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Kaplan et al.

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(54) **FIBROUS PROTEIN FUSIONS AND USE THEREOF IN THE FORMATION OF ADVANCED ORGANIC/INORGANIC COMPOSITE MATERIALS**

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C07K 14/00 (2006.01)

(52) **U.S. Cl.**
USPC **435/41**; 530/350; 530/352

(58) **Field of Classification Search**
None
See application file for complete search history.

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(57) **ABSTRACT**

The claimed invention provides a fusion polypeptide comprising a fibrous protein domain and a mineralization domain. The fusion is used to form an organic-inorganic composite. These organic-inorganic composites can be constructed from the nano- to the macro-scale depending on the size of the fibrous protein fusion domain used. In one embodiment, the composites can also be loaded with other compounds (e.g., dyes, drugs, enzymes) depending on the goal for the materials, to further enhance function. This can be achieved during assembly of the material or during the mineralization step in materials formation.

19 Claims, 15 Drawing Sheets

Fusion proteins of spider silk and dentin matrix protein 1 (DMP1)

1. Spider silk (CRGD-15mer) - full length DMP1 (Mw = 92.9 kDa)



CRGD-15mer (506 aa) full length DMP1 (474 aa)

2. Spider silk (CRGD-15mer) -C-terminal DMP1 (Mw = 58.9 kDa)



CRGD-15mer (506 aa) C-terminal DMP1 (156 aa)

3. Spider silk (CRGD-15mer) – (CD1 + pA + pB + CD2)



CRGD-15mer (506 aa) (37aa)

FIG. 1A

Fusion protein of spider silk and bone sialoprotein domain (BSP)

1. Spider silk (15mer)- sialoprotein domain (Mw = 42.7 kDa)



15mer (495 aa) BSP (48 aa)

2. Spider silk (CRGD-15mer) – sialoprotein domain (Mw = 48.1 kDa)



CRGD-15mer (499 aa) BSP (48 aa)

FIG. 1B

FIG. 2

MASMTGGCQMGRCRGDTSGRGGLGGGQAGAAAAAGCAGCGGGYGGGLGSGQTS
GRGGLGGGQAGAAAAAGGAGCGGGYGGGLGSGQTSGRGGLGGGQAGAAAAAGC
AGCGGGYGGGLGSGQTSGRGGLGGGQAGAAAAAGGAGCGGGYGGGLGSGQTSGRG
GLGGGQAGAAAAAGGAGCGGGYGGGLGSGQTSGRGGLGGGQAGAAAAAGCAGC
GGYGGGLGSGQTSGRGGLGGGQAGAAAAAGGAGCGGGYGGGLGSGQTSGRGGL
GGQAGAAAAAGGAGCGGGYGGGLGSGQTSGRGGLGGGQAGAAAAAGGAGCGGGY
GGGLGSGQTSGRGGLGGGQAGAAAAAGCAGCGGGYGGGLGSGQTSGRGGLGGGQ
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GSGQTSGRGGLGGGQAGAAAAAGGAGCGGGYGGGLGSGQTSGRGGLGGGQAG
AAAAAGGAGCGGGYGGGLGSGQTSGRGGLGGGQAGAAAAAGGAGCGGGYGGGLGSG
GTSRGDCGSLFVARYONTESSESSEERTGNLAQSPPPMANS DHTDSSSESGE
ELGSDRSQYRPAGGLSKSAGMDADKEEDEDSDGDDTFGDEDNGPGPEERQW
GGPSRLDSDDESDATTTQSSSEDSTSQENSAQDTPSDSKDHHSDEADSRPEAG
DSTQDSESEEYRVGGGSGEGSSHGDGSEFFDDEGMQSDDPGSTRSDRGHTRM
SSADISSEESKGDHEPTSTQSDSDSQDVEFSSRKSFRRSRVSEEDDRGELA
DSNSRETQSVSTEDFRSKEESRSETQEDTAETQSQEDSPEGQDPSSSESEE
AGEPSQESSSESQEGVASESRGDNPNQTSQTDGDRDESSEEDRLNTFSS
ESQSTEEQGDSESNESLSLEESQESAQDEDSSSQEGLSQSASRESRSQ
SQSEEDSRSEENRSDSDSQDSSRSKEESNSTGSTSSSEEDNHFKNIEADNRK
LIVDAYHNKPIGDQDQDNDQDGY (SEQ ID NO: 10)

FIG. 3

15mer-BSP (SEQ ID NO: 11): a fusion protein of (SEQ ID NO: 4) (light grey highlight) and (SEQ ID NO: 8) (dark grey highlight)

```
MASMTGGQQMGRGSAMASGRGGLGGQGAGAAAAAGGAGQGGYGGLGSQGTS
GRGGLGGQGAGAAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAAGG
AGQGGYGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGLGSQGTSGRG
GLGGQGAGAAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQ
GGYGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGLGSQGTSGRGGLG
GGAGAAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGY
GGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGLGSQGTSGRGGLGGQG
AGAAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGL
GSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGA
AAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGLGSQ
GTSEFFVQSSSDSSEENGNGDSSEEEEEEEENENNEENNEENEDSDGNEIX
LHHHHHH (SEQ ID NO: 11)
```

FIG. 4

CRGD-15mer-BSP (SEQ ID NO: 12): a fusion protein of (SEQ ID NO: 5) (light grey highlight) and (SEQ ID NO: 8) with linker

MASMTGGQQMGRGSCRGDTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQG
TSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAA
GGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSG
RGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGA
QGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGL
GGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGG
YGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGG
QGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGG
GLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGA
GAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLG
SQGTSGRGDCGSESEYFVQSSSDSSRENGNDSESESESESESESESESESESE
NEDSDGNEQKLHHHHHH (SEQ ID NO: 12)

FIG. 5

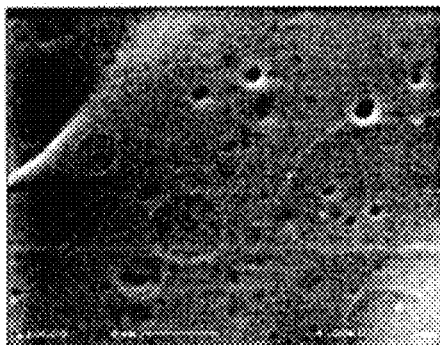
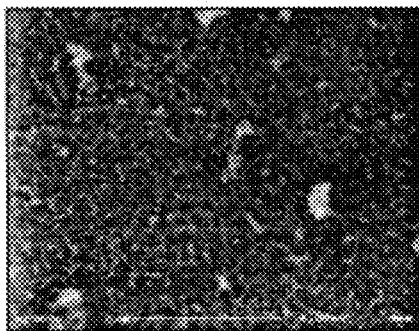
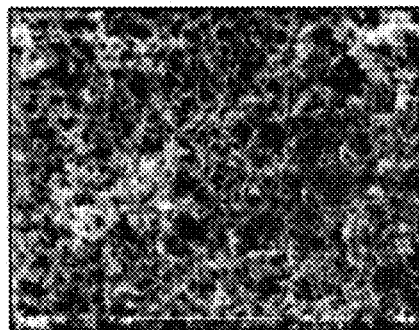


FIG. 6A



ROUND 1

FIG. 6B



ROUND 3

FIG. 6C

CRGD15mer-R5

MASMTGGQOM GRGS**CRG**DTs GRGGLGGQGA GAAAAAGGAG QGGYGGLGSQ GTSGRGGGLG 60
QGAGAAAAAG GAGQGGYGGL GSQGTSGRGG LGGQAGAGAA AAGGAGQGGY GGLGSQGTSG 120

RGGGGLGGQAG AAAAAAGGAGQ GYGGLGSQ TSGRGGGLGGQ GAGAAAAAGG AGQGGYGGLG 180
SQGTSGRGG L GGQAGAGAAA AGGAGQGGY GLGSQGTSGR GGLGGQGAGA AAAAGGAGQG 240
GYGGLGSQGT SGRGGLGGQ AGAAAAAGGA GQGGYGGLGS QGTSGRGG L GQAGAGAAAA 300
GGAGQGGYGG LGSQGTSGR GLGQAGAGAA AAAGGAGQGG YGGLGSQGTs GRGGLGGQGA 360
GAAAAAGGAG QGGYGGLGSQ GTSGRGG L GGQAGAGAAA GAGQGGYGG L GSQGTSGRGG 420
LGGQAGAGAA AAGGAGQGGY GGLGSQGTSG RGGGQGGAG AAAAAAGGAG GYGGLGSQ 480
TSGRGGGLGGQ GAGAAAAAGG AGQGGYGGLG SQGTs**RGDCG** SEFSSKKSGS YSGSKGSKRR 540
ILCGR**HHHHH** H (SEQ ID NO: 24) 551

FIG. 7A**15mer-R5**

M**HHHHH**SSG LVPRGSGMKE TAAAKFERQH MDSPDLGTDD DDKAMASGRG GLGGQAGAGAA 60
AAAGGAGQGG YGGLGSQGTs GRGGLGGQGA GAAAAAGGAG QGGYGGLGSQ GTSGRGG LGG 120
QGAGAAAAAG GAGQGGYGG L GSQGTSGRGG LGGQAGAGAA AAGGAGQGGY GGLGSQGTSG 180
RGGGGLGGQAG AAAAAAGGAGQ GYGGLGSQ TSGRGGGLGGQ GAGAAAAAGG AGQGGYGGLG 240
SQGTSGRGG L GGQAGAGAAA AGGAGQGGY GLGSQGTSGR GGLGGQGAGA AAAAGGAGQG 300
GYGGLGSQGT SGRGGLGGQ AGAAAAAGGA GQGGYGGLGS QGTSGRGG L GQAGAGAAAA 360
GGAGQGGYGG LGSQGTSGR GLGQAGAGAA AAAGGAGQGG YGGLGSQGTs GRGGLGGQGA 420
GAAAAAGGAG QGGYGGLGSQ GTSGRGG L GGQAGAGAAA GAGQGGYGG L GSQGTSGRGG 480
LGGQAGAGAA AAGGAGQGGY GGLGSQGTSG RGGGQGGAG AAAAAAGGAG GYGGLGSQ 540
TSSSKKSGSY SSKGSKRRI L (SEQ ID NO: 25) 561

