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(54) **Polypeptide useful as biomaterial and process for producing the same**
Polypeptid zur Verwendung als Biomaterial und Prozess zur Herstellung davon
Polypeptide utile comme biomaterieau et procédé de préparation

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Remarks:

The file contains technical information submitted after the application was filed and not included in this specification

Description

FIELD OF THE INVENTION

5 **[0001]** The present invention relates to a polypeptide and a process for producing the same. More specifically, the present invention relates to a polypeptide useful as biomaterial which is free from a risk of an infection by a pathogenic organism (or a causative factor) or an undesirable side effect, and which has a high safety, and a process for producing the same. Such a biomaterial or biocompatible material includes, for example, a medical material such as a carrier or support for a tissue engineering, a carrier or support for a regenerative medical treatment, a tissue binding agent or an antiadhesive material, a suture for a surgical operation, a hemostatic material and a contact lens; a raw material for a pharmaceutical preparation; a raw material for a cosmetic preparation; and others.

BACKGROUND OF THE INVENTION

15 **[0002]** A collagen is a fibrous protein found in all multicellular organisms. The collagen is a main component of skins or bones, and occupies 25% of total proteins in mammals. A typical collagen molecular has a rope-like superhelical structure, which is referred to as a triple helical structure, comprising three collagen polypeptide chains. The collagen is particularly rich in proline (Pro) and glycine (Gly). These two amino acid residues are important to form a stable triple helical structure of the collagen.

20 **[0003]** As methods for using a collagen as a biomaterial, there may be mentioned, for example, a method of grafting or transplanting an intact or lyophilized skin tissue derived from a pig on a skin area damaged by a burn or scald, a method of removing cellular components from a tissue with enzyme treatment, and a method of using a collagen which is solubilized by a treatment with an acidic solution or an enzyme to reconstitute a desirable form. A common preparation method and a common qualitative method are described in Methods Enzymol., Vol. 82, pp. 33 to 64, 1982.

25 **[0004]** There are various suggestions to utilize a collagen. For example, Japanese Patent Application Laid-Open No. 08-027192 (JP-08-027192A) discloses a production process of a collagen derivative for imparting moisture and smoothness to skin, which comprises esterifying and modifying an animal tissue containing a collagen with alcohol, and extracting the modified collagen, as well as a cosmetic base material using the collagen derivative.

30 **[0005]** Japanese Patent Application Laid-Open No. 07-097454 (JP-07-097454A) discloses a production process of a water-soluble crosslinked collagen which shows a high regeneration rate of a triple helical structure after thermal denaturation, and the process comprises subjecting a water-soluble collagen to a crosslinking treatment with a bifunctional alkylene diimidate cross-linker having imide ester groups at both ends of the methylene chain.

35 **[0006]** Japanese Patent Application Laid-Open No. 08-053548 (JP-08-053548A) discloses a matrix of a collagen and a synthetic polymer (a collagen-synthetic polymer matrix) which has a low immunogenicity and is useful for preparation of biocompatible implants utilized for various medical applications, and a production process of the matrix comprises reacting a collagen with a first synthetic hydrophilic polymer to form a collagen-synthetic polymer matrix, and further reacting the collagen-synthetic polymer matrix with a reactant such as a second synthetic hydrophilic polymer, a biologically active substance, a glycosaminoglycan and a derivative thereof, a chemical crosslinking agent, an esterifying agent, an amidating agent, anacylating agent, an amino acid, a polypeptide, or others.

40 **[0007]** Japanese Patent Application Laid-Open No. 07-278312 (JP-07-278312A) discloses a unit material containing a hydrophilic synthetic polymer covalently bonded to a chemically modified collagen which is substantially a nonfibrous form at pH 7. The literature discloses that the unit material is particularly useful for ophthalmological devices and optically transparent, and that the unit material has a biocompatibility.

45 **[0008]** Japanese Patent Application Laid-Open No. 05-000158 (JP-05-000158A) discloses a production process of a collagenic membrane-like substance, which comprises crushing a collagen matrix, centrifuging the crushed matrix under a high centrifugal field, homogenizing the resultant precipitate to obtain a paste, casting the paste, and drying the cast paste at a temperature of not higher than 37°C. The literature also discloses that the collagen membrane-like substance has a biocompatibility and a non-inflammatory property, and is useful for repairing a tissue as an artificial implantation matter.

50 **[0009]** Japanese Patent Application Laid-Open No. 05-125100 (JP-05-125100A) discloses a soluble fish scale collagen having high-purity and a production process thereof, and the process comprises pepsinating an intact or deashed fish scale to a pepsin treatment.

55 **[0010]** Japanese Patent Application Laid-Open No. 06-228506 (JP-06-228506A) discloses a production process of a dry particulate or powdery soluble collagen, which comprises injecting a collagen solution through a nozzle into 70 to 90% ethanol medium to form a strand-like or membranous product, drying the product, and chopping or grinding the dried product.

[0011] Japanese Patent Application Laid-Open No. 08-276003 (JP-08-276003A) discloses use of an unbaked single-crystal hydroxyapatite as a material for repairing a biological hard tissue (such as a bone), through attaching the single

crystal to at least part of a low antigenic collagen fiber.

[0012] Japanese Patent Application Laid-Open No. 08-041425 (JP-08-041425A) discloses a method which comprises removing fragments of cells or tissues in a collagen solution and subjecting the residue to an alkali treatment, for removing prion in a collagen derived from an animal or human being, and discloses a collagen obtained by this method.

[0013] Moreover, regarding methods for chemical synthesis of collagen analogues, it has been reported that a soluble polyamide having a molecular weight of 16,000 to 21,000 is obtained by dissolving p-nitrophenyl ester of Pro-Ser-Gly or p-nitrophenyl ester of Pro-Ala-Gly in dimethylformamide, and adding triethylamine thereto, and allowing to stand the mixture for 24 hours (J. Mol. Biol., Vol. 63, pp. 85 to 99, 1972). In this literature, the soluble polyamide is estimated to form a triple helical structure based on the circular dichroism spectra. However there are not referred to properties of the obtained polymer.

[0014] It also has been reported that a method for obtaining a polyamide, which comprises dissolving a 50-mer peptide containing the sequence Val-Pro-Gly-Val-Gly derived from elastin in dimethylsulfoxide, adding 2 equivalents of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide, 1 equivalent of 1-hydroxybenzotriazole and 1.6 equivalents of N-methylmorpholine thereto, allowing to stand the mixture for 14 days, and dialyzing the resultant mixture with a dialysis membrane (molecular weight cutoff: 50,000) (Int. J. Peptide Protein Res., Vol. 46, pp. 453 to 463, 1995).

[0015] Meanwhile, as described in the above-mentioned Japanese Patent Application Laid-Open No. 08-041425 (JP-08-041425A), a causative substance of sheep tremor or bovine spongiform encephalopathy is an infectious protein called as prion, and the infectious protein is considered as one of causes of human Creutzfeldt-Jakob disease infection. Prion is a protein, and it is indicated that prion is hard to deactivate with a conventional pasteurization or sterilization method, further that prion is infectious over species (Nature Review, Vol. 2, pp. 118 to 126, 2001).

[0016] Nat. Prod. Rep. 2002, Vol. 19, 49-59 discloses polypeptides such as (Pro-Pro-Gly)₁₀, (Pro-Hyp-Gly)₁₀ which represent peptides with a lower molecular weight.

[0017] J. Biomed. Mater. Res., 2000, Vol. 51, 612-624 discloses peptoid-containing collagen mimetics with cell binding activity and discloses polypeptides containing Nleu, Gly, Pro and the like. As the polypeptide, (Gly-Pro-Nleu)_n-NH₂ (n=1-7), (Gly-Pro-Nleu)_n-Gly-Pro-NH₂ (n=10), (Gly-Pro-Hyp)_n-NH₂ (n=10) are disclosed. Said reference explicitly discloses (Gly-Pro-Hyp)_n-NH₂ (tri=10).

[0018] Biochemistry, 1994, Vol. 33, 7948-7956 relates to electrostatic interactions in collagen-like triple-helical peptides (title) and discloses a CD spectrum of (Pro-Hyp-Gly)₁₀. Said reference discloses that the polypeptides are synthesized by a standard t-Boc N-protection method with use of a resin. Said reference explicitly discloses (Pro-Hyp-Gly)₁₀.

[0019] J. Org. Chem., 1972. Vol. 27, 4377-4380 discloses a polymer molecular weight determinations and a polymer of Pro-Hyp(H)-Gly having a weight average molecular weight of "45,000^g" with "g" having the meaning that "Other preparations had mol.wt 9800 and 23,000".

[0020] Biochem. Biophys. Res. Comm., 1979, Vol. 86, 654-660 refers to a collagen helix stabilization by hydroxyproline in (Ala-Hyp-Gly)_n and discloses optical properties of (Pro-Hyp-Gly)_n made via the p-nitrophenyl ester as well as via tetraethylpyrophosphate and that 4-Hyp indeed contributes to stability and collagen-like optical properties even when an adjacent Pro residue is absent. Said reference further discloses that (Pro-Hyp-Gly)_n (refractionated, average molecular weight 5000) made via the Onp ester (10) is compared with (Pro-Hyp-Gly)_n (refractionated, average molecular weight 2700) made via TEPP according to Scheme 2 in said reference. Further, said reference discloses CD spectra of (Pro-Hyp-Gly)_n with a molecular weight fraction of 5500 to 4500.

[0021] In general, a collagen derived from bovine or pig is frequently used as a raw material for medical kits (devices) or pharmaceutical preparations, and cosmetic preparations. Accordingly, there have been always existed the risk of an infection (or a transmission) to pathogenic organisms or a causative factor such as prion which cannot be removed by conventional pasteurization or sterilization.

[0022] Moreover, since various cell adhesion sites are found in a naturally occurring collagen, the naturally occurring collagen cannot exert cell selectivity for any applications. For example, in the case using a collagen as a material for inducing a neural axon, migration or growth rate of surrounding fibroblast is more than elongation rate of the axon resulting in forming scarring tissue, and the axon cannot be elongated. It is therefore necessary to take a step to cover around the collagen with a material for protecting migration of fibroblast, or others.

SUMMARY OF THE INVENTION

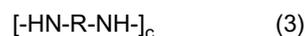
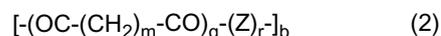
[0023] It is, therefore, an object of the present invention to provide a polypeptide which is free from a risk of an infection (or a transmission) by a pathogenic organism (or a causative factor) or an undesirable side effect, and a process for producing the same.

[0024] It is another object of the present invention to provide a polypeptide useful as a highly safe biomaterial or biocompatible material, and a process for producing the same.

[0025] It is further object of the present invention to provide a process for efficiently producing a polypeptide having above-mentioned properties with inhibiting dimerization or cyclization reaction.

[0026] The inventors of the present invention made intensive studies to achieve the above objects and finally found that a collagen-like polypeptide can be produced without cyclization by condensation of a specific peptide component. The present invention was accomplished based on the above findings.

