

Processes for the Manufacture of Macrocyclic Depsipeptides and New Intermediates

Summary of the Invention

The invention relates to a method or process for the manufacture of macrocyclic depsipeptides, to new intermediates and their manufacture, as well as related invention embodiments.

Background of the Invention

Cyclic depsipeptides have numerous uses in pharmacology. As an example, the depsipeptides disclosed in WO2009/024527 are useful for treatment of various diseases. For example, the compound of formula II mentioned in WO2009/024527 is useful for the treatment and prevention of inflammatory and/or hyperproliferative and pruritic skin diseases such as atopic dermatitis, psoriasis, pustular psoriasis, rosacea, keloids, hypertrophic scars, acne, Netherton's syndrome or other pruritic dermatoses such as prurigo nodularis, unspecified itch of the elderly as well as other diseases with epithelial barrier dysfunction such as aged skin.

Nostopeptin BN920, formerly isolated from the cyanobacterium Nostoc, was isolated also from Microcystis. Nostopeptin BN920 inhibited chymotrypsin with an IC₅₀ value of 31 nM (see J. Nat. Prod. 68(9), 1324-7 (2005)).

These compounds can be produced by fermentation (using chondromyces croactus, myxobacteria) along with other depsipeptides comprising the so-called ahp-substructure (ahp: 3-amino-6-hydroxy-piperidin-2-one) and the corresponding dehydro-ahp substructure (dehydro-ahp: 3-amino-3,4-dihydro-1H-pyridin-2-one), also called "dehydrate" herein, respectively. Therefore, the yield of fermentation with regard to any single of these compounds is rather low.

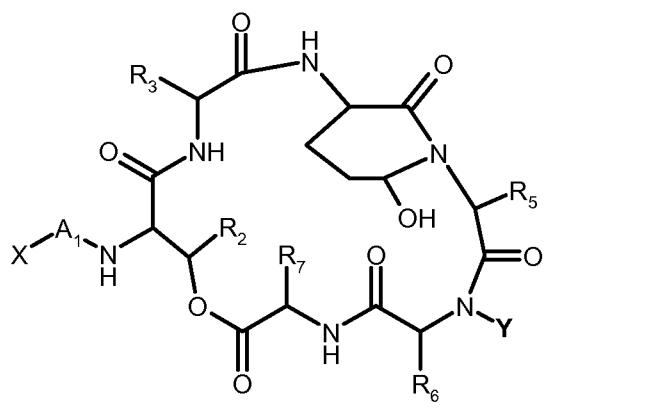
The present invention relates to processes or methods that allow obtaining such cyclic depsipeptides with increased yield and/or in good purity.

In view of the many risks, such as epimerization, tautomerization and the like in the synthesis of a complex molecule with many possible isomers, it has been possible to find

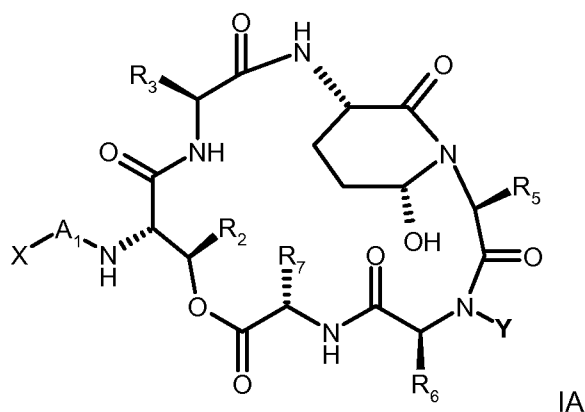
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a manufacturing process, preferably comprising a mixture of solid phase peptide synthesis and reactions in solution, that allows to produce cyclic depsipeptides of formula I in good yield and/or the required stereoisomeric purity, especially both. It is possible to reduce the amount of by-products, and even to improve yield, by converting such by-products, especially the dehydro-ahp substructure and/or an analogue of the desired ahp-comprising products with a five-membered ring instead of the ahp, into the desired final products. This allows to further increase yield. No synthesis has so far come to our attention making use of solid phase peptide synthesis in this field.

(i/a) In a first embodiment, the invention relates to a method or process for the preparation of a cyclic depsipeptide compound of the formula I,



especially of the formula IA



wherein

A₁ is a bivalent moiety of an amino acid with a terminal carboxy or carbamoyl group, especially asparagine or glutamine, and is bound at its right hand side in formula I (corresponding to the C-terminus) via a carbonyl (preferably the carbonyl of an α-

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carboxyl group thereof) to the rest of the molecule; or is C₁₋₈-alkanoyl or phosphorylated hydroxy-C₁₋₈-alkanoyl;

X is bound via an N of A₁ and is acyl, or is absent if A₁ is C₁₋₈-alkanoyl or phosphorylated hydroxy-C₁₋₈-alkanoyl;

R₂ is C₁₋₈-alkyl, especially methyl;

R₃ is the side chain of an amino acid, especially of leucine, isoleucine or valine;

R₅ is the side chain of an amino acid, preferably of phenylalanine, leucine, isoleucine or valine;

R₆ is the side chain of a hydroxy amino acid, especially of tyrosine;

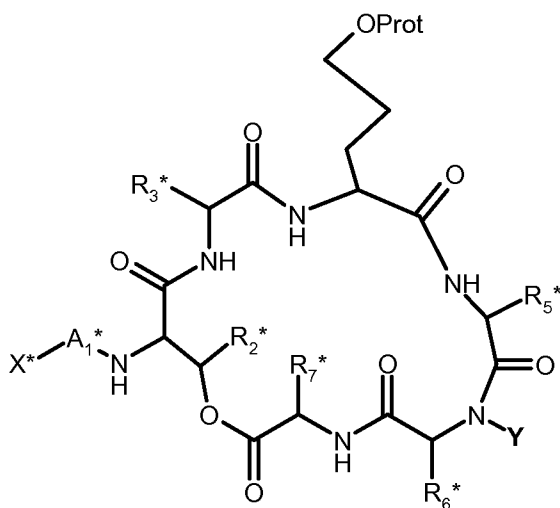
R₇ is the side chain of an amino acid, preferably of the amino acid leucine, isoleucine or valine; and

Y is hydrogen or C₁₋₈-alkyl;

or a salt thereof,

said method comprising

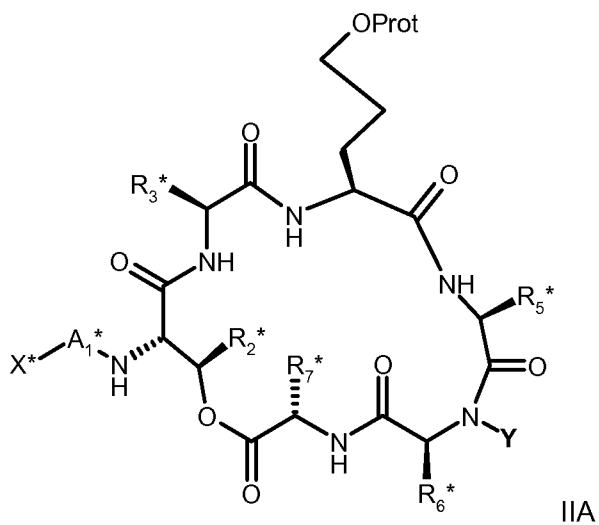
selectively deprotecting a compound of the formula II



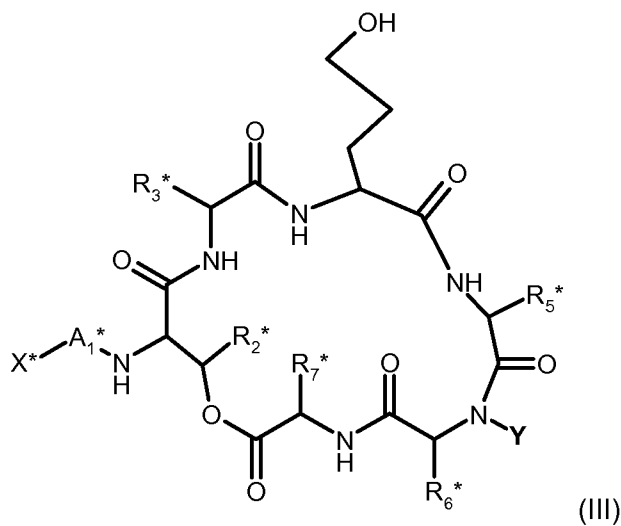
II

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especially of the formula IIA

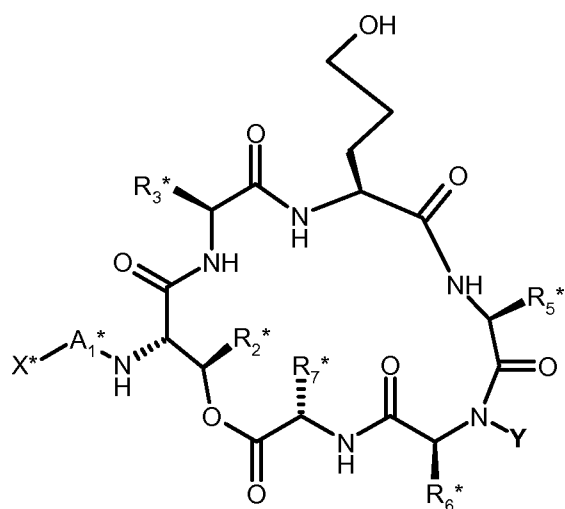


wherein Prot is a protecting group, Y is as defined for a compound of the formula I and X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* correspond to X, A_1 , R_2 , R_3 , R_5 , R_6 , and R_7 in formula I, respectively, but with the proviso that reactive functional groups on these moieties (such as amino, imino, hydroxy, carboxy, sulfhydryl, amidino, guanidino, O-phosphono ($-O-P(=O)(OH)_2$)) are present in protected form at least if they could participate in undesired side reactions, to result in a compound of the formula III,



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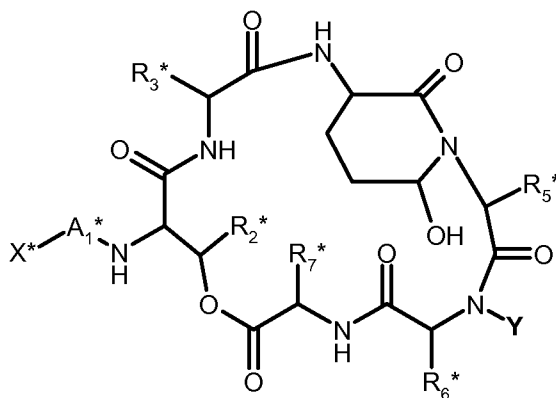
especially of the formula IIIA,



IIIA

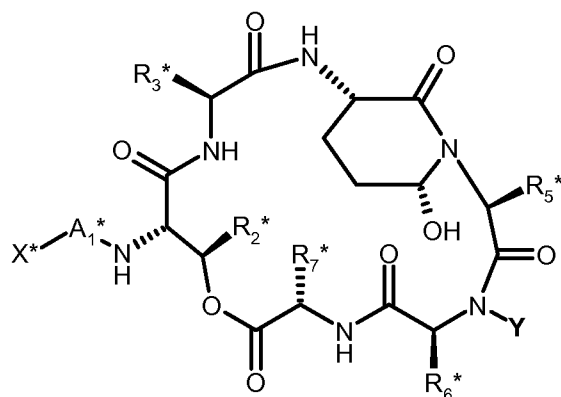
wherein X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* have the meanings just defined,

reacting the free hydroxyl group under oxidizing conditions to form a compound of the formula IV



IV

especially IVA,



IVA

and removing remaining protecting groups to yield a compound of the formula I, or a salt thereof,

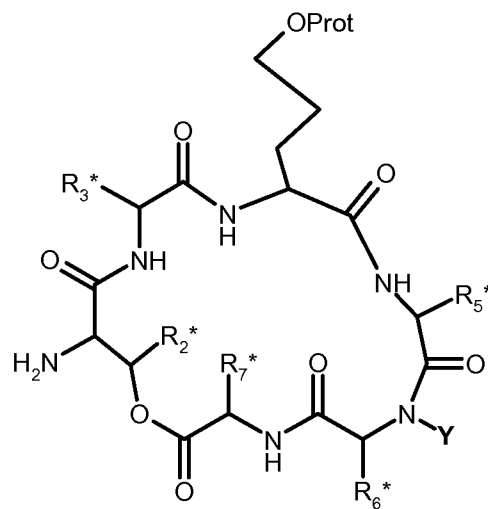
and, if desired, converting a free compound of the formula I, or especially IA, into a salt, a salt of a compound of the formula I into a different salt of a compound of the formula I, or especially IA, or into the free compound of the formula I, or especially IA, and/or converting a dehydrate analogue and/or five ring analogue of a compound of the formula I, or especially IA, into the corresponding compound of the formula I, or especially IA.

Suitable oxidizing conditions for the oxidation of a compound of the formula III or especially IIIA are usually using IBX in DMSO (J. Org. Chem. 1995, 60, 7272-7276); Pyridinium dichromate or Pyridinium chlorochromate (Tetrahedron Lett. 1979, 5, 399-402); oxalyl chloride, dimethyl sulfoxide and a tertiary amine (J. Peptide Sci. 2006, 12, 140-146), oxoammonium salts (J. Org. Chem. 1985, 50, 1332-1334); alkali hypochlorites catalyzed by oxoammonium salts (J. Org. Chem. 1989, 54, 2970-2972); oxoaminium salts (Tetrahedron Lett. 1988, 29, 5671-5672), $\text{RuCl}_2(\text{PPh}_3)_3$ (Tetrahedron Lett. 1981, 22, 1605-1608); TEMPO (1 mol%) in the presence of sodium hypochlorite (Tetrahedron Lett. 1990, 31, 2177-2180); NaIO_4 , TEMPO, NaBr (Tetrahedron 2006, 62, 8928-8932); SiO_2 supported vanadium(IV)oxide and t-BuOOH (Advanced Synthesis & Catalysis 2007, 349, 846-848). Preferably, the reaction is performed with IBX in DMSO or preferably in an inert solvent, such as tetrahydrofuran, in the presence of DMSO, at a temperature between 0-50°C, preferably between 20-25 °C.

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(ii/a) A further embodiment of the invention refers to the method or process described above, in addition comprising manufacturing the compound of the formula IV or especially IVA by a combination of Solid Phase Peptide Synthesis (especially For synthesis of the precursor XX or especially XXA given below for the oligopeptide precursor of the formula VIII or especially VIIIA given below, or of the oligopeptide precursor of the formula XXIV or especially XXIVA given below for the oligopeptide precuros of the formula XXV or especially XXVA given below) and Solution Phase synthesis (especially from the compounds just mentioned to the final product) from the corresponding starting amino acids and side chain precursors.

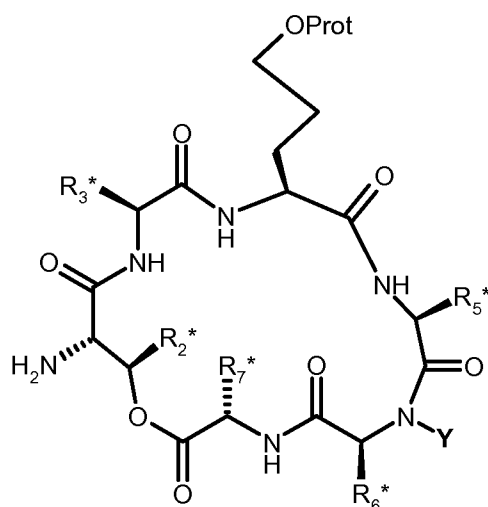
(iii/a) Yet a further embodiment of the invention relates to a method or process as described above, further comprising, for the synthesis of a compound of the formula II above, reacting a compound of the formula VI,



VI

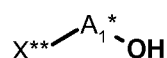
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especially VIA,



VIA

wherein Prot is a protecting group, Y is as defined for a compound of the formula I and R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* are as defined for a compound of the formula II above, with an acid of the formula VII



VII

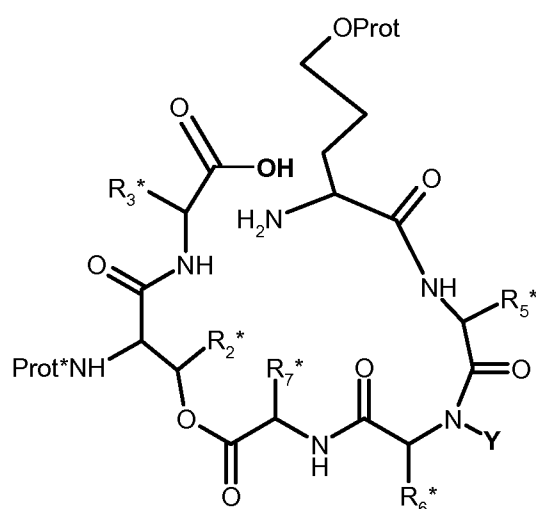
or a reactive derivative thereof,

wherein X^{**} is an amino protecting group or is X^* , and wherein X^* and A_1^* are as defined for a compound of the formula II above; and, if X^{**} is an amino protecting group, removing said amino protecting group X^{**} to yield the derivative of formula II (especially IIA) wherein, instead of X^* , H (a hydrogen) is present and coupling the resulting amino group with an acyl group X^* using the corresponding acid $X^* - OH$ wherein X^* is as defined for a compound of the formula II defined above, or a reactive derivative thereof.

(iv/a) Another embodiment of the invention relates to the methods or processes described above, especially in the preceding paragraph, further comprising cyclization under lactamization of a linear, that is, not yet cyclic, precursor peptide of the compound of the formula VI, carrying an N-terminal amino group and a C-terminal carboxy group, under reaction conditions that allow for the formation of an amide bond from said amino and said carboxy group, preferably using Solution Phase chemistry.

Lactamizations in solutions are usually carried out at very low concentrations of the substrate in order to avoid oligomerizations and polymerizations. This requires huge amounts of solvents and very large reactors to carry out the reactions. For example, the macrolactamization of an oligopeptide is performed at a concentration of 2 mMols/liter in reference Yokokawa et al., Tetrahedron 2005, 61, 1459-1480. This difficulty can be circumvented by dissolving the tertiary base and the coupling reagent and, in a controlled way adding a solution of the oligopeptide to this solution. The controlled, especially slow, addition of the oligopeptide-solution generates permanently low concentrations of the activated oligopeptide in solution and thus prevents oligomerization and polymerization. The addition rate of the oligopeptide solution can be adjusted according to the reaction rate for the macrocyclization: if the macrocyclization is a fast reaction, the solution of the oligopeptide can be added fast. If the macrocyclization is slow, the addition of the solution must be slow to ensure permanent low concentration of the activated oligopeptide. Thus the controlled addition of the oligopeptide enables to work with much less solvent amounts and still maintaining the concentration of the activated oligopeptide below 10^{-3} mM, e.g. in the range from 10^{-4} to 10^{-6} mM or even lower. This variant of controlled addition of the oligopeptide to the coupling reagent solution is an embodiment of the invention.

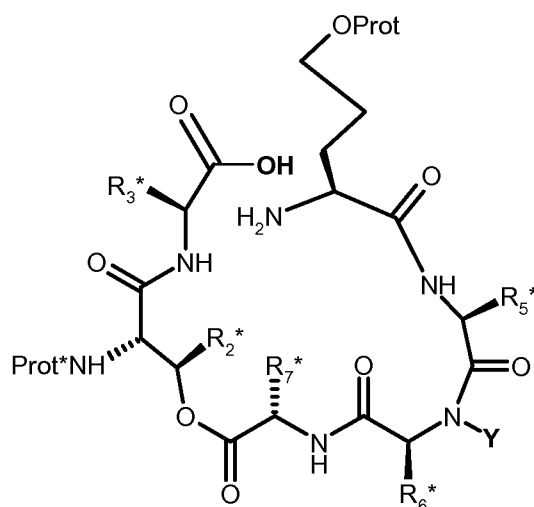
(v/a) In yet a further embodiment, the invention relates to the method or process as described above, especially in the preceding paragraph, where the linear (this term where used meaning not yet cyclic) precursor peptide is of the formula VIII,



VIII,

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especially VIIIA



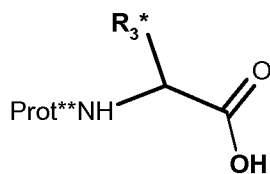
VIIIA

wherein Prot* is a protecting group that can be cleaved off selectively without affecting other protecting groups present and is stable during deprotection steps during synthesis of the linear precursor peptide (e.g. allyloxycarbonyl) and R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* are as defined for a compound of the formula VI as described above, further comprising, after cyclisation of the compound of the formula VIII, especially VIIIA, removing the protecting group Prot* *in situ* to yield the compound of the formula VI, especially VIA.

(vi/a) In another embodiment, the invention relates to the method or process described above, especially in the preceding paragraph, where the linear precursor peptide of the formula VIII, especially VIIIA, is synthesized from the corresponding amino acids by solid phase peptide synthesis and subsequent cleavage from the employed solid support.

(vii/a) An embodiment of the invention further relates to the method or process as described above (especially in the preceding paragraph (vi/a)), further comprising either

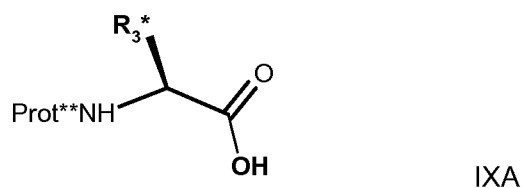
in a variant a), coupling an amino acid of the formula IX,



IX

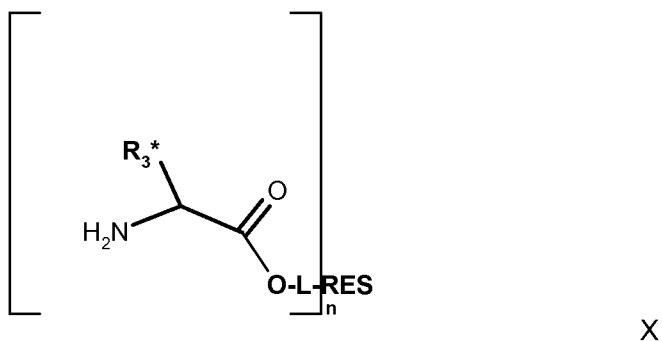
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especially of the formula IXA,

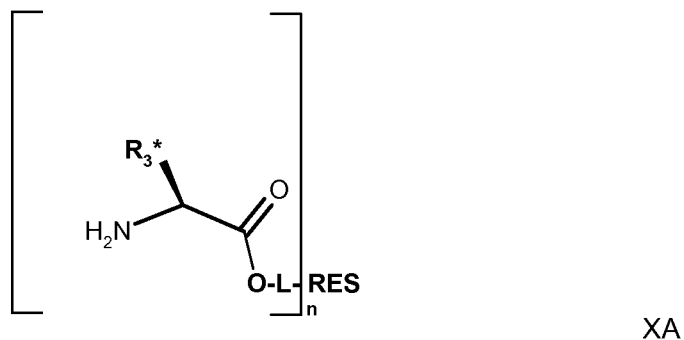


wherein R_3^* is as defined for a compound of the formula II above and Prot** is an amino protecting group that can be removed on the resin without cleaving other bonds, or a reactive derivative of said amino acid, via an oxygen to a cleavable linker L which is bound to a solid resin RES (e.g. by reaction with a resin of the formula $(X-L)_z-RES$ wherein L and RES are as just defined, X is e.g. halo, e.g. chloro, and z is a number larger than zero, e.g. a natural number) and removing the protecting group Prot**;

coupling the obtainable resin bound amino acid symbolized by the formula X,

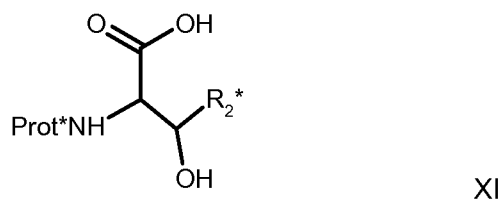


especially XA,

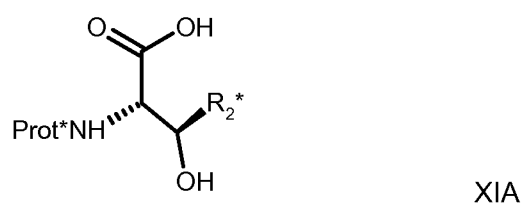


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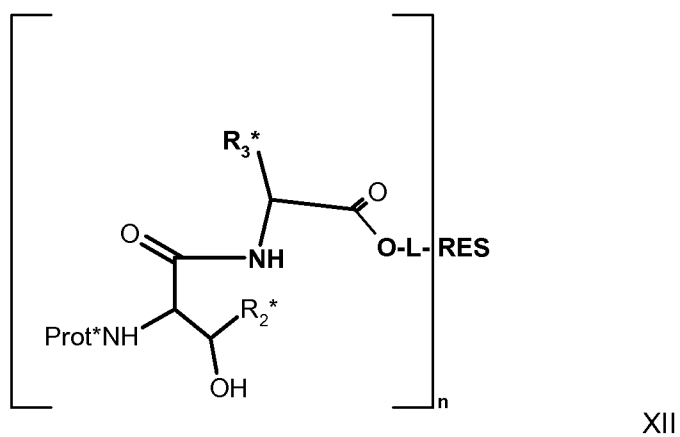
in which RES and R_3^* are as defined for a compound of the formula IX, n is a (e.g. natural) number larger than zero and L is a cleavable linker, with an amino acid of the formula XI,



especially XIA,

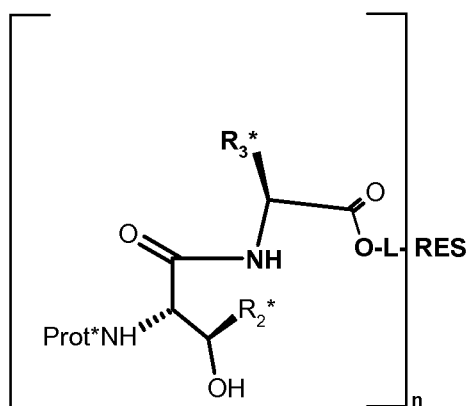


wherein Prot* is as defined for a compound of the formula VIII above and R_2^* is as defined for a compound of the formula II above, or a reactive derivative of said amino acid, coupling the obtainable resin bound dipeptide symbolized by the formula XII,



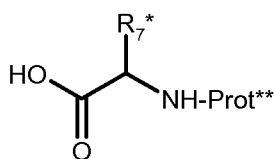
- 13 -

especially XIIA;



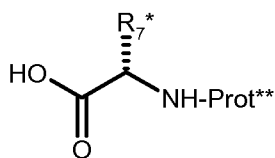
XIIA

in which Prot* is as defined for a compound of the formula VIII above, R_2^* and R_3^* are as defined for a compound of the formula II above, and n , L and RES are as defined for a compound of the formula X, via the free hydroxy group with an amino acid of the formula XIII,



XIII

especially XIII A,

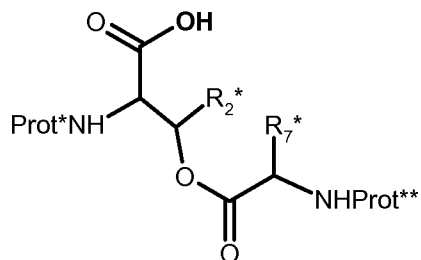


XIII A

wherein Prot** is as defined for a compound of the formula IX and R_7^* is as defined for a compound of the formula II above, or a reactive derivative of said amino acid, and removing the protecting group Prot**;

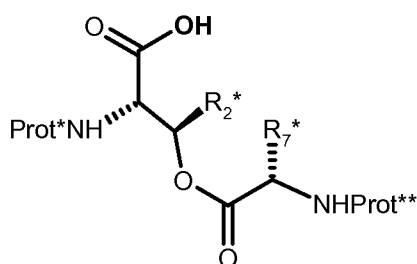
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or, in a variant b), coupling a dipeptide of the formula XXVII,



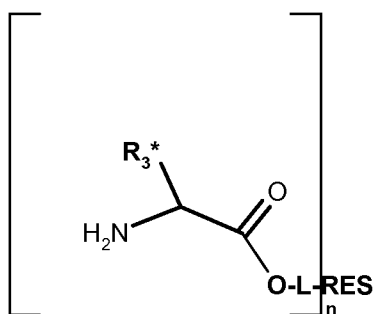
XXVII

especially of the formula XXVIIA,



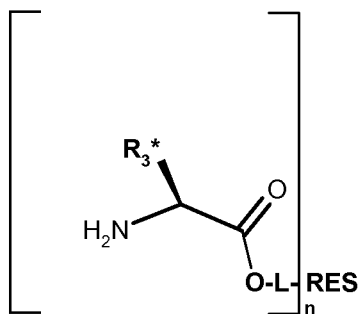
XXVIIA

wherein R_3^* and Prot^{**} are as described for a compound of the formula IX, especially IXA, and Prot^* is as defined for a compound of the formula VIII above, or a reactive derivative of said dipeptide, to an amino acyl moiety, bound via an oxygen to a cleavable linker L which is bound to a solid resin RES, having the formula X,



X

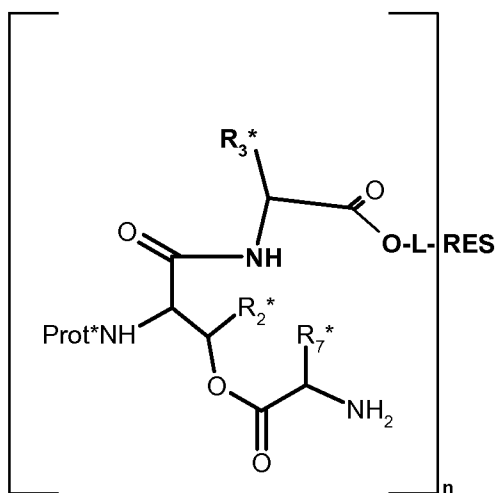
especially XA,



XA

that can be obtained as described under variant a), in which RES, and R_3^* are as defined for a compound of the formula IX and L and RES are as just defined; and removing the protecting group Prot^{**} ;

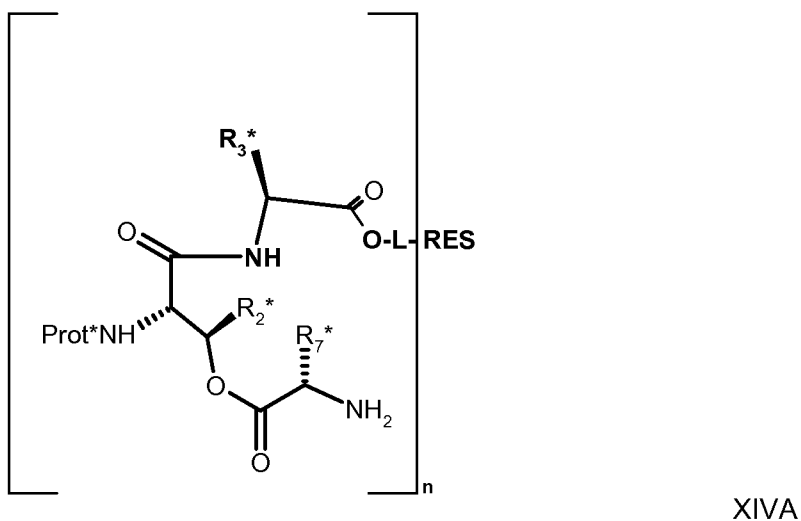
and, after the reactions of variant a) or of variant b),
(viii/a) coupling the obtainable compound of the formula XIV,