FIG. 7B

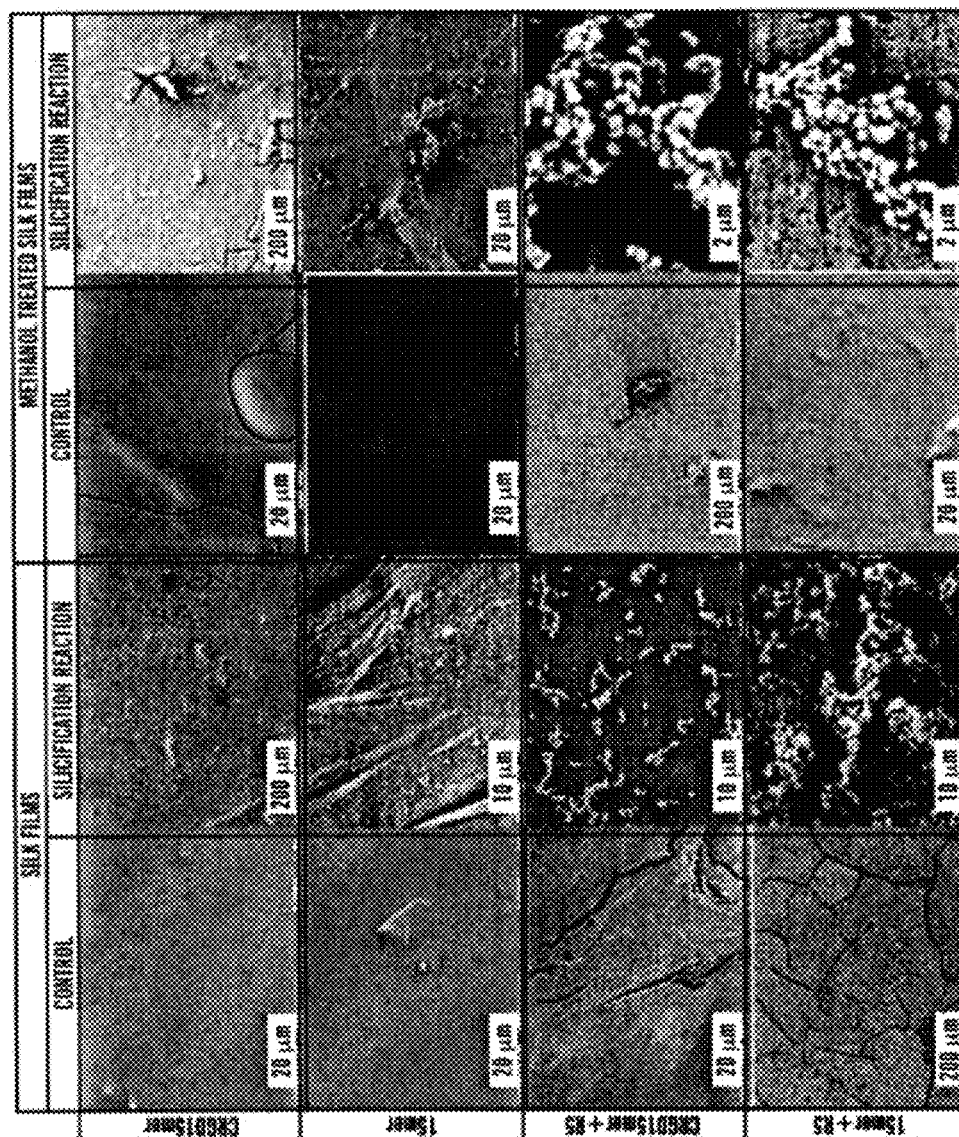


FIG. 8



FIG. 9E

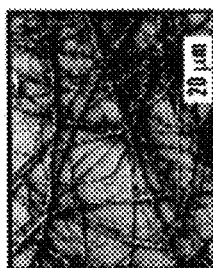


FIG. 9D

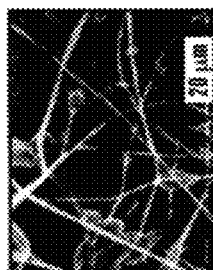


FIG. 9C



FIG. 9B

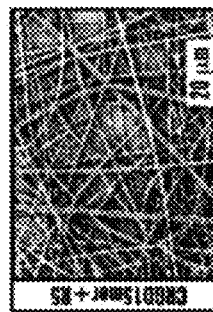
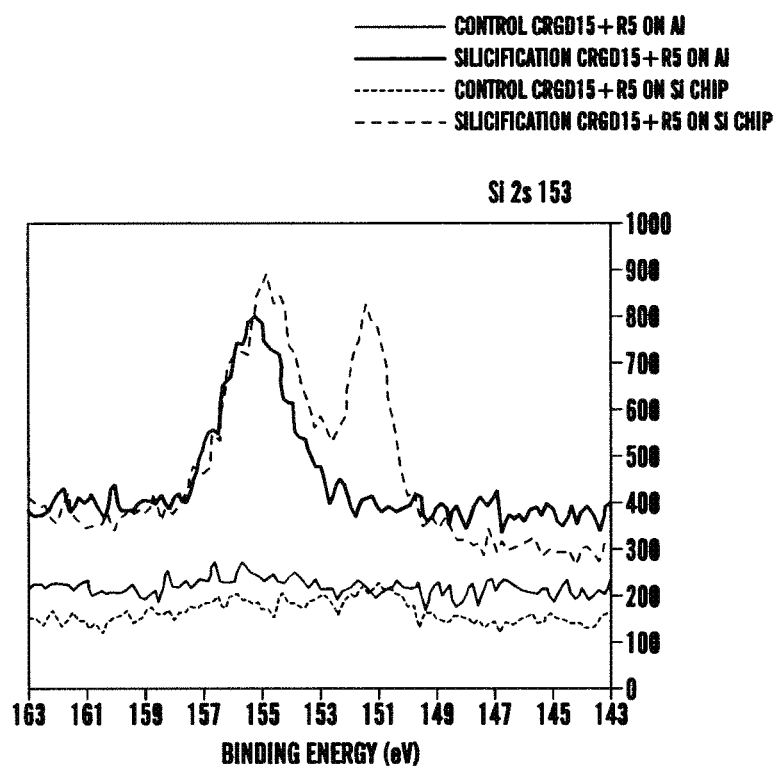
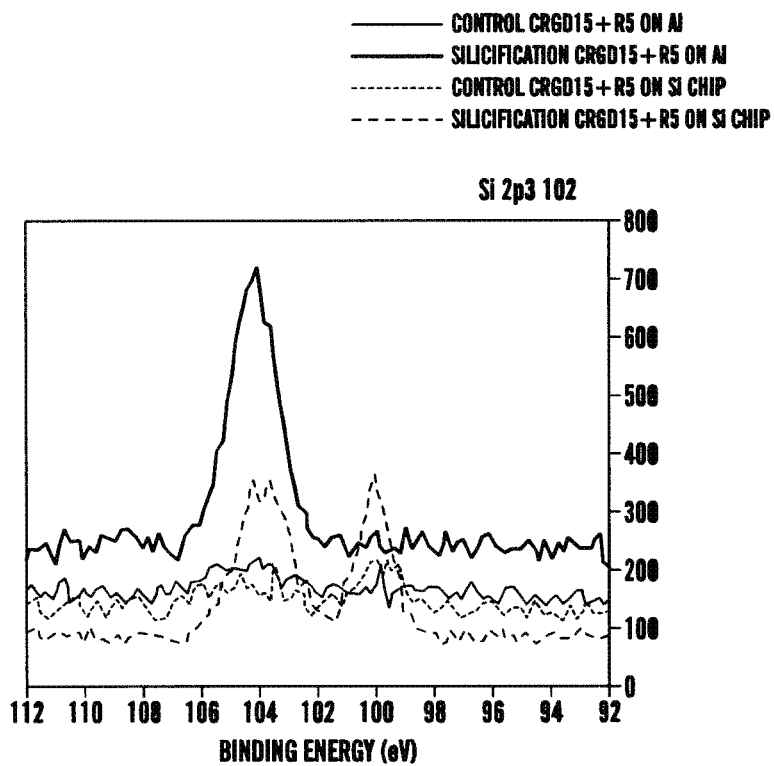
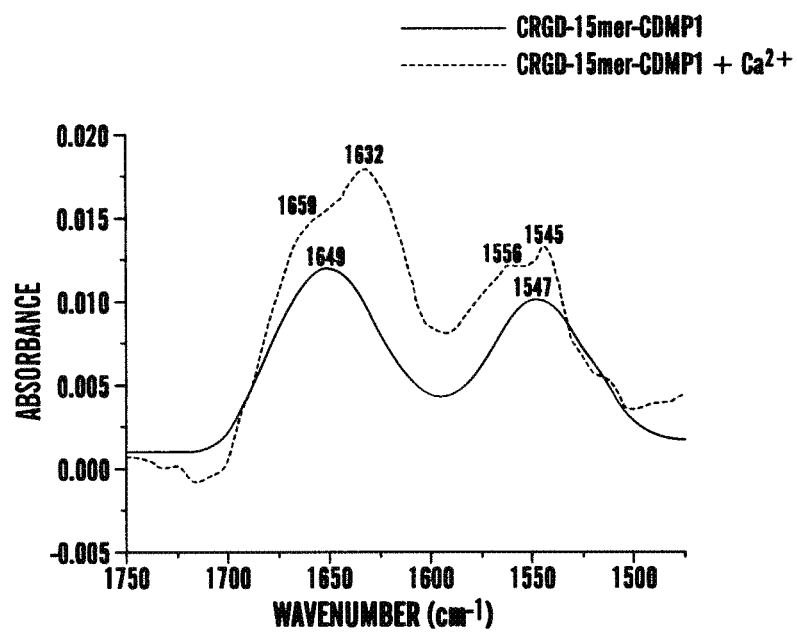
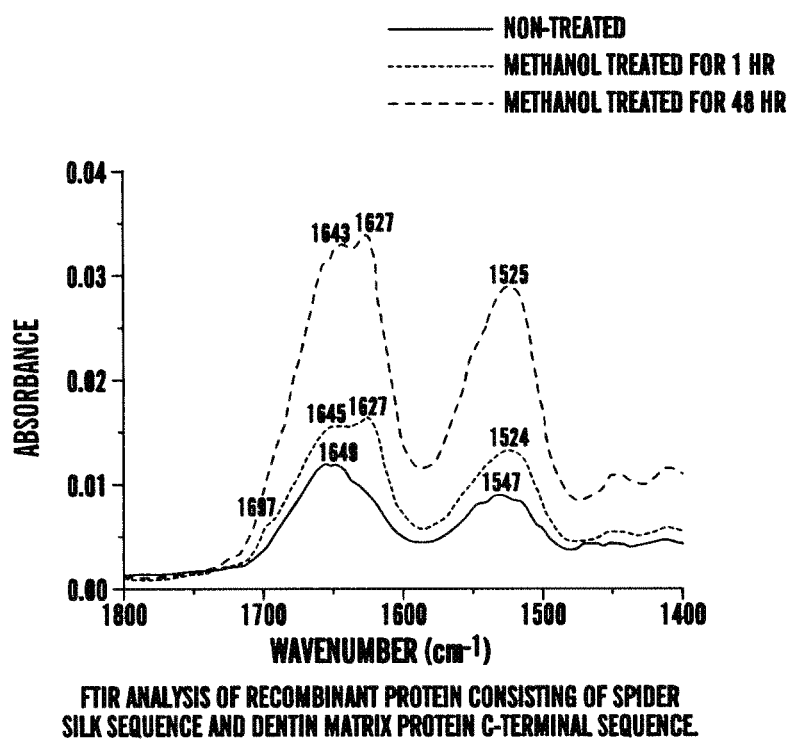


FIG. 9A

**FIG. 10A**

**FIG. 10B**

**FIG. 11**

**FIG. 12**

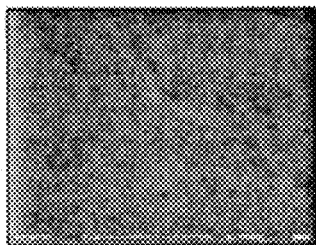
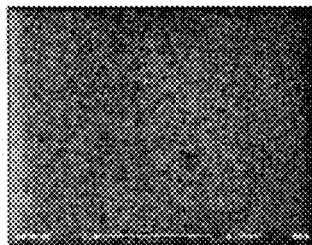
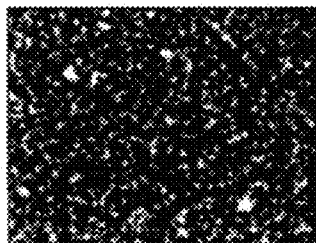
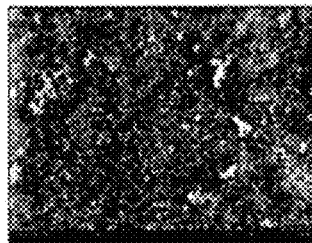
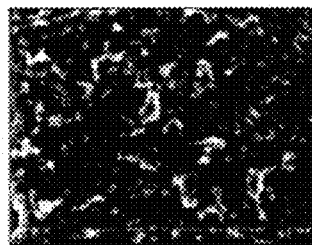
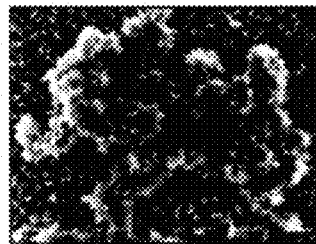
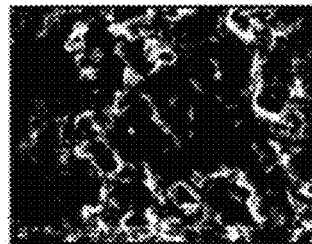
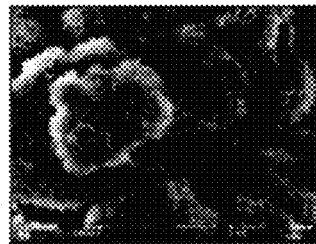
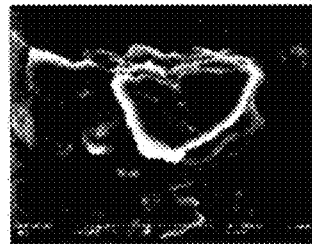
**FIG. 13A1****FIG. 13B1****FIG. 13A2****FIG. 13B2****FIG. 13A3****FIG. 13B3****FIG. 13A4****FIG. 13B4****FIG. 13A5****FIG. 13B5**