[0027] That is, the polypeptide of the present invention comprises units represented by the following formulae (1) to (3):



wherein "m" denotes an integer of 1 to 18, "p" and "q" are the same or different, each representing 0 or 1, "Y" represents Pro or Hyp, and "n" denotes an integer of 1 to 20; "Z" represents a peptide chain comprising 1 to 10 amino acid residue (s), "r" denotes an integer of 1 to 20, and "R" represents a linear or branched alkylene group; the molar ratio of "a" relative to "b" [a/b] is 100/0 to 30/70;

(i) when p=1 and q=0, c=a=1,

(ii) when p=0 and q=1, c=b=1,

(iii) when p=1 and q=1, c=a+b=2, and

(iv) when p=0 and q=0, c=0

wherein the polypeptide shows a peak corresponding to the molecular weight in a range from 1×10^4 to 100×10^4 in terms of a globular protein by means of an aqueous gel permeation chromatography.

[0028] Preferred embodiments of the invention are apparent from the dependent claims.

[0029] In the polypeptide comprising the units represented by the above formulae (1) to (3), usually, "m" denotes an integer of 2 to 12; "n" denotes an integer of 2 to 15; "Z" is a peptide chain comprising at least an amino acid residue or a peptide residue selected from the group consisting of Gly, Sar, Ser, Glu, Asp, Lys, His, Ala, Val, Leu, Arg, Pro, Tyr, and Ile. Moreover, in the above formula, usually, "r" denotes an integer of 1 to 10; and "R" represents a C_{2-12} alkylene group.

[0030] For example, the polypeptide of the present invention may comprise the following repeating unit (i), (ii) or (iii):

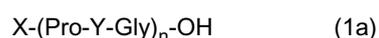
(i) a repeating unit comprising the unit $[-(\text{Pro-Y-Gly})_n-]_a$ (wherein "Y" and "n" have the same meanings as defined above) and the unit $[-(\text{Z})_r-]_b$ (wherein "Z" and "r" have the same meanings as defined above) in a ratio "a/b" of 100/0 to 40/60 (molar ratio); or

(ii) a repeating unit comprising the unit $[-(\text{OC}-(\text{CH}_2)_m-\text{CO})-(\text{Pro-Y-Gly})_n-]_a$ (wherein "m", "n" and "Y" have the same meanings as defined above) and the unit $[-\text{HN-R-NH-}]_c$ (wherein "R" has the same meaning as defined above) in a ratio "a/c" of substantial 1/1 (molar ratio); or

(iii) a repeating unit comprising the unit $[-(\text{OC}-(\text{CH}_2)_m-\text{CO})-(\text{Pro-Y-Gly})_n-]_a$ (wherein "m", "n" and "Y" have the same meanings as defined above), the unit $[-(\text{OC}-(\text{CH}_2)_m-\text{CO})-(\text{Z})_r-]_b$ (wherein "m", "r" and "Z" have the same meanings as defined above) and the unit $[-\text{HN-R-NH-}]_c$ (wherein "R" has the same meaning as defined above) in a ratio "a/b" of 100/0 to 40/60 (molar ratio) and in a ratio "(a+b)/c" of substantial 1/1 (molar ratio).

[0031] The polypeptide of the present invention usually shows positive Cotton effect at a wavelength in range of 220 to 230 nm and negative Cotton effect at a wavelength in range of 195 to 205 nm in a circular dichroism spectrum. At least part (part or all) of the polypeptide, therefore, forms a triple helical structure. Moreover, the polypeptide of the present invention is capable of forming a collagenous tissue (collagen tissue or collagen-like tissue). The polypeptide of the present invention may show a peak corresponding to the molecular weight in a range from 1×10^4 to 100×10^4 .

[0032] According to the present invention, the polypeptide may be produced by reacting a peptide component (A) represented by the following formula (1a) with a peptide component (B) represented by the following formula (2a) and a compound (C) represented by the following formula (3a), as an optional component:



wherein "X" represents H or the group $\text{HOOC}-(\text{CH}_2)_m-\text{CO}$ - ("m" denotes an integer of 1 to 18), "Y" represents Pro or Hyp, and "n" denotes an integer of 1 to 20, $\text{X}-(\text{Z})_r-\text{OH}$ (2a)

wherein "X" represents H or the group $\text{HOOC}-(\text{CH}_2)_m-\text{CO}$ - ("m" denotes an integer of 1 to 18), "Z" represents a peptide chain comprising 1 to 10 amino acid residue(s), and "r" denotes an integer of 1 to 20,



wherein "R" represents a linear or branched alkylene group,
 wherein the ratio of the peptide component (A) relative to the peptide component (B) [the former (A)/the latter (B)] is
 100/0 to 30/70 (mol%); provided that

5 in the case where "X" represents the group $\text{HOOC}-(\text{CH}_2)_m-\text{CO}-$ ("m" has the same meaning as defined above) in the
 formula (1a) and/or (2a), the amount of the compound (C) is substantially 1 mol relative to 1 mol of the total molar amount
 of the peptide component (A) and/or the peptide component (B);

10 in the case where "X" represents H in the formula (1a) and/or (2a), the reaction is conducted without the compound (C)
 wherein the polypeptide shows a peak corresponding to the molecular weight in a range from 1×10^4 to 100×10^4 in
 terms of a globular protein by means of an aqueous gel permeation chromatography.

15 **[0033]** The reaction may be usually carried out by condensing each component (1) to (3) in a solvent (water and/or
 an organic solvent) in the presence of at least a dehydrating and condensing agent (e.g., a carbodiimide-series condensing
 agent, a fluorophosphate-series condensing agent, and a diphenylphosphorylazide). Moreover, the reaction may be
 carried out in the presence of the dehydrating and condensing agent and a condensing auxiliary (or a dehydrating
 auxiliary) [for example, an N-hydroxypolycarboxylic imide, an N-hydroxytriazole (e.g., an N-hydroxybenzotriazole such
 as 1-hydroxybenzotriazole), a triazine, ethyl ester of 2-hydroxyimino-2-cyanoacetic acid]. In the case where the reaction
 is conducted in a water-free solvent system, the proportion of the dehydrating and condensing agent may be 0.7 to 5
 mol relative to 1 mol of the total molar amount of the reaction components (1a), (2a) and (3a). In the case where the
 reaction is conducted in a water-containing solvent system, the proportion of the dehydrating and condensing agent
 20 may be about 2 to 500 mol relative to 1 mol of the total molar amount of the reaction components (1a), (2a) and (3a).
 The amount of the condensing auxiliary may be 0.5 to 5 mol relative to the total molar amount of the above reaction
 components (1a), (2a) and (3a).

[0034] The present invention also includes use of the polypeptide for applying to a tissue of a subject.

25 BRIEF DESCRIPTION OF THE DRAWINGS

[0035]

30 Fig. 1 is a scanning electron micrograph showing a film obtained in Example 3.

DETAILED DESCRIPTION OF THE INVENTION

[0036] In the present invention, amino acid residues are abbreviated to the following condensation codes.

35 Ala: L-alanine residue
 Arg: L-arginine residue
 Asn: L-asparagine residue
 Asp: L-aspartic acid residue
 Cys: L-cysteine residue
 40 Gln: L-glutamine residue
 Glu: L-glutamic acid residue
 Gly: glycine residue
 His: L-histidine residue
 Hyp: L-hydroxyproline residue
 45 Ile: L-isoleucine residue
 Leu: L-leucine residue
 Lys: L-lysine residue
 Met: L-methionine residue
 Phe: L-phenylalanine residue
 50 Pro: L-proline residue
 Sar: sarcosine residue
 Ser: L-serine residue
 Thr: L-threonine residue
 Trp: L-tryptophan residue
 55 Tyr: L-tyrosine residue
 Val: L-valine residue

[0037] Moreover, in this specification, amino acid sequences of peptide chains are represented in accordance with

conventional expression that, in an amino acid residue, N-terminus and C-terminus are drawn at the left and the right, respectively.

[0038] It is essential that a unit (1) $[-(\text{OC}-(\text{CH}_2)_m-\text{CO})_p-(\text{Pro-Y-Gly})_n-]$ constituting the polypeptide of the present invention contains a repeating sequence Pro-Y-Gly. When the repeating number of the sequence Pro-Y-Gly is smaller, the polypeptide is deteriorated in stability of the triple helical structure. When the repeating number is too large, it is difficult to synthesize the peptide. The repeating number "n" is, therefore, 1 to 20, preferably 2 to 15 (e.g., 3 to 15), and more preferably 5 to 15.

[0039] In the above formula (1), "Y" may be either Pro or Hyp. From the viewpoint of stability of the triple helical structure, the preferred "Y" is Hyp. Incidentally, Hyp is usually 4Hyp (e.g., trans-4-hydroxy-L-proline) residue.

[0040] Further, the number "m" representing the repeating number of a methylene chain (CH_2) needs only to be selected from a range that the polypeptide is adversely affected physical and biological properties. The number "m" is usually 1 to 18, preferably 2 to 12, and more preferably 2 to 10 (particularly 2 to 6). The number "p" is 0 or 1.

[0041] In the above-mentioned unit (2) $[-(\text{OC}-(\text{CH}_2)_m-\text{CO})_q-(\text{Z})_r-]$, "Z" represents a peptide chain which has an arbitrary sequence comprising 1 to 10 amino acid residue(s). "Z" may be any sequence as far as the resulting polypeptide is not deteriorated in physical and biological properties. In order that the polypeptide shows useful physical and biological properties, for example, the peptide chain "Z" usually comprises at least one amino acid or peptide residue selected from the group consisting of Gly, Sar, Ser, Glu, Asp, Lys, His, Ala, Val, Leu, Arg, Pro, Tyr and Ile. In particular, the peptide chain "Z" practically comprises at least one amino acid or peptide residue selected from the group consisting of Gly, Ser, Glu, Asp, Lys, Arg and Pro. The peptide chain "Z" preferably comprises the amino acid residue or the sequence such as Gly, Sar, Ser, Glu, Asp, Lys, Arg-Gly-Asp, Tyr-Ile-Gly-Ser-Arg, Ile-Lys-Val-Ala-Val, Val-Pro-Gly-Val-Gly, Asp-Gly-Glu-Ala, Gly-Ile-Ala-Gly, His-Ala-Val, Glu-Arg-Leu-Glu, Lys-Asp-Pro-Lys-Arg-Leu, or Arg-Ser-Arg-Lys.

[0042] The number "r" representing the repeating number of the peptide chain "Z" needs only to be selected from a range that the resulting polypeptide demonstrates physical and biological properties. When the repeating number "r" is too large, it is difficult to synthesize the peptide, and the physical properties of the resulting polypeptide is apt to vary. The repeating number "r" is, therefore, usually 1 to 20, preferably 1 to 10, and more preferably 1 to 5.

[0043] The number "m" representing the repeating number of the methylene chain (CH_2) is 1 to 18, preferably 2 to 12, and more preferably 2 to 10 (particularly about 2 to 6) as the same manner as in the above formula (1). The number "q" is 0 or 1.

[0044] In the formulae (1) and (2), in the case where at least one of "p" and "q" is 0, the polypeptide contains the unit $[-\text{HN-R-NH}-]$ represented by the formula (3). In the unit represented by the formula (3), a linear or branched alkylene group represented by "R" needs only be any group as far as the polypeptide is not deteriorated in physical and biological properties. For example, as the group "R", there may be mentioned a C_{1-18} alkylene group such as methylene, ethylene, propylene, trimethylene, and tetramethylene groups. The alkylene group "R" may be a linear methylene chain $(\text{CH}_2)_s$ ("s" represents an integer of 1 to 18). The group "R" is preferably a C_{2-12} alkylene group, more preferably a C_{2-10} alkylene group, in particular a C_{2-6} alkylene group.