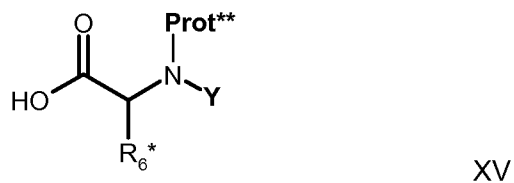
XIV

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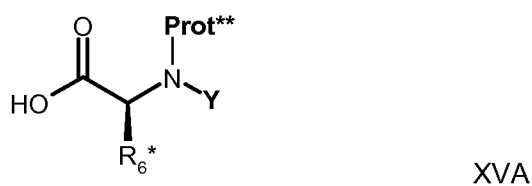
especially XIVA,



wherein R_2^* , R_3^* and R_7^* are as defined for a compound of the formula II above, Prot^* is as defined for a compound of the formula VIII above and n , L and RES are as defined for a compound of the formula X, with an amino acid of the formula XV,

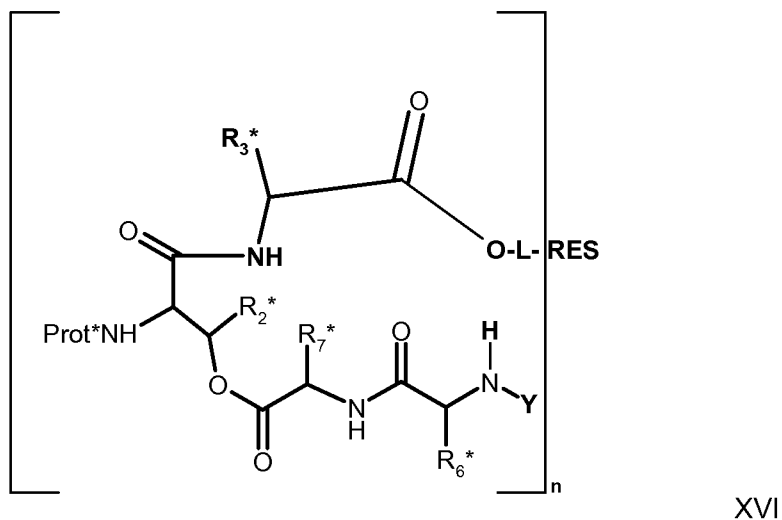


especially the formula XVA,

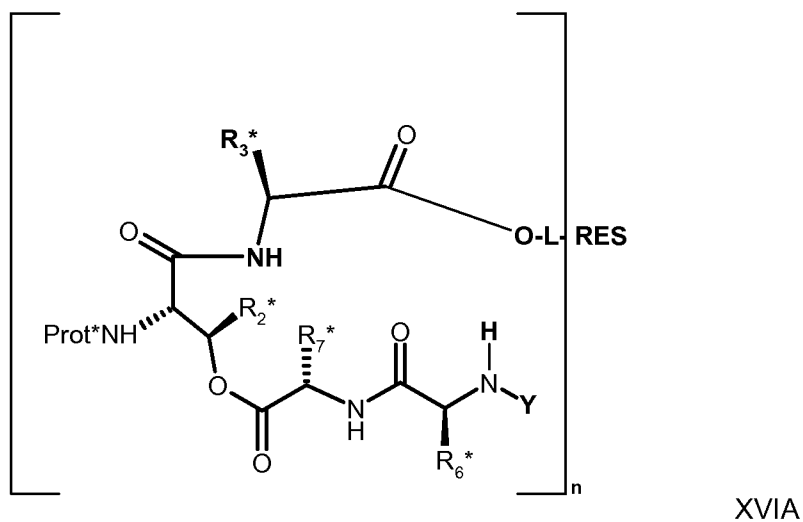


in which R_6^* and Y are as defined for a compound of the formula II above and Prot^{**} is as defined for a compound of the formula IX above, or a reactive derivative of said amino acid, and removing the protecting group Prot^{**} ;

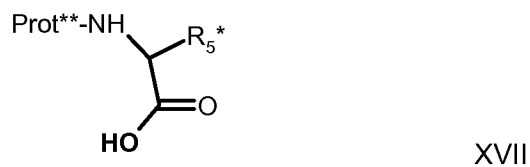
(ix/a) preferably coupling the obtainable compound of the formula XVI



especially the formula XVIA,

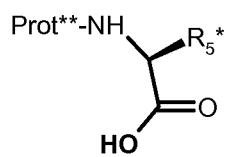


wherein Y, R₂^{*}, R₃^{*}, R₇^{*} and R₆^{*} are as defined for a compound of the formula II above, Prot^{*} is as defined for a compound of the formula VIII above and n, L and RES are as defined for a compound of the formula X, with an amino acid of the formula XVII,



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especially formula XVIIA,

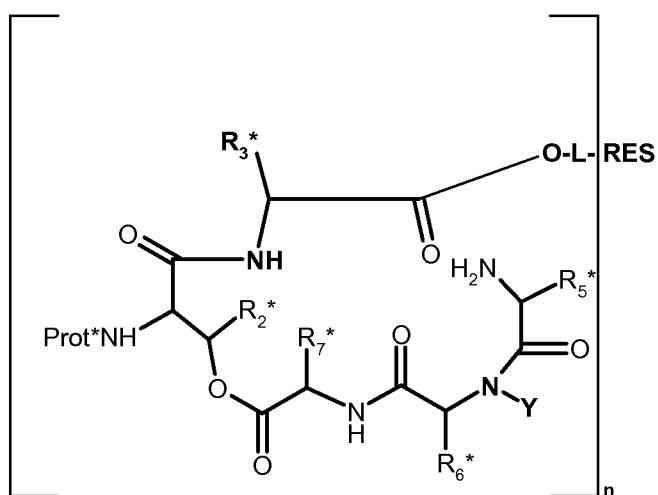


XVIIA

wherein R_5^* is as defined for a compound of the formula II above and Prot^{**} is as defined for a compound of the formula IX, or a reactive derivative of said amino acid, and removing the protecting group Prot^{**} ,

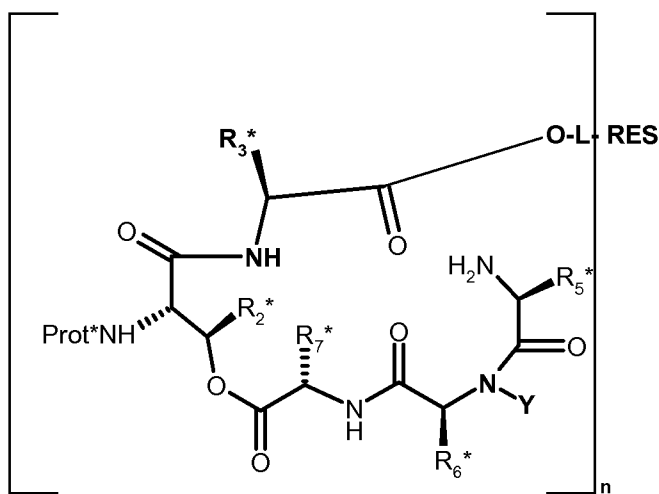
and preferably

(x/a) finally coupling the resulting compound of the formula XVIII,



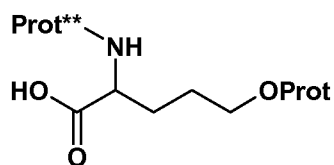
XVIII

especially XVIIIA,



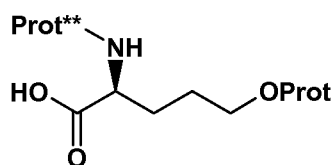
XVIII A

wherein Y, R₂^{*}, R₃^{*}, R₇^{*}, R₆^{*} and R₅^{*} are as defined for a compound of the formula II above, Prot^{*} is as defined for a compound of the formula VIII above and n, L and RES are as defined for a compound of the formula X above, to an unnatural amino acid (= synthon) of the formula XIX,



XIX

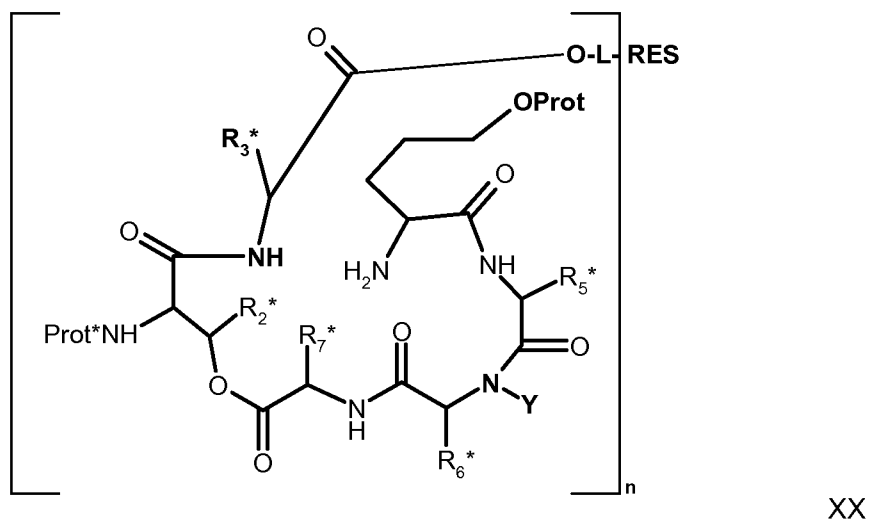
especially the formula XIXA,



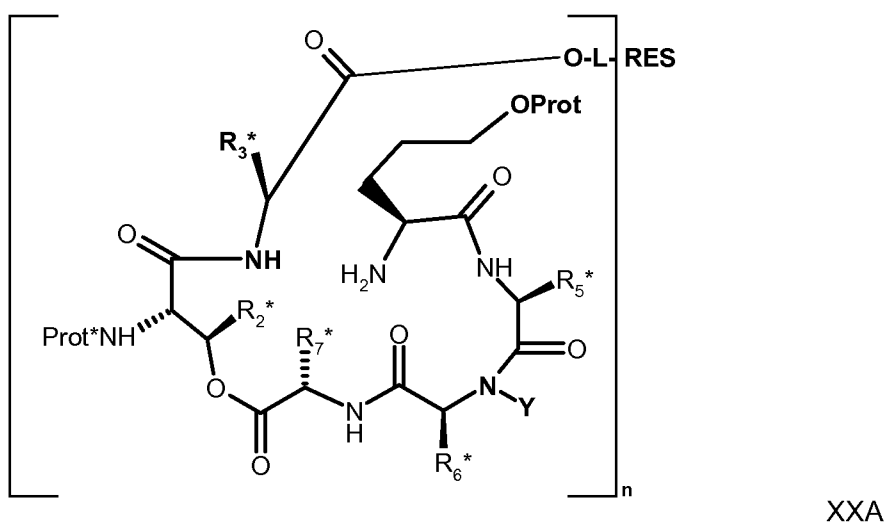
XIXA

wherein Prot is as defined for a compound of the formula II above and Prot^{**} is as defined for a compound of the formula IX, or an activated derivative of said synthon, and removing the protecting group Prot^{**} to yield a compound of the formula XX,

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especially XXA,



wherein Prot, Y, R_2^* , R_3^* , R_7^* , R_6^* and R_5^* are as defined for a compound of the formula II above, Prot* is as defined for a compound of the formula VIII above and n, L and RES are as defined for a compound of the formula X,

and

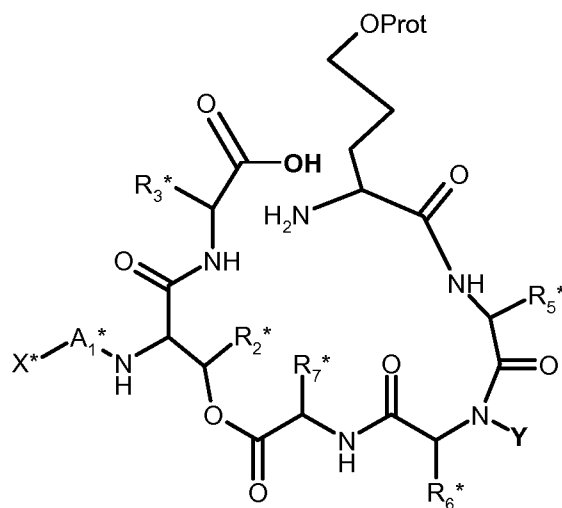
(xi/a) cleaving the solid phase bound peptide in formula XX off the solid phase L-RES to yield the corresponding compound of the formula VIII, especially VIIIA, as shown above.

Another embodiment of the invention relates to the synthesis of a compound of the formula II as given above according to section (i/a), preferably preceded by the reaction

according to section (ii/a) or more preferably according to section (iii/a); preferably preceded by the reaction according to section (iv/a) or preferably (v/a), preferably preceded by the reaction according to section (vi/a), preferably preceded by the reaction according to section (x/a), preferably preceded by the reaction according to section (ix/a), preferably preceded by the reaction according to section (viii/a), preferably preceded by the reaction according to section (vii/a).

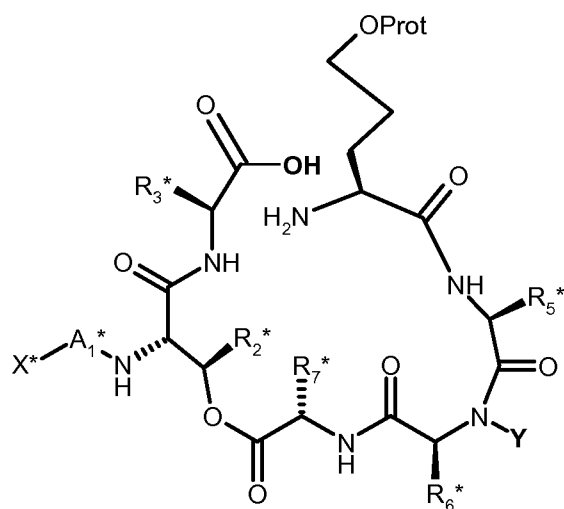
(i/b) Another embodiment of the invention relates to a method or process above, comprising, for the synthesis of the compound of the formula II given above, cyclization under lactamization of a linear, not yet cyclic, precursor peptide of the compound of the formula II, carrying an N-terminal amino group and a C-terminal carboxy group, under reaction conditions that allow for the formation of an amide bond from said amino and said carboxy group, preferably using Solution Phase chemistry.

(ii/b) A further embodiment of the invention relates to the method or process according to the preceding paragraph (i/b), where the linear precursor peptide is of the formula XXV,



XXV

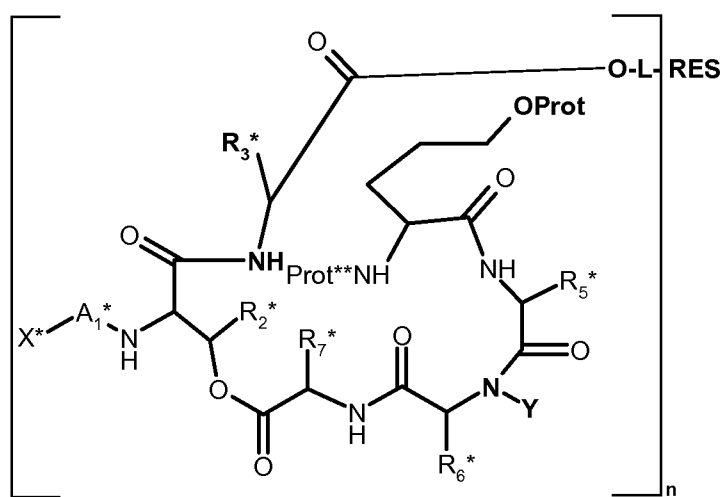
especially XXVA,



XXVA

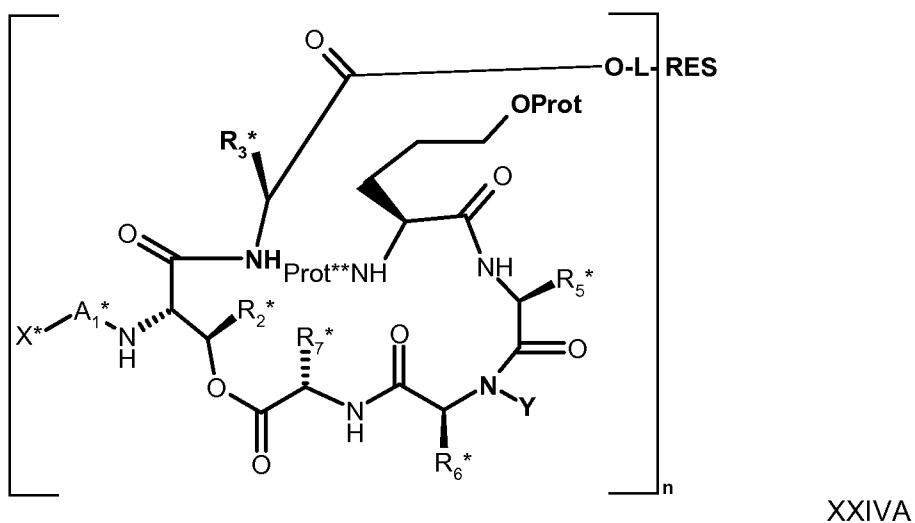
wherein X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , R_7^* and Prot are as defined for a compound of the formula II above.

(iii/b) Another embodiment refers to the method or process according to the preceding paragraph (ii/b), further comprising, for the synthesis of the compound of the formula XXV, cleaving a compound of the formula XXIV,



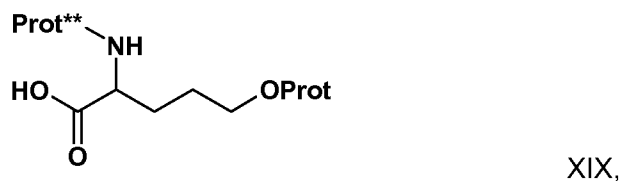
XXIV

especially XXIVA,

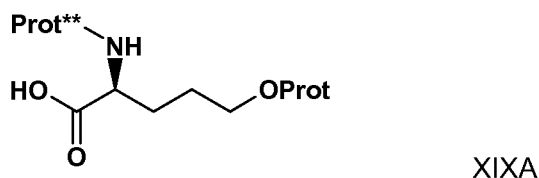


wherein X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , R_7^* and Prot are as defined for a compound of the formula II above, L is a cleavable linker, RES is a solid resin, n is a natural number and Prot** is an amino protecting group that can be removed without parallel removal of the protecting group Prot and with the product remaining on the resin, and (before the cleavage, in parallel or subsequently to it) removing the protecting group Prot** to yield the compound of the formula XXV.

(iv/b) A further embodiment of the invention relates to the method or process according to the preceding paragraph (iii/b), further comprising, for the synthesis of the compound of the formula XXIV, coupling an amino acid of the formula XIX,

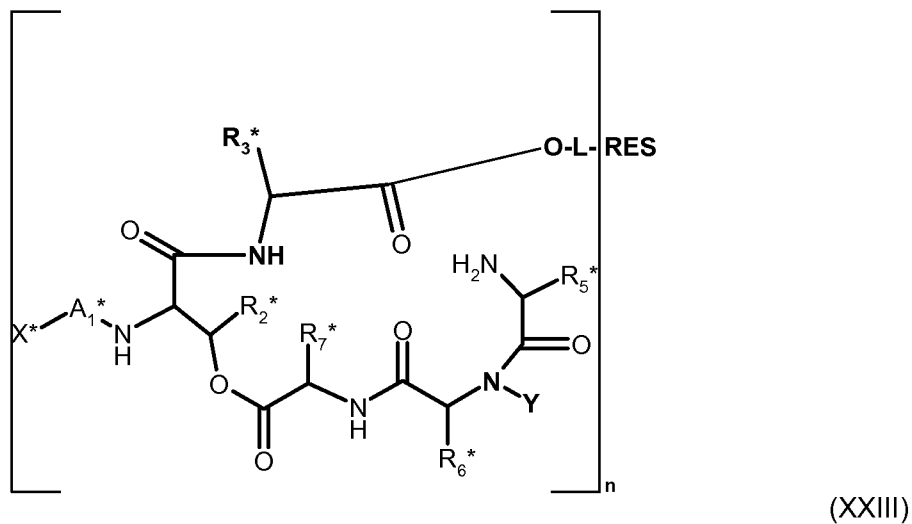


especially XIXA,

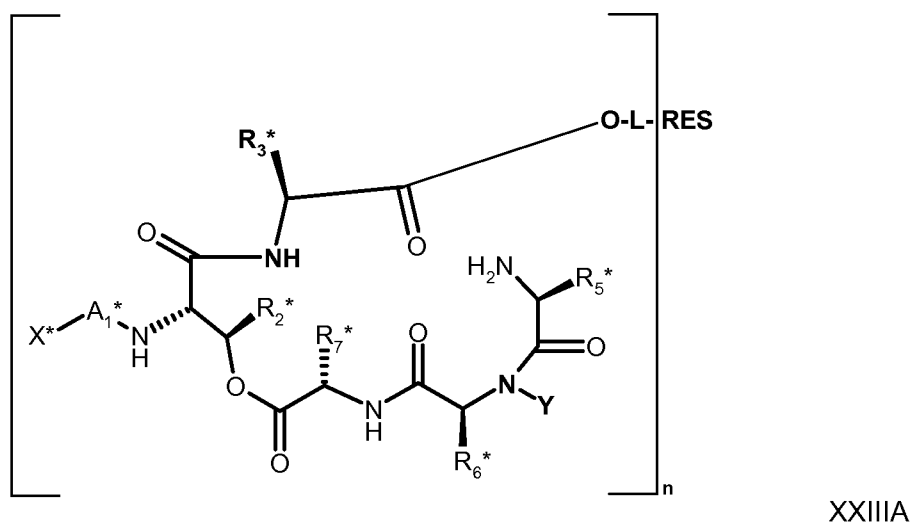


- 24 -

wherein Prot is as defined for a compound of the formula II above and Prot** is as defined for a compound of the formula XXIV above, or an activated derivative of said amino acid, with a compound of the formula XXIII,



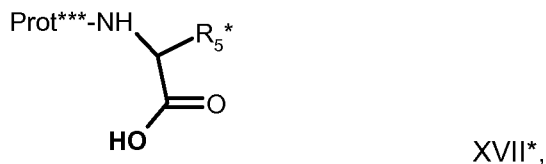
especially XXIIIA,



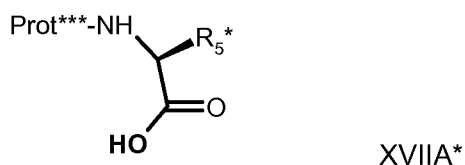
wherein X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* and R_7^* are as defined for a compound of the formula II above, L is a cleavable linker, RES is a solid resin, and n is a natural number.

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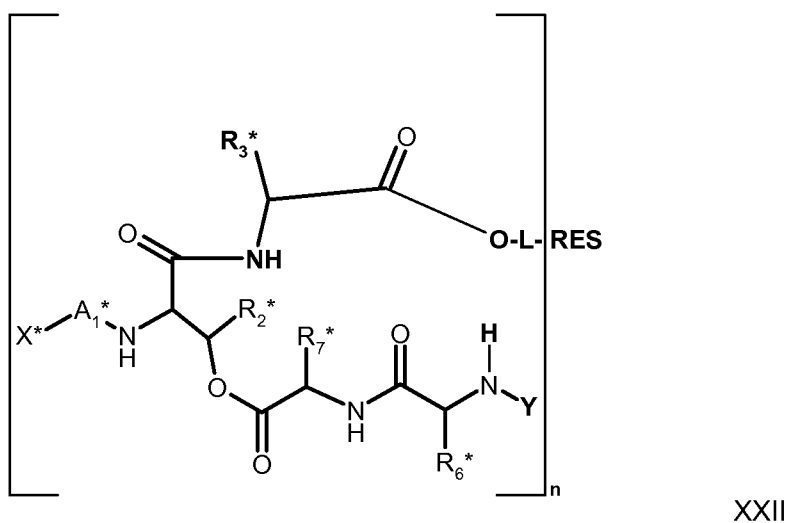
(v/b) Yet a further embodiment of the invention relates to the method or process according to the preceding paragraph (iv/b), further comprising, for the synthesis of the compound of the formula XXIII, coupling an amino acid of the formula XVII*



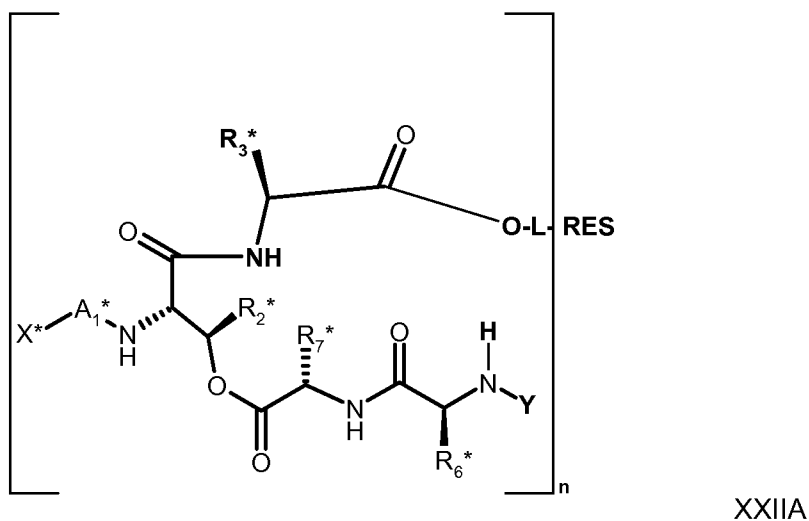
especially XVIIA*,



wherein R_5^* is as defined for a compound of the formula II above and Prot^{***} is an amino protecting group that can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin, or a reactive derivative of said amino acid, with a compound of the formula XXII,

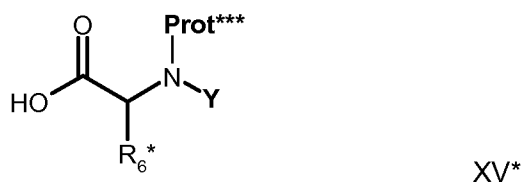


especially XXIIA

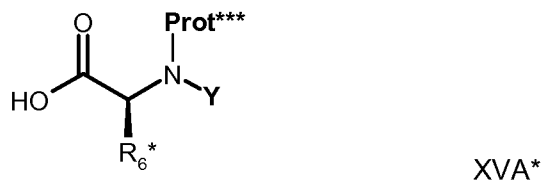


wherein X^* , A_1^* , R_2^* , R_3^* , R_6^* and R_7^* are as defined for a compound of the formula II above, L is a cleavable linker, RES is a solid resin, and n is a natural number, and removing the protecting group Prot***.

(vi/b) In yet a further embodiment, the invention relates to the method or process according to the preceding paragraph (v/b), further comprising, for the synthesis of the compound of the formula XXII, coupling an amino acid of the formula XV*,



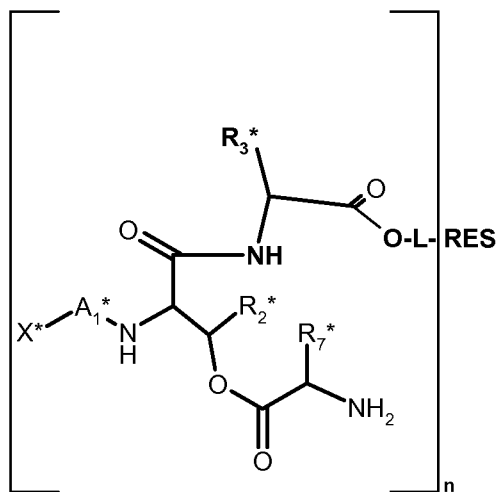
especially XVA*



in which R_6^* and Y are as defined for a compound of the formula II above and Prot*** is an amino protecting group that can be cleaved off selectively without affecting other

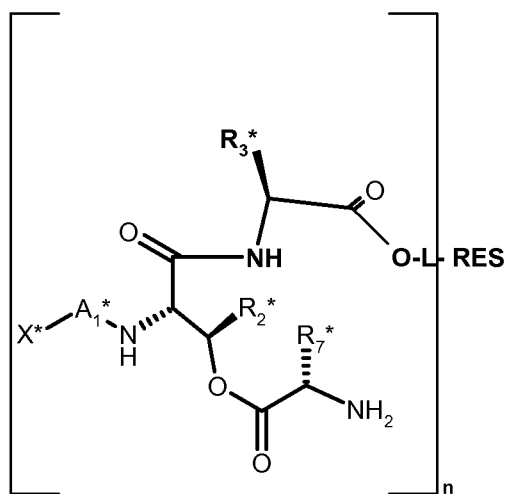
- 27 -

protecting groups present and with the product remaining on the resin, or a reactive derivative of said amino acid, with a compound of the formula XXI,



XXI,

especially XXIA,

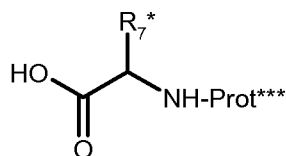


XXIA

wherein X^* , A_1^* , R_2^* , R_3^* and R_7^* are as defined for a compound of the formula II above, L is a cleavable linker, RES is a solid resin, and n is a natural number, and removing the protecting group Prot***.

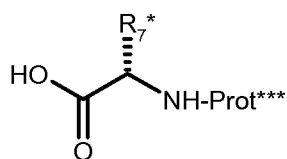
- 28 -

(vii/b) Another embodiment of the invention relates to the method or process according to the preceding paragraph (vi/b), further comprising, for the synthesis of a compound of the formula XXI, reacting an amino acid of the formula XIII*,



XIII*

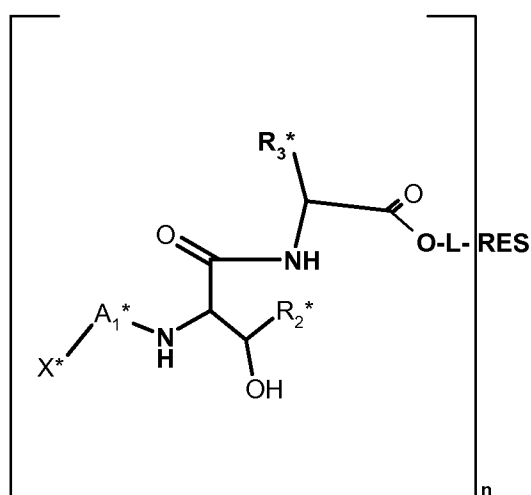
especially XIII A*,



XIII A*

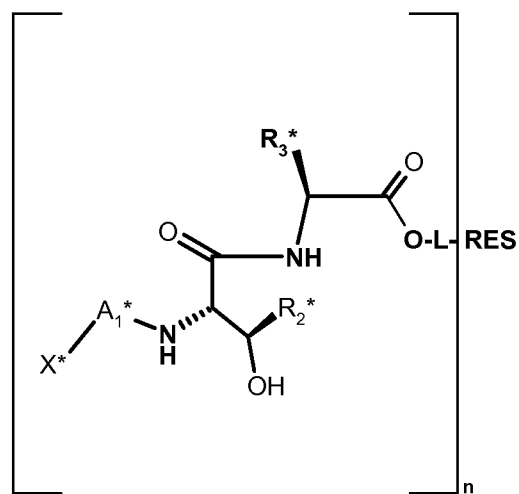
wherein Prot*** is an amino protecting group that can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin, and R₇* is as defined for a compound of the formula II above, or a reactive derivative of said amino acid,

with the hydroxyl group of a compound of the formula XXVI,



XXVI

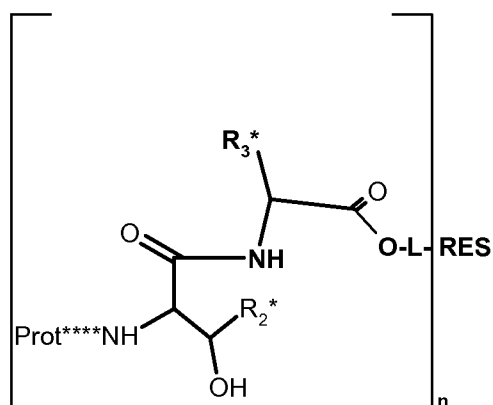
especially XXVIA,



XXVIA

wherein X^* , A_1^* , R_2^* and R_3^* are as defined for a compound of the formula II above, L is a cleavable linker, RES is a solid resin, and n is a natural number; and removing the protecting group Prot***.