FIG. 14A

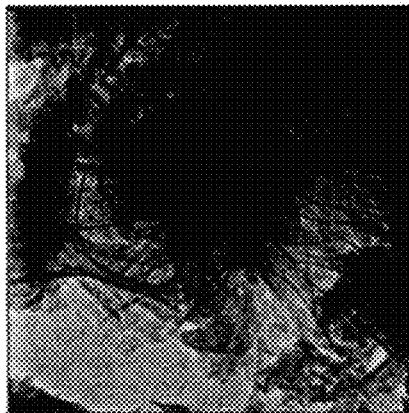


FIG. 14B

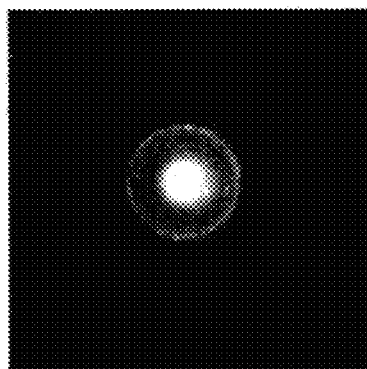


FIG. 14C



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FIBROUS PROTEIN FUSIONS AND USE THEREOF IN THE FORMATION OF ADVANCED ORGANIC/INORGANIC COMPOSITE MATERIALS

CROSS REFERENCED APPLICATIONS

This application is a continuation of U.S. Ser. No. 13/097, 538 filed Apr. 29, 2011, now issued as U.S. Pat. No. 8,129, 141, issued on Mar. 6, 2012, which is a continuation of U.S. Ser. No. 11/794,934 filed on Jan. 14, 2008, now U.S. Pat. No. 7,960,509, issued Jun. 14, 2011, which is a 371 National Phase Entry Application of International Application No. PCT/US2006/01536, filed Jan. 17, 2006, which designates the U.S., and which claims the benefit under 35 U.S.C. §119 (e) of U.S. Provisional Patent Application No. 60/644,264, filed Jan. 14, 2005, the contents of all of which are incorporated herein by reference in their entireties.

GOVERNMENT SUPPORT

This invention was supported by FA9550041-0363 awarded by the U.S. Air Force and EB003210-01 awarded by the National Institutes of Health. The government has certain rights in the invention.

BACKGROUND OF THE INVENTION

Many biomedical procedures require the provision of healthy tissue to counteract the disease process or trauma being treated. This work is often hampered by the tremendous shortage of tissues available for transplantation and/or grafting. Tissue engineering may ultimately provide alternatives to whole organ or tissue transplantation. In order to generate engineered tissues, various combinations of biomaterials and living cells are currently being investigated. Although attention is often focused on the cellular aspects of the engineering process, the design characteristics of the biomaterials also constitute a major challenge in this field.

In recent years, the ability to regenerate tissues and to control the properties of the regenerated tissue have been investigated by trying to specifically tune the mechanical or chemical properties of the biomaterial scaffold (Kim et al., 1997; Kohn et al. 1997). The majority of work has involved the incorporation of chemical factors into the material during processing, or the tuning of mechanical properties by altering the constituents of the material.

The foregoing methods have been used in an attempt to utilize chemical or mechanical signaling to affect changes in the proliferation and/or differentiation of cells during tissue regeneration. Despite such efforts there remains in the art a need for improved biomaterials, including composite materials, particularly those with a better capacity to support complex tissue growth in vitro (in cell culture) and in vivo (upon implantation).

SUMMARY OF THE INVENTION

The claimed invention provides a fusion polypeptide comprising a fibrous protein domain and a mineralization domain.

In one embodiment the fibrous protein domain is obtained from silk, collagens, coiled-coiled leucine zipper proteins, elastins, keratins, actins, and tubulins.

For example, in one preferred embodiment, the fibrous protein domain comprises an amino acid sequence from the silk protein Spidroin 1, such as the fibrous protein domains indicated by (SEQ ID NO: 1) or (SEQ ID NO: 3), which are

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derived from Spiroidin 1 of *Nephila clavipes*. A recombinant fibrous protein domain can be generated, which has multiple repeats of a fibrous protein domain.

In one embodiment, the fibrous domain sequence of (SEQ ID NO: 1) is repeated 15 times throughout the fibrous protein domain, which is used in the fusion proteins of the invention, i.e., the fibrous protein domain sequence indicated by (SEQ ID NO: 4), referred to herein as 15mer.

In one embodiment, a fibrous domain sequence of (SEQ ID NO: 1) is repeated 15 times throughout the fibrous protein domain that is used in the fusion proteins of the invention and has a CRGD (SEQ ID NO: 2) linker sequence, i.e. the fibrous protein domain sequence indicated by (SEQ ID NO: 5), referred to herein as CRGD-15mer.

In one embodiment, the fusion polypeptide of the invention has a mineralizing domain that is capable of inducing the formation of hydroxyapatite, silica, cadmium sulfide or magnetite.

In one embodiment, the mineralization domain is obtained from dentin matrix protein 1 (DMP1), bone sialoprotein (BSP), or silaffin-1 (Sil1) protein.

In one embodiment, the mineralization domain is derived from dentin matrix protein 1 (DMP1) and is (SEQ ID NO: 6) or (SEQ ID NO: 7).

In one embodiment, the mineralization domain is derived from bone sialoprotein (BSP) and is (SEQ ID NO: 8).

In one embodiment, the mineralizing domain is selected from the group consisting of the 19 amino-acid R5 peptide of the Sil1 protein (SEQ ID NO: 17), the R2 peptide of the Sil1 protein (SEQ ID NO: 18), the 19 amino-acid R3 peptide of the Sil1 protein (SEQ ID NO: 19), the 19 amino-acid R6 peptide of the Sil1 protein (SEQ ID NO: 20) and the 15 amino-acid R1 peptide of the Sil1 protein (SEQ ID NO: 21).

The invention also provides for the following fusion polypeptides that comprise a fibrous protein domain and a mineralization domain: the fusion polypeptide (SEQ ID NO: 9), the fusion polypeptide (SEQ ID NO: 10), the fusion polypeptide (SEQ ID NO: 11), the fusion polypeptide (SEQ ID NO: 12), a fusion polypeptide comprising a fusion of the 15 mer silk fibrous protein domain (SEQ ID NO: 4) and the R5 peptide of the Sil1 protein (SEQ ID NO: 17) and a fusion polypeptide comprising a fusion of the CRGD-15 mer silk fibrous protein domain (SEQ ID NO: 5) and the R5 peptide of the Sil1 protein (SEQ ID NO: 17).

A method for forming a fibrous protein inorganic-composite material is also provided. The method comprises (a) contacting the fusion proteins of the invention with an inorganic material capable of mineralizing for a sufficient period of time to allow mineralization of the inorganic material.

In one preferred embodiment, the fusion proteins of the invention are formed into a silk film, foam or sponge prior to deposition of inorganic material (mineralization).

In one embodiment, the inorganic material is capable of forming hydroxyapatite or silica.

In one embodiment, the fiber, film, or sponge further comprises an agent.

In one embodiment, the inorganic coating formed on the fibrous protein comprises an agent.

In one embodiment, the agent is selected from the group consisting of a protein, peptide, nucleic acid, PNA, aptamer, antibody or a small molecule.

The claimed invention also provides for biomaterial products produced by the methods of the invention.

BRIEF DESCRIPTION OF FIGURES

FIGS. 1A and 1B show diagrammatic illustrations of fibrous protein domain-mineralization domain fusion pro-

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teins. FIG. 1A shows three claimed fusion proteins composed of a spider silk fibrous protein domain ((CRGD-15mer) (See (SEQ ID NO: 5)) and a dentin matrix protein 1 (DMP1) mineralization domain. The fusion proteins contain a mineralization domain of either 1) the full length amino acid sequence of DMP1 (See (SEQ ID NO: 7)), 2) the C-terminal end of DMP1, (See (SEQ ID NO: 6)) or a composite sequence based on DMP1, which contains the collagen binding domain 1 (CD1) of DMP1, two acidic clusters in DMP1 that are responsible for hydroxyapatite nucleation (pA and pB) and the collagen binding domain 2 (CD2) of DMP1. FIG. 1B shows a diagram of two fusion proteins, 1) fusion of a spider silk fibrous protein domain (CRGD-15mer (See (SEQ ID NO: 5)) and a bone sialoprotein mineralizing domain (BSP) (See (SEQ ID NO: 8)) and 2) fusion of a spider silk fibrous protein domain 15-mer (without CRGD linker) (See (SEQ ID NO: 4)) and a bone sialoprotein mineralizing domain (BSP) (See (SEQ ID NO: 8)).

FIG. 2 shows the sequence of the fusion protein CRGD-15mer-CDMP1 (SEQ ID NO: 9). CRGD-15mer-CDMP1 is a fusion protein of a spider silk fibrous protein domain ((CRGD-15mer) (See, SEQ ID NO: 5), which is indicated by a dark grey highlight, and the C-terminal end of DMP1 mineralizing domain (See, SEQ ID NO: 6), which is indicated with light grey highlight.

FIG. 3 shows the sequence of the fusion protein CRGD-15mer-DMP1 (SEQ ID NO: 10). CRGD-15mer-DMP1 is a fusion protein of spider silk fibrous protein domain ((CRGD-15mer) (See, SEQ ID NO: 5), which is indicated by a dark grey highlight, and the full length sequence of DMP1 mineralizing domain (See, SEQ ID NO: 7), which is indicated with light grey highlight.

FIG. 4 shows the sequence of the fusion protein 15mer-BSP (SEQ ID NO: 11). 15mer-BSP is a fusion protein of spider silk fibrous protein domain ((15mer) (See, SEQ ID NO: 4), which is indicated by light grey highlight, and a mineralizing domain of bone sialoprotein (BSP) (SEQ ID NO: 8), which is indicated with dark grey highlight.

FIG. 5 shows the sequence of the fusion protein CRGD-15mer-BSP (SEQ ID NO: 12). CRGD-15mer-BSP a fusion protein of spider silk fibrous protein domain ((CRGD-15mer) (See, SEQ ID NO: 5), which is indicated by light grey highlight, and a mineralizing domain of bone sialoprotein (BSP) (SEQ ID NO: 8), which is indicated with dark grey highlight. A linker sequence between the two domains is also present.

FIGS. 6A to 6C show scanning electron microscope (SEM) images of recombinant silk film made from CRGD-15mer-CDMP1 (SEQ ID NO: 9), See Example 1. FIG. 6A, image of silk film morphology prior to mineralization (HFIP+MeOH). FIG. 6B, image of silk film morphology after one round of mineralization. FIG. 6C, image of silk film morphology after three rounds of mineralization.

FIGS. 7A to 7B show the sequence of the fusion proteins CRGD15mer-R5 (SEQ ID NO: 25) and 15mer-R5 (SEQ ID NO: 25) as described in Example II. The underlined sequence represents the monomeric repeat unit selected and used in the design of the recombinant proteins based on the consensus sequence of spidroin1 (Masp1) native sequence of *Nephila clavipes* (Accession #P19837).

FIG. 8 shows a grid of SEM of untreated and methanol treated silk films formed from the four different genetically engineered silk proteins CRGD15mer (SEQ ID NO: 5), 15mer (SEQ ID NO: 4), CRGD15mer-R5 (SEQ ID NO: 24), and 15mer-R5 (SEQ ID NO: 25) as described in Example 2. Images of the control films and the films that underwent silicification reactions are shown.

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FIGS. 9A to 9E show SEM images of untreated and methanol treated electropun CRGD15mer-R5 (SEQ ID NO: 24) silk fibers before, during and after silicification reactions, as described in Example 2. FIG. 9A, untreated electropun CRGD15mer-R5 (SEQ ID NO: 24) silk fibers. FIG. 9B, electropun CRGD15mer-R5 (SEQ ID NO: 24) silk fibers methanol treated before silification. FIG. 9C, 9D and 9E, electropun CRGD15mer-R5 (SEQ ID NO: 24) silk fibers during silification at 20 μ m and 2 μ m scale.