[0045] The ratio ["a/b"] of the unit represented by the formula (1) relative to the unit represented by the formula (2) is 100/0 to 30/70 (molar ratio), preferably 100/0 to 40/60 (molar ratio), and more preferably 100/0 to 50/50 (molar ratio).

[0046] Further, the ratio of the unit represented by the formula (3) may be selected depending on the value "p" of the formula (1) and the value "q" of the formula (2). When "p"=1 and "q"= 0, "c"=a (=1). When "p"=0 and "q"=1, "c"=b (=1). Moreover, when "p"=1 and "q"=1, "c"=a+b (=2). When "p"=0 and "q"=0, "c"=0.

[0047] That is, the polypeptide of the present invention includes the following polypeptides:

- (a) a polypeptide comprising a repeating unit of the unit $[-(\text{Pro-Y-Gly})_n-]_a$ ("P" is 0 in the formula (1));
- (b) a polypeptide which comprises a repeating unit containing the unit $[-(\text{Pro-Y-Gly})_n-]_a$ ("p" is 0 in the formula (1)) and the peptide unit $[-(\text{Z})_r-]_b$ ("q" is 0 in the formula (2)) in a ratio of "a/b" (mol%);
- (c) a polypeptide which comprises a repeating unit containing the unit $[-(\text{OC}-(\text{CH}_2)_m-\text{CO})-(\text{Pro-Y-Gly})_n-]_a$ ("p" is 1 in the formula (1)) and the unit $[-\text{HN-R-NH}-]_c$ represented by the formula (3) in a ratio of "a/c" (mol%); and
- (d) a polypeptide which comprises a repeating unit containing the unit $[-(\text{OC}-(\text{CH}_2)_m-\text{CO})-(\text{Pro-Y-Gly})_n-]_a$ ("p" is 1 in the formula (1)), the unit $[-(\text{OC}-(\text{CH}_2)_m-\text{CO})-(\text{Z})_r-]_b$ ("q" is 1 in the formula (2)) and the unit $[-\text{HN-R-NH}-]_c$ represented by the formula (3) in a ratio of "a/b/c" (mol%).

[0048] Such a polypeptide forms a linear polypeptide without forming a ring such as a six-membered ring by cyclization, and is soluble in a solvent (a hydrophilic solvent, e.g., water, an alcohol such as ethanol and propanol, a ketone such as acetone, a cyclic ether such as dioxane and tetrahydrofuran, a sulfoxide such as dimethyl sulfoxide, or a mixed solvent thereof). The polypeptide shows a peak corresponding to the molecular weight in a range from 1×10^4 to 100×10^4 in terms of a globular protein by means of an aqueous GPC (gel permeation chromatography).

[0049] Further, the polypeptide of the present invention is capable of forming (formable) a collagenous tissue (collagen tissue or collagen-like tissue). That is, the polypeptide of the present invention shows positive Cotton effect at a wavelength

in range of 220 to 230 nm and negative Cotton effect at a wavelength in range of 195 to 205 nm in circular dichroism spectra. At least one part (that is, part or all) of the polypeptide of the present invention is, accordingly, capable of forming a triple helical structure, and the polypeptide of the present invention forms a collagenous (collagen-like) polypeptide. Incidentally, Cotton effect means a phenomenon caused by difference between an absorption coefficient relative to a right circularly polarized light and that relative to a left at a specific wavelength in an optical rotatory substance.

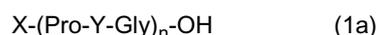
[0050] The polypeptide of the present invention can be obtained by subjecting the above-mentioned components (1a) to (3a) to a condensation reaction. The peptide chain of the present invention can be synthesized in accordance with a conventional peptide synthesis method. Peptides may, for example, be prepared based on a solid-phase synthesis method or a liquid-phase synthesis method, and the solid-phase synthesis method is operationally convenient [for example, see "Zoku Seikagaku Jikken Kouza 2, Tanpakushitsu no Kagaku (Supplemental Handbook of Biochemical Experiments, Chemistry of Protein) (the second volume)" edited by The Japanese Biochemical Society (issued by Tokyo Kagaku Dozin Co., Ltd., May 20, 1987), pp. 641 to 694]. For the peptide synthesis, a conventional manner may be utilized, and the manner includes, for example, a coupling method using a condensing agent, an active esterification method (e.g., a phenyl ester such as p-nitrophenyl ester (ONp) and pentafluorophenyl ester (Opfp), an N-hydroxydicarboxylic imide ester such as N-hydroxysuccinimide ester (ONSu), and 1-hydroxybenzotriazole ester (Obt)), a mixed acid anhydride method, an azide method, and others. In the preferred manner, at least a condensing agent (preferably a condensing agent as mentioned below, in particular a combination of a condensing agent as mentioned below with a condensing auxiliary as mentioned below) may be practically used.

[0051] Furthermore, in the peptide synthesis, protection of an amino group, a carboxyl group, and other functional group (e.g., a guanidino group, an imidazolyl group, a mercapto group, a hydroxyl group, an ω -carboxyl group) with a protective group, and elimination or removal of the protective group with a catalytic reduction or a strong acid treatment (e.g., anhydrous hydrogen fluoride, trifluoromethanesulfonic acid, trifluoroacetic acid) are repeatedly conducted depending on a species of amino acids or peptide fragments. For example, as a protective group for an amino group, there may be utilized benzyloxycarbonyl group (Z), p-methoxybenzyloxycarbonyl group (Z(OMe)), 9-fluorenylmethoxycarbonyl group (Fmoc), t-butoxycarbonyl group (Boc), 3-nitro-2-pyridinesulfonyl group (Npys), and the other groups. As a protective group for a carboxyl group, there may be utilized benzyloxy group (OBzl), phenacyloxy group (OPac), t-butoxy group (OBu), methoxy group (OMe), ethoxy group (OEt), and the other groups. Incidentally, an automatic synthesis apparatus may be utilized for the peptide synthesis.

[0052] More specifically, the preparation of the peptide chain of the present invention with the solid-phase synthesis method may be carried out in accordance with a conventional manner. As a solid-phase resin (or a carrier), there may be utilized a polymer insoluble to a reaction solvent, for example, a styrene-divinylbenzene copolymer (such as a chloromethylated resin, a hydroxymethyl resin, a hydroxymethylphenylacetamidemethyl resin, a 4-methylbenzhydrylamine resin).

[0053] In the solid-phase synthesis method, a peptide can be usually produced by the following steps: a step forming a peptide chain corresponding to an objective peptide, and a step comprising (iv) detaching the peptide chain from the polymer (resin) and eliminating the protective group(s) from the protected functional group(s) to obtain the objective peptide, and purifying the resulting peptide. The peptide chain-forming step comprises (i) bonding an amino acid or peptide fragment to the above polymer (resin) from C-terminal to N-terminal of the objective peptide, in which the amino acid or peptide fragment has a free α -COOH group and a functional group(s) (e.g., at least an α -amino group of the N-terminal) protected with a protective group(s), (ii) eliminating the protective group of an α -amino group forming a peptide bond among the bonded amino acid or peptide fragment, and (iii) sequentially repeating the above bonding operation and the eliminating operation to elongate the peptide chain for the formation of the object peptide. In the operation (i) for bonding the amino acid or peptide fragment, an amino acid which is corresponding to the C-terminal of the peptide chain and has a free α -COOH group, and in which at least the N-terminal is protected with a protective group (for example, a Fmoc-amino acid, a Boc-amino acid) is used. Incidentally, from the viewpoint of inhibiting a side reaction, detachment of the peptide chain from the polymer is preferably carried out concurrently with elimination of the protective group with the use of trifluoroacetic acid. Moreover, the resulting peptide may be purified by utilizing a separation and purification means (e.g., a reversed phase liquid chromatography, and a gel-permeation chromatography).

[0054] In the present invention, a polypeptide is prepared by condensing at least a peptide component (A) represented by the following formula (1a):



wherein "X" represents H or $\text{HOOC-(CH}_2\text{)}_m\text{-CO-}$ ("m" has the same meaning as defined above), "Y" and "n" have the same meanings as defined above.

[0055] The polypeptide of the present invention may be prepared by co-condensing the peptide component (A) represented by the above formula (1a) with a peptide component (B) represented by the following formula (2a):



wherein X represents H or $\text{HOOC-(CH}_2\text{)}_m\text{-CO-}$ ("m" has the same meaning as defined above), "Z" and "r" have the same meanings as defined above.

5 **[0056]** Incidentally, as the compound in which the above group "X" is $\text{HOOC-(CH}_2\text{)}_m\text{-CO-}$, there may be mentioned, for example, a C_{3-20} aliphatic dicarboxylic acid such as malonic acid, succinic acid, glutaric acid, adipic acid, pimelic acid, suberic acid, azelaic acid and sebacic acid, or an acid anhydride thereof. These compounds may be used singly or in combination. These compounds may be also subjected to a reaction according to a conventional amide bond-forming method (for example, a reaction using a tertiary amine described later as a catalyst) or the above-mentioned peptide synthesis method to obtain the compounds represented by the above formulae (1a) and (2a).

10 **[0057]** The ratio of the peptide component (A) relative to the peptide component (B) [the former (A)/the latter (B)] is, for example, 100/0 to 30/70 (mol%), preferably 100/0 to 40/60 (mol%), and more preferably 100/0 to 50/50 (mol%).

15 **[0058]** Further, in the above formulae (1a) and/or (2a), if the group "X" is $\text{HOOC-(CH}_2\text{)}_m\text{-CO-}$ ("m" has the same meaning as defined above), at least one of the peptide component (A) and the peptide component (B) is subjected to a co-condensation reaction with the compound (C) represented by the following formula (3a) for forming an amide group. If the group "X" is H, it is unnecessary to use the compound (C).



20 "R" has the same meaning defined above.

[0059] As the compound represented by the above formula (3a), there may be exemplified a diamine corresponding to the above formula (3), e.g., a C_{1-18} alkylenediamine such as ethylenediamine, trimethylenediamine, propylenediamine, tetramethylenediamine and hexamethylenediamine, a polyalkylenepolyamine such as diethylenetriamine and hexamethylenetetramine, and others. These compounds may be used singly or in combination.

25 **[0060]** The amount of the diamine compound (C), for example, may be substantially 1 mol (for example, 0.95 o 1.05 mol) relative to 1 mol of a total molar amount of the peptide component (A) and/or the peptide component (B). Incidentally, in the case where the group "X" is $\text{HOOC-(CH}_2\text{)}_m\text{-CO-}$ ("m" has the same meaning as defined above) in either the peptide component (A) or (B), the amount of the diamine compound (C) may be substantially 1 mol (for example, 0.95 o 1.05 mol) relative to 1 mol of the peptide component having such group.