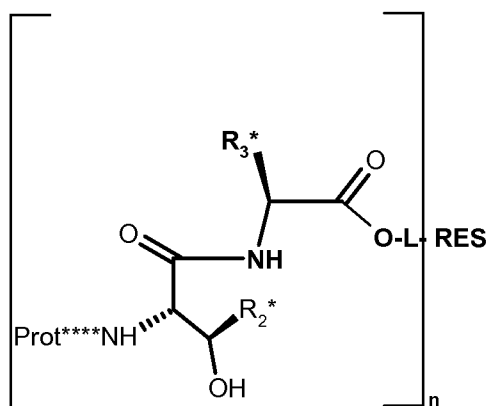
(viii/b) In a further embodiment, the invention relates to the method or process according to the preceding paragraph (vii/b), further comprising, for the synthesis of a compound of the formula XXVI, especially XXVIA, coupling a resin bound dipeptide symbolized by the formula XII*,



XII*

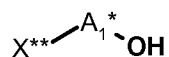
- 30 -

especially XIIA*



XIIA*

in which Prot**** is a protecting group that can be cleaved off selectively without affecting other protecting groups present in a compound of the formula II as defined above and with the product remaining on the resin, R_2^* and R_3^* are as defined for a compound of the formula II above, L is a cleavable linker, RES is a solid resin, and n is a natural number, after removal of the protecting group Prot**** via the thus obtainable free amino group, with an acid of the formula VII



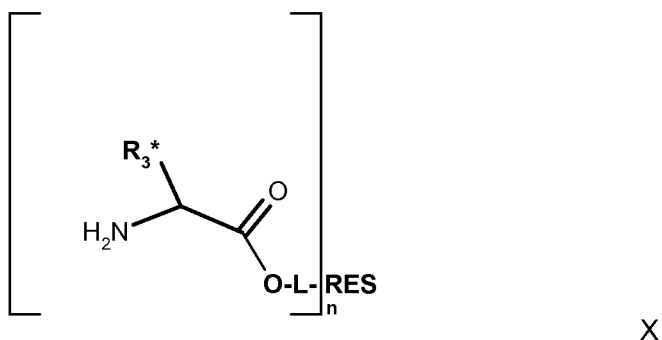
VII

wherein X^{**} is an amino protecting group or is X^* , and wherein X^* and A_1^* are as defined for a compound of the formula II above, or a reactive derivative of said acid;

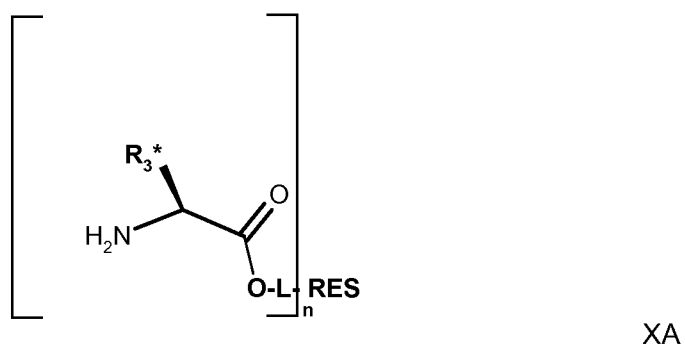
and, if X^{**} is an amino protecting group, removing said amino protecting group X^{**} to yield the derivative of formula II wherein, instead of X^* , H is present and coupling the resulting amino group with an acyl group X^* using the corresponding acid X^*-OH wherein X^* is as defined for a compound of the formula II above, or a reactive derivative of said acid.

(ix/b) A yet further embodiment of the invention relates to the method or process according to the preceding paragraph (viii/b), further comprising, for the synthesis of a compound of the formula XII, coupling a resin bound amino acid symbolized by the formula X,

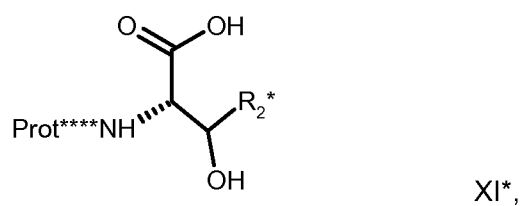
- 31 -



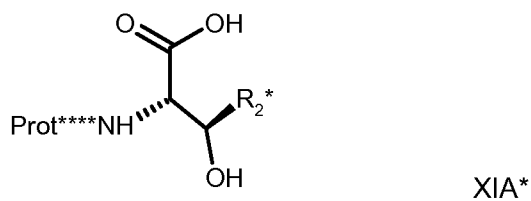
especially XA,



wherein R_3^* is as defined for a compound of the formula II above, L is a cleavable linker, RES is a solid resin, and n is a natural number, with an amino acid of the formula XI*,



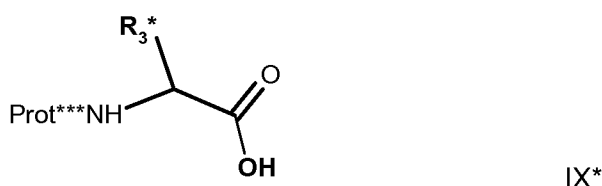
especially XIA*,



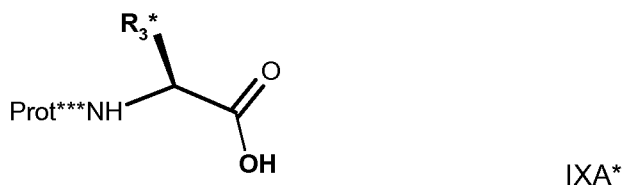
wherein Prot^{****} is a protecting group that can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin, and R_2^* is

as defined for a compound of the formula II above, or a reactive derivative of said amino acid.

(x/b) A further embodiment of the invention relates to the method or process according to the preceding paragraph (ix/b), further comprising, for obtaining the resin bound amino acid of the formula X, coupling an amino acid of the formula IX*,



especially IXA*,



wherein R_3^* is as defined for a compound of the formula II in claim 1 and Prot*** is an amino protecting group can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin ; or a reactive derivative of said amino acid of the formula IX, to a cleavable linker L which is bound to a solid resin RES, and removing the protecting group Prot***.

(i/c) Another embodiment of the invention relates to the method or process according to any one of the preceding paragraphs (i/a) to (x/b) where the symbols A_1 , R_2 , R_3 , R_5 , R_6 , R_7 , X and Y or the corresponding unprotected or protected moieties R_2^* , R_3^* , R_5^* , R_6^* , R_7^* , X^* and Y_3 are selected so that, in the resulting compound of the formula I, or a salt thereof,

A_1 is the bivalent radical of L-glutamine bound via the carbonyl of its α -carboxy group to the amino group at the right of A_1 in formula I and via its α -amino group to X, or is 2S-(2-hydroxy-3-phosphonooxy)-propionyl;

R_2 is methyl;

R_3 is isopropyl, isobutyl (2-methyl-n-propyl wherever used) or benzyl, especially isobutyl;

R_5 is sec-butyl or benzyl, especially sec-butyl;

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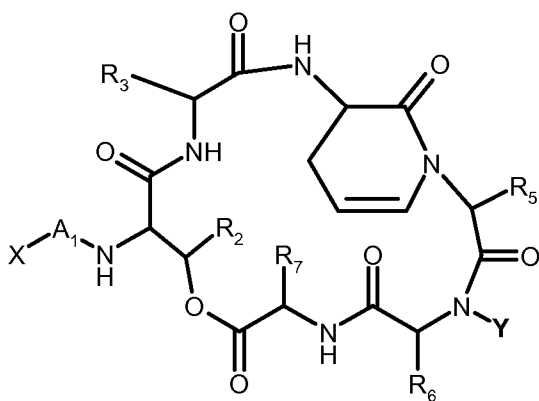
R₆ is 4-hydroxybenzyl;

R₇ is isopropyl or sec-butyl (1-methyl-n-propyl wherever used), especially sec-butyl;

X is acetyl or isobutyryl, or is absent if A₁ is 2S-(2-hydroxy-3-phosphonoxy)-propionyl and

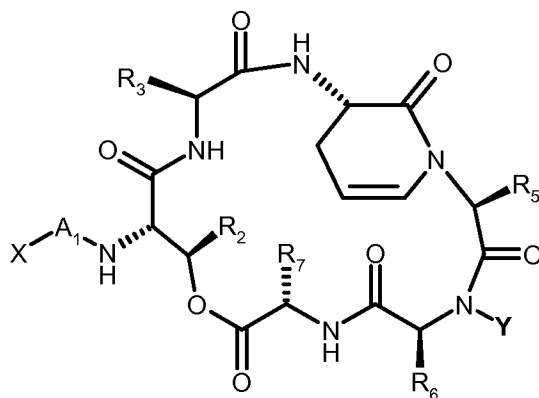
Y is methyl. This paragraph is also named paragraph a) below.

(i/d) In another particular embodiment, the invention relates to a method or process for converting a dehydrate of a compound of the formula I given above or in particular with the substituents as defined in the preceding paragraph (i/c) into the corresponding compound of the formula I, where the dehydrate has the formula V,



V,

especially VA,

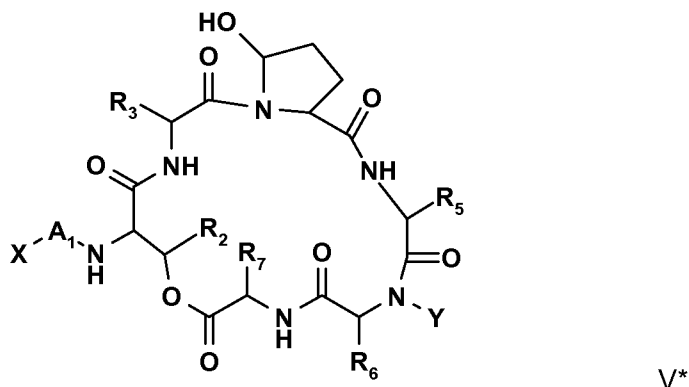


VA

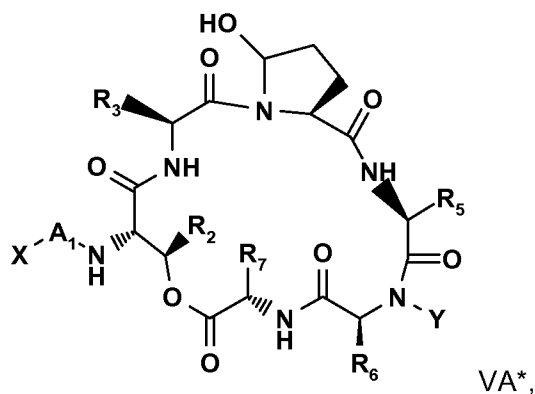
in which Y, X, A₁, R₂, R₃, R₅, R₆ and R₇ are as defined for a compound of the formula I above;

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or especially a method or process for shifting the equilibrium of a mixture of a compound of the formula I and its corresponding dehydrate, and/or its corresponding hemiaminal analogue with a five-ring instead of the ahp structure in formula I which may also be formed as byproduct and has the formula V*,



especially the formula VA*,



in which Y, X, A₁, R₂, R₃, R₅, R₆ and R₇ are as defined for a compound of the formula I above, respectively;

in favor of the compound of the formula I,

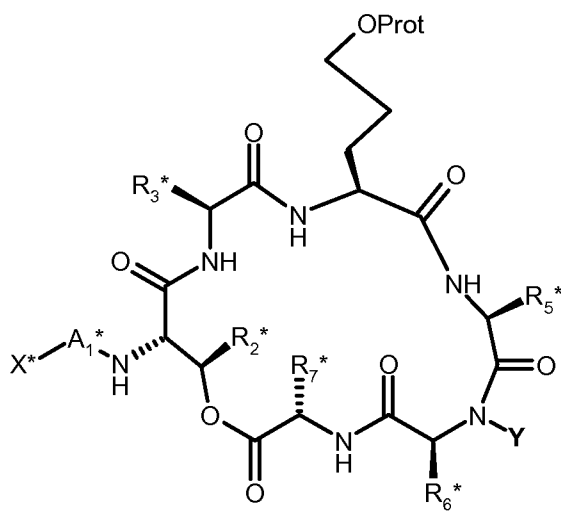
said method or process comprising using an aqueous acid as reactive solvent to drive the reaction. This method can be used independently (e.g. also for the product of fermentation or biosynthesis) or in addition to the other processes or methods described above and below to increase the yield or to re-convert a compound of the formula V,

especially VA, and/or the analogue with a five-membered ring instead of the ahp structure in formula I, into the corresponding compound of the formula I.

The method described for the conversion of the dehydrate and /or the five ring analogue (always regarding the desired ahp ring) into the desired compound of the formula I or especially IA, e.g. of Compound A -dehydrate from Example 3 B into Compound A, enables a straight-forward synthesis of this class of compounds. Up to now, an acidic treatment as final step had to be circumvented in order to avoid the dehydration of the product.

(i/e) A further embodiment of the invention relates to the method according to the preceding paragraph (i/d), wherein the acid is a carboxylic acid, especially a halo substituted C₁₋₈alkanoic acid, more especially trifluoroacetic acid or trichloroacetic acid.

(i/f) The invention, in yet a further embodiment, relates to a compound of the formula II,



II

wherein Prot is a protecting group, Y is as defined for a compound of the formula I in the first instance above or in particular as in claim 17 and X*, A₁*, R₂*, R₃*, R₅*, R₆*, and R₇* correspond to X, A₁, R₂, R₃, R₅, R₆, and R₇ in formula I as defined in claim 1 or in paragraph (ia) given above, respectively, however with the proviso that reactive functional groups on these moieties are present in protected form.

(i/g) In a further embodiment, the invention relates to a novel compound selected from the group consisting of compounds of the formula II, III, IV, V, VI, VIII, X, XII, XIV, XVI,

XVIII, XIX, XX, XXI, XXII, XXIII, XXIV, XXV, XXVI and XXVIII, and especially of the formula IIA, IIIA, IVA, VA, VIA, VIIIA, XA, XIIA, XIVA, XVIA, XVIIIA, XIXA, XXA, XXIA, XXIIA, XXIIIA, XXIVA, XXVA, XXVIA and XXVIIIA, yet more especially to the group consisting of the following compounds given in the examples: From Scheme 1: , compound 2, compound 3, compound 4, synthon 1; from Scheme 2: compound 5; Fmoc-Leu-Linker-Resin according to Example 1B(2); Fmoc-Thr-Leu-Linker-Resin according to Example 1B(3); Fmoc-Gln(Trt)-Thr-Leu-Linker-Resin according to Example 1B(4); Isobutyryl-Gln(Trt)-Thr-Leu-Linker-Resin according to Example 1B(5); Isobutyryl-Gln(Trt)-Thr(Ile-Fmoc)-Leu-Linker-Resin according to Example 1B(6); the product of Example 1B(7) = Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Fmoc)-Leu-Linker-Resin (previously named: Isobutyryl-Gln(Trt)-Thr(Ile-N-me-Tyr(tBu)-Fmoc)-Leu-Linker-Resin); Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Fmoc)-Leu-Linker-Resin (previously named: Isobutyryl-Gln(Trt)-Thr(Ile-N-me-Tyr(tBu)-Ile-Fmoc)-Leu-Linker-Resin) according to Example 1B(8); Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-Linker-Resin (previously named: Isobutyryl-Gln(Trt)-Thr(Ile-N-Me-Tyr(tBu)-Ile-Synthon 1-H)-Leu-Linker-Resin) according to example 1B(9). Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-OH (previously named: Isobutyryl-Gln(Trt)-Thr(Ile-N-Me-Tyr(tBu)-Ile-Synthon 1-H)-Leu-OH)according to example 1B(10) and Scheme 3; H-Thr-Leu-Resin according to example 1B(12); H-Gln(Trt)-Thr-Leu-Resin according to example 1B(13); Isobutyryl-Gln(Trt)-Thr-Leu-Resin according to example 1B(14); Isobutyryl-Gln(Trt)-Thr(Ile-H)-Leu-Resin according to example 1B(15); Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-H)-Leu-Resin according to example 1B(16); Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-H)-Leu-Resin according to example 1B(17); Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-Resin according to example 1B(18); from Scheme 3, Compound 6 and/or 7 (the latter preferred); from Scheme 4: Compound 8 and the 5-ring hemiaminal-isomer; from Scheme 5, precursor peptide 2, compound 9, compound 10 (this one being preferred in the present enumeration regarding Scheme 5), and/or compound 11; Fmoc-Thr-Leu-Trt-Tentagel-S from Example 2A(1); Fmoc-Gln(Trt)-Thr-Leu-Trt-Tentagel-S according to Example 2A(2); Ac-Gln(Trt)-Thr-Leu-Trt-Tentagel-S according to Example 2A(3); Ac-Gln(Trt)-Thr(Val-Fmoc)-Leu-Trt-Tentagel-S according to Example 2A(4); Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Fmoc)-Leu-Trt-Tentagel-S (previously named: Ac-Gln(Trt)-Thr(Val-N-Me-Tyr(tBu)-Fmoc)-Leu-Trt-Tentagel-S) according to Example 2A(5); Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Phe-Fmoc)-Leu-Trt-Tentagel-S (previously named: Ac-Gln(Trt)-Thr(Val-N-Me-Tyr(tBu)-Phe-Fmoc)-Leu-Trt-Tentagel-S)

according to Example 2A(6); and Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Phe-Synthon1-H)-Leu-OH (previously named: Ac-Gln(Trt)-Thr(Val-N-Me-Tyr(tBu)-Phe-Synthon1-H)-Leu-OH) (precursor peptide 2) according to Example 2A(7).

The following definitions (or also definitions already included above) can replace more general terms used in invention embodiments above and below in order to define further embodiments of the invention, with either one, two or more or all general terms being replaceable by the more specific terms in order to define such invention embodiments:

A bivalent moiety of an amino acid with a terminal carboxy or carbamoyl group is preferably an alpha-carbamoyl or carboxyl- C_{1-8} -substituted amino acid, especially the bivalent moiety of asparagine or glutamine, and is bound at its right hand side in formula I via a carbonyl (preferably the carbonyl of its α -carboxyl group) to the rest of the molecule.

C_{1-8} -alkanoyl or phosphorylated hydroxy- C_{1-8} -alkanoyl (C_{1-8} -alkanoyl carrying both a hydroxyl and a phosphono ($-O-P(=O)(OH)_2$) group) A_1 is e.g. 2,3-dihydroxy-propanoyl (preferably in S-form) or 2-hydroxy-3-phosphono-propanoyl (preferably in S-form).

R_2 and R_2^* are C_{1-8} -alkyl, especially methyl wherever mentioned.

R_3 is the side chain of an amino acid, especially of a natural amino acid. Preferably, it is C_{1-8} alkyl which may be branched or linear. Most especially, C_{1-8} alkyl is n-(2-methyl)propyl (isobutyl), n-(1-methylpropyl (sec-butyl) or methyl, that is, the amino acid carrying the moiety is leucine, isoleucine or valine.

R_3^* is the corresponding side chain in protected form if a functional group is present that has to be hindered to participate in a reaction. Preferably, it is C_{1-8} alkyl which may be branched or linear, especially as defined in the preceding paragraph.

A "side chain of an amino acid" may be selected from any moiety, e.g. a mono- or polycyclic, linear, saturated, unsaturated (e.g. with conjugated double bonds) or partially saturated organic moiety, e.g. with up to 20 carbon atoms and 0 to 5 heteroatoms in the basis structure independently selected from N, O and S replacing the corresponding

number of carbon atoms, and may be substituted by up to three moieties selected from amino, imino, hydroxy, carboxy, carbamoyl, sulfhydryl, amidino, guanidino, O-phosphono ($-O-P(=O)(OH)_2$). Preferably, the side chains are selected from those of the 20 standard alpha-amino acids arginine, histidine, lysine, aspartic acid, glutamic acid, serine, threonine, asparagine, glutamine, cysteine, glycine, alanine, leucine, isoleucine, methionine, phenylalanine, tryptophan, tyrosine, valine and further proline (then with internal cyclization including the alpha-amino group).

For the amino acids, either their names or the customary three letter codes are used in the present disclosure, in accordance with the following table:

Amino acid	Three letter code
Alanine	Ala
Arginine	Arg
Asparagine	Asn
Aspartic acid	Asp
Asparagine or aspartic acid	Asx
Cysteine	Cys
Glutamic acid	Glu
Glutamine	Gln
Glutamine or glutamic acid	Glx
Glycine	Gly
Histidine	His
isoleucine	Ile
Leucine	Leu
Lysine	Lys
Methionine	Met
Phenylalanine	Phe
Proline	Pro
Serine	Ser
Threonine	Thr
Tryptophan	Try
Tyrosine	Tyr
Valine	Val

R₅ is the side chain of an amino acid, preferably a standard amino acid. Preferably, it is C₁₋₈alkyl which may be branched or linear and which is unsubstituted or substituted by phenyl. Most especially it is benzyl, n-(2-methyl)propyl, isobutyl or methyl, that is, the amino acid carrying the moiety is phenylalanine, leucine, isoleucine or valine.

R₆ is the side chain of a hydroxy amino acid, especially of tyrosine.

R₇ is the side chain of an amino acid, especially of a natural amino acid. Preferably, it is C₁₋₈alkyl which may be branched or linear. Most especially it is n-(2-methyl)propyl (isobutyl), n-(1-methyl)propyl (sec-butyl) or methyl, that is, the amino acid carrying the moiety is leucine, isoleucine or valine.

C₁₋₈-alkyl can be linear or branched one or more times; for example, it can be n-(2-methyl)propyl, n-(1-methyl)propyl or methyl.

All of the compounds can, where salt-forming groups such as basic groups, e.g. amino or imino, or acidic groups, e.g. carboxyl or phenolic hydroxyl, are present, be used in free form or as salts or as mixtures of salts and free forms. Thus where ever a compound is mentioned, this includes all these variants. For example, basic groups may form salts with acids, such as hydrohalic acids, e.g. HCl, sulfuric acid or organic acids, such as acetic acid, while acidic groups may form salts with positive ions, e.g. ammonium, alkylammonium, alkali or alkaline-earth metal salt cations, e.g. Ca, Mg, Na, K or Li cations, or the like.

“Or the like” or “and the like”, wherever used in this disclosure, refers to the fact that other alternatives to those mentioned preceding such expression are known to the person skilled in the art and may be added to those expressions specifically mentioned; in other embodiments, “or the like” and “and the like” may be deleted in one or more or all invention embodiments.

The protecting groups Prot, Prot*, Prot**, Prot***, Prot**** and any further protecting groups present on the moieties A*, R₂*, R₃*, R₅*, R₆*, R₇*, X*, where ever mentioned throughout the present description and claims, are selected so that they allow for orthogonal protection.

Orthogonal protection is a strategy allowing the deprotection of multiple protective groups one (or more but not all) at the time where desired each with a dedicated set of reaction conditions without affecting the other protecting group(s) or bonds to resins, e.g. via linkers on solid synthesis resins. In other terms: The strategy uses different classes of protecting groups that are removed by different chemical mechanisms, also using appropriate linkers in the case of solid phase peptide synthesis (where the linker-resin bond might together be considered as a carboxy protecting group).

Preferably, the protecting groups are selected as follows:

The protecting group Prot is preferably selected so as to resist removal of any other protecting groups used or present during the synthesis according to the invention of a depsipeptide, e.g. able to resist mild bases (see Prot*), but removable with fluoride ion (especially under anhydrous conditions), e.g. $\text{Bu}_4\text{N}^+\text{F}^-$ (also if created *in situ*, e.g. using $\text{Bu}_4\text{N}^+\text{Cl}^-$ with $\text{KF} \cdot \text{H}_2\text{O}$, KF with 18-crown-6, LiBr with 18-crown-6, $\text{BF}_3 \cdot \text{diethylether}$, pyridine-HF, HF in urea, $\text{Et}_3\text{N}(\text{HF})_3$ (wherein Et is ethyl) or the like, where the solvent is e.g. selected from the group consisting of N,N-dimethylformamide, acetonitrile, chloroform and tetrahydrofuran.

Preferably, Prot is an ether protecting group, especially selected from the group consisting of silyl protecting groups in which the silyl moiety carries up to three organic moieties bound via a carbon (optionally via a further Si atom), such as tert-butyldiphenylsilyl, trimethylsilyl, triisopropylsilyl, tert-butyldimethylsilyl, triphenylsilyl, diphenylmethylsilyl, ti-tert-butyldimethylsilyl, tert-butylmethoxyphenylsilyl, tris(trimethylsilyl)silyl or the like.

Prot* is a protecting group that can be cleaved off selectively without affecting other protecting groups present, and also without affecting the depsipeptide forming ester bond or a linker to a resin RES, and is stable during deprotection steps during synthesis of the linear precursor peptide (e.g. removal of allyloxycarbonyl); it is preferably a protecting group removable by specific triphenylphosphin complexes in the presence of metal hydrides or other reductants, e.g. $(\text{PH}_3\text{P})_4\text{Pd}$ preferably in combination with di-n-butyl tin hydride or tri-n-butyl tin hydride, phenylsilane, sodium borohydride or dimesitylene, in an appropriate solvent, e.g. tetrahydrofuran, and is preferably not cleavable under

conditions that allow for the removal of a protecting group Prot^{**}; for example, and Prot^{*} is selected from the group consisting of C₃-C₈alk-2-enyloxycarbonyl moieties, e.g. allyloxycarbonyl (Alloc), 1-isopropylallyloxycarbonyl, 4-nitrocinnamyloxycarbonyl and 3-(3'-pyridyl)prop-2-enyloxycarbonyl.

Prot^{**} is a protecting group that can be removed on the resin without cleaving other bonds (no cleavage of an amino acid or peptide bound via the carbonyl of its (especially α -carboxyl group to the binding via a linker L mentioned below; also without cleaving off the protecting group Prot once present), especially a protecting group removable without cleavage of an ester (instead of an amide) bond in a depsipeptide or depsipeptide precursor and under conditions other than those for the protecting groups Prot^{*} and Prot, while preserving the binding via the linker to a resin RES where present; it is preferably removable by a mild base e.g. piperidine, morpholine, dicyclohexylamine, p-dimethylamino-pyridine, diisopropylamine, piperazine, tris-(2-aminoethyl)amine in an appropriate solvent, e.g. N,N-dimethylformamide, methylene chloride; Prot^{**} is, e.g., selected from the group consisting of fluoren-9-ylmethoxycarbonyl (Fmoc); 2-(2' or 4'-pyridyl)ethoxycarbonyl and 2,2-bis(4'-nitrophenyl)ethoxycarbonyl.

Prot^{***} (an amino protecting group that can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin) and Prot^{****} (a protecting group that can be cleaved off selectively without affecting other protecting groups present in a compound of the formula II as defined above and below and with the product remaining on the resin) are preferably protecting groups that can be removed as Prot^{**} and are, e.g., selected from those mentioned for R^{**}, e.g. fluoren-9-ylmethoxycarbonyl (Fmoc).

The preferred orthogonal synthesis method in this case makes use of the Fmoc method known in general for peptide synthesis using solid phase peptide synthesis combined with solution phase macrolactamization and further chemical conversions.

Alternatively, e.g. the Boc protecting group might be used instead of Fmoc Prot^{**}, Prot^{***} and Prot^{****}.

However, this will require different side chain protecting groups and also the hydroxy-group of N-methyl-Tyr will then have to be protected in a different way to maintain orthogonality of the protection groups.

Other protecting groups present as well as the binding linker to a resin RES where present are preferably not removable under conditions under which Prot* and Prot** can be removed, e.g. in A*, the amide can be N-protected e.g. with trityl (triphenylmethyl) (cleavage e.g. with trifluoro acetic acid (TFA); in R₆* a tyrosine hydroxy can be protected as t-butyl-ether, or protected by tert-butyldimethylsilyl, methoxymethyl, Boc (tert-butoxycarbonyl) or arylacetate (cleavage with TFA).

Appropriate protecting groups are known in the art, as well methods for their introduction and removal. For example, the protecting groups, their introduction and removal methods may be selected from those described in standard textbooks such as "Protective Groups in Organic Synthesis", 3rd ed., T.W. Green and P.G.M. Wuts (Eds.). J. Wiley & Sons, Inc., New York etc. 1999.

The protecting groups Prot, Prot*, Prot**, Prot***, Prot**** and other protecting groups are thus not limited to those mentioned above – rather they should fulfill conditions that make them appropriate for orthogonal protection, e.g. as described above or below.

It is recommended to avoid too basic conditions (though the bases described for Fmoc cleavage, such as piperidine, are usually allowable) to avoid cleavage of the depsipeptide (ester) bond.

Among the possible solid support for Solid Phase Peptide Synthesis (SPPS), the following may be mentioned:

- Gel-type supports without or with spacer: These are highly solvated polymers with an equal distribution of functional groups. This type of support is the most common, and includes:

Polystyrene: Styrene cross-linked with e.g. 1-2% divinylbenzene; Polyacrylamide or polymethacrylamide: as hydrophilic alternative to polystyrene; Polyethylene glycol (PEG): PEG-Polystyrene (PEG-PS) is more stable than polystyrene and spaces the site

of synthesis from the polymer backbone; PEG-based supports: Composed of a PEG-polypropylene glycol network or PEG with polyamide or polystyrene (these already include a spacer, PEG);

- Surface-type supports: Materials developed for surface functionalization, including controlled pore glass, cellulose fibers, and highly cross-linked polystyrene.
- Composites: Gel-type polymers supported by rigid matrices.

Usually these gels carry reactive groups to which a linker L as mentioned for various precursors above and below can be bound. For example, such groups include aminomethyl groups, polyethyleneglycol groups with a terminal hydroxy, and the like.

Any such support can be used in the embodiments of the present invention.

Gel type supports are used in another special embodiment of the invention, Among these, polystyrene (divinylbenzene crosslinked); polyacrylamide and polymethacrylamide resins are especially preferred.

Among the possible linkers, all commonly known and appropriate may be used.

Examples in possible embodiments of the invention are the 2-methoxy-4-benzyloxy-benzyl alcohol linker (a Sasrin-Linker, Sasrin stands for superacid sensitive resin, binds the amino acids or peptides via alcoholic OH); the trityl linker family (e.g. Trityl, 2Cl-Trityl, which bind the amino acids or peptides via OH); the 4-(2,4-dimethoxyphenylhydroxymethyl)phenoxymethyl-Linker (Rink-Acid-Linker, binds the amino acids or peptides via OH); or tris(alkoxy)benzyl ester linkers (HAL-Linker, binds the amino acids or peptides via OH).

Where reactive derivatives of acids, especially amino acids, or peptides, e.g. dipeptides, are mentioned, they may be formed *in situ* or may be used as such.

Reactive (or active) derivatives used as such include the acyl-halides, e.g. acyl-chlorides, -fluorides or -nitrophenyl esters, e.g. the 2,4-dinitrophenyl esters, or acid anhydrides (symmetric or e.g. with acetic acid) of the carboxy groups of the acids to be reacted.