FIGS. 10A to 10B show XPS analysis, as described in Example 2, of CRGD15mer-R5 (SEQ ID NO: 24) and silicified CRGD15mer-R5 (SEQ ID NO: 24) on Al foil and on silicon chip at the characteristic binding energies of (FIG. 10A) 153 eV and (FIG. 10B) 102 eV for electrons found in the 2s and 2p₃ electron shells of the silicon atom respectively.

FIG. 11 shows Fourier transform infrared spectroscopy (FTIR) analysis of the structure of CRGD-15mer-CDMP1 (SEQ ID NO: 9) before and after Ca ion binding, as described in Example 1.

FIG. 12 shows FTIR analysis of structure of CRGD-15mer-CDMP1 (SEQ ID NO: 9) before and after treatment with methanol, as described in Example 1.

FIGS. 13(A1) to 13(A5) and 13(B1) to 13(B5) show Scanning Electron Microscopy (SEM) surface morphologies of recombinant spider silk films after soaking in 1.5 \times SBF for various periods of time, as described in Example 1. FIG. 13(A1)-FIG. 13(A5), SS15m-CDMP1 soaked in 1.5 \times SBF for 0, 3, 7, 14 and 21 days: FIG. 13(A1)-FIG. 13(A5) respectively. FIG. 13(B1)-FIG. 13(B5), SS15m soaked in 1.5 \times SBF for 0, 3, 7, 14 and 21 days: FIG. 13(B1)-FIG. 13(B5) respectively. Scale bars are 2 μ m.

FIGS. 14A to 14C show Transmission Electron Microscopy (TEM) images of crystals grown on CRGD-15mer-CDMP1 (SEQ ID NO: 9) after soaking in 1.5 \times SBF for 21 days, as described in Example 1. (FIG. 14A) image of flake-like apatite; (FIG. 14B) electron diffraction pattern; (FIG. 14C) high-resolution image of apatite crystal.

DESCRIPTION OF THE INVENTION

Fibrous proteins represent an important category of proteins in biology, forming native structural entities both internally in organisms (e.g., collagens) and externally (such as spun silk fibers for webs and cocoons). As an example, silk proteins from spiders and insects provide a number of valuable materials features. When combined with functional domains, then the materials properties available from silks become significantly expanded. In accordance with the present invention, various domains from biology known to promote the formation of inorganic structures, e.g., promote formation of minerals, have been combined with a fibrous protein domain. Thus, these new fusion (composite) materials offer the benefits of the fibrous protein material features (e.g., silks) plus the benefit of the inorganic mineralization domains (e.g., hardness).

In one embodiment, the present invention provides a fusion polypeptide comprising a fibrous protein domain and a mineralization domain. The fusion is used to form an organic-inorganic composite. These organic-inorganic composites can be constructed from the nano- to the macro-scale depending on the size of the fibrous protein fusion domain used. In one embodiment, the composites can also be loaded with other compounds (e.g., dyes, drugs, enzymes) depending on the goal for the materials, to further enhance function. This can be achieved during assembly of the material or during the mineralization step in materials formation.

The fusion polypeptides of the present invention can be used for production of silk biomaterials, e.g., fibers, films, foams and mats. See, WO 03/022909. An all-aqueous process may be used. See, WO 03/022909.

In another embodiment, the invention provides a method for forming a composite material. The method comprises contacting a template formed from a fusion polypeptide of the invention with a suitable inorganic material for a sufficient period of time to allow mineralization of the inorganic material thus forming an inorganic coating on the template.

In one embodiment the template is in the form of a fiber, film, or sponge.

In an additional embodiment, growth factors, biological components or others agents, including therapeutic agents, are incorporated into the inorganic coating. The growth factors, biological components or other agents, can be incorporated during the formation of the template or during the crystallization process. Such composites can be used to deliver such components to cells and tissues. The ability to incorporate growth factors, biological regulators, enzymes, therapeutic agents, or cells in the construct of the present invention provides for stabilization of these components for long term release and stability, as well as better control of activity and release.

The products produced by these methods offer new options in the formation of scaffolds for biomaterials, tissue engineering applications and drug delivery. While the templates are useful in and of themselves, the ability to form inorganic coatings with controlled thickness leads to control of mechanical properties (e.g., stiffness) and biological interactions, such as for bone formation. Furthermore, the ability to control these processes allows one to match structural and functional performance of scaffolds for specific tissue targets and needs.

In another embodiment, the underlying template (e.g., fusion polypeptide formed into a film etc.) can be removed or etched away to generate porous networks, tubes, or lamellar sheets of inorganic material. These materials are useful directly as biomaterial scaffolds, for control of cell and tissue growth (e.g., as nerve conduits, bone conduits) and for non-biological applications (e.g., filtration and separation media, catalysis, decontamination (directly or if filled with appropriate chemical or enzymes), radar chaff, coatings in general, and many related needs, for example, inorganic fillers to toughen materials that can also be filled with a second component).

The fusion polypeptides may be created, for example, by chemical synthesis, or by creating and translation a polynucleotide in which the peptide regions are encoded in the desired relationship.

It is to be understood that while the invention has been described in conjunction with the preferred specific embodiments thereof that the foregoing description as well as the examples that follow are intended to illustrate and not limit the scope of the invention. Other aspects, advantages and modifications within the scope of the invention will be apparent to those skilled in the art to which the invention pertains.

Fibrous Protein Domains

Silks are unique in their self-assembly, formation of robust materials in the form of fibers, films and 3D matrices, and are polymorphic (the structure can be controlled among many assembly states). For example, silk fibers are the strongest known fibers and rival even high performance fibers. They are also effective in resisting compression. In 3D porous matrices the compressive resistance exceeds other commonly used organic polymers as biomaterial matrices. Inorganic domains, including silica and hydroxyapatite are prominent in biological systems as key inorganic components found

associated with proteins. These mineral phases dominate skeletal components of most systems.

The fibrous protein domain can include short to long versions, with the size in part influencing the scale of the composite from the nano to the macro level. For example, the size can range from a single sized hydrophobic block of 50 amino acids, up to multiple blocks as large as desired including up to proteins of sizes in the 100,000s of Daltons. In addition, the fibrous protein fusion domain could include other proteins such as collagens, coiled-coiled leucine zipper proteins, elastins, keratins, actins, and tubulins.

The silk protein suitable for use in the present invention is preferably fibroin or related proteins (i.e., silks from spiders). The silkworm silk is obtained, for example, from *Bombyx mori*. Spider silk may be obtained from *Nephila clavipes* See, for example, WO 97/08315 and U.S. Pat. No. 5,245,012.

Inorganic Domains/Mineralizing Domains

Mineralizing domains include those that can induce the formation of hydroxyapatite (Example 1), silica (Example 2), cadmium sulfide, and magnetite. In one embodiment, a mineralizing domain that can induce hydroxyapatite nucleation is obtained from dentin matrix protein 1 or bone sialoprotein (G. He, T. Dahl, A. Veis and A. George. Connective Tissue Res. 2003, 44 (Suppl. 1): 240-245; and G. He, T. Dahl, A. Veis and A. George. Nature materials. 2003, 2(8): 552-558; Stubbs et al. J Bone Miner Res. 1997 August; 12(8):1210-22). The full molecule or a minimum portion that can induce mineralization can be used, for example, a 37 amino acid segment from DMP (C-terminus) will induce controlled mineralization of hydroxyapatite. In one example, the mineralization domains were fused to a sequence derived from spider silk having the following silk fibrous domain repeated 15 times (SGRG-GLGGQGAGAAAAAGGAGQGGYGGLGSQGT) (SEQ ID NO: 1) in order to generate functional recombinant silks that possess both robust mechanical properties of the spider silk and the ability to promote mineralization. The recombinant proteins were cloned in pET21a(+) vector and expressed in *E. coli*. As demonstrated below, the identities of these recombinant silks were confirmed by amino acid composition analysis and the mineralization study was carried out on the surfaces of the cast functional recombinant silk films using recombinant protein containing only spider silk sequence as control. It was demonstrated that the functional recombinant silks exhibit an ability to promote the nucleation of hydroxyapatite. The functional recombinant silks (silk fusion proteins) of the present invention have potential application in biomaterials, tissue engineering, advanced material composites and biosensors.

Preferably, the mineralizing domain is capable of inducing the formation of hydroxyapatite, silica, cadmium sulfide, magnetite and other metal salts pending choice of peptide domain utilized.

Preferred mineralizing domains include, for example, dentin matrix protein 1 (DMP1), bone sialoprotein, and fragments of these proteins, and the 19 amino-acid R5 peptide of the Sil1 protein.

Alternative fibrous proteins and mineralizing domains can also be included in this system using the template provided. For example, peptides identified from other native proteins or identified by combinatorial screening methods can be used. Screening methods can involve any mineralization assay known in the art, for example the hydroxyapatite and silicification assays described herein.

Preparation of Fusion Proteins; Vectors, Host Cells and Expression

The fusion proteins containing a mineralization domain and a fibrous protein domain can be made using standard

molecular biology methods well known to those skilled in the art (See for example, Sambrook et al., *Molecular Biology: A laboratory Approach*, Cold Spring Harbor, N.Y. 1989; Ausubel, et al., *Current protocols in Molecular Biology*, Greene Publishing, Y, 1995).

In some embodiments, the fusion proteins of the invention are made using linker sequences. As used herein, the term "linker sequence" refers to a short (e.g., about 1-20 amino acids) sequence of amino acids that is not part of the sequence of either of two polypeptides being joined. For example, a linker sequence is attached on its amino-terminal end to one polypeptide or polypeptide domain and on its carboxyl-terminal end to another polypeptide or polypeptide domain.

The manipulation of nucleic acids that encode the protein domains in the present invention is typically carried out in recombinant vectors. Herein, both phagemid and non-phagemid vectors can be used. As used herein, vector refers to a discrete element that is used to introduce heterologous DNA into cells for the expression and/or replication thereof. Methods by which to select or construct and, subsequently, use such vectors are well known to one of skill in the art. Numerous vectors are publicly available, including bacterial plasmids, bacteriophage, artificial chromosomes, episomal vectors and gene expression vectors. A vector of use according to the invention may be selected to accommodate a polypeptide coding sequence of a desired size. A suitable host cell is transformed with the vector after in vitro cloning manipulations. Host cells may be prokaryotic, such as any of a number of bacterial strains, or may be eukaryotic, such as yeast or other fungal cells, insect or amphibian cells, or mammalian cells including, for example, rodent, simian or human cells. Each vector contains various functional components, which generally include a cloning (or "polylinker") site, an origin of replication and at least one selectable marker gene. If given vector is an expression vector, it additionally possesses one or more of the following: enhancer element, promoter, transcription termination and signal sequences, each positioned in the vicinity of the cloning site, such that they are operatively linked to the gene encoding a polypeptide repertoire member according to the invention.

Both cloning and expression vectors generally contain nucleic acid sequences that enable the vector to replicate in one or more selected host cells. Typically in cloning vectors, this sequence is one that enables the vector to replicate independently of the host chromosomal DNA and includes origins of replication or autonomously replicating sequences. Such sequences are well known for a variety of bacteria, yeast and viruses. For example, the origin of replication from the plasmid pBR322 is suitable for most Gram-negative bacteria, the 2 micron plasmid origin is suitable for yeast, and various viral origins (e.g. SV 40, adenovirus) are useful for cloning vectors in mammalian cells. Generally, the origin of replication is not needed for mammalian expression vectors unless these are used in mammalian cells able to replicate high levels of DNA, such as COS cells.

Advantageously, a cloning or expression vector may contain a selection gene also referred to as a selectable marker. This gene encodes a protein necessary for the survival or growth of transformed host cells grown in a selective culture medium. Host cells not transformed with the vector containing the selection gene will therefore not survive in the culture medium. Typical selection genes encode proteins that confer resistance to antibiotics and other toxins, e.g. ampicillin, neomycin, methotrexate or tetracycline, complement auxotrophic deficiencies, or supply critical nutrients not available in the growth media.

Since the replication of vectors according to the present invention is most conveniently performed in *E. coli* (e.g., strain TB1 or TG1), an *E. coli*-selectable marker, for example, the β -lactamase gene that confers resistance to the antibiotic ampicillin, is of use. These can be obtained from *E. coli* plasmids, such as pBR322 or a pUC plasmid such as pUC18 or pUC19, or pUC119.