30 **[0061]** The reaction of these components (1a), (2a) and (3a) is usually carried out in a solvent. The solvent may be capable of dissolving or suspending (partly or wholly dissolving) the peptide components and the compound, and there may be usually employed water and/or an organic solvent. As the solvent, there may be mentioned, for example, water, an amide (e.g., dimethylformamide, dimethylacetamide, hexamethylphosphoramide), a sulfoxide (e.g., dimethylsulfoxide), a nitrogen-containing cyclic compound (e.g., N-methylpyrrolidone, pyridine), a nitrile (e.g., acetonitrile), an ether (e.g., dioxane, tetrahydrofuran), an alcohol (e.g., methyl alcohol, ethyl alcohol, propyl alcohol), and a mixed solvent thereof. Among these solvents, water, dimethylformamide, or dimethylsulfoxide is practically used.

35 **[0062]** The reaction of these components (1a), (2a) and (3a) may be usually carried out in the presence of at least a dehydrating agent (a dehydrating and condensing agent). The reaction with these components in the presence of a dehydrating and condensing agent and a condensing auxiliary (synergist) produces smoothly a polypeptide with inhibiting dimerization or cyclization.

40 **[0063]** The dehydrating and condensing agent is not particularly limited as far as the agent can conduct dehydration and condensation efficiently in the above-mentioned solvent. For example, the dehydrating and condensing agent (the dehydrating agent) includes a carbodiimide-series condensing agent [e.g., diisopropylcarbodiimide (DIPC), 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDC=WSCI), 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride (WSCl·HCl), dicyclohexylcarbodiimide (DCC)], a fluorophosphate-series condensing agent [e.g., O-(7-azabenzotriazole-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate, O-benzotriazole-1-yl-N,N,N',N'-tetramethyluronium hexafluorophosphate, benzotriazole-1-yl-oxy-tris-pyrrolidinophosphonium hexafluorophosphate, a salt of benzotriazole-1-yl-tris(dimethylamino)phosphonium hexafluorophosphate (BOP)], diphenylphosphorylazide (DPPA), and others. The dehydrating and condensing agent(s) may be used singly, or used as a mixture in combination thereof. The dehydrating and condensing agent is preferably a carbodiimide-series condensing agent [e.g., 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide, 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride].

45 **[0064]** The condensing auxiliary is not particularly limited as the condensing auxiliary can facilitate the reaction of the condensing agent. For example, there may be mentioned an N-hydroxypolycarboxylic imide [e.g., an N-hydroxydicarboxylic imide such as N-hydroxysuccinic imide (HONSu) and N-hydroxy-5-norbornene-2,3-dicarboxylic imide (HONB)]; an N-hydroxytriazole [e.g., an N-hydroxybenzotriazole such as 1-hydroxybenzotriazole (HOBt)]; a triazine such as 3-hydroxy-4-oxo-3,4-dihydro-1,2,3-benzotriazine (HOObt); ethyl ester of 2-hydroxyimino-2-cyanoacetic acid; and others. The condensing auxiliary may be also used singly or in combination. The condensing auxiliary is preferably an N-

hydroxydicarboxylic imide [e.g., HONSu], an N-hydroxybenzotriazole or an N-hydroxybenzotriazine [e.g., HOBt].

[0065] The dehydrating and condensing agent may be suitably used in combination with the condensing auxiliary. As a combination of the dehydrating and condensing agent with the condensing auxiliary, there may be mentioned, for example, DCC-HONSu (HOBt or HOOBt), WSCI-HONSu (HOBt or HOOBt), and other combinations.

[0066] The amount of the dehydrating and condensing agent is, in a water-free solvent system, usually 0.7 to 5 mol, preferably 0.8 to 2.5 mol, and more preferably 0.9 to 2.3 mol (e.g., 1 to 2 mol) relative to 1 mol of the total molar amount of the reaction components (1a), (2a) and (3a). In a water-containing solvent system, since the dehydrating and condensing agent may be deactivated by water, the amount of the dehydrating and condensing agent is usually 2 to 500 mol (e.g., 2 to 50 mol), preferably 5 to 250 mol (e.g., 5 to 25 mol), and more preferably 10 to 125 mol (e.g., 10 to 20 mol) relative to 1 mol of a total molar amount of the reaction components (1a), (1b) and (1c). The amount of the condensing auxiliary is, for example, 0.5 to 5 mol, preferably 0.7 to 2 mol, and more preferably 0.8 to 1.5 mol relative to 1 mol of a total molar amount of the reaction components (1a), (1b) and (1c) irrespective of a kind or species of solvent.

[0067] In the condensation reaction of the present invention, the pH of the reaction system may be adjusted, or a base being inert for the reaction may be added to the system. The pH may be usually adjusted with an inorganic base [e.g., sodium hydroxide, potassium hydroxide, sodium carbonate, sodium hydrogen carbonate], an organic base, an inorganic acid [e.g., hydrochloric acid], or an organic acid. The pH of the reaction mixture is usually adjusted to approximately neutral pH (pH = 6 to 8). As the base being inert for the reaction, there may be exemplified a tertiary amine (e.g., a trialkylamine such as trimethylamine, triethylamine and diisopropylethylamine, a heterocyclic tertiary amine such as N-methylmorpholine and pyridine), and others. The amount of the base may be usually selected within a range from one to two times as much as the total molar amount of amino groups in the peptide components and the compound.

[0068] In the present invention, both the formation of the peptide chain represented by the above formula X-(Pro-Y-Gly)_n-OH, and the formation of a triple helical structure in the resulting polypeptide can be usually proved by measuring circular dichroism spectra or gel-permeation chromatography for a solution of the peptide chain or the polypeptide. In particular, regarding circular dichroism spectra, it has been reported that a naturally-occurring collagen and peptide chain forming a triple helical structure distinctively shows positive Cotton effect at a wavelength in range of 220 to 230 nm and negative Cotton effect at a wavelength in range of 195 to 205 nm (J. Mol. Biol., Vol. 63 pp. 85 to 99, 1972). Moreover, in accordance with gel-permeation chromatography, since a peak corresponding to a molecular weight of a trimer forming a triple helical structure is observed away from a peak corresponding to that of a monomer, the proportion of the peptide forming the triple helical structure can be determined.

[0069] The polypeptide of the present invention is capable of forming a collagenous tissue, free from a risk of an infection of a pathogenic organism or a transmission of a causative factor (e.g., a protein converted into a pathological protein, such as abnormal prion) or an undesirable side effect, and has high safety. Moreover, the polypeptide is excellent in cytophilic property and biocompatibility. The polypeptide is, therefore, useful as a biomaterial or a biocompatible material, for example, an artificial collagen. The polypeptide of the present invention may be applied to a tissue (e.g., an epidermal tissue and a dermal tissue) of a subject. The subject includes human beings, and nonhumans (such as monkeys, sheep, bovines, horses, dogs, cats, rabbits, rats, and mice). Moreover, the polypeptide of the present invention may be used for inhibiting or preventing an infection (or a transmission) originated from (caused by) a polypeptide (e.g., an infection or a transmission of a pathogenic organism or a causative factor present in a polypeptide). Accordingly, the polypeptide of the present invention may be effectively utilized in a damaged area [for example, a diseased area or an injured area (e.g., an area damaged by a scratch and a burn or scald)] or a dissected (or an incised) area [for example, a dissected area such as a surgical cut]. For example, the polypeptide is utilized as a medical material such as a carrier for a tissue engineering, a carrier for a regenerative medical treatment (e.g., an artificial skin), a tissue binding agent or an antiadhesive material, a suture for a surgical operation, a hemostatic material and a contact lens; a raw material (or base material) for a pharmaceutical preparation; a raw material (or base material) for a cosmetic preparation; a food additive; and others.

[0070] The polypeptide of the present invention can be shaped or molded by a conventional manner depending on various applications. The form to be used of the polypeptide may be a liquid form (e.g., a solution, a suspension), a particulate form, a two-dimensional form (e.g., a film, a sheet), or a three-dimensional form. For example, a sheet or film of the polypeptide may be obtained by casting a solution or suspension of the polypeptide on a releasable substrate (support) (e.g., a sheet made from a fluorine-containing resin (polytetrafluoroethylene)) and drying the cast substrate. Moreover, a fibrous substance is obtained by extruding a solution or suspension of the polypeptide through a nozzle in a solution containing a salt of high concentration or in a solvent incapable of dissolving the polypeptide. Further, a gelatinous substance may be obtained by allowing to stand an aqueous solution or suspension of the polypeptide, or if necessary, with adding a polyvalent crosslinkable reagent (e.g., glutaraldehyde) thereto. Further, a sponge-like porous substance may be obtained by lyophilizing the resultant gelatinous substance. Furthermore, a porous substance can be also obtained by stirring the aqueous solution or suspension of the polypeptide to foam, and drying.

[0071] Further, the polypeptide of the present invention may be used as a coating agent. For example, a surface of a substrate may be coated with the polypeptide of the present invention by coating or spraying with the solution or

suspension of the polypeptide and drying the coated or sprayed layer. The substrate may be a shaped article made of various materials such as a metal, a ceramic, a plastic, a natural polymer, or the like. The form of the shaped article may be a two-dimensional structure or a three-dimensional structure, e.g., a particulate form, a linear or fibrous form, a film or sheet form. Further, the polypeptide may be held or supported by impregnating, into the solution or suspension of the polypeptide, a porous substance (e.g., a particulate porous substance, a two-dimensional porous substance such as a paper made from a cellulose fiber (cellulosic paper), a nonwoven or woven fabric, a three-dimensional porous substance such as a cylindrical substance).

[0072] In the case using the polypeptide of the present invention for a medical purpose, the polypeptide is preferably sterilized or pasteurized. A sterilization or pasteurization method includes various methods, for example, pasteurization with steam such as a heated and damp steam, pasteurization with a gamma ray, pasteurization with ethylene oxide gas, sterilization with a pharmaceutical preparation, sterilization with an ultraviolet ray, and others. Among these methods, pasteurization with a gamma ray and pasteurization with ethylene oxide gas are preferred from the viewpoint of pasteurization efficiency and low (or light) adverse effects on a material to be used.

[0073] The polypeptide of the present invention has no fear of a risk of an infection by a pathogenic organism (or a causative factor) or an undesirable side effect, with a high safety, and a high cytophilic property. Further, the polypeptide can be produced by a simple operation such as a condensation reaction without inhibition of dimerization or cyclization.

EXAMPLES

[0074] The following examples are intended to describe this invention in further detail and should by no means be interpreted as defining the scope of the invention.

Example 1

[0075] A peptide represented by the formula H-(Pro-Pro-Gly)₁₀-OH (Sequence ID No.1) (manufactured by Peptide Institute, Inc.) was dissolved in water containing 50% methyl alcohol at a concentration of 0.25 mg/mL, and a circular dichroism spectrum thereof was measured with a spectrometer (J-725, manufactured by JASCO Corp., light path: 1 mm). The result showed that positive Cotton effect was observed at a wavelength of 227 nm and negative Cotton effect at a wavelength of 197 nm were observed, and confirmed that the peptide formed a triple helical structure. Moreover, with respect to the same solution, the molecular weight was measured by gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex Peptide PE 7.5/300, flow rate: 0.25 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl). The result confirmed two peaks: one peak showing a molecular weight of about 9000 corresponding to a triple helical structure, and another peak showing a molecular weight of about 3000 corresponding to a monomer. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit (manufactured by Amersham Bioscience K.K.), human insulin (manufactured by Sigma Corporation) and glycine (manufactured by Wako Pure Chemical Industries, Ltd.) as reference materials. The proportion of the peptide forming the triple helical structure was about 50% on the basis of two peak areas. Incidentally, in the case using a solution dissolving the peptide H-(Pro-Pro-Gly)₁₀-OH in water at a concentration of 0.25 mg/mL, weak positive Cotton effect at a wavelength of 227 nm and negative Cotton effect at a wavelength of 198 nm were observed. However, a peak corresponding to a trimer was inappreciably found in gel-permeation chromatography, and the proportion of the peptide forming the triple helical structure was almost 0%.