For *in situ* amino acid activation, customary coupling agents may be applied. Such reagents are known to the person skilled in the art and can be purchased conveniently from many sources, e.g. Aldrich ChemFiles – Peptide Synthesis (Aldrich Chemical Co., Inc., Sigma-Aldrich Corporation, Milwaukee, WI, USA) Vol. 7 No. 2, 2007 (see http://www.sigmaaldrich.com/etc/medialib/docs/Aldrich/Brochure/al_chemfile_v7_n2.Par.0001.File.tmp/al_chemfile_v7_n2.pdf). Among the possible coupling agents for amide and ester bond synthesis the following may be mentioned:

Triazoles, uronium or hexafluorophosponium derivatives, e.g. 1-hydroxy-benzotriazole (HOBt), 1-hydroxy-7-aza-benzotriazole (HOAt), ethyl 2-cyano-2-(hydroxyimino)acetate, 2-(1H-7-Azabenzotriazol-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate methanaminium (HATU), benzotriazol-1-yl-oxytripyrrolidinophosphonium hexafluorophosphate (PyBOP), Benzotriazole-1-yl-oxy-tris-(dimethylamino)-phosphonium hexafluorophosphate (BOP), 1-(mesitylene-2-sulfonyl)-3-nitro-1,2,4-triazole (MSNT), 2-(1H-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium-hexafluorophosphate (HBTU), 2-(1H-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium-hexafluoroborate (TBTU), 2-succinimido-1,1,3,3-tetramethyluronium-tetrafluoroborate (TSTU), 2-(5-norbornen-2,3-dicarboximido)-1,1,3,3-tetramethyluronium -tetrafluoroborate (TNTU), O-[(cyano(ethoxycarbonyl)-methyliden)amino]-1,1,3,3-tetramethyluronium-tetrafluoroborate (TOTU), O-(benzotriazol-1-yl)-1,3-dimethyl-1,3-dimethylene uronium hexafluorophosphate (HBMDU), O-(benzotriazol-1-yl)-1,1,3,3-bis(tetramethylene)uronium hexafluorophosphate (HBPYU), O-(benzotriazol-1-yl)-1,1,3,3-bis(pentamethylene)uronium hexafluorophosphate (HBPipU), 3-hydroxy-4-oxo-3,4-dihydro-1,2,3-benzotriazine (HODhbt), 1-hydroxy-7-aza-benzotriazole and its corresponding uronium or phosphonium salts, designated HAPyU and AOP, 1-cyano-2-ethoxy-2-oxoethylideneaminoxy-dimethylamino-morpholino-carbenium hexafluorophosphate (COMU), chlorotripyrrolidinophosphonium hexafluorophosphate (PyCloP), or the like;

Carbodiimides, e.g. dicyclohexylcarbodiimide, N-(3-dimethylaminopropyl)-N'-ethylcarbodiimide, 1-tert-butyl-3-ethylcarbodiimide, N-cyclohexyl-N'-2-morpholinoethyl)carbodiimide or diisopropylcarbodiimide (especially for ester formation via O-acyl urea formation of the carboxylic group); or

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active ester forming agents, e.g. 2-mercaptobenzothiazole (2-MBT),

azide forming agents, e.g. diphenyl phosphoryl azide,

acid anhydrides, such as propane phosphonic acid anhydride,

acid halogenation agents, e.g. 1-chloro-N,N,2-trimethyl-1-propenylamine, chloro-N,N,N',N'-bis(tetramethylene)formamidinium tetrafluoroborate or hexafluorophosphate, chloro-N,N,N',N'-tetramethylformamidinium hexafluorophosphate, fluoro-N,N,N',N'-tetramethylformamidinium hexafluorophosphate, fluoro-N,N,N',N'-bis(tetramethylene)formamidinium hexafluorophosphate,

or the like, or mixtures of two or more such agents.

Also for the ester coupling of compounds of the formula XII or XIIA with those of the formula XIII or XIII A, respectively, or of compounds of the formula XIII* or XIII A* with those of the formula XXVI or XXVIA, respectively, the corresponding reactive carboxyl compounds can be used or formed *in situ*. Here, especially MSNT is preferred as coupling agent as this allows for the maintenance of high stereospecificity.

The reaction may, where appropriate, be conducted in the presence of a mild base (e.g. N-methylmorpholine, a trialkylamine, e.g. ethyldiisopropylamine, a di-(alkyl)aminopyridine, such as N,N-dimethylaminopyridine, or the like (taking care that the conditions are not so basic as to allow for the hydrolysis of ester groups, e.g. the depsipeptide ester group, present in precursors of the compound of the formula I), where appropriate or required in the presence of an appropriate solvent or solvent mixture, e.g. an N,N dialkylformamide, such as dimethylformamide, a halogenated hydrocarbon, e.g. dichloromethane, N-alkylpyrrolidones, such as N-methylpyrrolidone, nitriles, e.g. acetonitrile, or further an aromatic hydrocarbon, e.g. toluene, or mixtures of two or more, where, provided an excess of coupling agent is present, also water may be present. The temperatures may be ambient temperature or lower or higher, e.g. in the range from -20 °C to 50 °C.

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The amino acids of the formula IX, IXA, XI, XIA, XIII, XIII A, XV, XVA, XVII, XVII A, XXVII (obtainable e.g. by Solution Phase synthesis), XVII*, XVII A*, XV*, XVA*, XIII*, XIII A*, XI*, XIA*, IX* and IXA* are known or they can be synthesized according to methods known in the art, they are commercially available, and/or they can be synthesized in analogy to methods known in the art.

Also the remaining starting materials, e.g. the acid of the formula XIX or VII, or the dipeptide of the formula XXVII or XXVII A, are known or they can be synthesized according to methods known in the art, they are commercially available, and/or they can be synthesized in analogy to methods known in the art.

For example, the synthon of the formula XIX can be prepared as described in Example 1 A(4) (which is a specific embodiment of the invention) or in analogy thereto. The synthesis of the intermediate compound 1 (Scheme 1) is described in Tetrahedron 61, 1459-1480 (2005).

The coupling reactions for dipeptides make use of the corresponding carboxylic groups of amino acids in free form or in activated form.

|

Examples: The following examples illustrate the invention without limiting its scope.

Abbreviations

aq.	aqueous
Boc/BOC	tert-Butoxycarbonyl
brine	sodium chloride solution in water (saturated at RT)
Bzl	benzyl
COMU	1-cyano-2-ethoxy-2-oxoethylideneaminooxy-dimethylamino-morpholino-carbenium hexafluorophosphate
DCM	dichloromethane
DIPEA	N,N-diisopropylethylamine
DMAP	4-Dimethylaminopyridine
DMF	N,N-dimethylformamide
Fmoc/FMOC	9-fluorenylmethoxycarbonyl
Et	ethyl
HATU	2-(1H-7-Azabenzotriazol-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate Methanaminium
HFIP	Hexafluoroisopropanol
HPLC	High Performance Liquid Chromatography
HR-MS	High Resolution Mass Spectroscopy
IPA	Isopropylacetate
IPC	In-Process Control
IR	Infrared Spectroscopy
IT	internal temperature
Kaiser test	Ninhydrin-based test to monitor deprotection in SPPS (see E. Kaiser, R. L. Colescott, C. D. Bossinger, P. I. Cook, Analytical Biochemistry 1970 , 34 595); if mentioned to be OK, this means successful deprotection.
me	methyl
MS	Mass Spectroscopy
MSNT	1-(Mesitylene-2-sulfonyl)-3-nitro-1,2,4-triazole
NMR	Nuclear Magnetic Resonance Spectroscopy
PyBOP	benzotriazol-1-yl-oxytripyrrolidinophosphonium hexafluorophosphate

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BOP	Benzotriazole-1-yl-oxy-tris-(dimethylamino)-phosphonium hexafluorophosphate	
RP	Reversed Phase	
RT/rt	room temperature	
SPPS	Solid Phase Peptide Synthesis	
TBME	tert-Butyl-methylether	
TFA	trifluoroacetic acid	2,2,6,6-Tetramethyl-1-
	piperidinyloxy, free radical.	
RBF	Round bottomed flask	

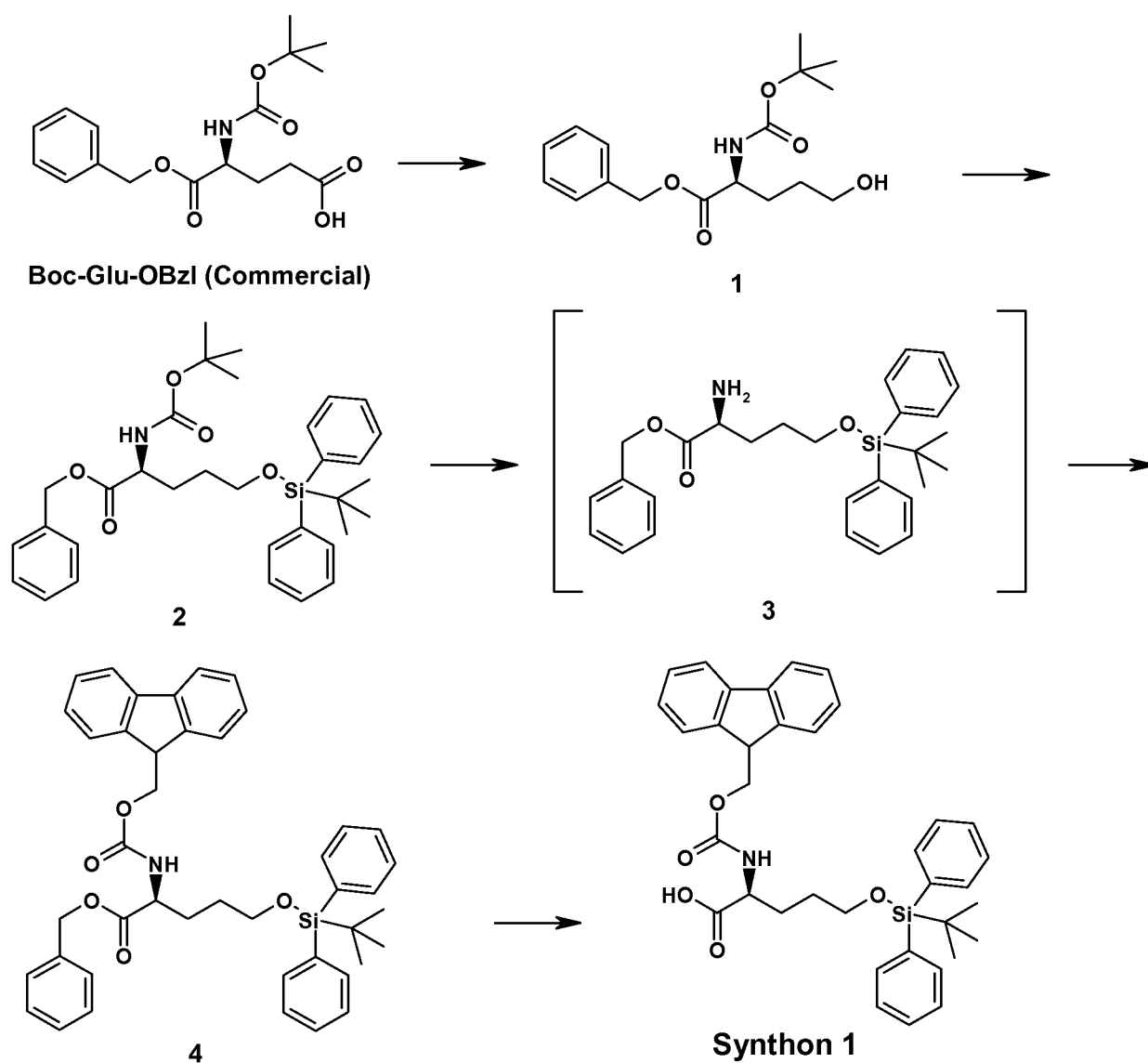
For amino acid abbreviations see the table above.

Compounds Names...

The names of the open chain oligopeptides have been derived according to recommendations of the Joint Commission on Biochemical Nomenclature ("International Union of Pure and Applied Chemistry" and "International Union of Biochemistry") published in Pure & Appl. Chem. 1984, 56, 595-624. Previously, a simple peptide naming convention was used for these compounds. The previous names are given in parenthesis.

Example 1: Synthesis of Compound A**1A Synthesis of Synthon 1****(i) Alternative 1:**

Reaction Scheme 1:



1A(1) Preparation of compound 1

(S)-Benzyl 2-((*tert*-butoxycarbonyl)amino)-5-hydroxypentanoate

Compound 1 was prepared by a procedure similar to the procedure reported in R. K. Olson, K. Ramasamy, T. Emery, *J. Org. Chem.* **1984**, 49, 3527. BOC-Glu-OBzl (50g, 148.2 mmol) was dissolved in tetrahydrofuran (800 mL) and triethylamine (47.3g, 467.4 mmol) was added. The solution was cooled down to IT= -10 °C. Ethyl-chloroformate (51.8g, 98% purity, 467.8 mmol) was added slowly, maintaining the temperature at IT = -10 to -15 °C. The suspension thus obtained was stirred for an additional hour. IPC (HPLC) indicated the disappearance of the starting material. The reaction mixture was allowed to warm-up to 0 °C and water (800 mL) was added at 0-5 °C within 25-30 minutes. Separation of the mixture into 2-phases became visible. Under intensive stirring, sodium-borohydride (11.8g, 299.4 mmol) was added in 10 portions at 0-5 °C. Caution was exercised as hydrogen gas evolved during addition of sodium-borohydride. The reaction mixture was stirred for additional 5 minutes at 0-5 °C and the temperature was increased to 20-25 °C within 30 minutes. Caution was required as hydrogen-gas continued to evolve during temperature increase. The reaction mixture was stirred for 15 minutes at 20-25 °C before work-up.

For work-up, water (1250 mL) was added to the reaction mixture, followed by the addition of ethyl acetate (1250 mL). The phases were separated and the aqueous phase was extracted with ethyl acetate (600 mL). The organic layers were combined and were washed with brine (2x 600 mL). The organic layer was dried over magnesium sulfate, the solvent was evaporated at reduced pressure and 40-45 °C to obtain 50.8g crude product. HPLC purity: 94 a%. The crude product was purified by flash chromatography on silica gel with hexane fraction/ethyl acetate (7:3 to 1:1) as mobile phase. Yield: 40.85g (85.2%). Purity: 98% (HPLC). MS and NMR confirmed the proposed structure.

1A(2) Preparation of compound 2: (S)-Benzyl 2-((*tert*-butoxycarbonyl)amino)5-((*tert*-butyldiphenylsilyl)oxy)pentanoate

The alcohol (compound 1) from the previous step (40.3g, 124.6 mmol) was dissolved in dimethyl formamide (200 mL) and imidazole (12.8g, 99.5% purity, 186.9 mmol) was added. The mixture was stirred at room temperature, until a solution was formed (5-10

min). *tert*-Butyl-diphenyl-silyl-chloride (41.9g, 98% purity, 149.5 mmol) was added dropwise within 10 minutes and stirring was continued for additional 15 minutes at room temperature. IPC (HPLC) indicated disappearance of the starting material (alcohol). For work-up, isopropyl acetate (400 mL) was added to the reaction mixture, followed by the slow addition of a half-saturated aq. sodium bicarbonate solution (400 mL). Caution was exercised as the addition was exothermic., gas evolution takes place. The phases were separated and the aqueous phase was extracted with isopropyl acetate (400 mL). The organic layers were combined and extracted with water (400 mL). The organic layer was dried on magnesium sulfate and the solvent was evaporated at 40-45 °C under reduced pressure. The residue was dried over night *in vacuo* at 25 °C to obtain 83.25g crude product as colorless oil. HPLC analysis indicated the presence of 81 a% desired product and 18.6 a% of the corresponding silanole. The crude product was used without further purification for the next step. NMR and HR-MS of a purified sample confirmed the proposed structure. HR-MS: calculated for $C_{33}H_{43}NO_5Si$: $[M+H]^+$: 562.29833; $[M+NH_4]^+$: 579.32488; $[M+Na]^+$: 584.28027. Found: $[M+H]^+$: 562.29848; $[M+NH_4]^+$: 579.32489; $[M+Na]^+$: 584.27995.

1A(3) Preparation of compound 4:

(*S*)-Benzyl 2-(((9*H*-fluoren-9-yl)methoxy)carbonyl)amino)-5-((*tert*-butyldiphenylsilyl)oxy)pentanoate

83.25g crude product from the previous step (corresponding to 70g theoretical yield of compound **2**, 124.6 mmol) was dissolved in dichloromethane (650 mL) and trifluoroacetic acid (358.8g, 99% purity, 3115 mmol) was added dropwise to the intensively stirred solution. IPC (HPLC) after 30 minutes indicated complete cleavage of the BOC protecting group. The reaction mixture (clear solution) was transferred into a 6 L 4-bottomed flask with mechanical stirrer and was diluted with 1000 mL of dichloromethane. Aqueous half saturated sodium carbonate solution (1800 mL) was slowly added to the intensively stirred solution. Caution was exercised as strong gas evolution was observed during addition of sodium carbonate. pH of the aqueous phase after completion of the addition: 9-10. The reaction mixture was stirred for additional 15 minutes and the phases were separated. The aqueous phase was extracted with dichloromethane (1000 mL) and the organic layers were combined to obtain a solution of the product in dichloromethane.

The solution was concentrated at 40-45 °C under reduced pressure to a final volume of ca. 650 mL. for the next step. HPLC indicated the presence of 80.5 % of compound **3** and 19.5 a% silanole in the solution.

For FMOCCylation, saturated aq. sodium bicarbonate solution (650 mL) was slowly added to the intensively stirred solution of compound **3**, followed by the addition of FMOCC-chloride (36.55g, 97% purity, 137 mmol). Caution was exercised as gas evolution was observed during the addition of FMOCC-Cl. Stirring was continued for 15 minutes at room temperature. An IPC (HPLC) of the organic layer indicated disappearance of the intermediate compound **3** and complete conversion into compound **4**. For work-up, the layers were separated and the aqueous phase was extracted with dichloromethane (650 mL). The organic layers were combined and were extracted with water (650 mL). The organic layer was dried over magnesium sulfate and the solvent was removed at 40-45 °C under reduced pressure to obtain 112.6g crude product.

The crude product was suspended in ethanol/isopropanol/water (89:5:6; 2400 mL) and the suspension was heated to IT 50 °C to obtain a solution. The solution was cooled down to 45 °C, seed crystals were added and the temperature was continued to cool down to 20-25 °C within 1 hour. Crystallization started at ca. 40 °C. The suspension was stirred at 20-25 °C over night, then cooled down to IT 0-5 °C within 30 minutes and stirring was continued for additional 2 hours at 0-5 °C. The product was isolated by filtration, the filter cake was washed with ethanol/isopropanol/water (89:5:6; 240 mL) and dried under reduced pressure to obtain pure compound **4** (100 a% purity according to HPLC). Yield: 66g (77.4%). The product was fully characterized by MS and NMR HR-MS: Calculated for C₄₃H₄₅NO₅Si: [M+H]⁺: 684.31398; [M+NaH₄]⁺: 701.34053; [M+Na]⁺: 706.29592. Found: [M+H]⁺: 684.31392; [M+NH₄]⁺: 701.34021; [M+Na]⁺: 706.29539. The mother liquor gave 30.6g of a foam comprising 30 a% product and 29 a% silanole according to HPLC.

1A(4) Preparation of Synthon 1 from compound 4:

(S)-2-(((9H-Fluoren-9-yl)methoxy)carbonyl)amino)-5-((*tert*-butyldiphenylsilyl)oxy)pentanoic acid

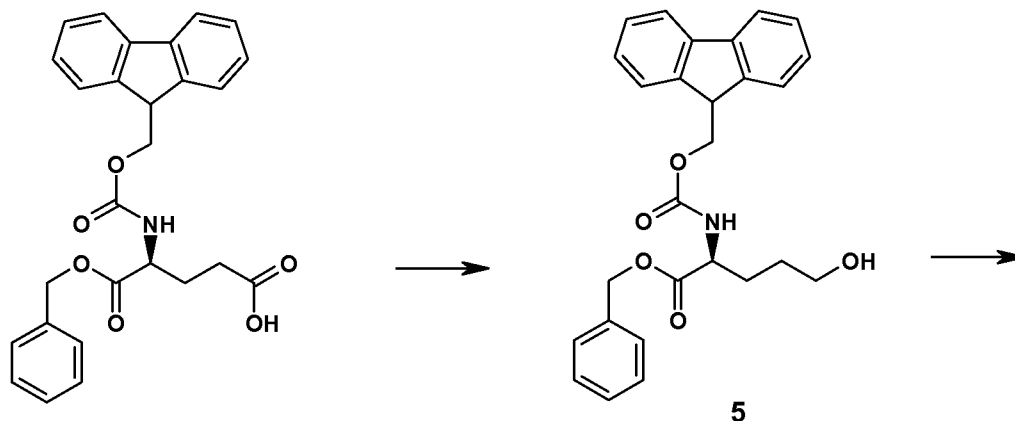
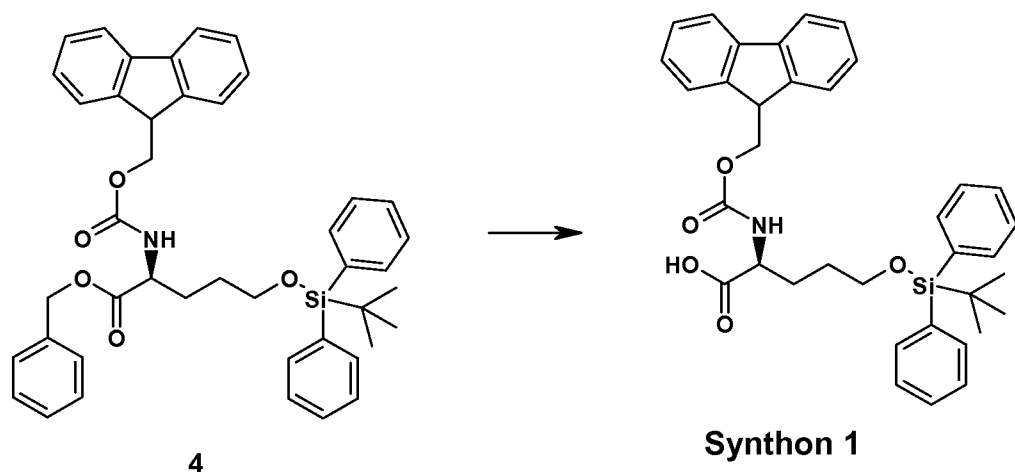
Compound **4** (66g, 96.6 mmol) was suspended in ethanol/isopropylalcohol/water (89:5:6; 3000 mL) and the suspension was heated to IT 45 °C to obtain a solution. The solution was cooled down to IT 30 °C. After inertization with Argon, palladium-catalyst (10% on

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barium sulfate; 6.6g) was added to the solution under an argon stream. The product was then hydrogenated under a hydrogen pressure slightly above the atmospheric pressure at 30-35 °C. The hydrogenation was completed after 1.5h according to HPLC. The reaction mixture was filtered over a cellulose based filter aid (Cellflock 40; cellulose based filtering aid) and the filter aid was washed with ethanol/isopropanol/water (89:5:6; 600 mL). Evaporation of the solvent under reduced pressure at 45-50 °C gave 59.58g foam as crude product. The crude product was purified by chromatography on silica gel in 2 portions (2x1 kg silica gel 60) using dichloromethane/methanol 95:5 to 80:20 as mobile phase. Yield: 52g (90.6%). The product was fully characterized by MS and NMR. HR-MS: Calculated for $C_{36}H_{39}NO_5Si$ $[M+H]^+$: 594.26703; $[M+NH_4]^+$: 611.29358; $[M+Na]^+$: 616.24897; Found: $[M+H]^+$: 594.26743; $[M+NH_4]^+$: 611.29385; $[M+Na]^+$: 616.24900.

(ii) Alternative 2:

Reaction Scheme 2:

**Fmoc-Glu-OBzl (Commercial)****1A(5) One-pot Preparation of compound 4 from Fmoc-Glu-OBzl via compound 5:**

Fmoc-Glu-OBzl (5g, 10.88 mmol) was dissolved in tetrahydrofuran (80 mL) and triethylamine (3.3g, 32.6 mmol) was added. The solution was cooled down to -15 °C and ethyl chloroformate (7.3g, 67.27 mmol) was added to the solution over 30 minutes at -12 °C. The suspension thus obtained was stirred for an additional hour at -10 to -15 °C and the temperature was elevated to 0 °C. Water (80 mL) was added dropwise to the reaction mixture, maintaining the temperature at 0 °C. Sodium borohydride (total 0.805 g, 21.27 mmol) was added in 3 portions (1 portion every 10 minutes) at 0 °C and the reaction mixture was stirred for additional 1h at 0 °C. Caution was exercised as

hydrogen gas evolved. The reaction mixture was diluted with water (100 mL) and extracted with isopropyl acetate (150 mL) and the layers were separated. The aqueous phase was extracted again with isopropyl acetate (100 mL) and the organic layers were combined. The combined organic phase was washed with half saturated sodium chloride solution (2x 50 mL) and the solution was concentrated at reduced pressure to a final volume of ca. 50 mL. The concentrated solution was clear-filtered and the filter residue was washed with isopropyl acetate (20 mL). The solution of compound **5** thus obtained was transferred into a round bottomed flask and imidazole (1.49g, 21.89 mmol) was added. The mixture was stirred for 15 min at Rt and tert.-butyl-diphenyl-silyl-chloride (4.45g, 16.19 mmol) was added. The reaction mixture was stirred for 15 h at rt. For work-up, the suspension was diluted with isopropyl acetate (20 mL) and extracted with water (3x50 mL). The organic layer was separated and the solvent was evaporated under reduced pressure to obtain 9.95g crude product. The crude product was purified by column chromatography on silicagel with isopropyl acetate/hexane (2:8) as mobile phase to obtain 6.5g of a solid, which was suspended in hexane and stirred for 3h at rt. The precipitate was isolated by filtration and dried at 50 °C under reduced pressure to obtain 5.6g of compound **4**. Yield: 75% over two steps. HR-MS: calculated for $C_{43}H_{45}NO_5Si$: $[M+H]^+$: 684.31398; $[M+NH_4]^+$: 701.34053; $[M+Na]^+$: 706.29592. Found: $[M+H]^+$: 684.31430; $[M+NH_4]^+$: 701.34073; $[M+Na]^+$: 706.29577.

1A(6) Alternative one-pot preparation of compound 4 from Fmoc-Glu-OBzl via compound 5 using isopropyl-chloroformate instead of ethyl-chloroformate:

Fmoc-Glu-OBzl (60g, 130.579 mmol) was dissolved in tetrahydrofuran (550 mL) and triethylamine (40.8g, 403.202 mmol) was added. A cloudy solution was obtained with some precipitate. This cloudy solution/suspension was transferred into a dropping funnel and was added to a pre-cooled solution of isobutyl-chloroformate (54.96g, 402.41 mmol) in tetrahydrofuran (300 mL) in a 4.5L reactor at -35 to -30 °C, maintaining this temperature during the addition. Residuals in the dropping funnel were washed with additional tetrahydrofuran (50mL) and the reaction mixture was stirred at -35 to -30 °C for another 2 hours. Water (960 mL) was added to the reaction mixture within 45 minutes, allowing the temperature to increase until 0 °C. A suspension was formed. Sodium borohydride (14.4g, 380.625 mmol) was added in 20 portions within 1h at 0°C and the reaction mixture was stirred for an additional hour at 0°C. Caution was exercised as hydrogen gas evolved.

The suspension was poured onto t-butyl-methylether (600 mL) and the reaction flask was washed with water (600 mL), which was added to the product mixture (2-phases). The phases were separated, the water phase was extracted with t-butyl-methylether (600 mL) and the organic phases were combined. The organic phase was washed with water (2x600 mL), dried over anhydrous magnesium sulfate (200g) and the solvent was removed under reduced pressure until a final volume of 1L was achieved. The solution was diluted with dimethyl-formamide (600g) and the solvent was evaporated under reduced pressure, until a final volume of 400 mL is achieved. The solution of compound **5** thus obtained was transferred into a round bottomed flask. Imidazole (14.4g, 211.524 mmol) was added to the DMF solution of compound **5** and the mixture was stirred for 5 minutes at rt. Finally, TBDPS-Cl (39.6g, 144.07 mmol) is added dropwise during 20 minutes at 20-25 °C and the reaction mixture is stirred for an additional hour at this temperature.

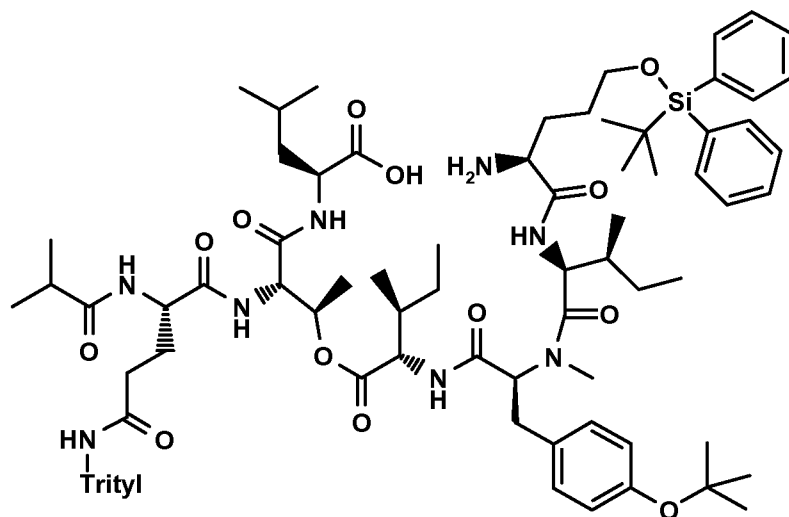
The reaction mixture was then poured onto ethyl acetate (1200 mL) and the mixture was extracted with water (700 mL). The layers were separated and the organic layer was washed with water (3x300 mL). Evaporation of the solvent under reduced pressure gave 106g crude product.

The crude product (106g) was dissolved in Ethanol/Isopropanol/Water (89:5:6; 1200 mL) at 40-50 °C and seed crystals (0.5g compound **4**) were added. The mixture was allowed to cool down to room temperature and stirred for 17 hrs at rt. The suspension was cooled to -20 °C and stirred for 2 hrs at -20 °C. The product was isolated by filtration, the filter cake was washed with the solvent mixture Ethanol/Isopropanol/Water (89:5:6; 3x200 mL) and dried at 40 °C under reduced pressure to obtain 66.5g of compound **4** (74.6% yield over 2 steps). HPLC indicated >99 a% purity for the product.

Additional product can be isolated from the mother liquor (34g after evaporation of the solvent), which contains ca. 30 a% compound **4** according to HPLC.

1B Synthesis of Precursor peptide 1 by SPPS

Precursor peptide 1: Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-OH



Precursor peptide 1

Precursor peptide 1 has been prepared using 2 different Solid Supports

Equipment:

Solid phase synthesis reactor with a filter cloth or sintered glass filter plate at the bottom. A nitrogen manifold allows to drain the reactor contents via filter cloth or sintered glass filter plate and bottom valve.

