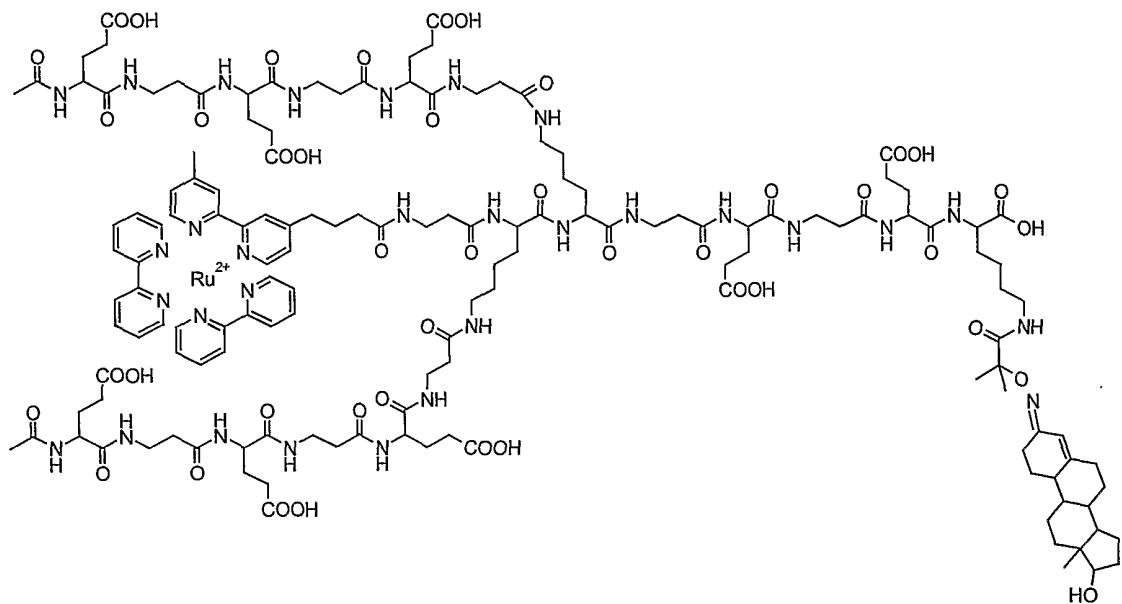
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Figure 18:Figure 19:

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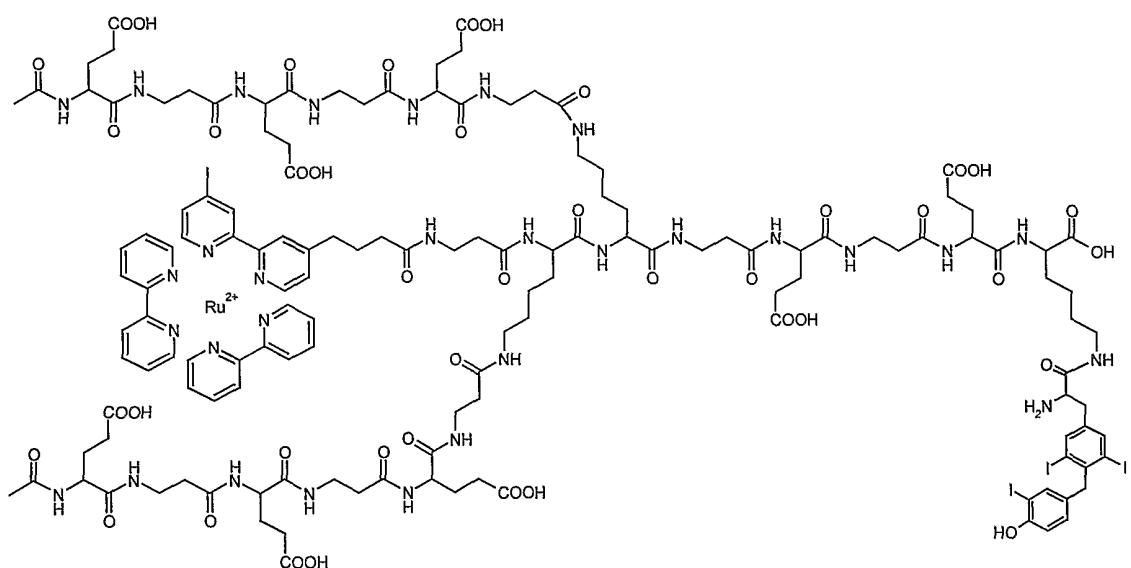