Expression vectors usually contain a promoter that is recognized by the host organism and is operably linked to the coding sequence of interest. Such a promoter may be inducible or constitutive. The term "operably linked" refers to a juxtaposition wherein the components described are in a relationship permitting them to function in their intended manner. A control sequence "operably linked" to a coding sequence is ligated in such a way that expression of the coding sequence is achieved under conditions compatible with the control sequences.

Promoters suitable for use with prokaryotic hosts include, for example, the β -lactamase and lactose promoter systems, alkaline phosphatase, the tryptophan (trp) promoter system and hybrid promoters such as the tac promoter. Promoters for use in bacterial systems will also generally contain a Shine-Delgarno sequence operably linked to the coding sequence. Preferred promoters for use in the present invention are the isopropylthiogalactoside (IPTG)-regulatable promoters.

In one preferred embodiment of the invention, the fusion protein of the invention further comprises a tag, e.g. Flag, His, Myc, HA, VSV, or V5, to aid in purification of the protein by standard means. After purification, the fusion protein can be lyophilized or suspended in aqueous solution for preparation of silk materials, such as films, foams or fibers. Formation of Silk Fibers, Films, Foams and Gels Using the Silk Fusion Protein of the Invention.

The silk fusion proteins of the invention can be processed into films, foams or fibers. As used herein, "silk fibroin" or "silk protein" refers to a silk fusion protein of the invention.

Fibers can be formed by electrospinning. Electrospinning can be performed by any means known in the art (see, for example, U.S. Pat. No. 6,110,590). Preferably, a steel capillary tube with a 1.0 mm internal diameter tip is mounted on an adjustable, electrically insulated stand. Preferably, the capillary tube is maintained at a high electric potential and mounted in the parallel plate geometry. The capillary tube is preferably connected to a syringe filled with silk/biocompatible polymer solution. Preferably, a constant volume flow rate is maintained using a syringe pump, set to keep the solution at the tip of the tube without dripping. The electric potential, solution flow rate, and the distance between the capillary tip and the collection screen are adjusted so that a stable jet is obtained. Dry or wet fibers are collected by varying the distance between the capillary tip and the collection screen.

A collection screen suitable for collecting silk fibers can be a wire mesh, a polymeric mesh, or a water bath. Alternatively and preferably, the collection screen is an aluminum foil. The aluminum foil can be coated with Teflon fluid to make peeling off the silk fibers easier. One skilled in the art will be able to readily select other means of collecting the fiber solution as it travels through the electric field. As is described in more detail in the Examples section below, the electric potential difference between the capillary tip and the aluminum foil counter electrode is, preferably, gradually increased to about 12 kV, however, one skilled in the art should be able to adjust the electric potential to achieve suitable jet stream.

The process of the present invention may further comprise steps of immersing the spun fiber into an alcohol/water solution to induce crystallization of silk. The composition of alcohol/water solution is preferably 90/10 (v/v). The alcohol

is preferably methanol, ethanol, isopropyl alcohol (2-propanol) or n-butanol. Methanol is most preferred. Additionally, the process may further comprise the step of washing the fibroin fiber in water.

In another embodiment, the biomaterial is a film. The process for forming the film comprises, for example, the steps of (a) preparing an aqueous silk fibroin solution comprising silk protein; (b) adding a biocompatible polymer to the aqueous solution; (c) drying the mixture; and (d) contacting the dried mixture with an alcohol (preferred alcohols are listed above) and water solution to crystallize a silk blend film. Preferably, the biocompatible polymer is poly(ethylene oxide) (PEO). The process for producing the film may further include step (e) of drawing or mono-axially stretching the resulting silk blend film to alter or enhance its mechanical properties. The stretching of a silk blend film induces molecular alignment in the fiber structure of the film and thereby improves the mechanical properties of the film.

In a further embodiment, the biomaterial is a foam. Foams may be made from methods known in the art, including, for example, freeze-drying and gas foaming in which water is the solvent or nitrogen or other gas is the blowing agent, respectively.

In one embodiment, the foam is a micropatterned foam. Micropatterned foams can be prepared using, for example, the method set forth in U.S. Pat. No. 6,423,252, the disclosure of which is incorporated herein by reference.

Formation of Organic-Inorganic Composites

The claimed invention provides a method for forming an organic-inorganic composite material. The method comprises contacting a template (such as recombinant silk films, sponges or fibers) formed from a fusion polypeptide of the invention with a suitable inorganic material for a sufficient period of time to allow mineralization of the inorganic material. Mineralized material is deposited onto the silk forming an inorganic coating on the template.

The procedure for mineralization of the coating is dependent upon the mineralization domain that is part of the fusion polypeptide of the invention. For example, the mineralizing domain of dentin matrix protein 1 (DMP1), an acidic phosphoprotein secreted into the extracellular matrix during the formation and mineralization of bone and dentin, is involved in the precipitation of hydroxyapatite, the main inorganic component in calcified hard tissue such as bone and teeth of vertebrates (Koutsopoulos, S. *J Biomed Mater Res.* 2002, 62: 600-612). Further, the mineralizing domain of Silaffin protein is involved in the precipitation of silica.

Those in the art are skilled to select the appropriate buffer for precipitation of inorganic material. The formation of inorganic coatings is further described in Examples 1 and 2.

The biomaterial products produced by the processes of the present invention may be used in a variety of medical applications such as wound closure systems, including vascular wound repair devices, hemostatic dressings, patches and glues, sutures, drug delivery and in tissue engineering applications, such as, for example, scaffolding, ligament prosthetic devices and in products for long-term or bio-degradable implantation into the human body. A preferred tissue engineered scaffold is a non-woven network of electrospun fibers.

Additionally, these biomaterials can be used for organ repair replacement or regeneration strategies that may benefit from these unique scaffolds, including but are not limited to, spine disc, cranial tissue, dura, nerve tissue, liver, pancreas, kidney, bladder, spleen, cardiac muscle, skeletal muscle, tendons, ligaments and breast tissues.

In another embodiment of the present invention, silk biomaterials can contain therapeutic agents. To form these mate-

rials, the polymer would be mixed with a therapeutic agent prior to forming the material or loaded into the material after it is formed. In addition, the agent can be incorporated into the inorganic coating formed by the silk fusion protein mineralization domain by mixing it with the mineralization solution. The variety of different therapeutic agents that can be used in conjunction with the biomaterials of the present invention is vast. In general, therapeutic agents which may be administered via the pharmaceutical compositions of the invention include, without limitation: anti-infectives such as antibiotics and antiviral agents; chemotherapeutic agents (i.e. anticancer agents); anti-rejection agents; analgesics and analgesic combinations; anti-inflammatory agents; hormones such as steroids; growth factors (bone morphogenic proteins (i.e. BMP's 1-7), bone morphogenic-like proteins (i.e. GFD-5, GFD-7 and GFD-8), epidermal growth factor (EGF), fibroblast growth factor (i.e. FGF 1-9), platelet derived growth factor (PDGF), insulin like growth factor (IGF-I and IGF-II), transforming growth factors (i.e. TGF-beta.1-III), vascular endothelial growth factor (VEGF)); and other naturally derived or genetically engineered proteins, polysaccharides, glycoproteins, or lipoproteins. These growth factors are described in *The Cellular and Molecular Basis of Bone Formation and Repair* by Vicki Rosen and R. Scott Thies, published by R. G. Landes Company hereby incorporated herein by reference.

Silk biomaterials containing bioactive materials may be formulated by mixing one or more therapeutic agents with the polymer used to make the material. Alternatively, a therapeutic agent could be coated on to the material preferably with a pharmaceutically acceptable carrier. Any pharmaceutical carrier can be used that does not dissolve the foam. The therapeutic agents, may be present as a liquid, a finely divided solid, or any other appropriate physical form. Typically, but optionally, the matrix will include one or more additives, such as diluents, carriers, excipients, stabilizers or the like.

The amount of therapeutic agent will depend on the particular drug being employed and medical condition being treated. Typically, the amount of drug represents about 0.001 percent to about 70 percent, more typically about 0.001 percent to about 50 percent, most typically about 0.001 percent to about 20 percent by weight of the material. Upon contact with body fluids the drug will be released.

The biocompatible polymer may be extracted from the biomaterial prior to use. This is particularly desirable for tissue engineering applications. Extraction of the biocompatible polymer may be accomplished, for example, by soaking the biomaterial in water prior to use.

The tissue engineering scaffolds biomaterials can be further modified after fabrication. For example, the scaffolds can be coated with bioactive substances that function as receptors or chemoattractors for a desired population of cells. The coating can be applied through absorption or chemical bonding.

Additives suitable for use with the present invention include biologically or pharmaceutically active compounds. Examples of biologically active compounds include cell attachment mediators, such as the peptide containing variations of the "RGD" integrin binding sequence known to affect cellular attachment, biologically active ligands, and substances that enhance or exclude particular varieties of cellular or tissue ingrowth. Such substances include, for example, osteoinductive substances, such as bone morphogenic proteins (BMP), epidermal growth factor (EGF), fibroblast growth factor (FGF), platelet-derived growth factor (PDGF), vascular endothelial growth factor (VEGF), insulin-like growth factor (IGF-I and II), TGF- and the like.

The scaffolds are shaped into articles for tissue engineering and tissue guided regeneration applications, including reconstructive surgery. The structure of the scaffold allows generous cellular ingrowth, eliminating the need for cellular pre-seeding. The scaffolds may also be molded to form external scaffolding for the support of in vitro culturing of cells for the creation of external support organs.

The scaffold functions to mimic the extracellular matrices (ECM) of the body. The scaffold serves as both a physical support and an adhesive substrate for isolated cells during in vitro culture and subsequent implantation. As the transplanted cell populations grow and the cells function normally, they begin to secrete their own ECM support.

In the reconstruction of structural tissues like cartilage and bone, tissue shape is integral to function, requiring the molding of the scaffold into articles of varying thickness and shape. Any crevices, apertures or refinements desired in the three-dimensional structure can be created by removing portions of the matrix with scissors, a scalpel, a laser beam or any other cutting instrument. Scaffold applications include the regeneration of tissues such as nervous, musculoskeletal, cartilaginous, tendinous, hepatic, pancreatic, ocular, integumentary, arteriovenous, urinary or any other tissue forming solid or hollow organs.

The scaffold may also be used in transplantation as a matrix for dissociated cells, e.g., chondrocytes or hepatocytes, to create a three-dimensional tissue or organ. Any type of cell can be added to the scaffold for culturing and possible implantation, including cells of the muscular and skeletal systems, such as chondrocytes, fibroblasts, muscle cells and osteocytes, parenchymal cells such as hepatocytes, pancreatic cells (including Islet cells), cells of intestinal origin, and other cells such as nerve cells, bone marrow cells, skin cells and stem cells, and combination thereof, either as obtained from donors, from established cell culture lines, or even before or after genetic engineering. Pieces of tissue can also be used, which may provide a number of different cell types in the same structure.

The cells are obtained from a suitable donor, or the patient into which they are to be implanted, dissociated using standard techniques and seeded onto and into the scaffold. In vitro culturing optionally may be performed prior to implantation. Alternatively, the scaffold is implanted, allowed to vascularize, then cells are injected into the scaffold. Methods and reagents for culturing cells in vitro and implantation of a tissue scaffold are known to those skilled in the art.

The biomaterials of the claimed invention may be sterilized using conventional sterilization process such as radiation based sterilization (i.e. gamma-ray), chemical based sterilization (ethylene oxide) or other appropriate procedures. Preferably the sterilization process will be with ethylene oxide at a temperature between 52-55° C. for a time of 8 hours or less. After sterilization the biomaterials may be packaged in an appropriate sterilize moisture resistant package for shipment and use in hospitals and other health care facilities.

The invention will be further characterized by the following examples which are intended to be exemplary of the invention.