1B(1) Coupling of the trityl-linker to the solid support:

200g of Aminomethyl-polystyrene resin (crosslinked with 1% divinyl benzene, loading of aminomethyl groups 1mmol/g) (supplier: Senn Chemicals AG, Dielsdorf/Switzerland) were stirred alternately with several portions of dimethylformamide (1600mL) and isopropanol (1600mL) After two final washes with dimethylformamide, the resin was treated with a previously prepared solution of 4-hydroxy-diphenylmethyl-benzoic acid (91.3g 300mmol), 1-hydroxy-benzotriazole monohydrate (45.9g, 300mmol) and diisopropylcarbodiimide (75.7g, 600mmol) in dimethylformamide (1600mL). The reaction mixture was stirred for 1.5h and a Ninhydrin test was performed. The test still showed free amino groups and thus diisopropylcarbodiimide (7.6g, 60mmol) was added and the

reaction stirred over night. A further ninhydrin test in the morning was negative and the reaction mixture was filtered off. The resin was washed with dimethylformamide and isopropanol alternatingly. The resin was dried *in vacuo* and yielded 257g of dry linker-resin. The material was used for the next synthesis step without further analysis.

1B(2) Coupling of Fmoc-Leu-OH

Preparation of Fmoc-Leu-Linker-Resin

Linker-Resin (190g, 147.8mmol) was swollen by stirring in toluene (1400mL). The solvent was filtered off and replaced by a solution of toluene (1400mL) and acetyl chloride (53mL, 1478mmol). This mixture was stirred for 2h, filtered off, replaced by an identical mixture which was stirred for another 2h before filtering off. The chlorinated resin was washed twice with toluene and three times with dichloromethane.

In a round bottom flask, a solution of Fmoc-Leu-OH (104.8g, 296mmol) and of N-Methylmorpholine (49mL, 444mmol) in dichloromethane (600mL) was prepared. This solution was added to the resin and stirred over night. In the morning, the solution was filtered off and the resin was washed with dichloromethane and isopropanol alternatingly. The resin was dried *in vacuo* and yielded 234.7g of dry Fmoc-Leu-linker-Resin. The loading with Fmoc-groups was determined at 0.787mmol/g what led to a yield of 185mmol (125% of theory). Amino acid analysis at an external contractor confirmed <0.1% D-Leu enantiomer.

1B(3) Coupling of Fmoc-Thr-OH

Preparation of Fmoc-Thr-Leu-Linker-Resin

Fmoc-Leu-Linker-Resin (140g, 109mmol) was swollen by stirring in two successive portions of dimethylformamide (1100mL) for 30min each.

Fmoc protecting group was cleaved by two subsequent washings of 20% piperidine in dimethylformamide for 5min and 15min respectively. The resin was washed by several alternating washes with dimethylformamide and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink colour confirmed successful removal of piperidine.

The resin was washed with tetrahydrofurane (1200mL) three times to prepare for the following coupling step.

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In a round bottomed flask a solution of Fmoc-Thr-OH (112.1g, 328mmol), hydroxybenzotriazole monohydrate (51.25g, 334mmol) and diisopropylcarbodiimide (51mL, 655mmol) in tetrahydrofuran (600mL) was prepared. The solution was added to the resin and the pH checked immediately (pH= 6.5). The reaction mixture was stirred for 1.5h until a ninhydrin test showed complete reaction. The solution was filtered off and the resin was washed with dimethylformamide and isopropanol alternatingly. A small sample of the resin was dried and sent for amino acid analysis (0.13% D-Leu, <0.1% D-Thr, <0.1% L-allo-Thr, <0.1% D-allo-Thr), the bulk of the material was subjected to the next step without further drying.

1B(4) Coupling of Fmoc-Gln(Trt)-OH

Preparation of Fmoc-Gln(Trt)-Thr-Leu-Linker-Resin

The Fmoc-Thr-Leu-Linker-Resin from the previous step was swollen by stirring in two subsequent portions of dimethylformamide (1100mL) for 30min each.

Fmoc protecting group was cleaved by two subsequent washings with 20% piperidine in dimethylformamide for 5min and 15min respectively. The resin was washed by several alternating washes with dimethylformamide and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink colour proved successful removal of piperidine.

The resin was washed with dimethylformamide (1100mL) three times to prepare for the following coupling step.

In a round bottomed flask a solution of Fmoc-Gln(Trt)-OH (138.6g, 226mmol), HATU (86.2g, 226mmol) and Ethyldiisopropylamine (58.4g, 452mmol) in dimethylformamide (400mL) was prepared.

The solution was added to the resin and the pH checked immediately (pH= 10). The reaction mixture was stirred for 3h until a ninhydrin test showed complete reaction. The solution was filtered off and the resin was washed with dimethylformamide and isopropanol alternatingly.

The resin was dried in vacuo and yielded 170g of dry Fmoc-Gln(Trt)-Thr-Leu-Linker-Resin. The loading with Fmoc-groups was determined at 0.60 mmol/g indicating a yield of 102mmol (94% of theory over the last two steps). Amino acid analysis at an external contractor led to the following values: (0.13% D-Leu, <0.1% D-Thr, <0.1% L-allo-Thr, <0.1% D-allo-Thr, <0.8% D-Gln).

1B(5) Coupling of Isobutyric acid**Preparation of Isobutyryl-Gln(Trt)-Thr-Leu-Linker-Resin**

Fmoc-Gln(Trt)-Thr-Leu-Linker-Resin (169g, 101mmol) was swollen by stirring in two subsequent portions of dimethylformamide (1300mL) for 30min each.

Fmoc protecting group was cleaved by two subsequent washings with 20% piperidine in dimethylformamide (1300mL) for 5min and 15min respectively. The resin was washed by several alternating washes with dimethylformamide and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink colour proved successful removal of piperidine.

The resin was washed with dimethylformamide (1100mL) three times to prepare for the following coupling step.

In a round bottom flask a solution of Isobutyric acid (17.9g, 203mmol), PyBOP (105.5g, 203mmol) and Ethyldiisopropylamine (52.4g, 406mmol) in dimethylformamide (550mL) was prepared.

The solution was added to the resin and the pH checked immediately (pH= 9.5). The reaction mixture was stirred for 2.5h until a ninhydrin test showed complete reaction. The solution was filtered off and the resin was washed with dimethylformamide and isopropanol alternatingly.

The batch was directly subjected to the next step without drying and further analysis.

1B(6) Coupling of Fmoc-Ile-OH (Esterification)**Preparation of Isobutyryl-Gln(Trt)-Thr(Ile-Fmoc)-Leu-Linker-Resin**

Isobutyryl-Gln(Trt)-Thr-Leu-Linker-Resin (wet from step above, 101mmol) was swollen by stirring in three subsequent portions of dichloromethane (1200mL) for 20min each.

The solvent was filtered off and MSNT (88g, 297mmol) and Fmoc-Ile-OH (105g, 297mmol) were added as solids. Dichloromethane (500mL) was added as well as a solution of N-methyl imidazole (18.2g, 223mmol) and ethyldiisopropylamine (51.2g, 396mmol) in dichloromethane (100mL). The reaction mixture was stirred for 2h until HPLC in process control showed complete reaction. The solution was filtered off and the resin was washed with three portions of dichloromethane, three portions of dimethylformamide and three portions of isopropanol subsequently. The resin was dried in vacuo and yielded 172g of dry Isobutyryl-Gln(Trt)-Thr(Ile-Fmoc)-Leu-Linker-Resin. Fmoc loading was determined to be 0.418mmol/g thus indicating a yield of 72mmol (71% over the last two steps).

1B(7) Coupling of Fmoc-N-methyl-Tyr(tBu)-OH**Preparation of Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Fmoc)-Leu-Linker-Resin
(Previously named: Isobutyryl-Gln(Trt)-Thr(Ile-N-me-Tyr(tBu)-Fmoc)-Leu-Linker-Resin)**

Isobutyryl-Gln(Trt)-Thr(Ile-Fmoc)-Leu-Linker-Resin (172g, 72mmol) was swollen by stirring in two subsequent portions of dimethylformamide (1300mL) for 30min each. Fmoc protecting group was cleaved by two subsequent washings with 20% piperidine in dimethylformamide (1400mL) for 5min and 15min respectively. The resin was washed by several alternating washes with dimethylformamide and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink colour proofed successful removal of piperidine.

The resin was washed with dimethylformamide (1100mL) three times to prepare for the following coupling step.

A solution of Fmoc-N-methyl-Tyr(tBu)-OH (68.7g, 144mmol) and HATU (55.1g, 144mmol) in dimethylformamide (700mL) was prepared and added to the peptide-resin, followed by the addition of a solution of Ethyl-diisopropylamine (37.5g 289mmol) in dimethylformamide (100mL) under stirring. pH checks immediately after addition of the coupling solution and after 1h of reaction gave the same result (pH 10) The solution was stirred for 2h until a ninhydrin test showed complete reaction. The solution was filtered of and the resin was washed with dimethylformamide and isopropanol alternatingly. The batch was directly subjected to the next step without drying and further analysis.

1B(8) Coupling of Fmoc-Ile-OH**Preparation of Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-IleFmoc)-Leu-Linker-Resin.
(Previously named: Isobutyryl-Gln(Trt)-Thr(Ile-N-me-Tyr(tBu)-Ile-Fmoc)-Leu-Linker-Resin.)**

The wet Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Fmoc)-Leu-Linker-Resin (72mmol) was swollen by stirring in two subsequent portions of dimethylformamide (1200mL and 1300mL) for 30min each. Fmoc protecting group was cleaved by two subsequent washings with 20% piperidine in dimethylformamide (1400mL) for 5min and 15min respectively. The resin was washed by several alternating washes with dimethylformamide and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink colour proved successful removal of piperidine.

The resin was washed with dimethylformamide (1100mL) three times to prepare for the following coupling step.

In a round bottom flask a solution of Fmoc-Ile-OH (103.9g, 294mmol) COMU (125.9g, 294mmol) and Ethyldiisopropylamine (76g, 588mmol) in dichloromethane (440mL) and dimethylformamide (440mL) was prepared.

The solution was added to the resin and the reaction mixture stirred for 20h. After that time a ninhydrin test was done, showing complete reaction. The solution was filtered off and the resin was washed with dimethylformamide and isopropanol alternatingly.

The resin was dried in vacuo and yielded 185.5g of dry Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Fmoc)-Leu-Linker-Resin. The loading with Fmoc groups was determined to be 0.40mmol/g. Thus a quantitative yield of 74mmol resulted.

1B(9) Coupling of Synthon 1 and cleavage of final Fmoc protecting group:

Preparation of Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-Linker-Resin

(Previously named: Isobutyryl-Gln(Trt)-Thr(Ile-N-me-Tyr-Ile-Synthon 1-H)-Leu-Linker-Resin)

Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Fmoc)-Leu-Linker-Resin (30g, 12mmol) was swollen by stirring in two subsequent portions of dimethylformamide (240mL and 250mL) for 30min each.

Fmoc protecting group was cleaved by two subsequent washings with 20% piperidine in dimethylformamide (250mL) for 5min and 15min respectively. The resin was washed by several alternating washes with dimethylformamide and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink colour proved successful removal of piperidine.

The resin was washed with dimethylformamide (250mL) three times to prepare for the following coupling step.

In a round bottom flask, a solution of of synthon 1 (14.6g, 24.6mmol) PyBOP (12.85g, 24.6mmol) and Ethyldiisopropylamine (6.4g, 49.2mmol) in dimethylformamide (120mL) was prepared.

The solution was added to the resin and the reaction mixture stirred for 3h. After that time a ninhydrin test confirmed complete reaction. The solution was filtered off and the resin was washed with dimethylformamide and isopropanol alternatingly.

The resulting peptide-resin was swollen by stirring in two subsequent portions of dimethylformamide (250mL) for 30min each.

Fmoc protecting group was cleaved by two subsequent washings with 20% piperidine in dimethylformamide (250mL) for 5min and 15min respectively. The peptide-resin was washed by several alternating washes with dimethylformamide and isopropanol.

Finally the peptide-resin was washed three times with dichloromethane (250mL) to prepare for the cleavage of the peptide.

1B(10) Cleavage of Precursor peptide 1 from solid Support

Preparation of Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-OH (= Precursor peptide 1)

(Previously named: Isobutyryl-Gln(Trt)-Thr(Ile-N-me-Tyr(tBu)-Ile-Synthon 1-H)-Leu-OH)

To the wet Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-Linker-Resin, a mixture of acetic acid (125mL) and dichloromethane (125mL) was added and the suspension stirred for 2h. The suspension was filtered and the filtrate collected in a round bottomed flask (filtrate 1). The resin was washed twice with dichloromethane (250mL) and the washes were combined with filtrate 1.

The resin was treated with a fresh portion of acetic acid (125mL) and dichloromethane (125mL) for another 2h. The suspension was filtered and the filtrate collected in a round bottomed flask (filtrate 2). The resin was washed twice with dichloromethane (250mL) and the washes combined with filtrate 2.

Acetic acid (200mL) and dichloromethane (50mL) were added to the resin and the suspension stirred over night. The suspension was filtered and the filtrate collected as filtrate 3.

The three filtrates were worked up separately to assess the effectiveness of the cleavage method. The filtrates were concentrated in a rotary evaporator and the residual acetic acid removed by azeotropic distillation with three portions of toluene (100mL). The oily residues were then dried in high vacuo at a lyophilisator. Yield: filtrate 1: 12.5g, filtrate 2: 2.1g, filtrate 3: 0.3g.

The crude material, Precursor peptide 1, was purified by RP-chromatography, the fractions concentrated in a rotary evaporator and the concentrate freeze dried. Purity: 98.8% Yield: 8.5g (48% for the last two steps). The product was characterized by, ¹H-

NMR, ^{13}C -NMR and HR-MS. The spectra confirmed the proposed structure. The NMR spectra indicated the presence of several conformations.

HR-MS: calculated for $\text{C}_{85}\text{H}_{116}\text{N}_8\text{O}_{13}\text{Si}$: $[\text{M}+\text{H}]^+$: 1485.85039. Found: $[\text{M}+\text{H}]^+$: 1485.84929.

Second SPPS-synthesis of Precursor peptide 1 using a chlorinated resin

1B(11) Immobilization of the first amino acid (Fmoc-Leu-OH)

2-Chlorotrityl chloride resin (30 g, $L = 1.2 \text{ mmol/g}$; Merck Novabiochem 855017) was swelled with 240 ml of dried DCM for 10 min and then the solvent was drained, the solid phase reactor was kept close under a soft nitrogen stream. In the meantime, in a round bottom flask were mixed Fmoc-Leu-OH (42.6g, 120.8 mmol) and 100 ml of dry 1,4-Dioxane. The solvent was evaporated under vacuum at 45°C , then a second portion of 1,4 dioxane was added. The solvent was evaporated again under vacuum at 45°C until a colorless oil residue was obtained. To this oily residue were then added sequentially dry DCM (155 mL) and DIPEA (40.4 mL), the solution was added in one single portion to the pre-swollen resin and then 5 min later a second portion of DIPEA (16.5 mL) was added. The suspension was stirred for 2h and the reaction medium was drained. In order to quench potential non-reacted chloride, 240 ml of a quenching solution of DCM/MeOH/DIPEA (70/15/15, v/v) was added and the mixture was stirred for 10 min. A loading of 0.95 mmol/g was measured.

General protocol for Fmoc cleavage:

To the pre-swollen resin were added 240 ml of a solution of 25% piperidine in DMF, the suspension was stirred for 5 minutes and then the reaction mixture was removed by filtration, then a second portion of the same solution of piperidine was added, the suspension was stirred for 15 minutes and then the solvent was removed by filtration.

Resin washing with drying down:

The resin was washed as follows:

250 ml of DMF (6x2 min)

250 ml of IPA (3x2 min)

250 ml of TBME (6x2 min)

The resin was dried overnight under vacuum at 40°C to obtain 35.7 g of H-Leu-Resin.

1B(12): SPPS synthesis of H-Thr-Leu-Resin H-Leu-Resin (30g) from the previous step was swollen with 270 ml of DMF for 30 min then the solvent was drained.

In a round bottomed flask (RBF) were mixed Fmoc-Thr-OH (25.0g, 73.2 mmol), BOP (40.4g, 91.5 mmol) and DMF (250 mL), the mixture was stirred for 2 min and then DIPEA (18.9g, 146.4 mmol) was added. The mixture thus obtained was added in one portion to the pre-swollen peptide-resin and the reaction mixture was stirred for 2.5h. The Kaiser test was positive and therefore a second coupling was done according to the conditions previously described. After 1.0h the Kaiser test was negative and the reaction was considered to be complete and the reaction medium was removed via filtration.

The resin was washed as follows:

250 ml DMF (4x2 min)

250 ml IPA (3x2min)

250 ml TBME (5x2 min).

The resin was dried over night under vacuum at 40 C° yielding 52.2 g of Fmoc-Thr-Leu-Resin ready for the Fmoc-cleaving step.

Fmoc cleavage:

The resin (52.2g) was suspended in 272 ml of DMF and stirred for 1.0h then the solvent was removed by filtration.

The Fmoc protecting group was cleaved in the same manner as described for 1B(11) and the peptide-resin was washed as follows:

250 ml DMF (6x2 min)

250 ml IPA (3x2min)

250 ml TBME (6x2 min)

The resin was dried overnight under vacuum at 40 C° yielding 39.9 g of H-Thr-Leu-Resin

1B(13): SPPS synthesis of H-Gln(Trt)-Thr-Leu-Resin

H-Thr-Leu-Resin from the previous step (39.9g) was swollen with 270 ml of DMF for 30 min then the solvent was drained.

In an RBF were mixed Fmoc-Gln(Trt)-OH (44.7g, 73.2 mmol), BOP (40.4g, 91.5 mmol) and 250 ml of DMF. The mixture was stirred for 2 min and then DIPEA (18.9g, 146.4 mmol) was added. The resulting mixture was added in one portion to the pre-swollen H-Thr-Leu-Resin and the reaction mixture was stirred for 2.5h, the Kaiser test was negative, then the reaction was considered to be complete and the reaction medium was removed via filtration.

The resin was washed as follows:

250 ml DMF (4x2 min)

250 ml IPA (3x2min)

250 ml DMF (4x2min)

The wet peptide-resin was used for the Fmoc-cleaving step without any further manipulation.

Fmoc cleavage:

The fmoc protecting group was cleaved according to the procedure described in step 1B(11). Then the resin was washed as follows:

250 ml DMF (4x2 min)

250 ml IPA (3x2min)

250 ml TBME (5x2 min)

The resin was dried overnight under vacuum at 40 C° yielding 54.4 g of H-Gln(Trt)-Thr-Leu-Resin.

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1B(14): SPPS synthesis of Isobutyryl-Gln(Trt)-Thr-Leu-Resin

The peptide-resin from the previous step (54.4g) was swollen with 270 ml of DMF for 30 min, then the solvent was drained.

In a RBF were mixed Isobutyric acid (6.4 g, 73.2 mmol), PyBop (38.1 g, 73.2 mmol) and 230 ml of DMF, the mixture was stirred for 2 min and then DIPEA (28.3 g, 219.6 mmol) was added. The solution was added in one portion to the pre-swollen peptide-resin and the reaction mixture was stirred for 1.5h. The Kaiser test was negative, then the reaction was considered to be complete and the reaction medium was removed via filtration.

The resin was washed as follows:

250 ml DMF (4x2 min)

250 ml IPA (3x2min)

250 ml TBME (5x2 min).

The resin was dried overnight under vacuum at 40 C° yielding 55.8 g of Isobutyryl-Gln(Trt)-Thr-Leu-Resin.

1B(15): SPPS synthesis of Isobutyryl-Gln(Trt)-Thr(Ile-H)-Leu-Resin

The peptide-resin coming from the previous step (55.8g) was suspended in 250 ml of dry DCM for 1.0h. Then the solvent was removed by filtration and the wet peptide-resin was kept under a soft stream of nitrogen and the temperature was adjusted to 0-5°C.

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In a RBF were mixed Fmoc-Ile-OH (51.7 g, 146.4 mmol), and 150 ml of dry dioxane. The solvent was evaporated under vacuum at 45°C until an oily residue was observed. The distillation process was repeated and then to the residue were added 150 ml of dried DCM, the solution was cooled to -10°C and MSNT (43.3 g, 146.4 mmol) was added, the suspension was stirred for 3 min and then N-methyl-imidazole (14.2 g, 173 mmol) was added, the mixture was stirred for 2 min and the solution was added dropwise in 10 min to the above prepared peptide-resin. After the addition was finished the suspension was stirred under a nitrogen atmosphere for 2.0h

The resin was washed as follows:

250 ml DMC (4x2 min)

250 ml DMF (3x2min)

The wet peptide-resin was used for the Fmoc-cleaving step without any further manipulation.

Fmoc cleavage:

The Fmoc protecting group was cleaved according to the general protocol described in 1B(11)

After the Fmoc cleavage was done, the resin was washed as follows:

250 ml DMF (5x2 min)

250 ml IPA (3x2min)

250 ml TBME (5x2 min)

The resin was dried overnight under vacuum at 40 C° yielding 57.2 g of Isobutyryl-Gln(Trt)-Thr(O-Ile-H)-Leu-Resin.

1B(16): SPPS synthesis of Isobutyryl-Gln(Trt)-Thr(O-Ile-Tyr(tBu)MeN)-Leu-Resin

The peptide-resin coming from the previous step (57.2g) was suspended in 250 ml DMF for 30 min. for swelling, then the solvent was removed by filtration.

In a RBF were mixed Fmoc-NMeTyr(tBu)-OH (52.0 g, 109.8 mmol), HATU (41.7 g, 109.8 mmol) and 230 ml of DMF and the mixture was stirred for 2 min. Then DIPEA (28.3 g, 219.6 mmol) was added and the solution was stirred for 2.0 min. The resulting solution was then added to the pre-swollen peptide-resin.

The reaction mixture was stirred for 1.0 h and a Kaiser-test was performed. The Kaiser test was negative. The reaction was considered to be complete and the reaction medium was removed via filtration.

The peptide-resin was washed as follows:

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250 ml DMF (4x2 min)

250 ml IPA (1x2min)

The wet peptide-resin was used for the Fmoc-cleaving step without any further manipulation.

Fmoc cleavage:

The Fmoc protecting group was cleaved according to the general protocol used in 1B(11).

After the Fmoc cleavage was done, the resin was washed as follows:

250 ml DMF (5x2 min)

250 ml IPA (1x2min)

250 ml DMF (5x2 min)

The obtained isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-H)-Leu-Resin was used for the next step without drying.

1B(17): SPPS synthesis of Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-H)-Leu-Resin

Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-H)-Leu-Resin from the previous step was swollen with 270 ml of DMF for 30 min, then the solvent was drained.

In a round bottomed flask were mixed Fmoc-Ile-OH (38.8 g, 109.8 mmol), HATU (41.7 g, 109.8 mmol) and DMF (250 mL), the mixture was stirred for 2 min and then DIPEA (28.3 g, 219.6 mmol) was added. The solution was added in one portion to the pre-swollen peptide-resin and the reaction mixture was stirred for 2.0h, The Chloranil test was positive and then a second coupling with Fmoc-Ile-OH was performed. 3h after the second coupling, the peptide-resin was isolated by filtration.

The peptide-resin was washed as follows:

250 ml DMF (4x2 min)

250 ml IPA (3x2 min)

250 ml DMF (4x2 min)

The wet peptide-resin was used for the Fmoc-cleaving step without any further manipulation.

Fmoc cleavage:

The Fmoc protecting group was removed in the same manner as described in 1B(11), and the peptide-resin was washed as follows.

250 ml DMF (5x2 min)

250 ml IPA (3x2 min)

250 ml TBME (4x2 min).

The peptide-resin was dried overnight under vacuum at 40 C° yielding 67.0 g of Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-H)-Leu-Resin.

1B(18): SPPS synthesis of Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-Resin

Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-H)-Leu-Resin from the previous step (33.5 g) was suspended in 150 ml of DMF for 30 min then the solvent was drained. In a RBF were mixed Fmoc-Synthon 1-OH (34.3 g, 57.8 mmol), PyBop (30.0 g, 57.7 mmol) and 113 ml of DMF, the mixture was stirred for 2 min and then DIPEA (14.9g, 115.2 mmol) was added. The solution was added in one portion to the pre-swollen peptide-resin and the reaction mixture was stirred for 2.0h. The reaction medium was removed by filtration. The peptide-resin was washed as follows:

150 ml DMF (4x2 min)

150 ml IPA (3x2min)

150 ml DMF (4x2 min)

The wet peptide-resin was used for the Fmoc-cleaving step without any further manipulation.

Fmoc cleavage:

The Fmoc protecting group was removed in the same way as in 1B(11) but using 150 ml of the piperidin solution, and the peptide-resin was washed as follows.

150 ml DMF (5x2 min)

150 ml IPA (3x2min)

150 ml TBME (4x2 min)

The peptide-resin was dried overnight under vacuum at 40 C° yielding 36.0 g of Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-Resin.

1B(19): Alternative SPPS synthesis: cleavage of peptide from solid support

Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-Resin from the previous step (29.75g) was treated with 350 ml of dried DCM for 1.0h, the solvent was removed by filtration and then 350 ml of a solution of 30% v/v HFIP in DCM was added. The mixture was stirred for 10 min and then the solvent was removed by filtration and kept aside. To the wet resin was added a second portion of the same HFIP solution and the solution was stirred for 10 min, then the solvent was removed by filtration and pooled with the

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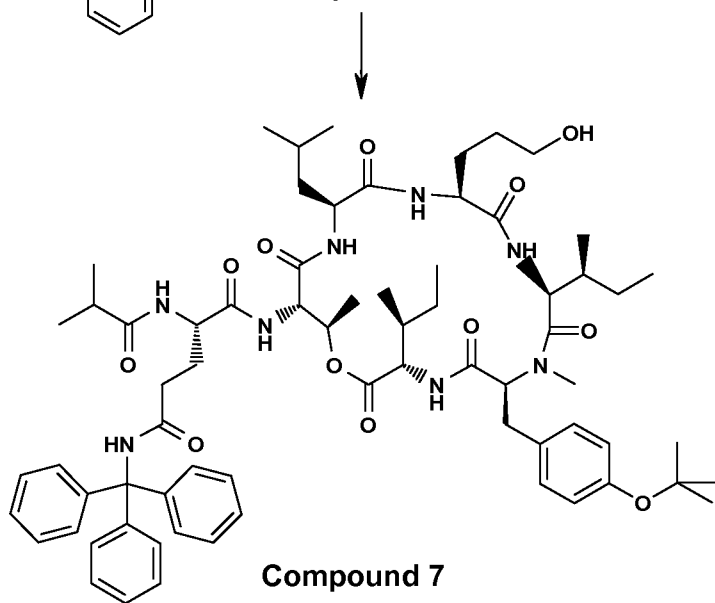
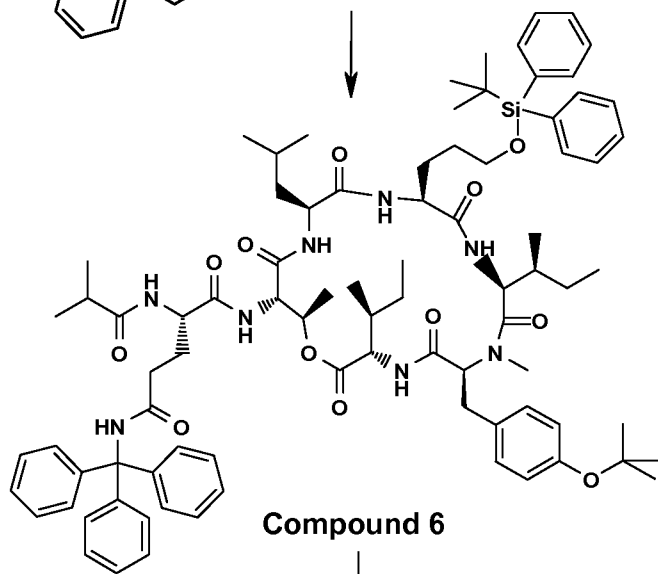
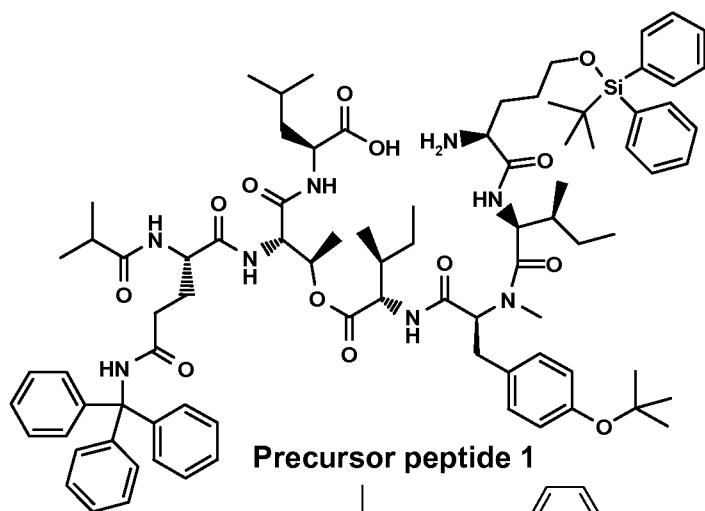
previous solution. The resin was washed 3 times with DCM (350 ml) and the washes were combined with the cleavage solutions.

The combined solution was concentrated under vacuum until an oily residue was observed and then 200 ml of toluene was added, the solvent was evaporated under reduced pressure at 45°C until an oily residue was obtained. 350 ml of hexane was added and the suspension was stirred for 2h. The solvent was removed by filtration and the filter cake was washed with hexane (50 ml), the wet cake was dried overnight under vacuum at 35°C to yield 19.24g of crude **precursor peptide 1** (= Isobutyryl-Gln(Trt)-Thr(Ile-Tyr(tBu)Me-Ile-Synthon 1-H)-Leu-OH).

Part of the crude precursor peptide 1 (6.0g) was purified by RP-chromatography, the product fractions were concentrated in a rotary evaporator and the concentrate was freeze dried to obtain 2.2g of pure precursor peptide 1 in 99.3 %a purity. Product containing side fractions were treated similarly to obtain additional 1.25g of less pure precursor peptide in 71.4 %a purity. Extrapolated for the whole amount of crude precursor peptide 1 (19.24g), this would correspond to 7.05g pure **precursor peptide 1** from major fractions and additional 4.01g **precursor peptide 1** (in 71.4 %a purity) from side fractions.

1C Solution Phase Synthesis of Compound 8

Reaction Scheme 3



1C(1) Synthesis of Compound 6

(*S*)-*N*¹-((3*S*,6*S*,9*S*,12*S*,15*S*,18*S*,19*R*)-6-(4-(*tert*-butoxy)benzyl)-3,9-di((*S*)-*sec*-butyl)-12-(3-((*tert*-butyldiphenylsilyl)oxy)propyl)-15-isobutyl-7,19-dimethyl-2,5,8,11,14,17-hexaoxo-1-oxa-4,7,10,13,16-pentaazacyclonadecan-18-yl)-2-isobutyramido-*N*⁵-tritylpentanediamide

A cloudy solution/suspension of Precursor peptide 1 (1.9 g, 1.28 mmol) in acetonitrile (120 mL) was added during 90 min to a stirred mixture of HATU (972 mg, 2.56 mmol) and DMAP (468.6 mg, 3.84 mmol) in acetonitrile (100 mL) at 35 °C. The dropping funnel was washed with acetonitrile (30 mL). An IPC (HPLC) after the addition of the suspension indicated the absence of the precursor peptide and the completion of the cyclization. For work-up, the solvent was evaporated under reduced pressure until a final volume of ca. 50 mL and the residue was diluted with isopropyl acetate (250). The organic phase was extracted with water (2x100 mL) and the solvent was evaporated under reduced pressure to obtain 2.3 g of crude product. The crude product was purified by flash-chromatography on silicagel with ethyl acetate as mobile phase to obtain 1.79 g (1.22 mmol) compound 6 as a foam. Yield: 95%. The product was characterized by, ¹H-NMR ¹³C-NMR and HR-MS. The spectra confirmed the proposed structure. NMR spectra indicated the presence of several conformations.