Figure 20: Ruthenium- and gp36-labeled hydrophilic (PEG 5000) linker

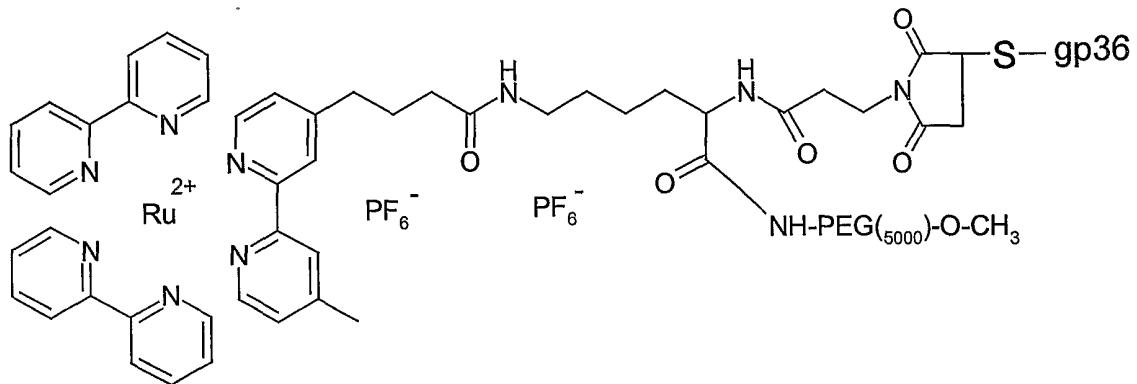


Figure 21: Solid-phase bound branched linker (with protective groups)

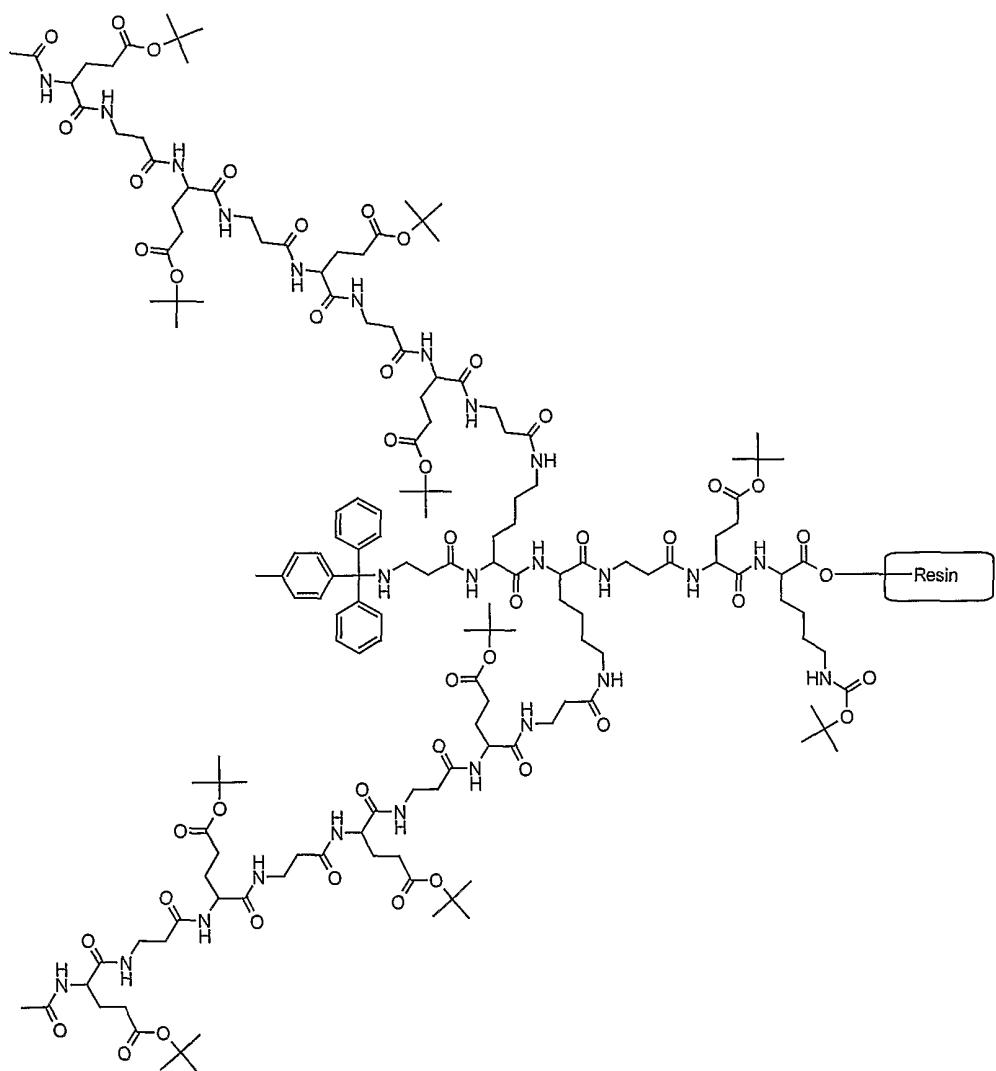


Figure 22: Branched linker with protected amino acid side chains and un-protected ϵ -amino-group of lysine