[0076] Five (5) mg (0.002 mmol) of the peptide H-(Pro-Pro-Gly)₁₀-OH was suspended in 2 mL dimethyl sulfoxide, and the mixture was stirred at a room temperature. To the mixture were added 0.31 mg (0.0024 mmol) of diisopropylethylamine, 0.32 mg (0.0024 mmol) of 1-hydroxybenzotriazole, and 0.46mg (0.0024 mmol) of 1-ethyl-3-(3-dimethylamino-propyl)-carbodiimide hydrochloride, and the resultant mixture was further stirred for 7 days at a room temperature.

[0077] The reaction mixture (solution) was diluted 20-fold with water, and the diluted solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex 200 HR 10/30, flow rate: 0.5 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl). The result confirmed a polypeptide peak in a molecular weight range of 40000 to 200000. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit and a Gel Filtration HMW Calibration Kit (manufactured by Amersham Bioscience K.K.) as reference materials.

[0078] The resulting reaction mixture was diluted 5-fold with water, and the diluted solution was dialyzed against water for 3 days to remove the reagent(s) such as the condensing agent and the unreacted monomer(s). The circular dichroism spectrum was measured in the obtained polypeptide, and the result showed that positive Cotton effect was observed at a wavelength of 227 nm and negative Cotton effect at a wavelength of 199 nm, and confirmed that the polypeptide formed a triple helical structure.

Example 2

[0079] A peptide chain represented by the formula H-(Pro-Pro-Gly)₅-OH (Sequence ID No. 2) was synthesized by a solid-phase synthesis with an automatic peptide synthesis machine. That is, with the use of 0.1 mmol of a particulate resin [HMP glycine, manufactured by Applied Biosystems (US)] which comprises a styrene-divinylbenzene copolymer containing 4-(N α -9-(fluorenylmethoxycarbonyl)-glycine)-oxymethyl-phenoxy-methyl group in a proportion of 0.65 mmol/g (resin) [component molar ratio of styrene relative to divinylbenzene: 99/1], the carboxyl terminal of one amino acid was sequentially linked (or bound) to the amino terminal of the other amino acid so as to obtain the object peptide. In this link reaction, 1 mmol of Na-9-(fluorenylmethoxycarbonyl)-L-proline [Fmoc-proline] and 1 mmol of N α -9-(fluorenylmethoxycarbonyl)-glycine [Fmoc-glycine] (manufactured by Applied Biosystems (US)) were used as amino acids in each linking step.

[0080] The peptide resin obtained in the foregoing manner was suspended in 10 mL of dimethylformamide, and 50 mg (0.5 mmol) of succinic anhydride and 13 mg (0.1 mmol) of diisopropylethylamine were added thereto. The mixture was subjected to a reaction at a room temperature for 12 hours. Thereafter, the reaction mixture was washed with methyl alcohol and dichloromethane alternately, and dried under a reduced pressure.

[0081] The peptide resin obtained by the above-mentioned manner was subjected to a treatment with 10 mL of trifluoroacetic acid containing 5% water for 3 hours. The resulting solution was added to diethyl ether to form a precipitate, and the precipitate was further washed with diethyl ether several times to deprotect the peptide and to eliminate the peptide from the resin. The resulting crude product was purified by a PD10 column (manufactured by Amersham Bioscience K.K.) to give a peptide. The purified peptide obtained in the foregoing manner was run through "AKTA explorer10XT" manufactured by Amersham Bioscience K.K. [column: "Nova-Pak C18", manufactured by Millipore Corporation, 3.9mm ϕ x 150mm, mobile phase: a mixed solvent of acetonitrile and water containing 0.05 vol.% of trifluoroacetic acid (concentration of acetonitrile was linearly increased from 5 to 50 vol.% for 30 minutes), flow rate: 1.0 mL/min.], and a single peak was shown at a retention time of 14.5 minutes. The molecular weight of the purified peptide was determined as 1375 based on FAB method mass spectrum (theoretical value: 1374.52).

[0082] The compound HOOC-(CH₂)₂-Co-(Pro-Pro-Gly)₅-OH obtained by the above-mentioned manner was dissolved in water or water containing 50% of methyl alcohol to prepare a solution having a concentration of 0.25 mg/mL, and a circular dichroism spectrum thereof was measured with a spectrometer (J-725, manufactured by JASCO Corp., light path: 1 mm). The results in both solutions showed that no positive Cotton effect was observed at a wavelength in range of 220 to 230 nm and only negative Cotton effect was observed at a wavelength in range of 200 to 202 nm, and confirmed that the peptide did not form a triple helical structure. Moreover, with respect to the same solution, the molecular weight was measured by gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex Peptide PE 7. 5/300, flow rate: 0.25mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl). The result confirmed one peak showing a molecular weight of not more than 2000 corresponding to a monomer. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit (manufactured by Amersham Bioscience K.K.), human insulin (manufactured by Sigma Corporation) and glycine (manufactured by Wako Pure Chemical Industries, Ltd.) as reference materials. The proportion of the peptide forming the triple helical structure was 0% on the basis of the peak area.

[0083] The compound HOOC-(CH₂)₂-CO-(Pro-Pro-Gly)₅-OH (1.4 mg (0.001 mmol)) and ethylenediamine (0.06 mg (0.001 mmol)) were suspended in 0.05 mL water, and to the suspended mixture were added 0.32 mg (0.0024 mmol) of 1-hydroxybenzotriazole and 4.6 mg (0.024 mmol) of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride. Then, the resulting mixture was shaken at a room temperature for 3 days.

[0084] The reaction mixture was diluted 100-fold with water, and the diluted solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex 200 HR 10/30, flow rate: 0.5 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl) to determine the molecular weight of the reaction product. The result confirmed a polypeptide peak in a molecular weight range of 30000 to 200000. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit and a Gel Filtration HMW Calibration Kit (manufactured by Amersham Bioscience K.K.) as reference materials.

[0085] The resulting reaction mixture was diluted 5-fold with water, and the diluted solution was dialyzed against water for 3 days to remove the reagent (s) such as the condensing agent and the unreacted monomer(s). The circular dichroism spectrum was measured in the obtained polypeptide, and the result showed that positive Cotton effect was observed at a wavelength of 228 nm and negative Cotton effect at a wavelength of 198 nm, and confirmed that the polypeptide formed a triple helical structure.

Example 3

[0086] A peptide represented by the formula H-(Pro-Hyp-Gly)₁₀-OH (Sequence ID No. 3) (manufactured by Peptide Institute, Inc.) was dissolved in water or water containing 50% of methyl alcohol to prepare a solution having a concen-

tration of 0.25 mg/mL, and a circular dichroism spectrum thereof was measured with a spectrometer (J-725, manufactured by JASCO Corp., light path: 1 mm). In water, the result showed that positive Cotton effect was observed at a wavelength of 225 nm and negative Cotton effect at a wavelength of 195 nm, and confirmed that the peptide formed a triple helical structure. Moreover, the same solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex Peptide PE 7.5/300, flow rate: 0.25 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl) thereby confirming one peak showing a molecular weight of about 9000 corresponding to a triple helical structure. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit (manufactured by Amersham Bioscience K.K.), human insulin (manufactured by Sigma Corporation) and glycine (manufactured by Wako Pure Chemical Industries, Ltd.) as reference materials. The proportion of the peptide forming the triple helical structure was about 100% on the basis of the peak area. Also, in the case of water containing 50% of methyl alcohol, almost the same results were given, and the proportion of the peptide forming the triple helical structure was about 100%.

[0087] Five (5) mg (0.0016 mmol) of the peptide H-(Pro-Hyp-Gly)₁₀-OH was suspended in 2 mL dimethyl sulf oxide, and the mixture was stirred at a room temperature. To the mixture were added 0.23 mg (0.0018 mmol) of diisopropylethylamine, 0.24 mg (0.0018 mmol) of 1-hydroxybenzotriazole, and 0.65 mg (0.0034 mmol) of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride, and the resultant mixture was further stirred for 7 days at a room temperature.

[0088] The reaction mixture was diluted 20-fold with water, and the diluted solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex 200 HR 10/30, flow rate: 0.5 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl). The result confirmed a polypeptide peak was observed in a molecular weight range of 60000 to not less than 200000. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit and a Gel Filtration HMW Calibration Kit (manufactured by Amersham Bioscience K.K.) as reference materials.

[0089] The resulting reaction mixture was diluted 5-fold with water, and the diluted solution was dialyzed against water for 3 days to remove the reagent(s) such as the condensing agent and the unreacted monomer(s). The circular dichroism spectrum was measured in the obtained polypeptide. The result showed that positive Cotton effect was observed at a wavelength of 225 nm and negative Cotton effect at a wavelength of 197 nm, and confirmed that the polypeptide formed a triple helical structure.

[0090] The water suspension containing the resulting polypeptide was cast on a sheet made from a fluorine-containing resin (polytetrafluoroethylene), and the cast sheet was air-dried to obtain a cast film. Gold was deposited on the film, and the deposited film was observed with the use of a scanning electron microscope. As a result, a fibrous structure was observed as shown in Figure 1.

Example 4

[0091] A peptide represented by the formula H-(Pro-Pro-Gly)₅-OH (manufactured by Peptide Institute, Inc.) was dissolved in water or water containing 50% of methyl alcohol to prepare a solution having a concentration of 0.25 mg/mL and a circular dichroism spectrum thereof was measured with a spectrometer (J-725, manufactured by JASCO Corp., light path: 1 mm). The results in both solutions showed that no positive Cotton effect was observed at a wavelength in range of 220 to 230 nm and only negative Cotton effect was observed at a wavelength in range of 200 to 202 nm, and confirmed that the peptide did not form a triple helical structure. Moreover, the same solutions were subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex Peptide PE 7.5/300, flow rate: 0.25 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl). The result confirmed one peak showing a molecular weight of not more than 2000 corresponding to a monomer. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit (manufactured by Amersham Bioscience K.K.), human insulin (manufactured by Sigma Corporation) and glycine (manufactured by Wako Pure Chemical Industries, Ltd.) as reference materials. The proportion of the peptide forming the triple helical structure was 0% on the basis of the peak area.

[0092] The peptide H-(Pro-Pro-Gly)₅-OH (3.5 mg (0.0026 mmol)) and the peptide H-(Val-Pro-Gly-Val-Gly)₂-OH (Sequence ID No. 4) (0.92 mg (0.0011 mmol)) synthesized in the same manner as in Example 2 were suspended in 1.5 mL dimethyl sulfoxide in a predetermined proportion, and the suspension was stirred at a room temperature. To the mixture were added 0.52 mg (0.0040 mmol) of diisopropylethylamine, 0.51 mg (0.0038 mmol) of 1-hydroxybenzotriazole, and 1.45 mg (0.0076 mmol) of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride, and the resultant mixture was further stirred for 7 days at a room temperature.