HR-MS: calculated for C₈₅H₁₁₄N₈O₁₂Si: [M+H]⁺:1467.83983; [M+NH₄]⁺:1484.86637; [M+Na]⁺: 1489.82177. Found: [M+H]⁺:1467.83984; [M+NH₄]⁺:1484.86555; [M+Na]⁺: 1489.82073.

Second example for the synthesis of Compound 6 (15g scale)

A solution of the precursor peptide 1 (15.0g) in t-butyl-methyl-ether (750 mL) was added slowly during 1.5h to a pre-cooled solution of DMAP (2.80g) and HATU (5.87g) in acetonitrile (375mL) at 0°C. The reaction mixture was stirred for additional 30 min at room temperature. The reaction mixture was then diluted with t-butyl-methyl-ether (750mL) and poured onto half-saturated aq. NaCl-solution (1500 mL). The phases were separated and the organic phase was extracted again with half-saturated aq. NaCl-solution (1500 mL). The organic layer was separated and the solvent was partly evaporated under reduced pressure to a final volume of ca. 175 mL. This solution was filtered over silica gel (column with 225g silica gel) using t-butyl-methyl-ether as mobile

phase. Evaporation of the solvent and drying in vacuo at 40-45 °C gave compound 6 in 97.53% purity according to HPLC. Yield: 14.19g (95.7%).

1C(2) Synthesis of compound 7

(S)-N¹-((3S,6S,9S,12S,15S,18S,19R)-6-(4-(*tert*-butoxy)benzyl)-3,9-di((S)-*sec*-butyl)-12-(3-hydroxypropyl)-15-isobutyl-7,19-dimethyl-2,5,8,11,14,17-hexaoxa-1-oxa-4,7,10,13,16-pentaazacyclononadecan-18-yl)-2-isobutyramido-N⁵-tritylpentanediamide

Compound 6 (1.79 g, 1.22 mmol) was dissolved in tetrahydrofurane (60 mL) and Et₃N(HF)₃ (6.62 g, 41.1 mmol) was added at room temperature. The reaction mixture was stirred at room temperature for 8.5 hours, then diluted with isopropyl acetate (200 mL), and the resulting solution/suspension was slowly added to an intensively stirred saturated aq. NaHCO₃-solution. The organic phase was separated and extracted with water (100 mL). Evaporation of the solvent under reduced pressure gave 1.86g crude product, which was purified by flash-chromatography on silica gel with ethyl acetate/isopropanol (95:5) as mobile phase to obtain 1.24 g (1.00 mmol) of compound 7. Yield: 82%.

The product was fully characterized by IR, NMR and MS. The spectra confirmed the proposed structure.

HR-MS: calculated for C₆₉H₉₆N₈O₁₂: [M+H]⁺:1229.72205; [M+NH₄]⁺: 1246.74860; [M+Na]⁺: 1251.70399. Found: [M+H]⁺: 1229.72128; [M+NH₄]⁺:1246.74780; [M+Na]⁺:1251.70310.

Second example for the preparation of compound 7 (14g scale)

Compound 6 (14.0g, 9.537 mmol) was dissolved in tetrahydrofuran (220 mL) and t-butyl-methyl-ether (116 mL) was added. The solution was treated with Et₃N(HF)₃ (23.05g) by slow addition within 10 minutes. The reaction mixture was stirred for 24h at room temperature. For work-up the reaction mixture was diluted with t-butyl-methyl-ether (570 mL) and the mixture was poured onto half saturated aq. NaHCO₃-solution (632 mL). The biphasic mixture was stirred for 30 min and the phases were separated. The organic phase was extracted with water (280 mL). Both aqueous phases were extracted with t-butyl-methyl-ether (380 mL) and the organic phases were combined. The organic phase was dried over anhydrous MgSO₄ (8.0g) and the solvent was partly evaporated under reduced pressure to a final volume of ca. 100 mL. Toluene (140 mL) was added and the

solvent was evaporated again to a final volume of 80 mL. This solution was diluted with t-butyl-methyl-ether (70 mL) and the product was precipitated by slow addition of heptanes (140 mL) during 30 min. The formed suspension was heated to 50-55 °C and stirred for 30 min at this temperature. The suspension was then cooled down to 0 °C within 30 min, stirred at 0 °C for 2h and the product was isolated by filtration. The product, a white precipitate, was dried in vacuo to obtain 11.08g of compound **7**. (94.5% yield). HPLC of the product indicated 97 a% purity.

1C(3) Synthesis of Compound 8 (see Reaction scheme 4 for structure)

(*S*)-*N*¹-((3*S*,6*S*,9*S*,12*S*,15*S*,18*S*,19*R*)-6-(4-(*tert*-butoxy)benzyl)-3,9-di((*S*)-*sec*-butyl)-15-isobutyl-7,19-dimethyl-2,5,8,11,14,17-hexaoxo-12-(3-oxopropyl)-1-oxa-4,7,10,13,16-pentaazacyclononadecan-18-yl)-2-isobutyramido-*N*⁵-tritylpentanediamide

Compound **7** (1.2g, 0.98 mmol) was dissolved in a mixture of tetrahydrofuran (190 mL) and dimethyl sulfoxide (62 mL). 1-Hydroxy-1,2-benziodoxol-3(1*H*)-one 1-oxide (IBX) (2.42g, 45% g/g, 3.9 mmol) was added and the solution was stirred for ca. 4.5 hours, after which time HPLC indicated disappearance of the starting material (compound **7**). The reaction mixture was then poured onto saturated, aq. NaHCO₃-solution (300 mL) and was extracted with dichloromethane (2x300 mL). The organic layers were combined and were washed with water (2x300 mL). Evaporation of the solvent under reduced pressure gave 2.31g of crude product as a foam. The crude product was purified by flash-chromatography on silicagel with ethyl acetate/isopropanol (95:5) to obtain 1.16g of a product mixture, comprising at least 2 products with the desired mass in LC-MS. The product mixture was used as such for the next step.

HR-MS (major isomer): calculated for C₆₉H₉₄N₈O₁₂: [M+H]⁺: 1227.70640; [M+NH₄]⁺: 1244.73295; [M+Na]⁺: 1249.68834 . Found: [M+H]⁺: 1227.70599; [M+NH₄]⁺: 1244.73200; [M+Na]⁺: 1249.68733.

HR-MS (minor isomer): calculated for C₆₉H₉₄N₈O₁₂: [M+H]⁺: 1227.70640; [M+NH₄]⁺: 1244.73295; [M+Na]⁺: 1249.68834 . Found: [M+H]⁺: 1227.70598; [M+NH₄]⁺: 1244.73198; [M+Na]⁺: 1249.68751.

Second example for the synthesis of compound 8 (10g scale)

Compound **7** (10.0g, 8.13 mmol) was dissolved in tetrahydrofuran (127 mL) and dimethylsulfoxide (42 mL). To this solution, 1-Hydroxy-1,2-benziodoxol-3(1*H*)-one 1-

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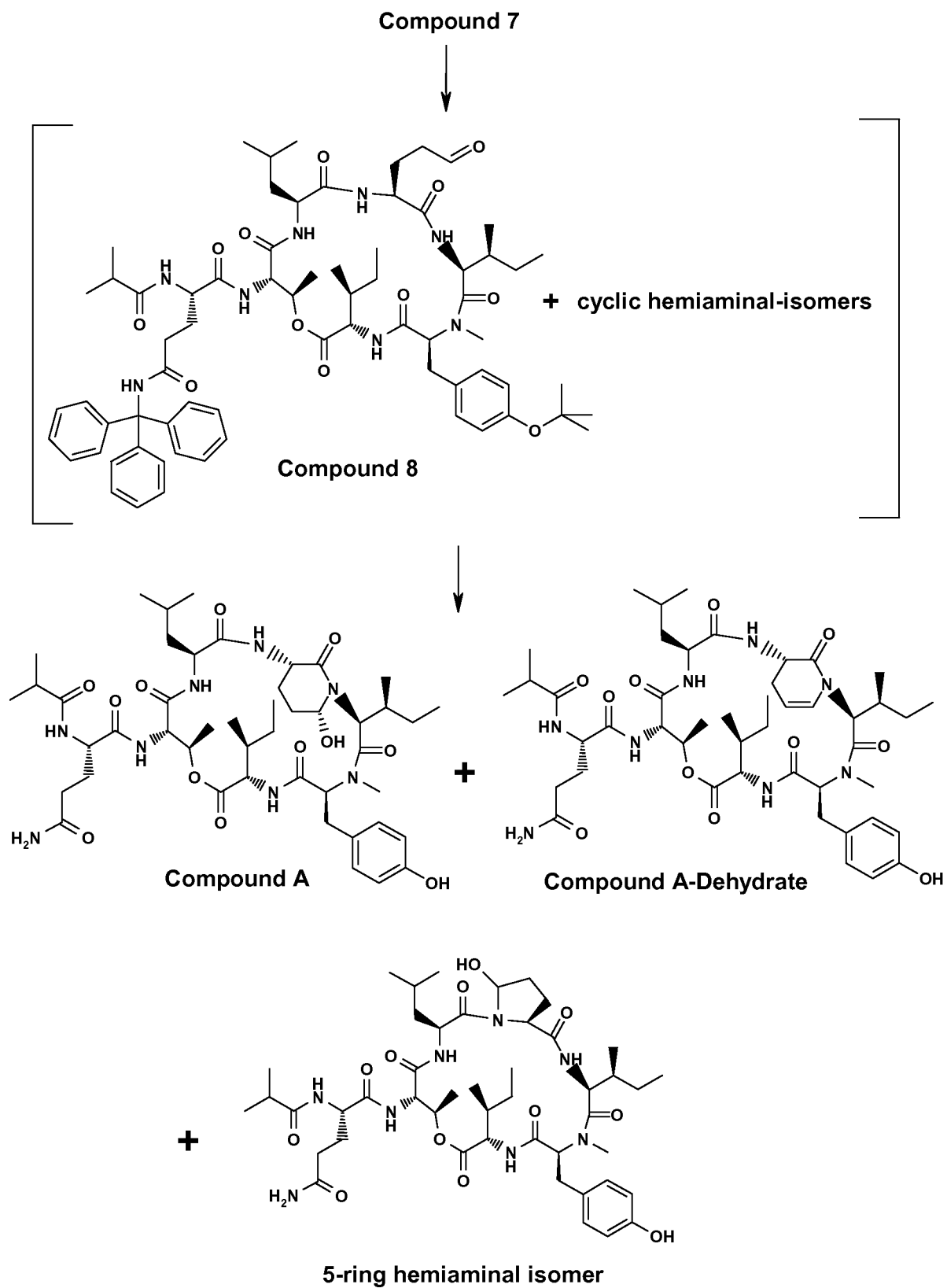
oxide (IBX) (15.18g, 45 %m/m, 24.4 mmol) was added under intense stirring at room temperature. The reaction mixture was stirred for 16h at room temperature. The reaction mixture was then poured onto aq. NaHCO₃-solution (500 mL) and ethyl acetate (250 mL) was added. The biphasic mixture was stirred for 30 minutes and the layers were separated. The organic phase was washed with half-saturated aq. NaCl-solution (250 mL) and the layers were separated. The water layers were extracted with ethyl acetate ((250 mL) and the organic phases were combined. The combined organic phase was dried on anhydrous magnesium sulfate and the solvent was partly evaporated to obtain ca. 50g solution. This solution was flashed over a silica gel column (100g silica gel) using ethyl acetate/methanol (98:2 v/v) as mobile phase. The product solution (537.4g) thus obtained was treated with toluene (135 mL) and the resulting solution was concentrated to 112g final weight by partly evaporation of the solvent at 45°C under reduced pressure. The resulting solution was cooled to 0 °C and heptanes (135 mL) was added during 30 min. The suspension thus obtained was stirred for 1h at 0°C, the product was isolated by filtration and washed with heptanes (2x30 mL).. Finally the product was dried in vacuo at 45 °C over night to obtain 8.824g of a mixture of compound **8** and its hemiaminal-isomers. Yield: 88.4%.

1D Synthesis of compound A

(S)-N¹-((2S,5S,8S,11R,12S,15S,18S,21R)-2,8-di-(S)-sec-butyl-21-hydroxy-5-(4-hydroxybenzyl)-15-isobutyl-4,11-dimethyl-3,6,9,13,16,22-hexaoxo-10-oxa-1,4,7,14,17-pentaazabicyclo[16.3.1]docosan-12-yl)-2-isobutyramidopentanediamide

Reaction Scheme 4:

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Compound 8 (2.0g) was dissolved in dichloromethane (400 mL) and the solution was cooled down to 0 °C. Trifluoroacetic acid (115.9g) was added to the stirred solution at 0 °C and the reaction mixture was stirred for 4h at 0 °C. Dichloromethane (400 mL) was added at this temperature, followed by the addition of water (20g). The reaction mixture was allowed to warm-up to room temperature and stirring was continued for additional 5 hours at room temperature. For work-up, the reaction mixture was poured onto a stirred solution of sodium acetate (165.1g) in water (800 mL) and ethyl acetate (400 mL) was added to obtain a solution. The upper layer (aqueous phase) was removed and the lower organic phase (dichloromethane phase) was washed with water (2x200 mL). The water layers were extracted with ethyl acetate (200 mL) and the organic layers were combined. The solvent was removed under reduced pressure to obtain crude Compound A as a mixture of 5- and 6-ring isomers (see Reaction Scheme 5), accompanied by trityl alcohol and other byproducts of the reaction.

The crude product was purified by RP-chromatography on Silica Kromasil 100-10-C8 with a gradient of acetonitrile/water as mobile phase. The product containing fractions were evaporated to remove acetonitrile, the precipitate was dissolved in ethyl acetate and the solvent was removed under reduced pressure to obtain 0.895g of compound A; yield: 59.1%. The product was fully characterized by NMR and MS and the spectra of the product were identical to Compound A from fermentation.

HR-MS: Calculated for $C_{46}H_{72}O_{12}N_8$: $[M+H]^+$: 929.53425; $[M+NH_4]^+$: 946.56080; $[M+Na]^+$: 951.51619. Found: $[M+H]^+$: 929.53445; $[M+NH_4]^+$: 946.56129; $[M+Na]^+$: 951.51624.

1H -NMR (600 MHz, d_6 -DMSO) δ_H : -0.11 (3H, d, J= 6.2Hz), 0.64 (4H, m), 0.77 (3H, d, J= 6.2Hz), 0.81 (3H, t, J= 7.3Hz), 0.84 (3H, d, J= 7.0 Hz), 0.88 (3H, d, J= 6.6Hz), 1.02 (3H, d, J= 6.7Hz), 1.02 (1H, m), 1.03 (3H, d, J= 6.7Hz), 1.09 (1H, m), 1.20 (3H, d, J= 6.2Hz), 1.24 (1H, m), 1.39 (1H, m), 1.51 (1H, m), 1.75 (6H, m), 1.83 (1H, m), 1.92 (1H, m), 2.12 (2H, m), 2.47 (1H, m), 2.58 (1H, m), 2.67 (1H, m), 2.71 (3H, s), 3.16 (1H, d, J= 14.2Hz), 4.30 (1H, m), 4.34 (1H, m), 4.42 (1H, d, J= 10.6Hz), 4.45 (1H, m), 4.61 (1H, d, J= 9.2Hz), 4.71 (1H, dd, J= 9.5, 5.5Hz), 4.93 (1H, s), 5.05 (1H, dd, J= 11.4, 2.6 Hz), 5.48 (1H, m), 6.07 (1H, d, J= 2.6Hz), 6.64 (2H, d, J= 8.4Hz), 6.73 (1H, s), 6.99 (2H, d, J= 8.4Hz), 7.25 (1H, s), 7.35 (1H, d, J= 9.2Hz), 7.64 (1H, d, J= 9.5Hz), 7.73 (1H, d, J= 9.2Hz), 8.01 (1H, d, J= 7.7Hz), 8.42 (1H, d, J= 8.8Hz), 9.17 (1H, s).

^{13}C -NMR (150 MHz, d_6 -DMSO) δ_C : 10.35, CH₃; 11.21 CH₃; 13.85, CH₃; 16.00, CH₃; 17.68, CH₃; 19.52, 2 x CH₃; 20.89, CH₃; 21.75, CH₂; 23.30, CH₃; 23.74, CH₂; 24.21, CH; 24.48, CH₂; 27.35, CH₂; 29.78, CH₂; 30.08, CH₃; 31.49, CH₂; 33.18, CH; 33.24, CH₂;

33.76, CH; 37.41, CH; 39.23, CH₂; 48.84, CH; 50.69, CH; 52.11, CH; 54.17 CH; 54.70, CH; 55.31, CH; 60.66, CH; 71.89, CH; 73.97, CH; 115.32, 2 x CH; 127.34, Cq; 130.37, 2 x CH; 156.27, Cq; 169.12, Cq; 169.29, Cq; 169.37, Cq; 169.79 Cq; 170.65, Cq; 172.40, Cq; 172.53, Cq; 173.87 Cq; 176.38, Cq.

Second example for the synthesis of compound A (8.5g scale)

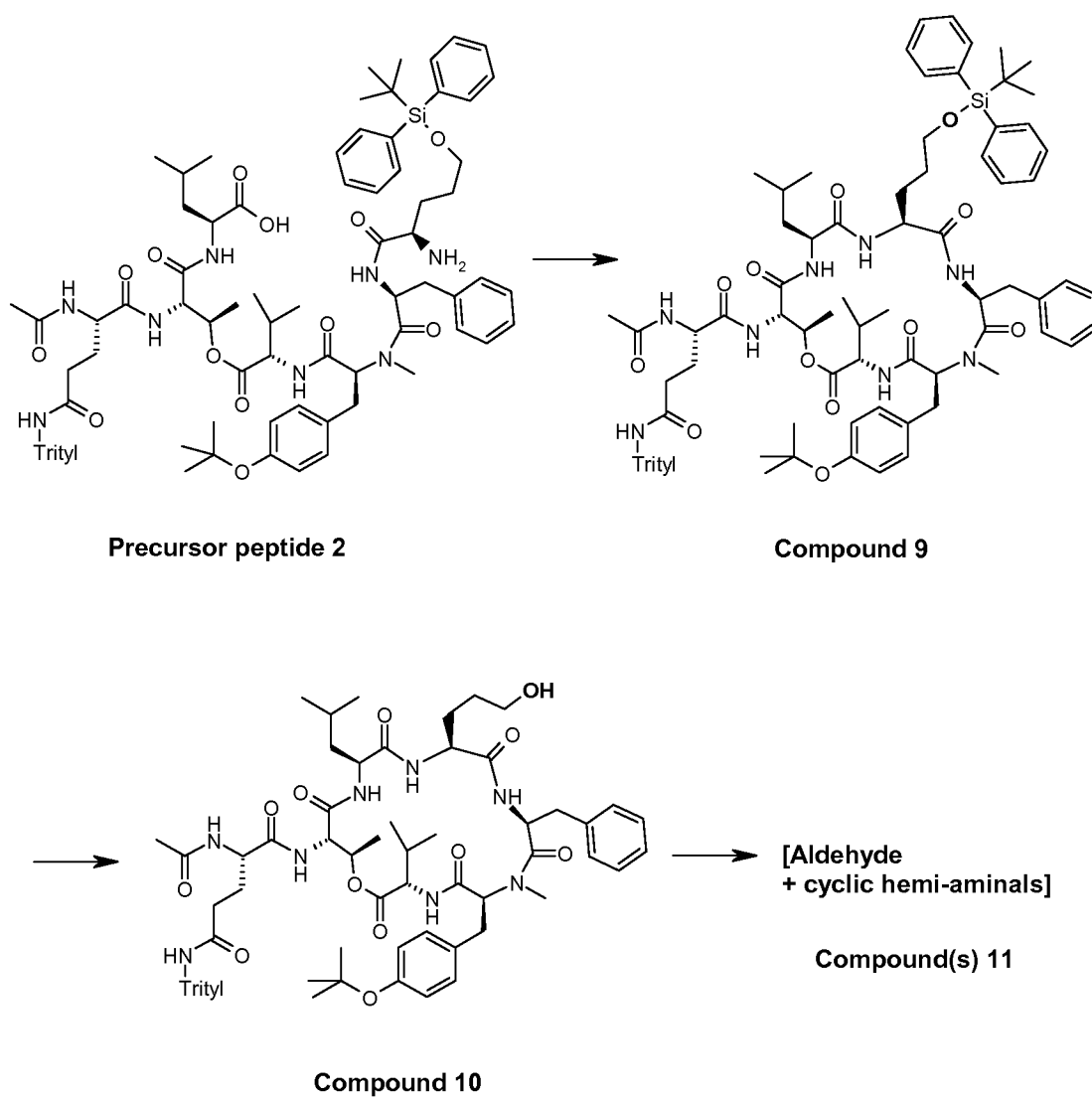
Compound **8** (8.5g, 6.924 mmol) was dissolved in dichloromethane (595 mL) and the solution was cooled to 0 °C. A solution of trifluoroacetic acid (127.5 mL) in dichloromethane (127.5 mL) was added to the cooled solution, maintaining the temperature at 0-5 °C. The reaction mixture was stirred for 5.5 h at 0 °C and was diluted with dichloromethane (850 mL), followed by addition of water (42.5 mL). The reaction mixture was allowed to warm up to room temperature and was stirred for 17.5h at room temperature. For work-up, the reaction mixture was poured onto a stirred, biphasic mixture of aqueous sodium acetate (169g sodium acetate in 722g water) and ethyl acetate (700 mL). The phases were separated and the organic phase was washed again with aqueous sodium acetate solution (169g sodium acetate in 722g water). The layers were separated and the organic phase was washed with water (2x850 mL). The individual aqueous phases were extracted with ethyl acetate (700 mL) and the organic layers were combined. The organic layer was dried over anhydrous magnesium sulfate and the solvent was partly evaporated at 40-45 °C under reduced pressure to obtain 187g of a thin suspension. Heptane (187g) was added within 15 minutes and the suspension formed was cooled to 0 °C. The suspension was stirred for 1h at 0 °C and the product was isolated by filtration. The crude product was dried in vacuo at 40 °C to obtain 5.65g crude compound **A**. Crude yield: 87.8%.

0.96g of the crude product was purified by RP-chromatography on Silica Kromasil 100-10-C8 with a gradient of acetonitrile/water as mobile phase. The product containing fractions were evaporated to remove acetonitrile, the precipitate was dissolved in ethyl acetate and the solvent was partly removed under reduced pressure to obtain 40g of a solution. Heptane (40g) was added to the stirred solution during 30 min at room temperature and the resulting suspension was stirred at room temperature for additional 2h. The product was isolated by filtration and washed with ethyl acetate/heptane (1:1, 2x10mL). The product was dried in vacuo at 40-45 °C for 16h to obtain 0.702g of compound **A**. Yield after RP-chromatography and precipitation: 64%.

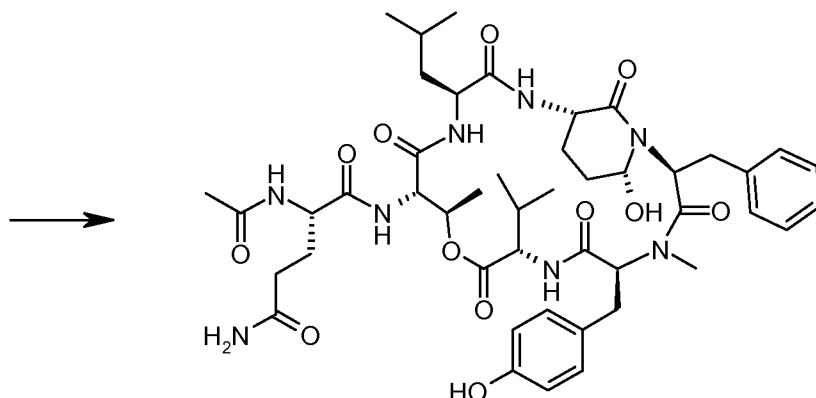
Example 2: Synthesis of Compound B (Nostopeptin BN920)

(*S*)-2-acetamido-*N*¹-((1*S*,2*S*,5*S*,8*S*,11*R*,12*S*,15*S*,18*S*,21*R*)-2-benzyl-21-hydroxy-5-(4-hydroxybenzyl)-15-isobutyl-8-isopropyl-4,11-dimethyl-3,6,9,13,16,22-hexaoxo-10-oxa-1,4,7,14,17-pentaazabicyclo[16.3.1]docosan-12-yl)pentanediamide

Reaction Scheme 5:



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**Compound B (Nostopeptin BN920)****2A Synthesis of Precursor Peptide 2 by SPPS:****Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Phe-Synthon1-H)-Leu-OH****Equipment:**

Peptide-synthesizer equipped with a 250 ml glass-reactor with frit and manifold for automatic solvent delivery, shaking and sucking off reagents.

2A(1) Synthesis of Fmoc-Thr-Leu-Trt-Tentagel-S

Fmoc-Leu-Trt-Tentagel-S-Resin (18.7g loading 0.37mmol/g (supplied by Rapp Polymere GmbH, Tübingen/Germany)) were swollen in DMF by shaking for 30 min.

The Fmoc protecting group was cleaved by two subsequent treatments with 20% piperidine in DMF for 5min and 15min respectively. After resin washing by several alternating washes with DMF and isopropanol the complete removal of bases was checked with the absence of a pink color after Phenolphthalein and water addition to the last washing step.

4.7g of Fmoc-Thr-OH, 5.26g of HATU and 1.8g of DIPEA were dissolved in 50ml of DMF. After 5min of stirring additional 1.8g of DIPEA were added. After checking the pH (>11) the mixture was added to the deprotected resin and shaken for 2h. The performed Kaiser test was OK and the resin was washed by several alternating washes with DMF and isopropanol. After drying the resin weight was 19.01g. A small sample was cleaved and checked by HPLC. A single main peak showed a successful conversion.

2A(2) Synthesis of Fmoc-Gln(Trt)-Thr-Leu-Trt-Tentagel-S

19.01g Fmoc-Thr-Leu-Trt-Tentagel-S-Resin (6.6 mmol) were pre-swollen in DMF and the Fmoc protecting group was cleaved by two subsequent treatments of 20% piperidine in DMF for 5min and 15min respectively. After resin washing by several alternating washes with DMF and isopropanol the complete removal of bases was checked with the absence of a pink color after Phenolphthalein and water addition to the last washing step.

8.07g of Fmoc-Gln(Trt)-OH, 5.01g of HATU and 3.4g of DIPEA were dissolved in 50ml of DMF. After checking the pH (>11) the mixture was added to the deprotected resin and shaken for 1.5h. The performed Kaiser test was OK and the resin was washed by several alternating washes with DMF and isopropanol. The resulting resin was directly used in the following step below, only a small sample was cleaved and checked by HPLC. A single main peak showed successful conversion.

2A(3) Synthesis of Ac-Gln(Trt)-Thr-Leu-Trt-Tentagel-S

The Fmoc-Gln(Trt)-Thr-Leu-Trt-Tentagel-S resin from above was re-swollen in DMF by shaking in DMF for 10min. The Fmoc protecting group was cleaved by two subsequent treatments with 20% piperidine in DMF for 5min and 15min respectively. The resin was washed by several alternating washes with DMF and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink color proved successful removal of piperidine.

0.774g of acetic-acid, 6.707g of PyBOP and 3.33g of DIPEA were dissolved in 50ml of DMF. After checking the pH (>11) the mixture was added to the deprotected resin and shaken for 2.5h. The performed Kaiser test was OK and the resin was washed by several alternating washes with DMF and isopropanol. The resulting resin was directly used in the following step below, only a small sample was cleaved and checked by HPLC. A single main peak showed successful conversion.

2A(4) Synthesis of Ac-Gln(Trt)-Thr(Val-Fmoc)-Leu-Trt-Tentagel-S (Previously named: Ac-Gln(Trt)-Thr(Val-Fmoc)-Leu-Trt-Tentagel-S) (side-chain esterification)

The Ac-Gln(Trt)-Thr-Leu-Trt-Tentagel-S resin from above was re-swollen in DMF by shaking in DMF for 10min.

8.7g of Fmoc-Val-OH and 8.3g of DIPEA were dissolved in 25ml of DCM. In parallel 2.76g of MSNT were dissolved in another 25ml of DCM. Both solutions were combined

and after 3min pre-activation put to the peptide resin and shaken for 2h. The resin was washed by several alternating washes with DMF and isopropanol. The resulting resin was directly used in the following step below, only a small sample was cleaved and checked by HPLC. A single main peak showed successful conversion.

**2A(5) Synthesis of Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Fmoc)-Leu-Trt-Tentagel-S
(Previously named: Ac-Gln(Trt)-Thr(Val-N-me-Tyr(tBu)-Fmoc)-Leu-Trt-Tentagel-S)**

The Ac-Gln(Trt)-Thr(Val-Fmoc)-Leu-Trt-Tentagel-S resin from above was re-swollen in DMF by shaking in DMF for 10min.

The Fmoc protecting group was cleaved by two subsequent treatments with 20% piperidine in DMF for 5min and 15min respectively. The resin was washed by several alternating washes with DMF and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink color proved successful removal of piperidine.

6.1g of Fmoc-N-Me-Tyr(tBu)-OH, 4.8g of HATU and 3.3g of DIPEA were dissolved in 50ml of DMF. After checking the pH (>11) the mixture was added to the deprotected resin and shaken for 2.5h. The performed Kaiser test was OK and the resin was washed by several alternating washes with DMF and isopropanol. The resulting resin was directly used in the following step below, only a small sample was cleaved and checked by HPLC. A single main peak showed successful conversion.

**2A(6) Synthesis of Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Phe-Fmoc)-Leu-Trt-Tentagel-S
(Previously named: Ac-Gln(Trt)-Thr(Val-N-me-Tyr(tBu)-Phe-Fmoc)-Leu-Trt-Tentagel-S)**

Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Fmoc)-Leu-Trt-Tentagel-S resin from above was re-swollen in DMF by shaking in DMF for 10min.

The Fmoc protecting group was cleaved by two subsequent treatments with 20% piperidine in DMF for 5min and 15min respectively. The resin was washed by several alternating washes with DMF and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink color proved successful removal of piperidine.

9.66g of Fmoc-Phe-OH, 9.48g of HATU and 6.4g of DIPEA were dissolved in 100ml of DMF. After checking the pH (>11) the mixture was added to the deprotected resin and shaken for 2h. The performed Kaiser test was OK and the resin was washed by several

alternating washes with DMF and isopropanol. The resulting resin was directly used in the following step below, only a small sample was cleaved and checked by HPLC. A single main peak showed successful conversion.

**2A(7) Synthesis of Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Phe-Synthon1-H)-Leu-OH
(Previously named: Ac-Gln(Trt)-Thr(Val-N-me-Tyr(tBu)-Phe-Synthon1-H)-Leu-OH)
(= Precursor peptide 2)**

Ac-Gln(Trt)-Thr(Val-Tyr(tBu)Me-Phe-Fmoc)-Leu-Trt-Tentagel-S resin from above was re-swollen in DMF by shaking in DMF for 10min.