EXAMPLES

Example 1

Formation of Hydroxyapatite Silk Fusions

Hydroxyapatite (HAP, $\text{Ca}_5(\text{PO}_4)_3(\text{OH})$) is the most stable calcium phosphate salt at normal temperature and pH between 4 and 12 (Aaron, S. Posner *Physiological review*,

1969, 49(4): 760-792). Hydroxyapatite is the main inorganic component in calcified hard tissue such as bone and teeth of vertebrates (Koutsopoulos, S. *J Biomed Mater Res.* 2002, 62: 600-612). In addition, hydroxyapatite also has many applications in protein chromatography, water treatment processes, fertilizer and pharmaceutical products (Bailliez, S.; Nzihou, A.; Beche, E.; Flamant, G. *Process Safety and Environmental Protection*, 2004, 82(B2): 175-180). For calcified hard tissue, hydroxyapatite contributes to its stiffness, while organic matrix contributes to its plasticity (Landis, W. J. *Bone*, 16(5): 533-544). The inherent strength and other mechanical properties of the skeletal system are thought to depend on an interaction between its organic and inorganic matrix strength, here hydroxyapatite. One example is the composite formed between the normal mineral salt, hydroxyapatite, and collagen, the principal organic component of the vertebrate skeleton, able to resist a wide range of compressive or tensile forces whereas either material alone can not (Chen, Q. Z.; Wong, C. T.; Lu, W. W.; Cheung, K. M. C.; Leong, J. C. Y. and Luk, K. D. K. *Biomaterials*, 2004, 25: 4243-4254). In the present disclosure, we demonstrate a proof of concept with dental matrix protein (DMP).

Dentin matrix protein 1 (DMP1) is an acidic phosphoprotein secreted into the extracellular matrix during the formation and mineralization of bone and dentin, therefore, it is believed that DMP1 plays an important role in the initiation of mineralization (J. Q. Feng, H. Huang, Y. Lu, L. Ye, Y. Xie, T. w. Tsutsui, T. Kunieda, T. Castranio, G. Scott, L. B. Bonewald, and Y. Mishina. *J Dent Res.* 2003, 82(10): 776-780; W. T. Butler, H. Ritchie. *Int J Dev Biol*, 1995, 39: 169-179; George, B. Sabsay, P. A. L. Simonian, and A. Veis. *J Biol Chem*, 1993, 268(17):12624-12630). It has been reported that DMP1 nucleates the formation of hydroxyapatite in vitro (G. He, T. Dahl, A. Veis and A. George. *Connective Tissue Res.* 2003, 44 (Suppl. 1): 240-245; and G. He, T. Dahl, A. Veis and A. George. *Nature materials*. 2003, 2(8): 552-558) by binding calcium ions and initiating mineral deposition. It has also been reported that DMP1 binds to the N-telopeptide region of type I collagen and the collagen-binding domains involved in this interaction have been identified (G. He and A. George. *J Biol. Chem.* 2004, 279(12): 11649-11656).

We generated various fibrous protein domain-mineralizing domain fusion proteins. See for example FIGS. 1-5. FIG. 2 shows the sequence of CRGD-15mer-CDMP1 (SEQ ID NO: 9), a fusion protein of a spider silk fibrous protein domain ((CRGD-15mer) (See, SEQ ID NO: 5), which is indicated by a dark grey highlight, and the C-terminal end of DMP1 mineralizing domain (See, SEQ ID NO: 6), which is indicated with light grey highlight.

FIG. 3 shows the sequence of the fusion protein CRGD-15mer-DMP1 (SEQ ID NO: 10). CRGD-15mer-DMP1 (SEQ ID NO: 10) is a fusion protein of spider silk fibrous protein domain ((CRGD-15mer) (See, SEQ ID NO: 5), which is indicated by a dark grey highlight, and the full length sequence of DMP1 mineralizing domain (See, SEQ ID NO: 7), which is indicated with light grey highlight.

FIG. 4 shows the sequence of the fusion protein 15mer-BSP (SEQ ID NO: 11). 15mer-BSP (SEQ ID NO: 11) is a fusion protein of spider silk fibrous protein domain ((15mer) (See, SEQ ID NO: 4), which is indicated by light grey highlight, and a mineralizing domain of bone sialoprotein (BSP) (SEQ ID NO: 8), which is indicated with dark grey highlight.

FIG. 5 shows the sequence of the fusion protein CRGD-15mer-BSP (SEQ ID NO: 12). CRGD-15mer-BSP (SEQ ID NO: 12) a fusion protein of spider silk fibrous protein domain ((CRGD-15mer) (See, SEQ ID NO: 5), which is indicated by

light grey highlight, and a mineralizing domain of bone sialoprotein (BSP) (SEQ ID NO: 8), which is indicated with dark grey highlight.

Construction of Expression Vector for Recombinant Spider Silks: The consensus repeat unit of spider silk (-SGRG-GLGGQGAGAAAAAGGAGQGGY GGLGSQGT-) (SEQ ID NO: 1) was derived from the native sequence of the spidroin 1 of *N. clavipes* (accession P19837). The construction of the expression vector pET30a (+) containing 15 repeats of the silk was previously described (Bini et al., 2005) and was termed pET30a (+)-15mer. Plasmid containing the rat dentin matrix protein 1 cDNA (pGEX-DMP1) was previously described (Feng et al., J. Dent. Res. 2003, 82(10):776-780). Primers with BamHI and XhoI restriction enzyme sites were designed in order to copy the C-terminal portion of rat DMP1 from pGEX-DMP1 by PCR: BamHI site, C-DMP1f: 5'-CAGGATCCAGGGGTGACAACCCAGAT-3' (SEQ ID NO: 26) and Xho I site, C-DMP1r: 5'-GCC TCGAGGTAGCCATCTTGCAATC-3' (SEQ ID NO: 27).

PCR products were double digested with BamHI and XhoI before running on a 1% agarose gel. The band of C-terminal DMP1 DNA were cut from gel and purified using a MinElute gel extraction kit. Expression vector pET21a (+) was also double digested with BamHI and XhoI before running on a 0.8% agarose gel. The band of linearized pET21a (+) was purified using a QIAquick gel extraction kit. After determining the DNA concentration of the vector pET21a (+) and insert of C-terminal DMP1 based on intensities of 1 µl of each in a 1% gel against 1 µl of high mass DNA ladder, the ligation reactions were performed using a Quick ligation Kit™ based on an insert to vector molar ratio of 5:1. Fifty µl of one Shot® Mach1™ T1 phage-resistant cells were transformed with 1 µl of ligation reaction. The correct clone of vector pET21a (+) containing C-terminal DMP1, named as pET21a (+)-CDMP1 was determined by DNA sequencing with forward and reverse T7 primers. pET21a (+)-CDMP1 was digested with BamHI and dephosphorylated with CIP before running on 0.8% agarose gel. The linearized vector was purified using a QIAquick gel extraction kit. pET30a (+)-15mer was digested with BamHI before running on 1% agarose gel and the DNA encoding the repeat unit of spider silk was purified using a MinElute gel extraction kit. The DNA concentrations of the vector pET21a (+)-CDMP1 and the insert: spider silk 15 mer were determined using the same method as described above. The spider silk 15 mer was then inserted into pET21a (+)-CDMP1 at the BamHI site by ligation reaction and the resulting vector was named as pET21a (+)-5515m-CDMP1. DNA sequencing with forward and reverse T7 primers was performed to check the sequences of the vector.

Protein Expression and Purification: Expression plasmid pET21a (+)-SS15m-CDMP1 was transformed into *Escherichia coli* RY-3041 strain, kindly provided by Professor Ry Young (Texas A&M University, College Station, Tex.). High cell density cultivation using a 1.25 liter jar fermentor (Bioflo 3000; New Brunswick Scientific Co., Edison, N.J.) was performed as described previously (Butler et al. Int. J. Dev. Biol. 1995, 39: 169-179). Expression was induced by adding isopropyl-β-D-thiogalactoside to a final concentration of 2 mM at the early log phase when A₆₀₀ was about 30. Cells were harvested after 3 h of induction by centrifuging the culture at 4° C., 6000×g, for 10 min. The cells were then sonicated on ice using sonicator equipped with a microtip. Ten second burst at 100 W with a 10 second cooling period between each burst was applied. The lysate was centrifuged at 10,000×g for 30 min at 4° C. to pellet the cellular debris. The recombinant silk protein was purified using Ni-NTA agarose resin from supernatant at 4° C. The purification fractions that contain the

target protein were dialyzed against distilled water for 2 days and dry proteins were prepared by lyophilizing the aqueous solution. Dialyzed purification fractions were also concentrated and desalted by using Centricon plus 70 (NMW=10 kDa).

Protein Characterization: SDS-PAGE was performed to analyze purification fractions and Western blot was performed using His-tag monoclonal antibody to further confirm the expression of the target proteins. Western blot analysis of CRGD-15mer-CDMP1 (SEQ ID NO: 9) showed expression of the protein (data not shown). The target protein bands on 4-12% Bis-Tris PAGE gel were analyzed at the Yale University W.M. Keck Biotechnology Resource Laboratory to determine amino acid compositions. A Bruker Proflex™ mass spectrometer (Bruker, Billerica, Mass.) was used for molecular weight determination.

Change of Protein Secondary Structure by Calcium Ion Binding: 0.2 mg/mL protein aqueous solution was prepared by the method described in protein expression and purification. Calcium chloride solution (1M) was added such that a molar ratio of protein to Ca ion was maintained at 1:1000. Protein secondary structures before and after adding Ca ion were studied by Fourier Transform Infrared Spectroscopy (FTIR). FTIR studies were performed using a Bruker Tensor 27 FTIR spectrometer with a BioATR accessory.

Film Formation: Recombinant proteins were dissolved in hexafluoroisopropanol (HFIP) to a final concentration of 2% w/v. One-hundred µl of silk HFIP solution was loaded onto a silica wafer and dried in hood. Dried silk film was treated with 90% methanol to induce the transition of secondary structure from random coil to beta sheet. Concentrated silk aqueous solution was then loaded onto silk film and incubated at 37° C. overnight to dry.

In vitro Mineralization: The silk films formed on silica wafer were then incubated in 1.5× simulated body fluid (SBF). 1.5×SBF was prepared as described previously (3) by dissolving reagent grade CaCl₂, KH₂PO₄, NaCl, KCl, MgCl₂·6H₂O, NaHCO₃, Na₂SO₄ in distilled water and buffering at pH 7.3 with tris-hydroxymethyl aminomethane and hydrochloric acid (HCl). A 1.5×SBF solution contains 1.5 times higher ion concentration than the SBF solution with ion concentrations close to human blood plasma. Fresh 1.5×SBF was prepared daily to replace the 1.5×SBF. After the silk coated silica wafers were soaked in 1.5×SBF for 3, 7, 14 and 21 days, they were removed, gently rinsed with distilled water and dried at room temperature.

Scanning Electron Microscopy: Morphological investigation of the dried films before and after incubating in 1.5×SBF was performed using LEO 982 Field Emission Scanning Electron Microscope (SEM) (LEO Electron Microscopy, Inc., Thornwood, N.Y.) at 3.0 kV. The sample was sputter coated with gold prior to examination.

Transmission Electron Microscopy: Samples for transmission electron microscopy (TEM) analysis were prepared by ultrasonication of the silk films after incubation in 1.5×SBF in 200 µL of absolute ethanol for 10 min and then depositing a few drops of the suspension onto a standard TEM copper grid with a holey carbon support film. TEM images were obtained with a JOEL 2100 TEM operating at 200 kV with a LaB₆ filament and recorded with a slow scan CCD camera. The diffraction patterns were obtained at calibrated camera lengths using a NiO_x test specimen as a reference.

Results
Protein Expression and Purification: CRGD-15mer (SEQ ID NO: 5), molecular weight of which is 48.56 kDa, migrated to the right position on SDS-PAGE gel, while the apparent molecular weight of CRGD-15mer-CDMP1 (SEQ ID NO: 9)

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indicated by SDS-PAGE electrophoresis was higher than the calculated molecular weight, which is 58.89 kDa. This is due to high content of acidic amino acids such as aspartic acid and glutamic acid in CRGD-15mer-CDMP1 (SEQ ID NO: 9). Western blot analysis of CRGD-15mer-CDMP1 (SEQ ID NO: 9) and CRGD-15mer-CDMP1 (SEQ ID NO: 9) was detected based on the His tag (data not shown).

Amino Acid Composition Analysis: Amino acid composition analysis confirmed the correct composition for the purified CRGD-15mer-CDMP1 (SEQ ID NO: 9).