[0093] The reaction mixture was diluted 20-fold with water, and the diluted solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex 200 HR 10/30, flow rate: 0.5 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl) to determine the molecular weight of the resulting polypeptide. The result confirmed a polypeptide peak in a molecular weight range of 80000 to 450000. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit and a Gel Filtration HMW Calibration Kit (manufactured by Amersham Bioscience K.K.) as reference materials.

[0094] The resulting reaction mixture was diluted 5-fold with water, and the diluted solution was dialyzed against water for 3 days to remove the reagent(s) such as the condensing agent and the unreacted monomer(s). The circular dichroism spectrum was measured in the obtained polypeptide. The result showed that positive Cotton effect was observed at a wavelength of 227 nm and negative Cotton effect at a wavelength of 198 nm, and confirmed that the polypeptide formed a triple helical structure.

[0095] The water suspension containing the resulting polypeptide was cast on a sheet made of a fluorine-containing resin (polytetrafluoroethylene), and the cast sheet was air-dried to obtain a cast film. The film was immersed in 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl to give a gel sheet. The gel sheet was transparent at a room temperature, and was reversibly clouded at a temperature of not lower than 40°C.

Example 5

[0096] A peptide represented by the formula H-(Pro-Hyp-Gly)₅-OH (Sequence ID No. 5) (manufactured by Peptide Institute, Inc.) was dissolved in water or water containing 50% of methyl alcohol to prepare a solution having a concentration of 0.25 mg/mL, and a circular dichroism spectrum thereof was measured with a spectrometer (J-725, manufactured by JASCO Corp., light path: 1 mm). In water, the result showed that positive Cotton effect was observed at a wavelength of 223 nm and negative Cotton effect at a wavelength of 201 nm, and confirmed that the peptide formed a triple helical structure. Moreover, the same solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex Peptide PE 7.5/300, flow rate: 0.25 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl) thereby confirming one peak showing a molecular weight of about 4100 corresponding to a triple helical structure. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit (manufactured by Amersham Bioscience K.K.), human insulin (manufactured by Sigma Corporation) and glycine (manufactured by Wako Pure Chemical Industries Ltd.) as reference materials. The proportion of the peptide forming the triple helical structure was about 100% on the basis of the peak area. Also, in the case of water containing 50% of methyl alcohol, almost the same results were given, and the proportion of the peptide forming the triple helical structure was about 100%.

[0097] Five (5) mg (0.0033 mmol) of the peptide H-(Pro-Hyp-Gly)₅-OH was suspended in 2 mL dimethyl sulfoxide, and the mixture was stirred at a room temperature. To the mixture were added 0.44 mg (0.0034 mmol) of diisopropylethylamine, 0.46 mg (0.0033 mmol) of 1-hydroxybenzotriazole, and 1.3 mg (0.0068 mmol) of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride, and the resultant mixture was further stirred for 14 days at a room temperature.

[0098] The reaction mixture was diluted 20-fold with water, and the diluted solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex 200 HR 10/30, flow rate: 0.5 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl). A polypeptide peak was observed in a molecular weight range of 40000 to not less than 100000. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit and a Gel Filtration HMW Calibration Kit (manufactured by Amersham Bioscience K.K.) as reference materials.

[0099] The resulting reaction mixture was diluted 5-fold with water, and the diluted solution was dialyzed against water for 3 days to remove the reagent(s) such as the condensing agent and the unreacted monomer(s). The circular dichroism spectrum was measured in the obtained polypeptide. The result showed that positive Cotton effect was observed at a wavelength of 224 nm and negative Cotton effect at a wavelength of 199 nm, and confirmed that the polypeptide formed a triple helical structure.

Example 6

[0100] Five (5) mg (0.0016 mmol) of a peptide represented by the formula: H-(Pro-Hyp-Gly)₁₀-OH (Sequence ID No. 3) (manufactured by Peptide Institute, Inc.) was dissolved in 0.5 mL of 10 mM phosphate buffer (containing 8.1 mM Na₂HPO₄·12H₂O, 1.5 mM KH₂PO₄ and 2.7 mM KCl, pH 7.4), and the solution was stirred at 20°C. To the solution were added 0.24 mg (0.0018 mmol) of 1-hydroxybenzotriazole and 31 mg (0.16 mmol) of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride, and the resultant mixture was further stirred for 24 hours at 20°C.

[0101] The reaction mixture was diluted 60-fold with water, and the diluted solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superose 6 HR 10/30, flow rate: 0.5 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl) to determine the molecular weight of the reaction product. The result confirmed a polypeptide peak corresponding to an average molecular weight of 400000. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit and a Gel Filtration HMW Calibration Kit (manufactured by Amersham Bioscience K.K.) as reference materials.

[0102] Moreover, the resulting reaction mixture was diluted 5-fold with water, and the diluted solution was dialyzed against water for 3 days to remove the reagent(s) such as the condensing agent and the unreacted monomer(s). The circular dichroism spectrum was measured in the obtained polypeptide, and the result showed that positive Cotton effect

was observed at a wavelength of 225 nm and negative Cotton effect at a wavelength of 197 nm, and confirmed that the polypeptide formed a triple helical structure.

Example 7

[0103] A peptide represented by the formula: H-(Pro-Hyp-Gly)₁-OH (manufactured by Peptide Institute, Inc.) was dissolved in water to prepare a solution having a concentration of 0.25 mg/mL, and a circular dichroism spectrum thereof was measured with a spectrometer (J-820, manufactured by JASCO Corp., light path: 1 mm). The results showed that positive Cotton effect was observed at a wavelength of 214 nm and negative Cotton effect at a wavelength of 196 nm, and confirmed that the peptide did not form a triple helical structure. Moreover, with respect to the same solution, the molecular weight was measured by gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex Peptide PE 7.5/300, flow rate: 0.25 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl). The result confirmed one peak showing a molecular weight of about 250 corresponding to a monomer. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit (manufactured by Amersham Bioscience K.K.), human insulin (manufactured by Sigma Corporation) and glycine (manufactured by Wako Pure Chemical Industries, Ltd.) as reference materials. The proportion of the peptide forming the triple helical structure was 0%.

[0104] Twenty-five (25) mg (0.088 mmol) of a peptide represented by the formula: H-(Pro-Hyp-Gly)₁-OH was dissolved in 2.5 mL of 10 mM phosphate buffer (pH 7.4), and the solution was stirred at 20°C. To the solution were added 12.1 mg (0.09 mmol) of 1-hydroxybenzotriazole and 168 mg (0.88 mmol) of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride, and the resultant mixture was further stirred for 24 hours at 20°C.

[0105] The reaction mixture was diluted 20-fold with water, and the diluted solution was subjected to a gel-permeation chromatography (AKTA purifier system, manufactured by Amersham Bioscience K.K., column: Superdex 200 HR 10/30, flow rate: 0.5 mL/min., eluent: 10 mM phosphate buffer (pH 7.4) containing 150 mM NaCl) to determine the molecular weight of the reaction product. The result confirmed a polypeptide peak in a molecular weight range of 20000 to 30000. The molecular weight was calculated with a Gel Filtration LMW Calibration Kit and a Gel Filtration HMW Calibration Kit (manufactured by Amersham Bioscience K.K.) as reference materials.

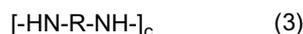
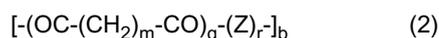
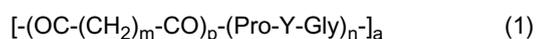
[0106] The resulting reaction mixture was diluted 5-fold with water, and the diluted solution was dialyzed against water for 3 days to remove the reagent(s) such as the condensing agent and the unreacted monomer(s). The circular dichroism spectrum was measured in the obtained polypeptide, and the result showed that positive Cotton effect was observed at a wavelength of 225 nm and negative Cotton effect at a wavelength of 199 nm, and confirmed that the polypeptide formed a triple helical structure.

Test Example

[0107] A cast film was made from each polypeptide manufactured in Examples 1 and 3, and the film was pasteurized with 25 kGy gamma irradiation. About 10,000 of mouse normal fibroblast strain NIH3T3 were dispersed on the film, and was incubated in an Eagle's Minimum Essential Medium containing 10% bovine fetal serum in the presence of 5% CO₂ at 37°C for 3 days. The NIH3T3 cell was well adhered to and proliferated on the film made from each polypeptide obtained in Examples 1 and 3, and was no abnormality in configuration or others.

Claims

1. A polypeptide comprising units represented by the following formulae (1) to (3):



wherein "m" denotes an integer of 1 to 18, "p" and "q" are the same or different, each representing 0 or 1, "Y" represents Pro or Hyp, and "n" denotes an integer of 1 to 20; "Z" represents a peptide chain comprising 1 to 10 amino acid residue(s), "r" denotes an integer of 1 to 20, and "R" represents a linear or branched alkylene group; the molar ratio of "a" relative to "b" [a/b] is 100/0 to 30/70;

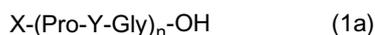
- (i) when p=1 and q=0, c=a=1,
- (ii) when p=0 and q=1, c=b=1,

(iii) when $p=1$ and $q=1$, $c=a+b=2$, and

(iv) when $p=0$ and $q=0$, $c=0$

wherein the polypeptide shows a peak corresponding to the molecular weight in a range from 1×10^4 to 100×10^4 in terms of a globular protein by means of an aqueous gel permeation chromatography.

2. A polypeptide according to claim 1, wherein "m" denotes an integer of 2 to 12; "n" denotes an integer of 1 to 15; "Z" is a peptide chain comprising at least an amino acid residue or a peptide residue selected from the group consisting of Gly, Sar, Ser, Glu, Asp, Lys, His, Ala, Val, Leu, Arg, Pro, Tyr, and Ile; "r" denotes an integer of 1 to 10; and "R" represents a C_{2-12} alkylene group.
3. A polypeptide according claim 1, which shows positive Cotton effect at a wavelength in range of 220 to 230 nm and negative Cotton effect at a wavelength in range of 195 to 205 nm in a circular dichroism spectrum.
4. A polypeptide according to claim 1 or 2, wherein at least part of the polypeptide forms a triple helical structure.
5. A polypeptide according to claim 1, which is in the form of a solution, a suspension, a particulate, a film, sheet, a fibrous substance, a gelatinous substance, or a sponge-like porous substance.
6. A polypeptide according to claim 1, which is in the form of a sponge-like porous substance obtained by lyophilizing the gelatinous substance.
7. A process for producing a polypeptide, which comprises reacting a peptide component (A) represented by the following formula (1a) with a peptide component (B) represented by the following formula (2a) and a compound (C) represented by the following formula (3a), as an optional component:



wherein "X" represents H or the group $\text{HOOC}-(\text{CH}_2)_m-\text{CO}-$ ("m" denotes an integer of 1 to 18), "Y" represents Pro or Hyp, and "n" denotes an integer of 1 to 20,



wherein "X" represents H or the group $\text{HOOC}-(\text{CH}_2)_m-\text{CO}-$ ("m" denotes an integer of 1 to 18), "Z" represents a peptide chain comprising 1 to 10 amino acid residue(s), and "r" denotes an integer of 1 to 20,



wherein "R" represents a linear or branched alkylene group,

wherein the ratio of the peptide component (A) relative to the peptide component (B) [the former (A)/the latter (B)] is 100/0 to 30/70 (mol%); provided that

in the case where "X" represents the group $\text{HOOC}-(\text{CH}_2)_m-\text{CO}-$ ("m" has the same meaning as defined above) in the formula (1a) and/or (2a), the amount of the compound (C) is substantially 1 mol relative to 1 mol of the total molar amount of the peptide component (A) and/or the peptide component (B);

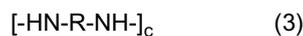
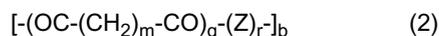
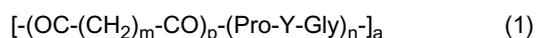
in the case where "X" represents H in the formula (1a) and/or (2a), the reaction is conducted without the compound (C) wherein the polypeptide shows a peak corresponding to the molecular weight in a range from 1×10^4 to 100×10^4 in terms of a globular protein by means of an aqueous gel permeation chromatography.