The Fmoc protecting group was cleaved by two subsequent treatments with 20% piperidine in DMF for 5min and 15min respectively. The resin was washed by several alternating washes with DMF and isopropanol. Phenolphthalein and water were added to a sample of the final wash solution. The absence of pink color proved successful removal of piperidine.

6.77g of Synthon1, 5.9g of PyBOP and 2.95g of DIPEA were dissolved in 100ml of DMF. After checking the pH (>11) the mixture was added to the deprotected resin and shaken for 2h. The performed Kaiser test was OK and the resin washed by several alternating washes with DMF and isopropanol. After taking a small sample for a HPLC check, the Fmoc protecting group was cleaved by two subsequent treatments with 20% piperidine in DMF for 5min and 15min respectively. The resin was washed by several alternating washes with DMF and isopropanol. Finally the resin was washed two times with DCM. Cleavage of the synthesized peptide from the resin was achieved by shaking in a mixture of 80 % of acetic acid in DCM over night. The resulting peptide solution was filtered off and the resin was washed two times with DCM. The combined filtrates were evaporated and finally purified by RP-chromatography using a gradient system. The collected fractions were analyzed by HPLC and the pure fractions were pooled, evaporated and finally lyophilized.

Yield: 3.54g (42%) Precursor peptide 2.

HR-MS: Calculated for $C_{85}H_{108}N_8O_{13}Si$: $[M+H]^+ = 1477.78779$; Found: $[M+H]^+ = 1477.78691$.

2B Synthesis of Compound B

(*S*)-2-acetamido-*N*¹-((1*S*,2*S*,5*S*,8*S*,11*R*,12*S*,15*S*,18*S*,21*R*)-2-benzyl-21-hydroxy-5-(4-hydroxybenzyl)-15-isobutyl-8-isopropyl-4,11-dimethyl-3,6,9,13,16,22-hexaoxo-10-oxa-1,4,7,14,17-pentaazabicyclo[16.3.1]docosan-12-yl)pentanediamide

2B(0) Synthesis of Compound 9

(*S*)-2-acetamido-*N*¹-((3*S*,6*S*,9*S*,12*S*,15*S*,18*S*,19*R*)-9-benzyl-6-(4-(*tert*-butoxy)benzyl)-12-(3-((*tert*-butyldiphenylsilyl)oxy)propyl)-15-isobutyl-3-isopropyl-7,19-dimethyl-2,5,8,11,14,17-hexaoxo-1-oxa-4,7,10,13,16-pentaazacyclononadecan-18-yl)-*N*⁵-tritylpentanediamide

The lactamization of Precursor peptide 2 to obtain Compound B was performed under similar conditions described for the preparation of Compound 6. The product, Compound 9, was fully characterized by IR, NMR and MS. The spectra confirmed the proposed structure.

HR-MS: Calculated for C₈₅H₁₀₆N₈O₁₂Si: [M+H]⁺: 1459.77723; [M+NH₄]⁺: 1476.80377.

Found: [M+H]⁺: 1459.77719; [M+NH₄]⁺: 1476.80291.

2B(1) Synthesis of Compound 10

(*S*)-2-acetamido-*N*¹-((3*S*,6*S*,9*S*,12*S*,15*S*,18*S*,19*R*)-9-benzyl-6-(4-(*tert*-butoxy)benzyl)-12-(3-hydroxypropyl)-15-isobutyl-3-isopropyl-7,19-dimethyl-2,5,8,11,14,17-hexaoxo-1-oxa-4,7,10,13,16-pentaazacyclononadecan-18-yl)-*N*⁵-tritylpentanediamide

The de-silylation reaction to obtain compound 10 was performed under similar conditions described for the preparation of compound 7. The product was fully characterized by IR, NMR and MS. The spectra confirmed the proposed structure.

HR-MS: Calculated for C₆₉H₈₈N₈O₁₂: [M+H]⁺: 1221.65945; [M+NH₄]⁺: 1238.6860;

[M+Na]⁺: 1243.64139. Found: [M+H]⁺: 1221.65894; [M+NH₄]⁺: 1238.68518; {M+Na}⁺: 1243.64001.

2B(2) Synthesis of Compound 11 (Aldehyde)

(*S*)-2-acetamido-*N*¹-((3*S*,6*S*,9*S*,12*S*,15*S*,18*S*,19*R*)-9-benzyl-6-(4-(*tert*-butoxy)benzyl)-15-isobutyl-3-isopropyl-7,19-dimethyl-2,5,8,11,14,17-hexaoxo-12-(3-oxopropyl)-1-oxa-4,7,10,13,16-pentaazacyclononadecan-18-yl)-*N*⁵-tritylpentanediamide

The oxidation of Compound 10 to obtain Compound 11 was performed under similar

conditions described for the preparation of Compound 8. LC-MS analysis showed several peaks with the desired mass, indicating the presence of a mixture of the aldehyde, 5-ring-hemiaminal and 6-ring-hemiaminal. The mixture was used as such for the next step. HR-MS (major peak): Calculated for $C_{69}H_{86}N_8O_{12}$: $[M+H]^+$: 1219.64380; $[M+NH_4]^+$: 1236.67035; $[M+Na]^+$: 1241.62574. Found: $[M+H]^+$: 1219.64404; $[M+NH_4]^+$: 1236.67053; $[M+Na]^+$: 1241.62524.

2B(3) Synthesis of Compound B (Nostopeptin BN920) from Compound 11

(S)-2-acetamido-*N*¹-((1*S*,2*S*,5*S*,8*S*,11*R*,12*S*,15*S*,18*S*,21*R*)-2-benzyl-21-hydroxy-5-(4-hydroxybenzyl)-15-isobutyl-8-isopropyl-4,11-dimethyl-3,6,9,13,16,22-hexaoxa-10-oxa-1,4,7,14,17-pentaazabicyclo[16.3.1]docosan-12-yl)pentanediamide

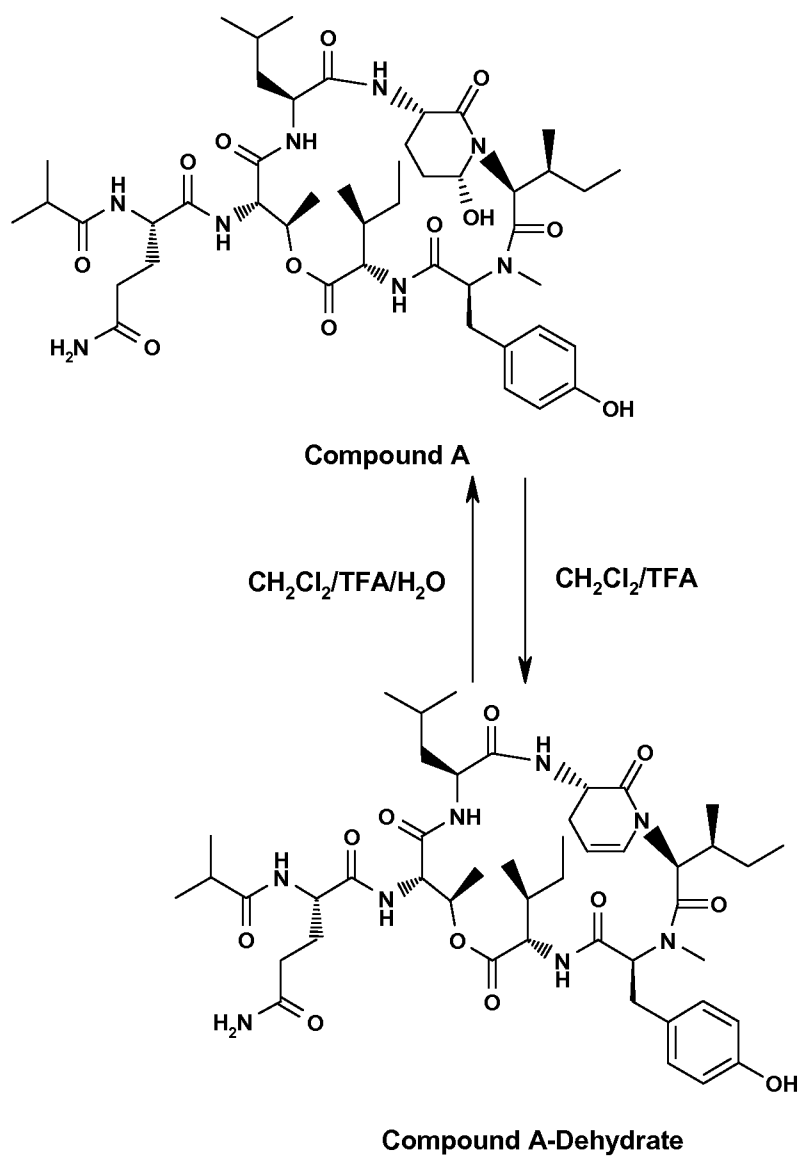
Compound(s) 11 (1.0g, 0.82 mmol) was dissolved in dichloromethane (200 mL). Trifluoroacetic acid (57.8g) was added at 15 to 25 °C during 10 minutes and the reaction mixture was stirred for 45 hours at room temperature. The reaction mixture was then diluted with dichloromethane (200 mL) and water (10 mL) was added. Stirring was continued for additional 24 hours at room temperature. The reaction mixture was poured onto a saturated aq. $NaHCO_3$ -solution (700 mL) within 20 minutes and dichloromethane (500 mL), isopropanol (50 mL) and water (500 mL) were added sequentially. The layers were separated and the aqueous layer was extracted with a solution of isopropanol (50 mL) in dichloromethane (500 mL). The layers were again separated and the aqueous layer was extracted several times with dichloromethane (5x300 mL). The organic layers were combined and the solvent was evaporated under reduced pressure to obtain crude product mixture (1.0 g), comprising the desired Nostopeptin BN920 along with trityl alcohol and other byproducts of the reaction. The crude product was purified by silica gel chromatography with dichloromethane/isopropanol as mobile phase to obtain pure Nostopeptin BN920 (212 mg, 97 % (A) purity). Collection of less pure fractions gave additional 402 mg Product with 90 % (A) purity. Yield from all fractions: 614 mg (81%). The product was fully characterized by IR, NMR and MS. The spectra confirmed the proposed structure.

HR-MS: Calculated for $C_{46}H_{64}N_8O_{12}$: $[M+H]^+$: 921.47165; $[M+NH_4]^+$: 938.49820; $[M+Na]^+$: 943.45359. Found: $[M+H]^+$: 921.47167; $[M+NH_4]^+$: 938.49861; $[M+Na]^+$: 943.45337.

Example 3: Shift of Equilibrium

Reaction Scheme 4 shows the presence of the dehydrate form of Compound A. It has now been discovered that this can be converted easily (back) into Compound A using a simple procedure for hydration of the dehydrate form depicted in the following reaction scheme:

Reaction Scheme 6:



This allows to improve the yield of Compound A in any type of synthesis (be it chemical as in the present disclosure or by use of fermentation as in WO2009/024527).

For example, during cleavage of acid sensitive protecting groups in a compound comprising the ahp-subunit, e.g. Compound A in Example 1, the formation of large amounts of the corresponding dehydrated byproduct is observed. This byproduct is usually separated e.g. by chromatography and disposed. This leads to loss of valuable product and to low yield for this step. For example, if the oxidation product(s) of Compound 8 are subjected to acidic conditions to cleave the trityl- and t-butyl protecting groups (scheme 4), significant amounts of Compound-A-dehydrate are formed as byproduct. Depending on the acid concentration and reaction conditions, Compound-A-dehydrate might be formed even as major product in this product mixture.

For example, a ratio Compound A/Compound A-dehydrate (1:2) was observed when trifluoroacetic acid/dichloromethane (5:95 v/v) was used to cleave the protecting groups after the oxidation step (Example 1).

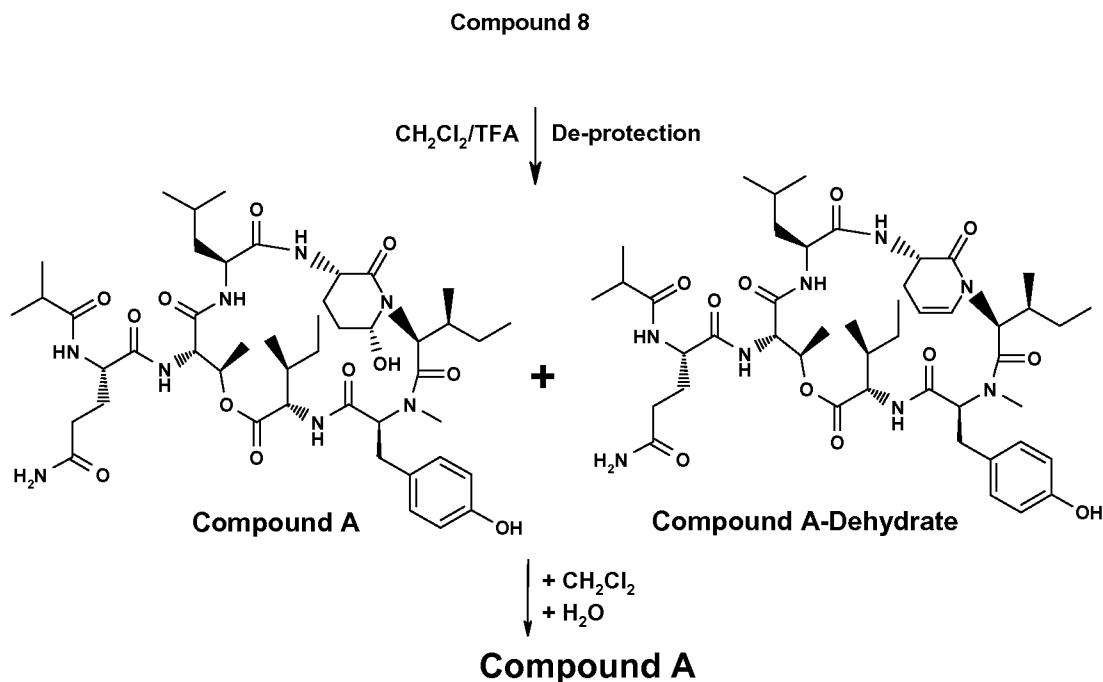
It was therefore searched for ways to convert the dehydrate byproduct into the desired product. It has now been found that this can be achieved by acid catalyzed equilibration of the product mixture in the presence of water under well defined conditions. Addition of water to the reaction mixture of example 1 and subsequent stirring at room temperature for 19h gave a product mixture with a ratio Compound A/Compound A-dehydrate of ca. 96:4. Thus, addition of water to the reaction mixture after the acid-catalyzed de-protection step (scheme 4) changed the ratio of Compound A / Compound A -dehydrate from (1:2) under water free conditions to (96:4) after water addition and equilibration.

The formation of Compound A -dehydrate from Compound A under acidic de-protection conditions was confirmed by conversion of pure Compound A into Compound A -dehydrate using trifluoroacetic acid in DCM. Treatment of Compound A with 33% (v/v) TFA in DCM for 2h at room temperature gave a product mixture of Compound A -dehydrate/ Compound A in a ratio of 78:22 according to HPLC. The dehydration could be driven to >95% conversion, when water absorbing agents, such as molecular sieves were added to the reaction mixture. Thus, stirring of pure Compound A in a 1:2 mixture of TFA/DCM in the presence of molecular sieves gave the dehydrated-product in

quantitative crude yield and ca. 96 area% HPLC purity (example 3 B). There were still ca. 4 area% of Compound A present in the crude product.

Conversion of Compound A -dehydrate from Example 3 B into Compound A was demonstrated by stirring Compound A-dehydrate in dichloromethane in the presence of trifluoroacetic acid and water (Example 3 C). The product thus obtained comprised 95.6 area% Compound A and only 4.4 area% Compound A -dehydrate according to HPLC.

Reaction Scheme 7:



Experimental details for Example 3

3A:

23 mg product mixture from oxidation (compound 8 and cyclic amins derived from it) was dissolved in DCM (5.7 mL) and TFA (0.3 mL) was added to the solution under intensive stirring. The reaction mixture was stirred for 5h at room temperature, until an IPC (HPLC) indicated complete cleavage of the protecting groups. Compound A and Compound A-dehydrate were present in the reaction mixture in a ratio of 1:2, along with additional byproducts. The reaction mixture was diluted with DCM (5.7 mL) and the

intensively stirred solution was treated with water (0.23 mL). HPLC after 19h stirring time at room temperature indicated a ratio of 96:4 for Compound A / Compound A -dehydrate.

For workup, the reaction mixture was poured onto a solution of sodium acetate (1.62g) in water (23 mL) and ethyl acetate (35 mL) was added. The phases were separated and the organic layer was extracted with water (2x25 mL). The water layers were extracted with ethyl acetate (35 mL) and the organic layers were combined. The solvent was evaporated at reduced pressure to obtain 22 mg of crude product as a foam. The crude product was purified by flash chromatography on silica gel with ethyl acetate/methanol 95:5 to 90:10 to obtain Compound A in 96.8 area% purity, accompanied by 1.8 area% of the 5-ring isomer.

3B:

Compound A (250 mg) was dissolved in dichloromethane (10 mL) and trifluoroacetic acid (5 mL) was added, followed by the addition of molecular sieves (1g). IPC (HPLC) after 1h indicated a ratio of 17:83 for Compound A /Compound A -dehydrate. The reaction mixture was stirred for a total of 72h at room temperature. For workup, the molecular sieves were removed by filtration and the solution was poured onto saturated aq. NaHCO₃-solution. The aqueous layer was extracted with ethyl acetate (50 mL). The organic phase was washed with water (20 mL) and the solvent was evaporated at reduced pressure and the residue was dried in vacuo to obtain quantitative yield (245 mg) crude product as a foam. HPLC analysis of the crude product indicated the presence of Compound A / Compound A-dehydrate at a ratio of ca. 4:96 area%.

HR-MS: calculated for C₄₆H₇₀N₈O₁₁ [M+H]⁺= 911.52368, [M+NH₄]⁺= 928.55023, [M+Na]⁺= 933.50563; found [M+H]⁺= 911.52372, [M+NH₄]⁺= 928.55029, [M+Na]⁺= 933.50538. The structure of Compound A-dehydrate was confirmed by ¹H-NMR.

¹H-NMR (600 MHz, d₆-DMSO) δ_H: 0.06 (3H, d, J= 6.6 Hz), 0.67 (3H, t, J= 7.3 Hz), 0.70 (3H, d, J= 7.0 Hz), 0.77 (3H, d, J= 6.6 Hz), 0.81 (1H, m), 0.87 (6H, m), 1.00 (3H, d, J= 7.0 Hz), 1.02 (3H, d, J= 7.0 Hz), 1.06 (1H, m), 1.16 (3H, d, J= 6.5 Hz), 1.17 (1H, m), 1.30 (1H, m), 1.41 (1H, m), 1.53 (1H, m), 1.74 (2H, m), 1.91 (2H, m), 2.01 (1H, m), 2.11 (2H, m), 2.45 (3H, m), 2.73 (3H, s), 2.74 (1H, m), 3.18 (1H, m), 4.32 (2H, m), 4.50 (1H, m), 4.54 (1H, m), 4.64 (1H, d, J= 9.5 Hz), 4.77 (1H, d, J= 11.0 Hz), 5.18 (1H, m), 5.26 (1H, m), 5.42 (1H, q, J= 6.6 Hz), 6.25 (1H, d, J= 7.3 Hz), 6.32 (1H, d, J= 7.7 Hz), 6.67 (2H, d,

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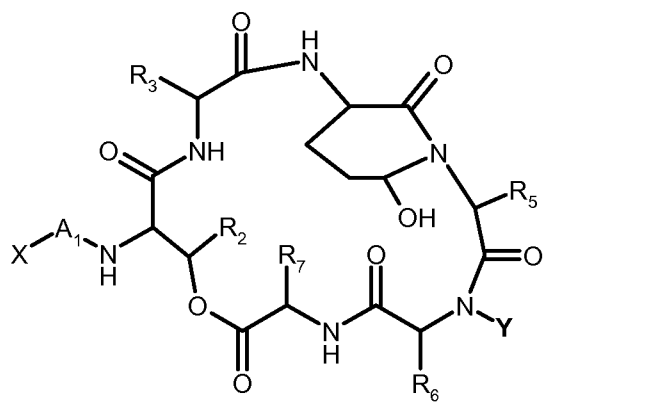
J= 8.4 Hz), 6.75 (1H, s), 7.04 (2H, d, J= 8.4 Hz), 7.25 (1H, s), 7.30 (1H, d, J= 8.8 Hz), 7.89 (1H, d, J= 9.2 Hz), 7.98 (1H, d, J= 8.1 Hz), 8.50 (1H, d, J= 8.4 Hz), 9.24 (1H, s).

3C:

Compound A-dehydrate (110 mg) from example 2 was dissolved in DCM (20 mL). TFA (1g) and water (0.2 mL) were added and the reaction mixture was stirred for 20h at room temperature. For workup, the reaction mixture was poured onto ethyl acetate (50 mL) and the ethyl acetate solution was extracted with a saturated aq. NaHCO₃-solution (50 mL). The organic layer was extracted with water (20 mL) and the solvent was evaporated. The residue was dissolved in ethyl acetate/isopropanol 9:1 (20 mL) and the solution was clear filtered over a 0.45 micrometer filter. The solvent was evaporated under reduced pressure and the residue was dried in vacuo at 45 °C to obtain 100mg crude product. HPLC analysis indicated the presence of ca. 95.6 area% Compound A and ca. 4.4 area% Compound A -dehydrate.

Claims

1. A method or process for the preparation of a cyclic depsipeptide compound of the formula I,



wherein

A₁ is a bivalent moiety of an amino acid with a terminal carboxy or carbamoyl group, especially asparagine or glutamine, and is bound at its right hand side in formula I via a carbonyl to the rest of the molecule; or is C₁₋₈-alkanoyl or phosphorylated hydroxy-C₁₋₈-alkanoyl;

X is bound via an N of A₁ and is acyl, or is absent if A₁ is C₁₋₈-alkanoyl or phosphorylated hydroxy-C₁₋₈-alkanoyl;

R₂ is C₁₋₈-alkyl, especially methyl;

R₃ is the side chain of an amino acid, especially of leucine, isoleucine or valine;

R₅ is the side chain of an amino acid, preferably of phenylalanine, leucine, isoleucine or valine;

R₆ is the side chain of a hydroxy amino acid, especially of tyrosine;

R₇ is the side chain of an amino acid, preferably of the amino acid leucine, isoleucine or valine; and

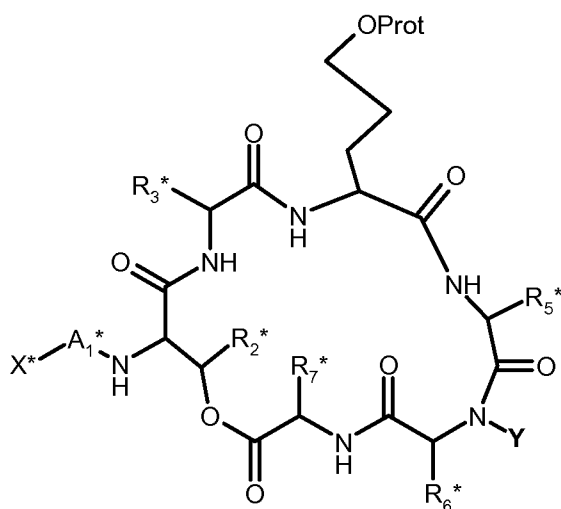
Y is hydrogen or C₁₋₈-alkyl;

or a salt thereof,

said method comprising

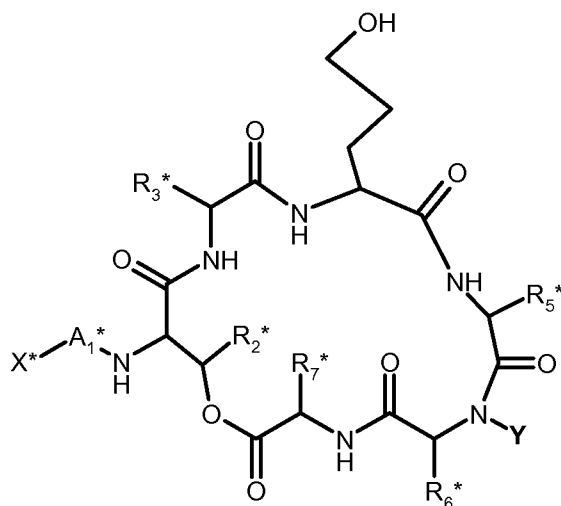
selectively deprotecting a compound of the formula II

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II

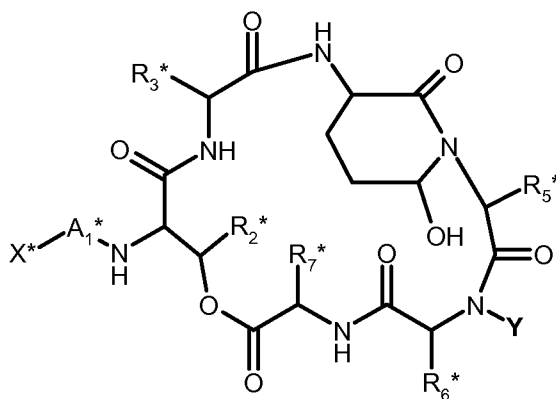
wherein Prot is a protecting group, Y is as defined for a compound of the formula I and X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* correspond to X, A_1 , R_2 , R_3 , R_5 , R_6 , and R_7 in formula I, respectively, but with the proviso that reactive functional groups on these moieties are present in protected form at least if they could participate in undesired side reactions, to result in a compound of the formula III,



(III)

wherein X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* have the meanings defined above,

reacting the free hydroxyl group under oxidizing conditions to form a compound of the formula IV



IV

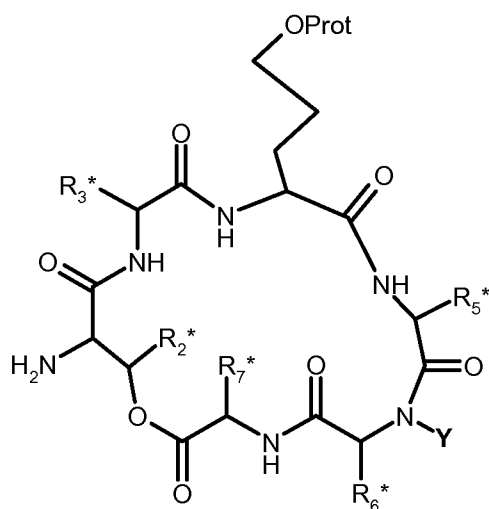
and removing remaining protecting groups to yield a compound of the formula I, or a salt thereof,

and, optionally, converting a free compound of the formula I into a salt, a salt of a compound of the formula I into a different salt of a compound of the formula I or into the free compound of the formula I and/or converting a dehydrated analogue of a compound of the formula I into the corresponding compound of the formula I.

2. The method or process according to claim 1, wherein the compound of the formula IV is synthesized by a combination of Solid Phase Peptide Synthesis and Solution Phase synthesis from the corresponding starting amino acids and side chain precursors.

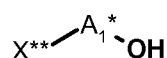
3. The method or process according to any one of the preceding claims, further comprising, for the synthesis of a compound of the formula II shown in claim 1, reacting a compound of the formula VI,

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VI

wherein Prot is a protecting group, Y is as defined for a compound of the formula I and R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* are as defined for a compound of the formula II in claim 1, with an acid of the formula VII



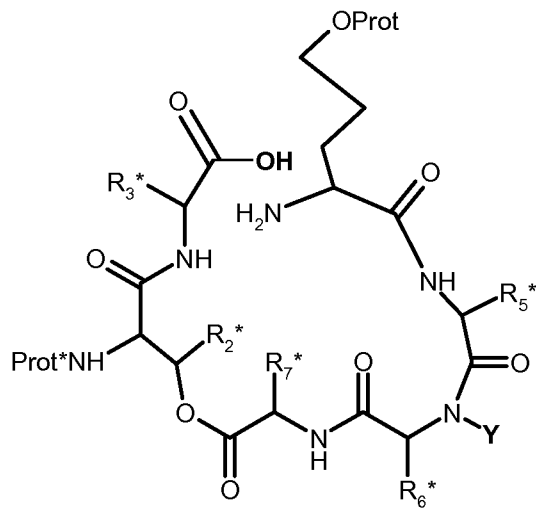
VII

or a reactive derivative thereof,

wherein X^{**} is an amino protecting group or is X^* , and wherein X^* and A_1^* are as defined for a compound of the formula II in claim 1; and, if X^{**} is an amino protecting group, removing said amino protecting group X^{**} to yield H instead of X^* and coupling the resulting amino group with an acyl group X^* using the corresponding acid X^*-OH wherein C^* is as defined for a compound of the formula II in claim 1, or a reactive derivative thereof.

4. The method or process according to claim 3, further comprising cyclization under lactamization of a linear precursor peptide of the compound of the formula VI, carrying an N-terminal amino group and a C-terminal carboxy group, under reaction conditions that allow for the formation of an amide bond from said amino and said carboxy group, preferably using Solution Phase chemistry.

5. The method or process according to claim 4, where the linear precursor peptide is of the formula VIII,

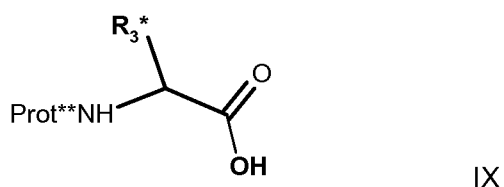


wherein Prot* is a protecting group that can be cleaved off selectively without affecting other protecting groups present and is stable during deprotection steps during synthesis of the linear precursor peptide (e.g. allyloxycarbonyl) and R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* are as defined for a compound of the formula VI in claim 4, further comprising, after cyclisation of the compound of the formula VIII, especially VIIIA, removing the protecting group Prot* *in situ* to yield the compound of the formula VI.

6. The method or process according to claim 5, where the linear precursor peptide of the formula VIII, especially VIIIA, is synthesized from the corresponding amino acids by solid phase peptide synthesis and subsequent cleaving-off from the employed solid phase.