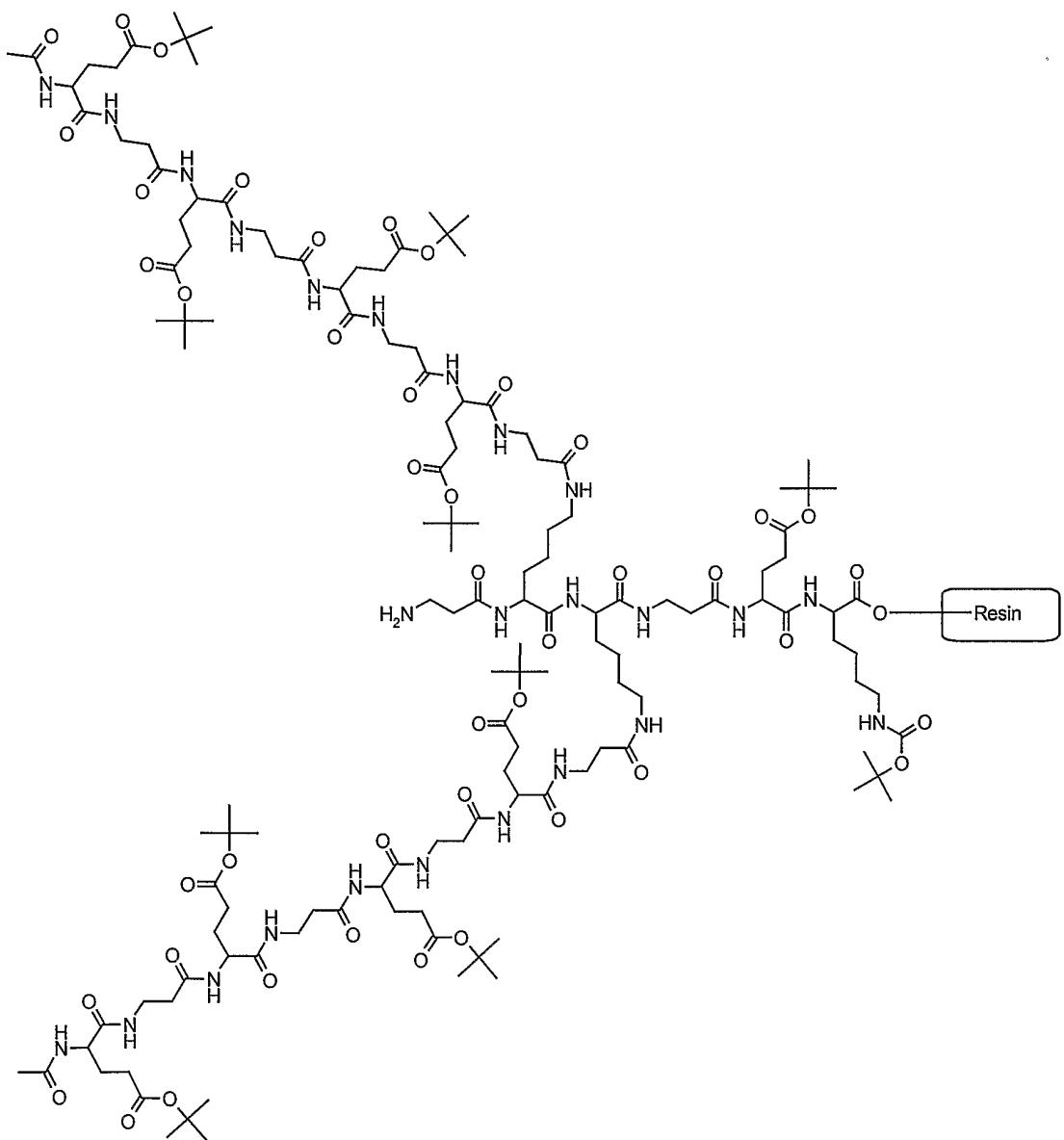


Figure 23: NHS-activated rhodamine-labeled branched linker

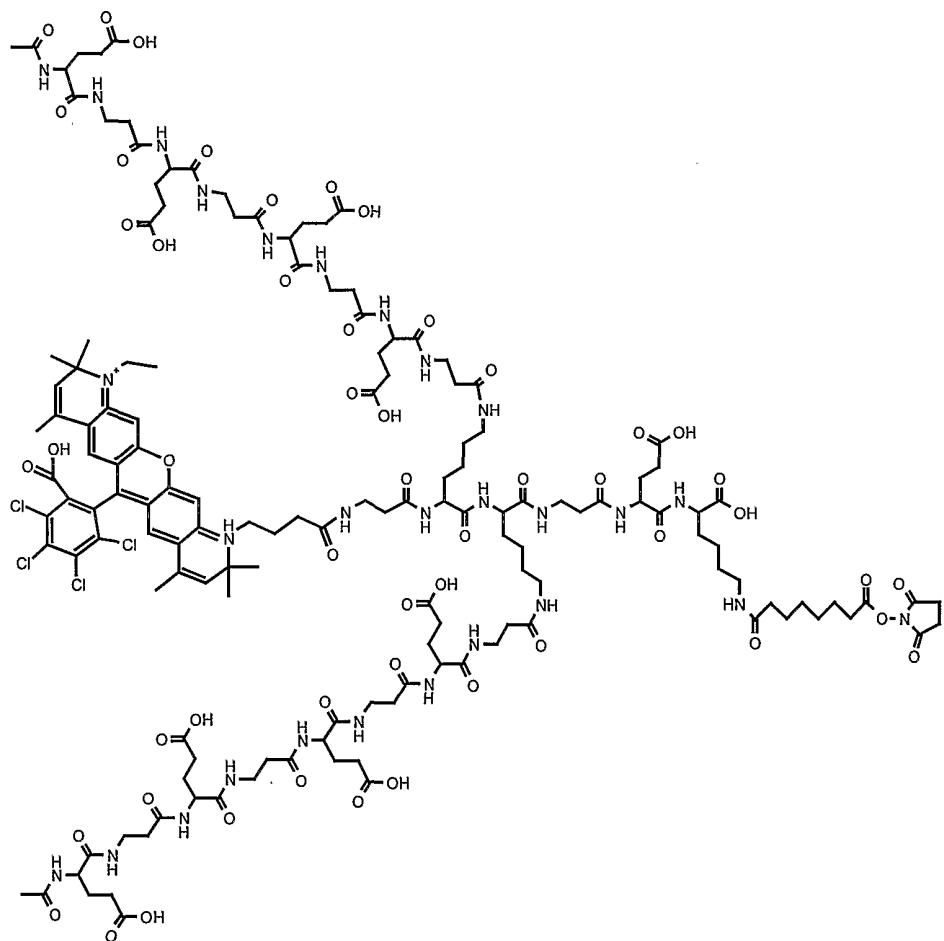