Functional Change of Protein Secondary Structure by Calcium Ion Binding: CRGD-15mer-CDMP1 (SEQ ID NO: 9) was unordered random coil in water before adding CaCl_2 , indicated by amide I peak at 1649 cm^{-1} and amide II peak at 1548 cm^{-1} . After Ca ion binding, α helix and β sheet structures showed up, demonstrated by amide I peak at 1659 cm^{-1} (α helix) and 1632 cm^{-1} (β sheet) (FIG. 11).

Film Formation: Silk secondary structure changed from random coil to β sheet after treated with 90% methanol (FIG. 12) so that the film was more robust in solution.

Functional Assessments of the Fusion Proteins

The apatite nucleation ability of the functional recombinant silks was studied using the method of alternate soaking process. Silk films of the recombinant fusion proteins were cast using standard methods. Lyophilized fusion proteins were dissolved in HFIP solvent at a concentration of 2.5% w/v overnight at 4°C . The silk-HFIP solutions were then pipetted into 24-well culture plates and left in the hood for about 3 hours for the silk films to form and dry out. The silk films were then treated with 90% v/v methanol to induce β -sheet formation and prevent resolubilization of the films in aqueous solutions.

Scanning Electron Microscopy: To assess the ability for mineralization, the cast recombinant silk film was first soaked in 200 mM aqueous calcium chloride solution buffered with tris(hydroxymethyl) aminomethane and HCl (pH 7.4) (Ca solution) for 1 hr at 37°C , then Ca solution was aspirated and the film was washed with abundant distilled water to wash away any unbound Ca^{2+} , then 120 mM aqueous disodium hydrogenphosphate (P solution) was added to immerse the silk film for 1 hr at 37°C . After soaking in P solution for 1 hr, the film was washed again by distilled water. This is one round of mineralization. Three rounds of the soaking were carried out. The film surface was then observed by SEM. See FIGS. 6A-6C, which show mineral deposition on silk film cast using CRGD-15mer-CDMP1 (SEQ ID NO: 9). Deposition began to occur as in round one (FIG. 6B) and continued through round three (FIG. 6C). Control films made from CRGD-15mer (SEQ ID NO: 4) and 15mer (SEQ ID NO: 5) showed no deposition (data not shown). The SEM images indicate that the functional fusion silk proteins are capable of inducing apatite nucleation and growth, while silk proteins that so not contain mineralizing domains do not promote apatite nucleation.

In addition, scanning electron microscopy (SEM) surface morphologies of recombinant spider silk films after soaking in $1.5\times\text{SBF}$ for various periods of time were assessed see FIG. 13. FIG. 13(A1)-FIG. 13(A5), SS15m-CDMP1 soaked in $1.5\times\text{SBF}$ for 0, 3, 7, 14 and 21 days: FIG. 13(A1)-FIG. 13(A5) respectively. FIG. 13(B1)-FIG. 13(B5), SS15m soaked in $1.5\times\text{SBF}$ for 0,3,7,14 and 21 days: FIG. 13(B1)-FIG. 13(B5) respectively.

The surfaces of the films made of CRGD-15mer-CDMP1 (SEQ ID NO: 9) and CRGD-15mer (SEQ ID NO: 5) were smooth after casting and there was no big difference between two films (FIG. 13A1 and FIG. 13B1). However, after

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immersing in $1.5\times\text{SBF}$ for 3 days, the surface morphologies of the films were different (Fig.13A2 and Fig.13B2).

Transmission Electron Microscopy: The high-magnification image of the crystals grew on CRGD-15mer-CDMP1 (SEQ ID NO: 9) (FIG. 14A) showed that the apatite, which appeared flake-like in the SEM, is composed of aggregates of nanocrystals with needle shapes 100-200 nm in length. The electron diffraction pattern of the nanocrystals (FIG. 14B) showed diffraction rings. The spacings of the rings agreed with the characteristic x-ray diffraction spacing of hydroxyapatite (JCPDS 09-0432). The rings could be assigned to the (002), (211) planes. In addition, a high-resolution TEM image (FIG. 14C) showed the lattice fringes of apatite nanocrystals. The average distance between fringes is 0.82 nm, which is consistent with the value of the (100) interplanar spacing in the apatite structure (0.817 nm).

Example 2

Formation of Silica Silk Fusions Introduction

Silica is widespread in biological systems and serves different functions, among which the most essential ones include support and protection in single-celled organisms, such as diatoms, to higher plants and animals. Despite its widespread occurrence and importance of function, little is known about biosilica and the mechanisms used to produce controlled microscopic and macroscopic silica structures with such nanoscale precision and symmetry. The remarkable control in vivo of the morphology of these beautiful intricate patterns at small length scales are species-specific and has attracted a great deal of interest in recent years as these features exceed the capabilities of present-day synthetic and technological approaches to materials engineering in vitro.

Silaffins^{1; 2; 3; 4; 5} form part of three families of proteins identified to date in the organic matrix of the cell wall of the diatom *Cylindrotheca fusiformis*. Silaffins are highly post-translationally modified peptides and have received the most attention because of their ability to induce and regulate silica precipitation at ambient temperature and pressure. Silaffins are low-molecular weight polypeptides: natSil-1A (6.5 kDa)⁴, natSil-1B (10 kDa)⁵, and natSil-2 (40 kDa)⁵, isolated by treating the diatom cell wall with ammonium fluoride. A gene sil 1 has been isolated from a *C. fusiformis* genomic library and it encodes a polypeptide of 265 amino acids. Seven repeated sequences were identified in this sequence and named R1 to R7¹. R1 corresponds to the precursor of Silaffin-1B, while R3 to R7 and R2 correspond to the precursors of the Silaffin-1A1 and Silaffin-1A2, respectively.

Controlled formation of biosilica structures by different proteins and peptides under various physical reaction environments has also been reported^{6; 7; 8; 9}. The 19 amino-acid R5 peptide of the Sil1 protein was utilized to obtain silica nanostructures with different morphologies including spheres, arch-shaped morphologies and even elongated fibers⁷. These results suggest that through careful manipulation of the environmental conditions and the application of a linear shear force, distinct morphologies can be attained. These opportunities for control of materials outcomes in biosilica morphology establish important links to device fabrication from these materials with regularity in process and outcomes to achieve technological relevance in the future, such as for silica based micro- and nano-devices.

Applicants have genetically cloned the R5 peptide unit of Sil1 protein to a 15mer spider silk clone of the consensus sequence of the golden orb spider *Nephila clavipes*, expressed the fusion protein and performed the same silicifi-

cation reactions on silk films made from the expressed protein. The purpose of fusing this silicification-inducing peptide unit to our genetically engineered silk is to combine the properties of our silk whether in the form of films or other such as spun fibers to the silica precipitating properties of R5 under ambient conditions to produce biomaterials with controlled silica morphologies on the surface. These biomaterials are especially useful in such fields as controlled drug delivery. Materials and Methods

Design and Cloning of Spider Silk Sequences

The repeat unit was selected used in the design of the 15mer and CRGD15mer (SEQ ID NO: 5) was based on the consensus sequence of spider silk native sequence of *Nephila clavipes* (accession P19837, UniProtKB) (-SGRGGLGGQ-GAGAAAAAGGAGQGGYGGGLGSQGT-) (SEQ ID NO: 1). Multimers encoding the repeat were cloned through the transfer of cloned inserts between two shuttle vectors based on pUC19 and pCR-Script (Novagen, Wis.), which were ampicillin and chloramphenicol resistant respectively^{10; 11}.

The expression vector pET-30a (Novagen, Madison, Wis.) was modified with a linker carrying the SpeI site flanked by sequences encoding the amino acids CRGD (SEQ ID NO: 2) to obtain pET30-link. The two complementary oligonucleotide sequences for the linker were: GGATCCTGTCGCGGTGACACTAGTCGCGGTGACTGTG (SEQ ID NO: 13) and GGATCCACAGTCACCGCGACTAGTGT-CACCGCGACAG (SEQ ID NO: 14). The restriction sites of BamHI and SpeI are underlined. The 15mer sequence obtained by multimerization was inserted into pET30-link to generate pET30-CRGD15mer. For the production of a spider silk protein without CRGD (SEQ ID NO: 2), the construct pET30-15mer was obtained by subcloning the NcoI-NotI fragment of pCR-15 into pET-30a vector.

To construct the fusion protein CRGD15mer-R5, the polylinker of pET-21a(+) shuttle vector was redesigned to contain a His-Tag and the CRGD15mer was digested with the restriction enzyme BamHI from pET-30a vector and inserted into the pET-21a(+)-link vector. Two complementary oligonucleotide sequences for the R5 peptide were designed with EcoRI and NotI sites at the 5' and 3' ends respectively:

(SEQ ID NO: 15)
5' aattcagcagcaaaaaagcggcagctattcgggcagcaaggca
gcaaacgcgcgcatcctcgc 3'
and

(SEQ ID NO: 16)
3' gtcgctgcttttttttcgcgctcgataagcccgctcgtttcgcgctt
tcggcgctaggagcgcgg 5'

These synthetic oligonucleotides were annealed and ligated into the pET-21a(+)-link vector right next to the CRGD15mer clone to create the fusion protein with the His-Tag at the C-terminus.

To construct the fusion protein 15mer-R5, the synthetic oligonucleotides were ligated into the NotI and XhoI restriction sites of pET-21a(+) vector right next to the 15mer clone. This fusion protein has a His-Tag at the N-terminus. The amino acid sequence of the R5 peptide of Sil1 protein is: SSKKSGSYSGSKGSKRRIL (SEQ ID NO: 17). Use of other Sil1 protein sequences are also contemplated, for example, the R2 peptide SSKKSGSYSGYSTKSGSKRRIL (SEQ ID NO: 18), the R3 peptide SSKKSGSYSGYSGKSKRRIL (SEQ ID NO: 19), the R4/R6 peptide SSKKSGSYSGYSGKSKRRNL (SEQ ID NO: 20), the R1 peptide SSKKSGSYSGYSGTKK (SEQ ID NO: 21).

Protein Expression and Purification

The constructs pET-21(a)-15mer-R5 and pET-30a-CRGD15mer-R5 were used to transform the *E. coli* host strains Ry-3041, a mutant strain defective in the expression of SlyD protein, for expression^{12; 13}. Cells were cultivated in LB broth at 37° C. Protein expression was induced by the addition of 1 mM IPTG (Fisher Scientific, Hampton, N.H.) when the OD₆₀₀ was between 0.6 and 0.8. After approximately 6 hours of protein expression, the cells were harvested by centrifugation at 9500 rpm. For large scale expression, *E. coli* was grown in a fermentor (Bioflo 3000, New Brunswick Scientific Co., Edison, N.J.) in minimal medium supplemented with 1% yeast extract. Ammonia was used as the base to maintain the pH at 6.8. When the pH exceeded 6.88, as a result of glucose exhaustion in the culture, a feed solution (50% glucose, 10% Yeast Extract, 2% MgSO₄·7H₂O) was added. Pure O₂ was also provided to the culture to sustain the level of dissolved oxygen above 40%. All culture media contained kanamycin (50 µg/ml) or ampicillin (100 µg/ml) for selectivity. For the fermentor grown cells, expression was induced when the absorbance was between 25 and 30 at OD₆₀₀.

The cell pellets were resuspended by adding denaturing buffer (100 mM NaH₂PO₄, 10 mM Tris HCl, 8 M urea, pH 8.0) containing 10 mM imidazole. The cells were lysed by stirring for 30 min and were then centrifuged at 9500 rpm at 4° C. for 30 min. His-tag purification of the proteins was performed by addition of Ni-NTA agarose resin (Qiagen, Valencia, Calif.) to the supernatant (batch purification) under denaturing conditions. After washing the column with denaturing buffer at pH 6.3, the proteins were eluted with denaturing buffer at pH 4.5 (without imidazole).