8. A process according to claim 7, wherein the reaction is carried out in the presence of at least a dehydrating and condensing agent.
9. A process according to claim 7, wherein the reaction is carried out in the presence of a dehydrating and condensing agent and a condensing auxiliary.
10. A process according to claim 8, wherein the dehydrating and condensing agent comprises at least one member selected from the group consisting of a carbodiimide-series condensing agent, a fluorophosphate-series condensing agent, and a diphenylphosphorylazide.

11. A process according to claim 8 which is conducted in a water-free solvent system, wherein the proportion of the dehydrating and condensing agent is 0.7 to 5 mol relative to 1 mol of the total molar amount of said reaction components (1a), (2a) and (3a).
- 5 12. A process according to claim 8 which is conducted in a water-containing solvent system, wherein the proportion of the dehydrating and condensing agent is 2 to 500 mol relative to 1 mol of the total molar amount of said reaction components (1a), (2a) and (3a).
- 10 13. A process according to claim 9, wherein the condensing auxiliary comprises at least one member selected from the group consisting of an N-hydroxypolycarboxylic imide, an N-hydroxytriazole, a triazine, and ethyl ester of 2-hydroxyimino-2-cyanoacetic acid.
14. A process according to claim 9, wherein the amount of the condensing auxiliary is 0.5 to 5 mol relative to the total molar amount of said reaction components (1a), (2a) and (3a).
- 15 15. A biomaterial or biocompatible material comprising the polypeptide according to any of claims 1 to 6.
16. A cosmetic preparation, which comprises the polypeptide according to claim any of claims 1 to 6.
- 20 17. A food additive, which comprises the polypeptide according to any of claims 1 to 6.
18. A coating agent for coating the surface of a substrate, consisting of the polypeptide of any of claims 1 to 6.
- 25 19. Polypeptide according to any of claims 1 to 6 for use as a biomaterial or biocompatible material, a medical material, a pharmaceutical raw or base material, a cosmetic raw or base material, a food additive, or a coating agent.
- 30 20. Polypeptide for use according to claim 19, wherein the medical material is a tissue engineering carrier, a regenerative medical treating carrier, a tissue binding agent, an antiadhesive material, a surgical operating suture, a hemostatic material or a contact lens.
- 35 21. Polypeptide according to any of claims 1 to 6 for use as a component for a pharmaceutical preparation or a component for a cosmetic preparation.
22. Use of the polypeptide according to any of claims 1 to 6 for the manufacture of a medicament for the treatment of a tissue of a subject.

Patentansprüche

- 40 1. Polypeptid, das Einheiten umfasst, die durch die folgenden Formeln (1) bis (3) dargestellt werden:



wobei "m" eine ganze Zahl von 1 bis 18 bezeichnet, "p" und "q" gleich oder verschieden sind und jeweils für 0 oder 1 stehen, "Y" für Pro oder Hyp steht und "n" eine ganze Zahl von 1 bis 20 bezeichnet, "Z" für eine Peptidkette steht, die 1 bis 10 Aminosäurereste umfasst, "r" eine ganze Zahl von 1 bis 20 bezeichnet und "R" für eine lineare oder verzweigte Alkylengruppe steht, wobei das Stoffmengenverhältnis von "a" zu "b" [a/b] 100/0 bis 30/70 beträgt; wobei gilt:

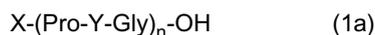
- 55 (i) wenn $p = 1$ und $q = 0$, dann $c = a = 1$;
 (ii) wenn $p = 0$ und $q = 1$, dann $c = b = 1$;
 (iii) wenn $p = 1$ und $q = 1$, dann $c = a + b = 2$; und
 (iv) wenn $p = 0$ und $q = 0$, dann $c = 0$;

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wobei das Polypeptid bei einer wässrigen Gelpermeationschromatographie einen Peak zeigt, der dem Molekulargewicht in einem Bereich von 1×10^4 bis 100×10^4 als globuläres Protein entspricht.

- 5 2. Polypeptid gemäß Anspruch 1, wobei "m" eine ganze Zahl von 2 bis 12 bezeichnet, "n" eine ganze Zahl von 1 bis 15 bezeichnet, "Z" eine Peptidkette ist, die wenigstens einen Aminosäurerest oder einen Peptidrest umfasst, der aus der Gruppe ausgewählt ist, die aus Gly, Sar, Ser, Glu, Asp, Lys, His, Ala, Val, Leu, Arg, Pro, Tyr und Ile besteht, "r" eine ganze Zahl von 1 bis 10 bezeichnet und "R" für eine C_{2-12} -Alkylengruppe steht.
- 10 3. Polypeptid gemäß Anspruch 1, das in einem Zirkulardichroismus-Spektrum einen positiven Cotton-Effekt bei einer Wellenlänge im Bereich von 220 bis 230 nm und einen negativen Cotton-Effekt bei einer Wellenlänge im Bereich von 195 bis 205 nm zeigt.
- 15 4. Polypeptid gemäß Anspruch 1 oder 2, wobei wenigstens ein Teil des Polypeptids eine Tripelhelixstruktur bildet.
5. Polypeptid gemäß Anspruch 1, das in Form einer Lösung, einer Suspension, von Teilchen, eines Films, einer Platte, einer faserigen Substanz, einer gelatinösen Substanz oder einer schwammartigen porösen Substanz vorliegt.
- 20 6. Polypeptid gemäß Anspruch 1, das in Form einer schwammartigen porösen Substanz vorliegt, die durch Lyophilisieren der gelatinösen Substanz erhalten wird.
7. Verfahren zur Herstellung eines Polypeptids, umfassend:

Umsetzen einer Peptidkomponente (A), die durch die folgende Formel (1a) dargestellt wird, mit einer Peptidkomponente (B), die durch die folgende Formel (2a) dargestellt wird, und einer Verbindung (C), die durch die folgende Formel (3a) dargestellt wird, als wahlfreier Komponente:



wobei "X" für H oder die Gruppe $\text{HOOC}-(\text{CH}_2)_m\text{-CO-}$ steht (und "m" eine ganze Zahl von 1 bis 18 bezeichnet), "Y" für Pro oder Hyp steht und "n" eine ganze Zahl von 1 bis 20 bezeichnet;



wobei "X" für H oder die Gruppe $\text{HOOC}-(\text{CH}_2)_m\text{-CO-}$ steht (und "m" eine ganze Zahl von 1 bis 18 bezeichnet), "Z" für eine Peptidkette steht, die 1 bis 10 Aminosäurereste umfasst, und "r" eine ganze Zahl von 1 bis 20 bezeichnet;



wobei "R" für eine lineare oder verzweigte Alkylengruppe steht;

wobei das Verhältnis der Peptidkomponente (A) zur Peptidkomponente (B) [erstere (A)/letzterer (B)] 100/0 bis 30/70 (Mol-%) beträgt; mit der Maßgabe, dass

im Fall, dass "X" in Formel (1a) und/oder (2a) für die Gruppe $\text{HOOC}-(\text{CH}_2)_m\text{-CO-}$ steht (und "m" dasselbe wie oben bedeutet), die Menge der Verbindung (C) im Wesentlichen 1 mol relativ zu 1 mol der Gesamtstoffmenge der Peptidkomponente (A) und/oder der Peptidkomponente (B) beträgt;

im Fall, dass "X" in Formel (1a) und/oder (2a) für H steht, die Reaktion ohne die Verbindung (C) durchgeführt wird, wobei das Polypeptid bei einer wässrigen Gelpermeationschromatographie einen Peak zeigt, der dem Molekulargewicht in einem Bereich von 1×10^4 bis 100×10^4 als globuläres Protein entspricht.

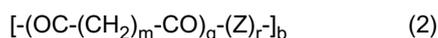
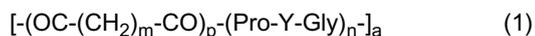
8. Verfahren gemäß Anspruch 7, wobei die Reaktion in Gegenwart von wenigstens einem Dehydratisierungs- und Kondensationsmittel durchgeführt wird.
9. Verfahren gemäß Anspruch 7, wobei die Reaktion in Gegenwart eines Dehydratisierungs- und Kondensationsmittels und eines Kondensationshilfsstoffs durchgeführt wird.
10. Verfahren gemäß Anspruch 8, wobei das Dehydratisierungs- und Kondensationsmittel wenigstens einen Vertreter umfasst, der aus der Gruppe ausgewählt ist, die aus einem Kondensationsmittel der Carbodiimid-Reihe, einem

Kondensationsmittel der Fluorophosphat-Reihe und einem Diphenylphosphorylazid besteht.