7. The method according to claim 6, comprising either

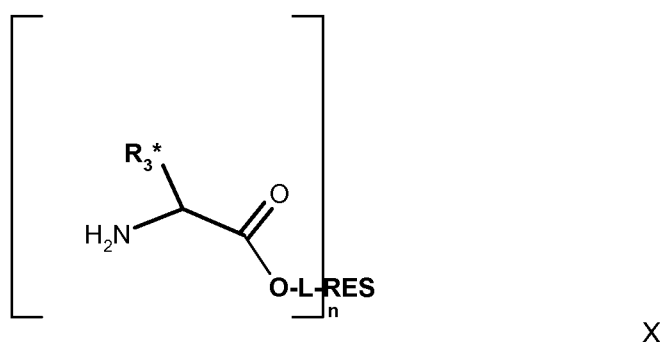
in a variant a), coupling an amino acid of the formula IX,



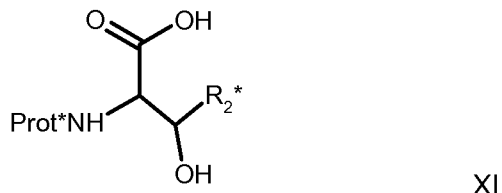
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wherein R_3^* is as defined for a compound of the formula II in claim 1 and $Prot^{**}$ is an amino protecting group that can be removed on the resin without cleaving other bonds, or a reactive derivative of said amino acid, via an oxygen to a cleavable linker L which is bound to a solid resin RES, and removing the protecting group $Prot^{**}$;

coupling the obtained resin bound amino acid symbolized by the formula X,

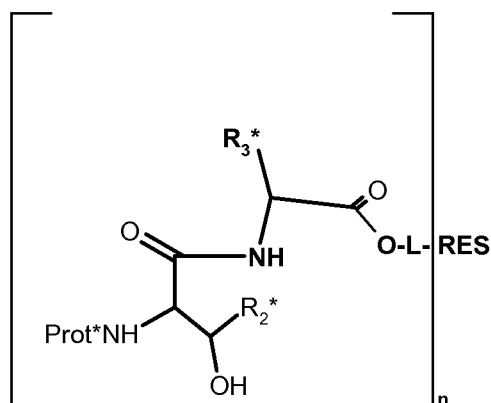


in which RES and R_3^* are as defined for a compound of the formula IX, n is a natural number and L is a cleavable linker, with an amino acid of the formula XI,



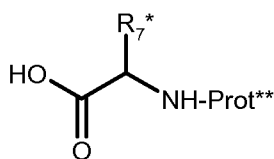
wherein $Prot^*$ is as defined for a compound of the formula VIII in claim 5 and R_2^* is as defined for a compound of the formula II in claim 1, or a reactive derivative of said amino acid, coupling the obtained resin bound dipeptide symbolized by the formula XII,

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XII

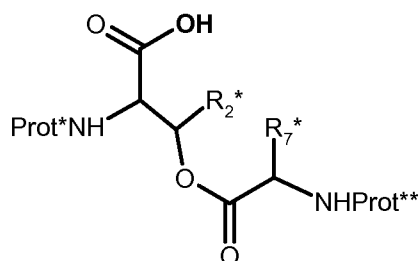
in which Prot* is as defined for a compound of the formula VIII in claim 5, R_2^* and R_3^* are as defined for a compound of the formula II in claim 1, and n , L and RES are as defined for a compound of the formula X, via the free hydroxy group with an amino acid of the formula XIII,



XIII

wherein Prot** is as defined for a compound of the formula IX and R_7^* is as defined for a compound of the formula II in claim 1, or a reactive derivative of said amino acid, and removing the protecting group Prot**;

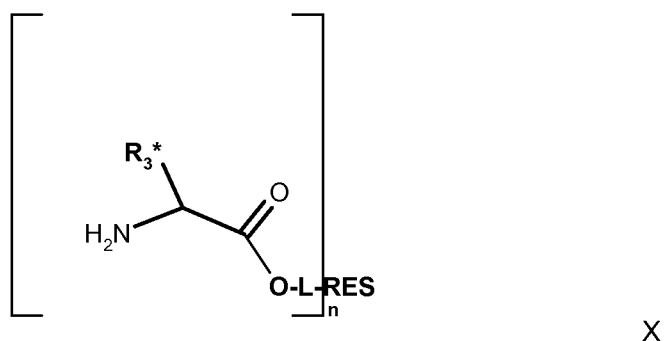
or, in a variant b), coupling a dipeptide of the formula XXVII,



XXVII

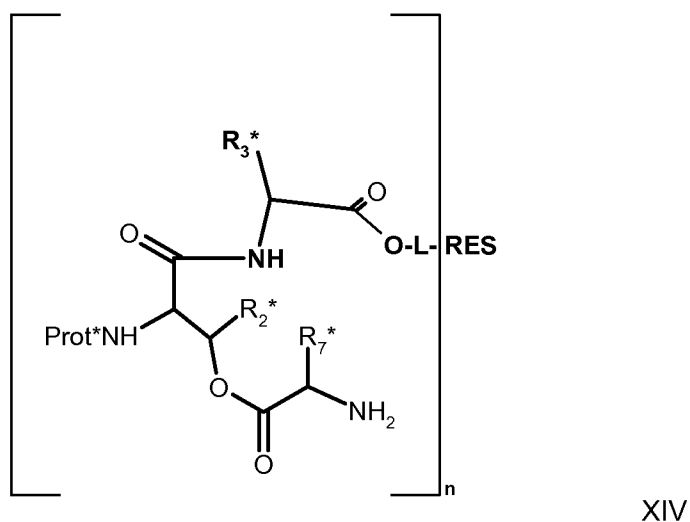
wherein R_3^* and Prot** are as described for a compound of the formula IX, especially IXA, and Prot* is as defined for a compound of the formula VIII in claim 5, or a reactive

derivative of said dipeptide, to an amino acyl moiety, bound via an oxygen to a cleavable linker L which is bound to a solid resin RES, having the formula X,



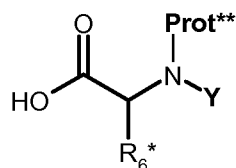
that can be obtained as described under variant a), in which RES, and R_3^* are as defined for a compound of the formula IX and L and RES are as just defined; and removing the protecting group Prot**;

and, after the reactions of variant a) or of variant b), coupling the obtained compound of the formula XIV,



wherein R₂^{*}, R₃^{*} and R₇^{*} are as defined for a compound of the formula II in claim 1, Prot^{*} is as defined for a compound of the formula VIII in claim 5 and n, L and RES are as defined for a compound of the formula X, with an amino acid of the formula XV,

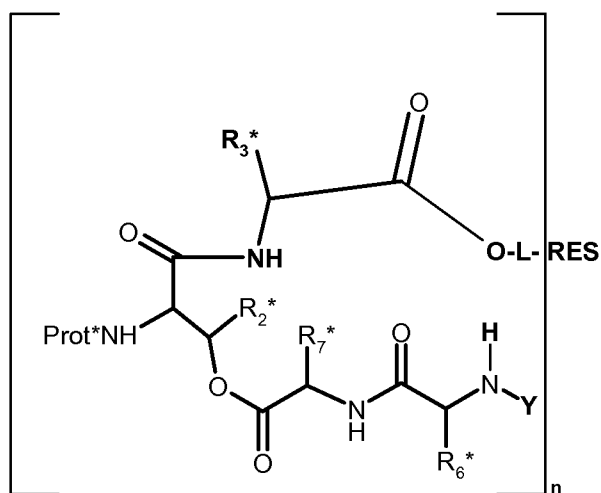
- 99 -



XV

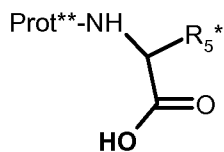
in which R_6^* and Y are as defined for a compound of the formula II in claim 1 and $Prot^{**}$ is as defined for a compound of the formula IX, or a reactive derivative of said amino acid, and removing the protecting group $Prot^{**}$;

coupling the obtained compound of the formula XVI



XVI

wherein Y, R_2^* , R_3^* , R_7^* and R_6^* are as defined for a compound of the formula II in claim 1, $Prot^*$ is as defined for a compound of the formula VIII in claim 5 and n, L and RES are as defined for a compound of the formula X, with an amino acid of the formula XVII,

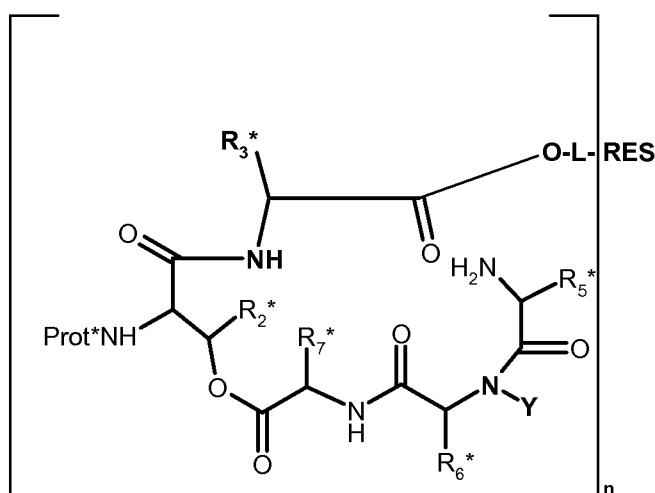


XVII

wherein R_5^* is as defined for a compound of the formula II in claim 1 and $Prot^{**}$ is as defined for a compound of the formula IX, or a reactive derivative of said amino acid, and removing the protecting group $Prot^{**}$,

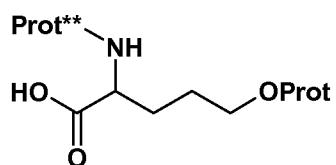
and finally coupling the resulting compound of the formula XVIII,

- 100 -



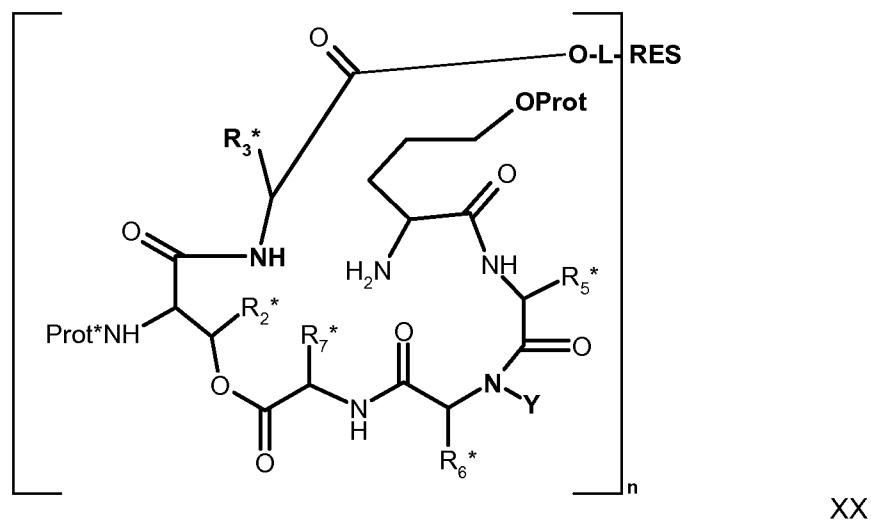
XVIII

wherein Y, R₂^{*}, R₃^{*}, R₇^{*}, R₆^{*} and R₅^{*} are as defined for a compound of the formula II in claim 1, Prot^{*} is as defined for a compound of the formula VIII in claim 5 and n, L and RES are as defined for a compound of the formula X, to a synthon of the formula XIX,



XIX

wherein Prot is as defined for a compound of the formula II in claim 1 and Prot^{**} is as defined for a compound of the formula IX, or an activated derivative of said synthon, and removing the protecting group Prot^{**} to yield a compound of the formula XX,



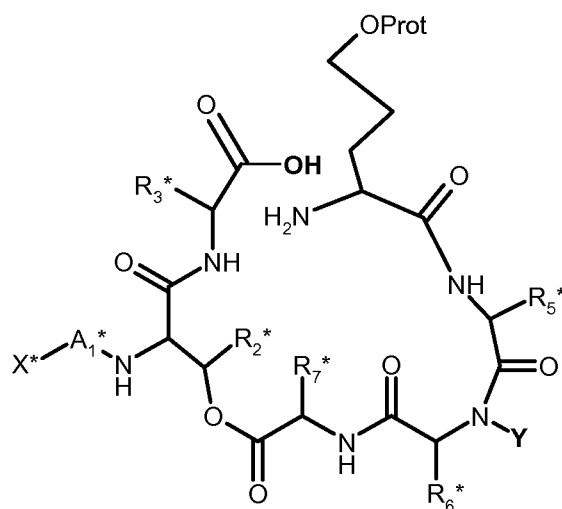
wherein Prot, Y, R_2^* , R_3^* , R_7^* , R_6^* and R_5^* are as defined for a compound of the formula II in claim 1, Prot* is as defined for a compound of the formula VIII in claim 5 and n, L and RES are as defined for a compound of the formula X,

and cleaving the solid phase bound peptide in formula XX off the solid phase L-RES to yield the corresponding compound of the formula VIII, especially VIIIA, as shown in claim 5.

8. The method or process according to claim 1 or claim 2, comprising, for the synthesis of the compound of the formula II of claim 1, cyclizing under lactamization of a linear precursor peptide of the compound of the formula II, carrying an N-terminal amino group and a C-terminal carboxy group, under reaction conditions that allow for the formation of an amide bond from said amino and said carboxy group, preferably using Solution Phase chemistry.

9. The method or process according to claim 8, where the linear precursor peptide is of the formula XXV,

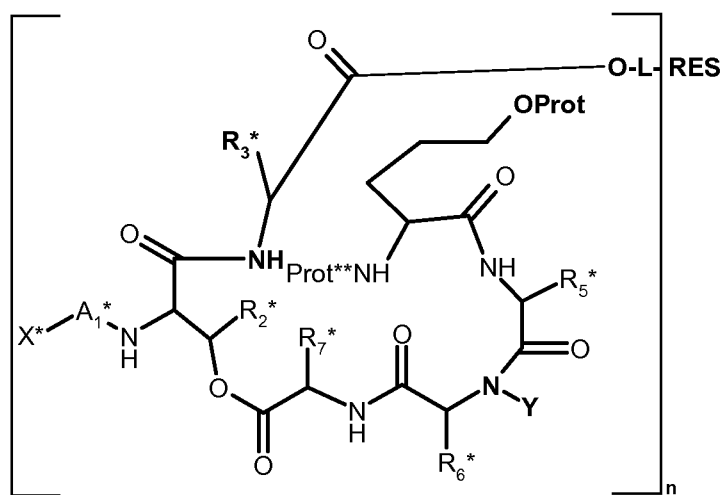
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XXV

wherein X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , R_7^* and Prot are as defined for a compound of the formula II in claim 1.

10. The method or process according to claim 9, further comprising, for the synthesis of the compound of the formula XXV, cleaving a compound of the formula XXIV,



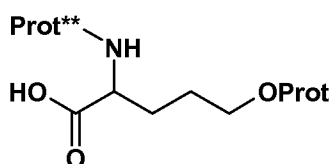
XXIV

wherein X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , R_7^* and Prot are as defined for a compound of the formula II in claim 1, L is a cleavable linker, RES is a solid resin, n is a natural number and Prot** is an amino protecting group that can be removed without parallel removal of the protecting group Prot and with the product remaining on the resin, and (before the

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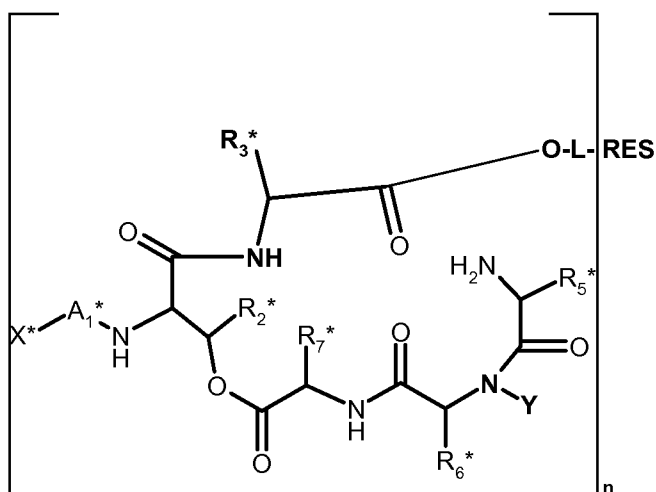
cleavage, in parallel or subsequently to it) removing the protecting group Prot** to yield the compound of the formula XXV.

11. The method or process according to claim 10, further comprising, for the synthesis of the compound of the formula XXIV, coupling an amino acid of the formula XIX,



XIX,

wherein Prot is as defined for a compound of the formula II in claim 1 and Prot** is as defined for a compound of the formula XXIV in claim 10, or an activated derivative of said amino acid, with a compound of the formula XXIII,

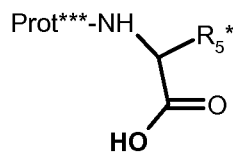


(XXIII)

wherein X*, A₁*, R₂*, R₃*, R₅*, R₆* and R₇* are as defined for a compound of the formula II in claim 1, L is a cleavable linker, RES is a solid resin, and n is a natural number.

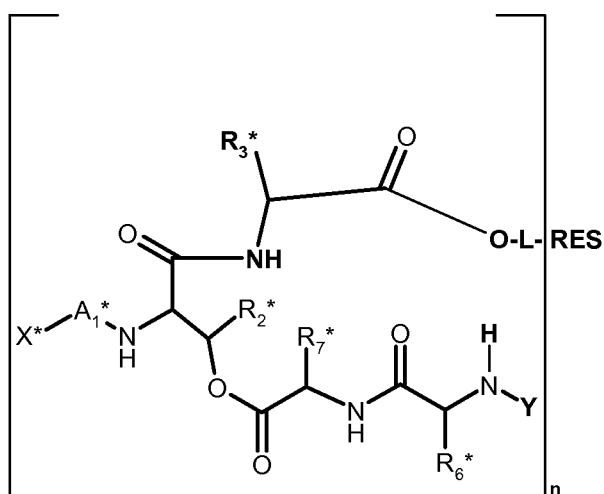
12. The method or process according to claim 11, further comprising, for the synthesis of the compound of the formula XXIII, coupling an amino acid of the formula XVII*

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XVII*,

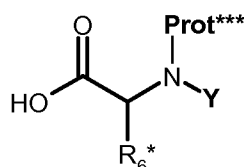
wherein R_5^* is as defined for a compound of the formula II in claim 1 and Prot^{***} is an amino protecting group that can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin, or a reactive derivative of said amino acid, with a compound of the formula XXII,



XXII

wherein X^* , A_1^* , R_2^* , R_3^* , R_6^* and R_7^* are as defined for a compound of the formula II in claim 1, L is a cleavable linker, RES is a solid resin, and n is a natural number, and removing the protecting group Prot^{***} .

13. The method or process according to claim 12, further comprising, for the synthesis of the compound of the formula XXII, coupling an amino acid of the formula XV*,

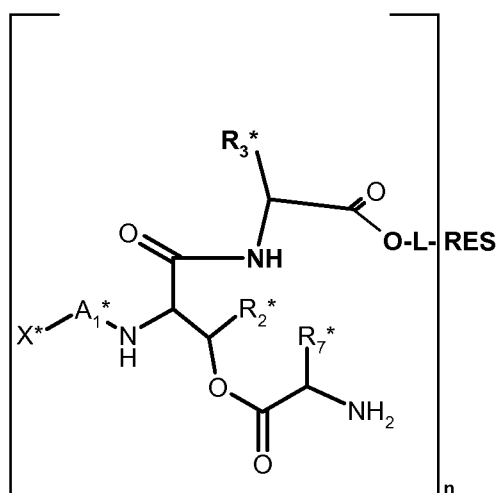


XV*

in which R_6^* and Y are as defined for a compound of the formula II in claim 1 and Prot^{***} is an amino protecting group that can be cleaved off selectively without affecting other

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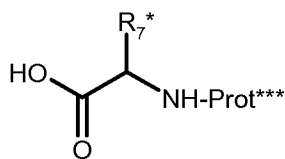
protecting groups present and with the product remaining on the resin, or a reactive derivative of said amino acid, with a compound of the formula XXI,



XXI,

wherein X^* , A_1^* , R_2^* , R_3^* and R_7^* are as defined for a compound of the formula II in claim 1, L is a cleavable linker, RES is a solid resin, and n is a natural number, and removing the protecting group Prot***.

14. The method or process according to claim 13, further comprising, for the synthesis of a compound of the formula XXI, reacting an amino acid of the formula XIII*,

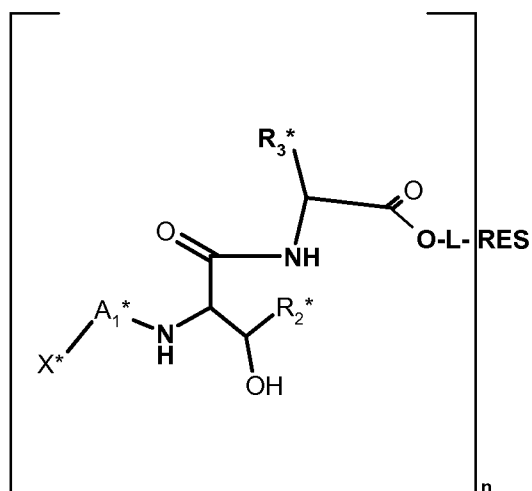


XIII*

wherein Prot*** is an amino protecting group that can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin, and R_7^* is as defined for a compound of the formula II in claim 1, or a reactive derivative of said amino acid,

with the hydroxyl group of a compound of the formula XXVI,

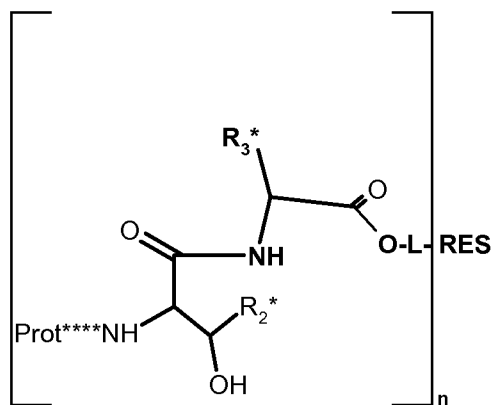
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XXVI

wherein X^* , A_1^* , R_2^* and R_3^* are as defined for a compound of the formula II in claim 1, L is a cleavable linker, RES is a solid resin, and n is a natural number; and removing the protecting group Prot***.

15. The method or process according to claim 14, further comprising, for the synthesis of a compound of the formula XXVI, coupling a resin bound dipeptide symbolized by the formula XII*,

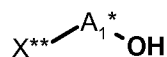


XII*

in which Prot**** is a protecting group that can be cleaved off selectively without affecting other protecting groups present in a compound of the formula II as defined in claim 1 and with the product remaining on the resin, R_2^* and R_3^* are as defined for a compound of the formula II in claim 1, L is a cleavable linker, RES is a solid resin, and n

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is a natural number, after removal of the protecting group Prot**** via the thus obtained free amino group, with an acid of the formula VII

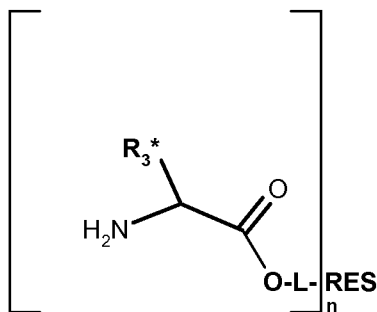


VII

wherein X** is an amino protecting group or is X*, and wherein X* and A₁* are as defined for a compound of the formula II in claim 1, or a reactive derivative of said acid;

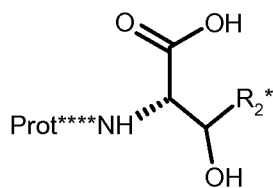
and, if X** is an amino protecting group, removing said amino protecting group X** to yield H instead of X* and coupling the resulting amino group with an acyl group X* using the corresponding acid X*-OH wherein X* is as defined for a compound of the formula II in claim 1, or a reactive derivative of said acid.

16. The method or process according to claim 15, further comprising, for the synthesis of a compound of the formula XII, coupling a resin bound amino acid symbolized by the formula X,



X

wherein R₃* is as defined for a compound of the formula II in claim 1, L is a cleavable linker, RES is a solid resin, and n is a natural number, with an amino acid of the formula XI*,

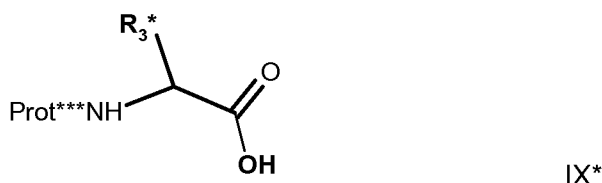


XI*,

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wherein Prot**** is a protecting group that can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin, and R₂* is as defined for a compound of the formula II in claim 1, or a reactive derivative of said amino acid.

17. The method or process according to claim 16, further comprising, for obtaining the resin bound amino acid of the formula X, coupling an amino acid of the formula IX*,



wherein R₃* is as defined for a compound of the formula II in claim 1 and Prot**** is an amino protecting group can be cleaved off selectively without affecting other protecting groups present and with the product remaining on the resin; or a reactive derivative of said amino acid of the formula IX, to a hydroxy group bound via a cleavable linker L which is bound to a solid resin RES, and removing the protecting group Prot****.

18. The method or process according to any one of claims 1 to 16, where the symbols A₁, R₂, R₃, R₅, R₆, R₇, X and Y or the corresponding unprotected or protected moieties R₂*, R₃*, R₅*, R₆*, R₇*, X* and Y₃ are selected so that, in the resulting compound of the formula I, or a salt thereof,

A₁ is the bivalent radical of L-glutamine bound via the carbonyl of its α-carboxyl group to the amino group at the right of A₁ in formula I and via its α-amino group to X, or is 2S-(2-hydroxy-3-phosphonooxy)-propionyl;

R₂ is methyl;

R₃ is isopropyl, isobutyl or benzyl, especially isobutyl;

R₅ is sec-butyl or benzyl, especially sec-butyl;

R₆ is 4-hydroxybenzyl;

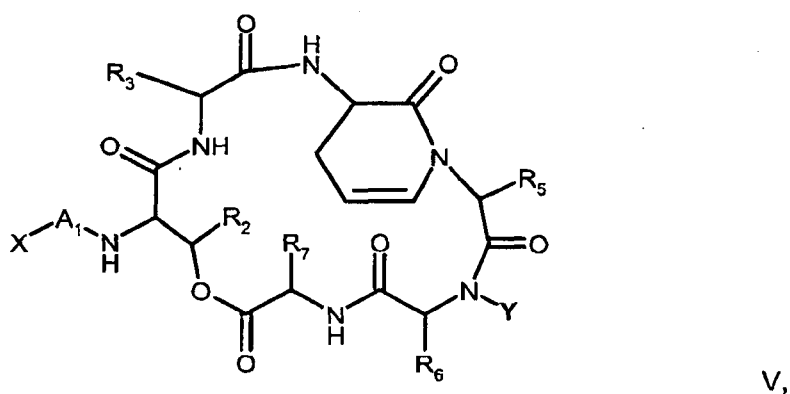
R₇ is isopropyl or sec-butyl, especially sec-butyl;

X is acetyl or isobutyryl, or is absent if A₁ is 2S-(2-hydroxy-3-phosphonooxy)-propionyl and

Y is methyl.

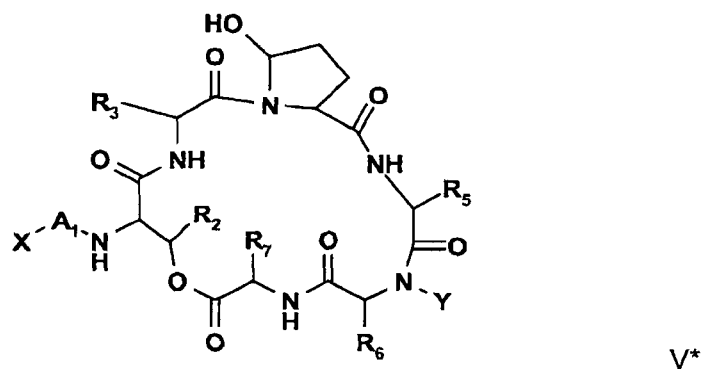
- 109 -

19. A method or process for converting a dehydrate of a compound of the formula I given in claim 1 or with the substituents as defined in claim 18 into the corresponding compound of the formula I, where the dehydrate has the formula V,



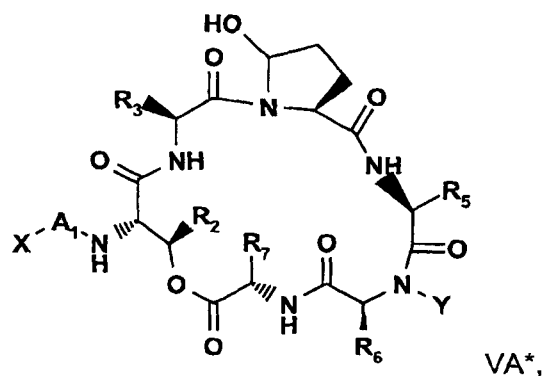
in which Y, X, A₁, R₂, R₃, R₅, R₆ and R₇ are as defined for a compound of the formula I in any one of claims 1 and 18;

and/or its corresponding hemiaminal analogue with a five-ring instead of the ahp structure in formula I which may also be formed as byproduct and has the formula V*,



especially the formula VA*,

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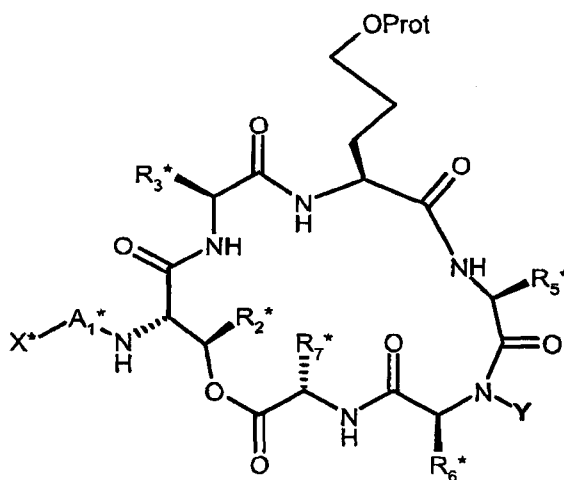
in which Y, X, A₁, R₂, R₃, R₅, R₆ and R₇ are as defined for a compound of the formula I in any one of claims 1 and 18, respectively;

or a method or process for shifting the equilibrium of a mixture of a compound of the formula I and its corresponding dehydrate and/or hemiaminal in favor of the compound of the formula I,

said method or process comprising using an aqueous acid as reactive solvent to drive the reaction.

20. The method according to claim 19, wherein the acid is a carboxylic acid, especially a halo substituted C₁₋₈alkanoic acid, more especially trifluoroacetic acid or trichloroacetic acid.

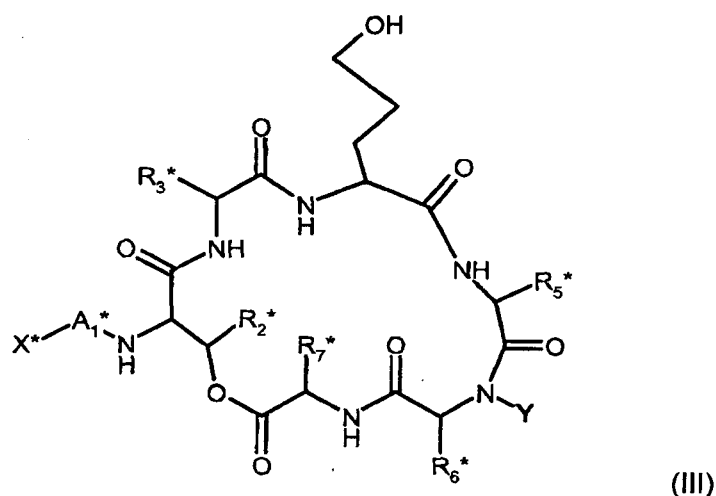
21. A compound of the formula II,



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wherein Prot is a protecting group, Y is as defined for a compound of the formula I in claim 1 or claim 18 and X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* correspond to X, A_1 , R_2 , R_3 , R_5 , R_6 , and R_7 in formula I as defined in claim 1 or in claim 18, respectively, however with the proviso that reactive functional groups on these moieties are present in unprotected or protected form.

22. A compound of the formula III,



wherein Y is as defined for a compound of the formula I in claim 1 or claim 18 and X^* , A_1^* , R_2^* , R_3^* , R_5^* , R_6^* , and R_7^* correspond to X, A_1 , R_2 , R_3 , R_5 , R_6 , and R_7 in formula I as defined in claim 1 or in claim 18, respectively, however with the proviso that reactive functional groups other than the hydroxy represented at the top in formula III on these moieties are present in unprotected or protected form.