Figure: 24: Branched linker labeled with an acridinium ester

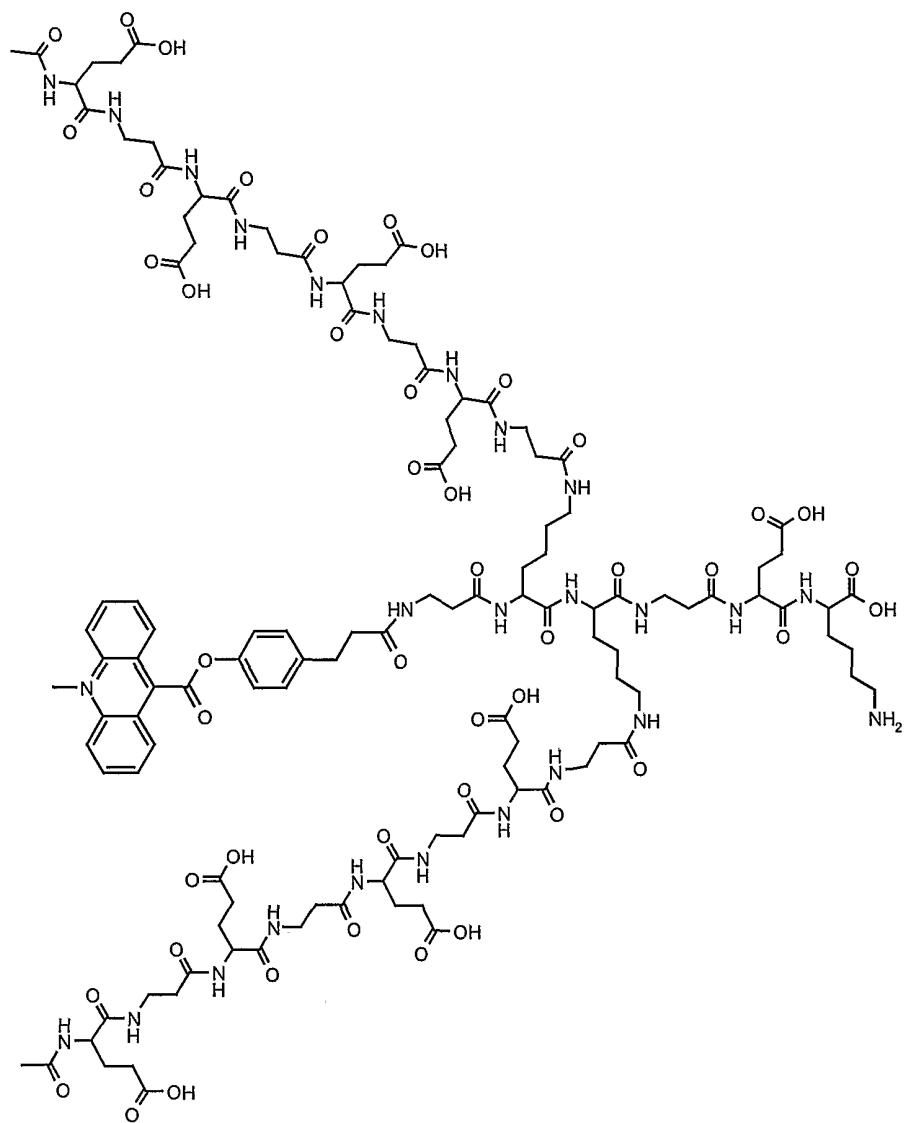
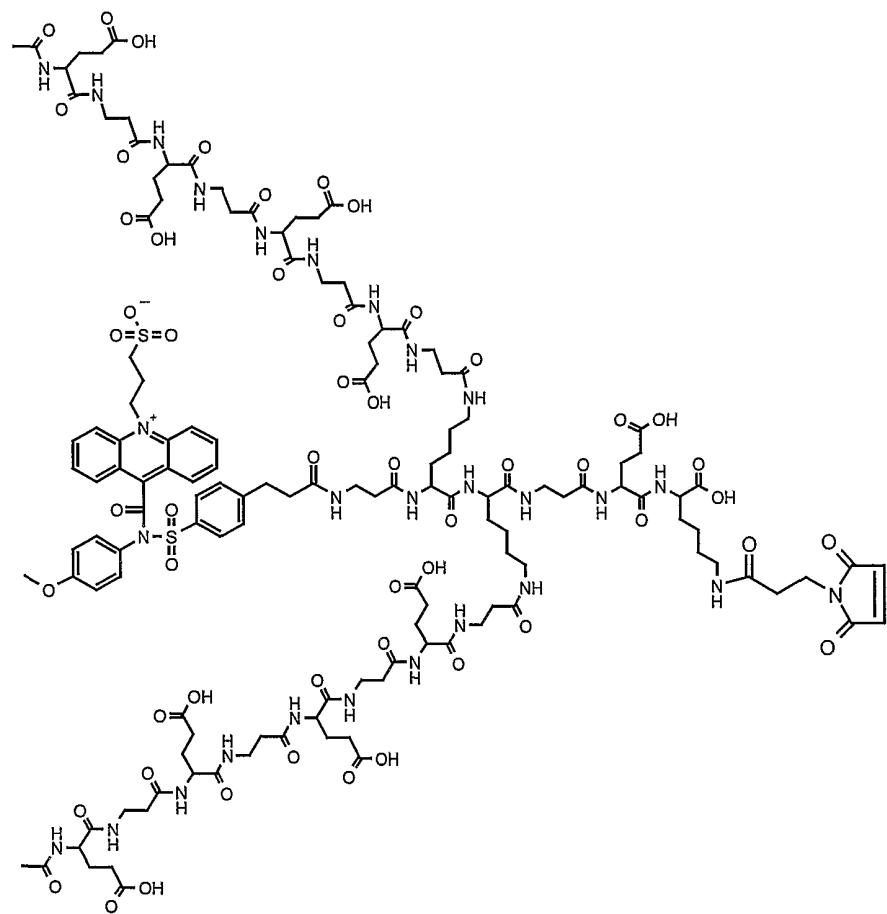


Figure 25: MPS-activated branched linker with an acridinium sulfonyl group as label



- 1 -

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Roche Diagnostics GmbH
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