- 5 11. Verfahren gemäß Anspruch 8, das in einem wasserfreien Lösungsmittelsystem durchgeführt wird, wobei der Anteil des Dehydratisierungs- und Kondensationsmittels 0,7 bis 5 mol relativ zu 1 mol der Gesamtstoffmenge der Reaktionskomponenten (1a), (2a) und (3a) beträgt.
- 10 12. Verfahren gemäß Anspruch 8, das in einem wasserhaltigen Lösungsmittelsystem durchgeführt wird, wobei der Anteil des Dehydratisierungs- und Kondensationsmittels 2 bis 500 mol relativ zu 1 mol der Gesamtstoffmenge der Reaktionskomponenten (1a), (2a) und (3a) beträgt.
- 15 13. Verfahren gemäß Anspruch 9, wobei der Kondensationshilfsstoff wenigstens einen Vertreter umfasst, der aus der Gruppe ausgewählt ist, die aus einem N-Hydroxypolycarboxylimid, einem N-Hydroxytriazol, einem Triazin und einem Ethylester von 2-Hydroxyimino-2-cyanoessigsäure besteht.
- 15 14. Verfahren gemäß Anspruch 9, wobei die Menge des Kondensationshilfsstoffs 0,5 bis 5 mol relativ zur Gesamtstoffmenge der Reaktionskomponenten (1a), (2a) und (3a) beträgt.
- 20 15. Biomaterial oder biokompatibles Material, das das Polypeptid gemäß einem der Ansprüche 1 bis 6 umfasst.
- 20 16. Kosmetikpräparat, das das Polypeptid gemäß einem der Ansprüche 1 bis 6 umfasst.
17. Lebensmittelzusatz, der das Polypeptid gemäß einem der Ansprüche 1 bis 6 umfasst.
- 25 18. Beschichtungsmittel zur Beschichtung der Oberfläche eines Substrats, wobei das Beschichtungsmittel aus dem Polypeptid gemäß einem der Ansprüche 1 bis 6 besteht.
- 30 19. Polypeptid gemäß einem der Ansprüche 1 bis 6 zur Verwendung als Biomaterial oder biokompatibles Material, medizinisches Material, pharmazeutischen Roh- oder Grundstoff, kosmetischen Roh- oder Grundstoff, Lebensmittelzusatz oder Beschichtungsmittel.
- 30 20. Polypeptid zur Verwendung gemäß Anspruch 19, wobei das medizinische Material ein Tissue-Engineering-Träger, ein Träger für eine regenerative medizinische Behandlung, ein Gewebepolymers, ein antiadhäsives Material, ein chirurgisches Nahtmaterial, ein hämostatisches Material oder eine Kontaktlinse ist.
- 35 21. Polypeptid gemäß einem der Ansprüche 1 bis 6 zur Verwendung als Komponente für ein pharmazeutisches Präparat oder als Komponente für ein kosmetisches Präparat.
- 40 22. Verwendung des Polypeptids gemäß einem der Ansprüche 1 bis 6 zur Herstellung eines Medikaments für die Behandlung eines Gewebes eines Patienten.

Revendications

- 45 1. Polypeptide comprenant des motifs représentés par les formules (1) à (3) suivantes:



où "m" désigne un nombre entier allant de 1 à 18, "p" et "q" sont identiques ou différents, chacun représentant 0 ou 1, "Y" représente Pro ou Hyp, et "n" désigne un nombre entier allant de 1 à 20; "Z" représente une chaîne peptidique comprenant de 1 à 10 restes d'acides aminés, "r" désigne un nombre entier allant de 1 à 20, et "R" représente un groupe alkylène linéaire ou ramifié; le rapport molaire de "a" relatif à "b" [a/b] est de 100/0 à 30/70;

55

(i) lorsque $p = 1$ et $q = 0$, alors $c = a = 1$;

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- (ii) lorsque $p = 0$ et $q = 1$, alors $c = b = 1$;
- (iii) lorsque $p = 1$ et $q = 1$, alors $c = a + b = 2$; et
- (iv) lorsque $p = 0$ et $q = 0$, alors $c = 0$;

5 dans lequel le polypeptide montre un pic correspondant au poids moléculaire dans la gamme de 1×10^4 à 100×10^4 en tant que protéine globulaire au moyen d'une chromatographie par perméation de gel aqueux.

10 **2.** Polypeptide selon la revendication 1, dans lequel "m" désigne un nombre entier allant de 2 à 12, "n" désigne un nombre entier allant de 1 à 15; "Z" est une chaîne peptidique comprenant au moins un reste d'acide aminé ou reste peptidique choisi dans le groupe consistant en Gly, Sar, Ser, Glu, Asp, Lys, His, Ala, Val, Leu, Arg, Pro, Tyr et Ile; "r" désigne un nombre entier allant de 1 à 10; et "R" représente un groupe alkylène en C_{2-12} .

15 **3.** Polypeptide selon la revendication 1, montrant un effet Cotton positif à une longueur d'onde comprise entre 220 et 230 nm et un effet Cotton négatif à une longueur d'onde comprise entre 195 et 205 nm dans un spectre de dichroïsme circulaire.

4. Polypeptide selon la revendication 1 ou 2, dans lequel au moins une partie du polypeptide forme une structure en triple hélice.

20 **5.** Polypeptide selon la revendication 1, qui est sous la forme d'une solution, d'une suspension, de particules, d'un film, d'une plaque, d'une substance fibreuse, d'une substance gélatineuse ou d'une substance poreuse spongieuse.

6. Polypeptide selon la revendication 1, qui est sous la forme d'une substance poreuse spongieuse obtenue par lyophilisation de la substance gélatineuse.

25

7. Procédé pour produire un polypeptide, comprenant l'étape consistant à:

30 faire réagir un composant peptidique (A) représenté par la formule (1a) suivante avec un composant peptidique (B) représenté par la formule (2a) suivante et un composé (C) représenté par la formule (3a) suivante en tant que composant optionnel:



35 où "X" représente H ou le groupe $\text{HOOC}-(\text{CH}_2)_m\text{-CO-}$ ("m" désignant un nombre entier allant de 1 à 18), "Y" représente Pro ou Hyp et "n" désigne un nombre entier allant de 1 à 20;



40 où "X" représente H ou le groupe $\text{HOOC}-(\text{CH}_2)_m\text{-CO-}$ ("m" désignant un nombre entier allant de 1 à 18), "Z" représente une chaîne peptidique comprenant de 1 à 10 restes d'acides aminés et "r" désigne un nombre entier allant de 1 à 20;



45 où "R" représente un groupe alkylène linéaire ou ramifié;

dans lequel le rapport entre le composant peptidique (A) et le composant peptidique (B) [le premier (A)/le dernier (B)] est de 100/0 à 30/70 (moles pour cent); à la condition que:

50 dans le cas où "X" représente le groupe $\text{HOOC}-(\text{CH}_2)_m\text{-CO-}$ ("m" ayant la même signification que celle définie ci-dessus) dans la formule (1a) et/ou (2a), la quantité du composé (C) est sensiblement 1 mole par mole de la quantité molaire totale du composant peptidique (A) et/ou du composant peptidique (B);

dans le cas où "X" représente H dans la formule (1a) et/ou (2a), la réaction est conduite en absence du composé (C), dans lequel le polypeptide montre un pic correspondant au poids moléculaire dans la gamme de 1×10^4 à 100×10^4 en tant que protéine globulaire au moyen d'une chromatographie par perméation de gel aqueux.

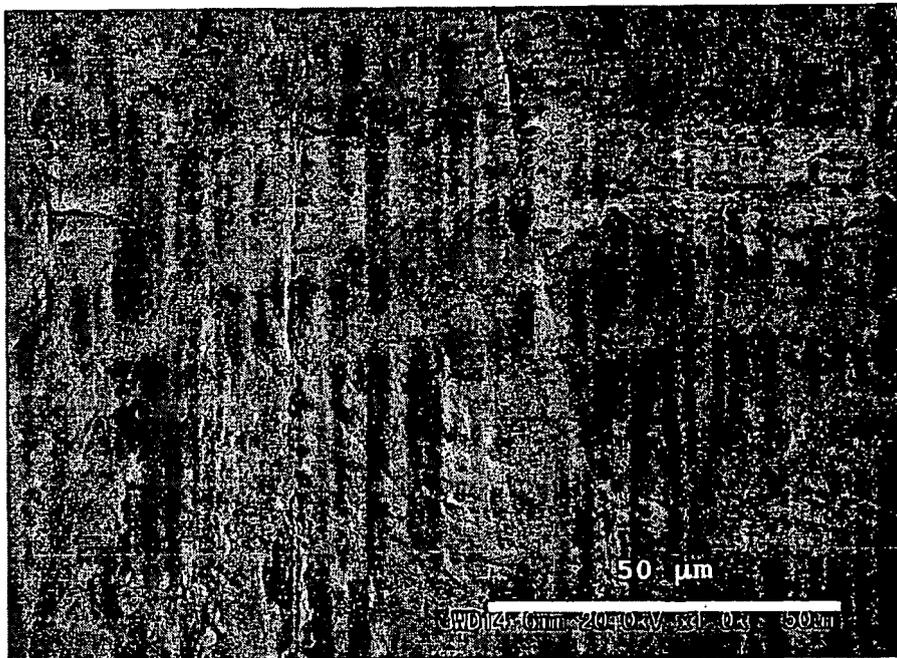
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8. Procédé selon la revendication 7, dans lequel la réaction est conduite en présence d'au moins un agent déshydratant et de condensation.

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9. Procédé selon la revendication 7, dans lequel la réaction est conduite en présence d'un agent déshydratant et de condensation et d'un auxiliaire de condensation.
- 5 10. Procédé selon la revendication 8, dans lequel l'agent déshydratant et de condensation comprend au moins un élément choisi dans le groupe consistant en un agent de condensation de la série des carbodiimides, un agent de condensation de la série des fluorophosphates et un diphénylphosphoryl-azide.
- 10 11. Procédé selon la revendication 8, qui est conduit dans un système de solvants anhydres, dans lequel la proportion de l'agent déshydratant et de condensation est de 0,7 à 5 moles par mole de la quantité molaire totale des composants réactifs (1a), (2a) et (3a).
- 15 12. Procédé selon la revendication 8, qui est conduit dans un système de solvants contenant de l'eau, dans lequel la proportion de l'agent déshydratant et de condensation est de 2 à 500 moles par mole de la quantité molaire totale des composants réactifs (1a), (2a) et (3a).
- 20 13. Procédé selon la revendication 9, dans lequel ledit auxiliaire de condensation comprend au moins un élément choisi dans le groupe consistant en un imide N-hydroxypolycarboxylique, un N-hydroxytriazole, une triazine et un ester éthyle de l'acide 2-hydroxyimino-2-cyanoacétique.
- 25 14. Procédé selon la revendication 9, dans lequel la quantité de l'auxiliaire de condensation est de 0,5 à 5 moles par rapport à la quantité molaire totale des composants réactifs (1a), (2a) et (3a).
- 30 15. Biomatériau ou matériau biocompatible comprenant le polypeptide selon l'une quelconque des revendications 1 à 6.
- 35 16. Préparation cosmétique comprenant le polypeptide selon l'une quelconque des revendications 1 à 6.
- 40 17. Additif alimentaire comprenant le polypeptide selon l'une quelconque des revendications 1 à 6.
- 45 18. Agent de revêtement pour revêtir la surface d'un substrat, l'agent de revêtement consistant en le polypeptide selon l'une quelconque des revendications 1 à 6.
- 50 19. Polypeptide selon l'une quelconque des revendications 1 à 6 destiné à être utilisé en tant que biomatériau ou matériau biocompatible, matériau médical, matière primaire pharmaceutique ou matériau pharmaceutique de base, matière primaire cosmétique ou matériau cosmétique de base, additif alimentaire ou agent de revêtement.
- 55 20. Polypeptide destiné à être utilisé selon la revendication 19, dans lequel ledit matériau médical est un transporteur de génie tissulaire, un transporteur de traitement médical régénératoire, un agent de liage tissulaire, un matériau antiadhésif, un matériau de suture chirurgical, un matériau hémostatique ou une lentille de contact.
21. Polypeptide selon l'une quelconque des revendications 1 à 6 destiné à être utilisé en tant que composant pour une préparation pharmaceutique ou composant pour une préparation cosmétique.
22. Utilisation du polypeptide selon l'une quelconque des revendications 1 à 6 pour la production d'un médicament pour le traitement d'un tissu d'un sujet.

Figure 1



REFERENCES CITED IN THE DESCRIPTION

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