SDS-polyacrylamide gel electrophoresis (PAGE) was performed using 4-12% precast NuPage Bis-Tris gels (Invitrogen, Carlsbad, Calif.). Electrophoresis was performed in MOPS buffer for 50 min at 200V. Purified samples were extensively dialyzed against several changes of H₂O. For dialysis, Snake Skin membranes (Pierce, Rockford, Ill.) with MWCO of 7000 or lower were used. The dialyzed samples were lyophilized using a LabConco lyophilizer. For determination of the amino acid composition, the samples were submitted to the W. M. Keck Foundation Biotechnology Resource Laboratory (Yale University, New Haven, Conn.). The samples were analyzed from bands of interest cut out from the gel or lyophilized powder after purification and dialysis. Determination of protein concentration was performed by BCA assay (Pierce, Rockford, Ill.) or the molar absorptivity at 280 nm. FIGS. 7A and 7B show the sequence of CRGD15mer-R5 (FIG. 7A) and 15mer-R5 (FIG. 7B) Silicification Reactions

The lyophilized fusion proteins were then dissolved in HFIP solvent at a concentration of 2.5% w/v overnight at 4° C. The silk-HFIP solutions were then pipetted into 24-well culture plates and left in the hood for about 3 hours for the silk films to form and dry out. The silk films were then either left as is or treated with 90% v/v methanol to induce β -sheet formation and prevent resolubilization of the films in aqueous solutions. 100 mM phosphate buffer at pH 5.5 was added to the silk films and leave to sit for about 30 minutes before 1M tetramethoxysilane (TMOS) dissolved in 1 mM hydrochloric acid was added to the mixture for about 10 minutes. The films were then washed with MilliQ water three times and left to dry overnight in the fume hood. These films were then analyzed using a LEO 982 Scanning Electron Microscope at the CIMS facility at Harvard University.

SEM Results

In order to exploit the self-assembling properties of silk in developing silk-silica nanocomposites, experiments were performed using TMOS alone as the precursor. Four genetically engineered variants of the spider silk protein (two controls, one with and one without RGD but both without R5, two chimeric versions of the controls but with R5) were cast into films that were either left untreated or were treated with methanol to induce a structural transition to β -sheet and thus decrease film solubility in aqueous buffer. Silicification reactions were performed on the films yielding spherical silica structures with diameters ranging from ~ 0.5 to $2.0 \mu\text{m}$ only when the silica precipitating domain, R5 peptide, was fused to the C-terminus of the silk proteins (FIG. 8). The silk proteins that did not contain R5, CRGD15mer and 15mer, did not yield significant changes in surface morphology of the films upon exposure to the silicification reactions.

Fusion proteins were assembled into fibers by electrospinning. SEM images of the electropun fibers formed from the chimeric proteins (FIGS. 9A-9E), and the morphological characteristics observed when the fibers were treated with methanol, were similar to those we observed previously for electrospun silk fibroin with polyethylene oxide¹⁵. Upon silicification on electrospun mats formed from the chimeric protein CRGD15mer+R5 (SEQ ID NO: 24) without methanol treatment, similar spherical silica structures were observed as in the reactions on the cast films. However, the dimensions of the silica spheres were slightly smaller, ranging from 200 to 400 nm (FIGS. 9B and 9C). When the electrospun fibers consisting of the chimera CRGD15mer+R5 (SEQ ID NO: 24) were not treated with methanol, the fibers fused together on the surface. Without the β -sheet inducing methanol treatment, the fibers are prone to partially solubilize on the surface yielding a thin film upon which the mineralization reaction takes place. However, upon treatment of the chimera CRGD15mer+R5 (SEQ ID NO: 24) electrospun mats with methanol before silicification, a thin film formed from the solubilized and then fused fibers at the surface and silica nanospheres did not form as in the sample above. Instead, the fibers fused with each other as expected¹⁵ and although mineralization occurred as confirmed by XPS, silica deposited around the fibers providing a non-uniform coating instead of the spheres (FIGS. 9B and 9C). When the chimera CRGD15mer+R5 (SEQ ID NO: 24) was electrospun during the silica polymerization process (concurrent processing), silica deposition was induced in and on the fibers and elliptically shaped silica particles fused to the fibers were observed (FIGS. 9D and 9E). XPS analysis of the resulting fibers confirmed the presence of elemental silicon (FIG. 10). Thus, the concurrent processing approach, fiber spinning and silicification reactions, resulted in a different morphology of the silica in terms of location within the fibers and shape, compared to the silicification reactions conducted post electro spinning.

Sequences
Fibrous protein domain sequence derived from Spidroin1 (the native sequence of the golden orb spider *Nephila clavipes*)

(SEQ ID NO: 1)
SGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGT

A linker sequence CRGD (SEQ ID NO: 2)

The consensus fibrous protein domain sequence derived from Spidroin1 (the native sequence of the golden orb spider *Nephila clavipes*) with CRGD linker

(SEQ ID NO: 3)
CRGDTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGT

15 mer: An amino acid sequence that contains a repeat of a fibrous protein domain sequence derived from Spidroin1 (the native sequence of the golden orb spider *Nephila clavipes*) The fibrous protein domain sequence is repeated 15 times.

(SEQ ID NO: 4)
MASMTGGQGMGRGSAMASGRGGLGGQGAGAAAAGGAGQGGYGGL
GSQGTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLG
GQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGG
AGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQ
GTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQG
AGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGGAGQ
GGYGGLGSQGTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGT
GRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGA
AAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGGAGQGGY
GGLGSQGTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRG
GLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAA
AGGAGQGGYGGLGSQGTSHHHHH

CRGD-15mer: an amino acid sequence that contains a repeat of a fibrous protein domain sequence derived from Spidroin1 (the native sequence of the golden orb spider *Nephila clavipes*) The fibrous protein domain sequence is repeated 15 times with a CRGD linking sequence

(SEQ ID NO: 5)
MASMTGGQGMGRGS CRGDTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQ
GTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAA
AAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQ
TSRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAA
AGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGT
SGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAA
GGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGT
GRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAG
GAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSG
RGGLGGQGAGAAAAGGAGQGGYGGLGSQGTSGRGGLGGQGAGAAAAGG
AGQGGYGGLGSQGTSGRGDCGSHHHHHH

CDMP1, the C-Terminal Sequence of DMP1, a Mineralizing domain

(SEQ ID NO: 6)
RGDNPNTSQTGDQRDSSESEEDRLNTFSSSESQSTEEQDSESNESSL
SEESQESAQDEDESSSQEGLQSQSASRESRSQESQSEEDSRSEENRSDSQ
DSSRSKEESNSTGSTSSSEEDNHPKNI EADNRKLI VDAYHNKPI GDQDDN
DCQDGY

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DMP1, the full length sequence of DMP1, a Mineralizing domain

(SEQ ID NO: 7)
 LPVARYQNTESSESSEERTGNLAQSPPPPMANS DHTDSSSESGEELGSDRSQ
 YRPAGGLSKSAGMDADKEEDEDSDGDDTFGDEDNGPGPEERQWGGPSRLD
 SDEDSADTTQSSSEDSTSQENSAQDTPSDSKDHHSDEADSRPEAGDSTQDS
 ESEFYRVGGGSEGESSEHGDSFEFDDGEMQSDDPGSTRSDRGHTRMSSADI
 SSEESKGDHEPTSTQSDSDSQDVEFSSRKSFRSRVSEEDDRGELADSN
 RETQSVSTEDFRSKEESRSETQEDTAETQSQEDSPEGQDPSSSESSEAGE
 PSQESSSESQEGVASESRGDNPDNTSQTGDQRDSSESSEEDRLNTFSSSES
 QSTEEQGDSESNESLSLSEESQESAQDESSSQEGLQSQSASRESRSQES
 QSEEDSRSEENRDSQSDSSRSKEESNSTGSTSSSEEDNHPKNI EADNRK
 LIVDAYHNKPIGDQDDNDCQDGY

BSP: Bone sialoprotein, a Mineralizing domain

(SEQ ID NO: 8)
 SEFPVQSSSDSSEENGNGDSSEEEEEEEENSNEEENNEENEDSDGNED

CRGD-15mer-CDMP1: (SEQ ID NO: 9): a fusion protein of (SEQ ID NO: 5) and (SEQ ID NO: 6)

(SEQ ID NO: 9)
 MASMTGGQMGRGCRGDTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGT
 SGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAA
 GGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGT
 GRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAG
 GAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSG
 RGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGG
 AGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGR
 GGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGA
 QGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRG
 GLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAG
 QGGYGGGLGSQGTSGRGGLGGQSDGSRGDNPDNTSQTGDQRDSSESSEEDRLNTFSS
 ESQSTEEQGDSESNESLSLSEESQESAQDESSSQEGLQSQSASRESRSQ
 ESQSEEDSRSEENRDSQSDSSRSKEESNSTGSTSSSEEDNHPKNI EADN
 RKLI VDAYHNKPIGDQDDNDCQDGY

CRGD-15mer-DMP1 (SEQ ID NO: 10): a fusion protein of (SEQ ID NO: 5) and SEQ ID NO: 7)

(SEQ ID NO: 10)
 MASMTGGQMGRGCRGDTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGT
 SGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAA
 GGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGT
 GRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAG
 GAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSG

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-continued

RGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGG
 AGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGR
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 QGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRG
 GLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAG
 10 QGGYGGGLGSQGTSGRGDCGSLPVARYQNTESSESSEERTGNLAQSPPPMAN
 SDHTDSSSESGEELGSDRSQYRPAGGLSKSAGMDADKEEDEDSDGDDTFGD
 EDNGPGPEERQWGGPSRLDSEDSADTTQSSSEDSTSQENSAQDTPSDSKD
 HHSDEADSRPEAGDSTQDSSEFYRVGGGSEGESSEHGDSFEFDDGEMQSD
 15 DPGSTRSDRGHTRMSADI SSEESKGDHEPTSTQSDSDSQDVEFSSRKSF
 RRSRVSEEDDRGELADSNRETQSVSTEDFRSKEESRSETQEDTAETQSQ
 EDSPGQDPSSSESSEAGEPSQESSSESQEGVASESRGDNPDNTSQTGDQ
 20 RDESSEEDRLNTFSSSESQSTEEQGDSESNESLSLSEESQESAQDESS
 SQEGLQSQSASRESRSQESQSEEDSRSEENRDSQSDSSRSKEESNSTGS
 TSSSEEDNHPKNI EADNRKLI VDAYHNKPIGDQDDNDCQDGY

15mer-BSP (SEQ ID NO: 11): a fusion protein of (SEQ ID NO: 4) and (SEQ ID NO: 8)

(SEQ ID NO: 11)
 MASMTGGQMGRGSAMASGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGT
 SGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAA
 GGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGT
 35 GRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAG
 GAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSG
 RGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGG
 40 AGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGR
 GGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGA
 QGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRG
 45 GLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAG
 QGGYGGGLGSQGTSEFPVQSSSDSSEENGNGDSSEEEEEEEENSNEEENNE
 ENEDSDGNEDKLHHHHH

CRGD-15mer-BSP (SEQ ID NO: 12): a fusion protein of (SEQ ID NO: 5) and (SEQ ID NO: 8) with linker

(SEQ ID NO: 12)
 55 MASMTGGQMGRGSCRGDTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQ
 GTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAA
 AAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQ
 60 TSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAA
 AGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGT
 SGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAA
 GGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGT
 65 GRGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGLGGQGAGAAAAAG

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GAGQGGYGGGLGSQGTSGRGGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSG
 RGGLGGQGAGAAAAAGGAGQGGYGGGLGSQGTSGRGGGLGGQGAGAAAAAGG
 AGQGGYGGGLGSQGTSGRGGCGSESEFPVQSSSDSSEENGNGDSSEEEEEEE
 ENSNEEENNEENEDSDGNEDKLHHHHHH

One of two complementary oligonucleotide sequences for
 the linker carrying the SpeI site flanked by sequences encod-
 ing the amino acids CRGD to obtain pET30-link

(SEQ ID NO: 13)
 GGATCCTGTGCGCGGTGACACTAGTCGCGGTGACTGTG

One of two complementary oligonucleotide sequences for
 the linker carrying the SpeI site flanked by sequences encod-
 ing the amino acids CRGD to obtain pET30-link

(SEQ ID NO: 14)
 GGATCCACAGTCACCGCGACTAGTGTACCGCGACAG

One of two complementary oligonucleotide sequences for
 the R5 peptide (contains the 19 amino-acid R5 peptide of the
 Sil1 protein) which were designed with EcoRI and NotI sites
 at the 5' and 3' ends respectively

(SEQ ID NO: 15)
 5' aattcagcagcaaaaaagcggcagctattcgggcagcaaaaggcagc
 aaacgcgcgcattcctcgc 3'

One of two complementary oligonucleotide sequences for
 the R5 peptide (contains the 19 amino-acid R5 peptide of the
 Sil1 protein) which were designed with EcoRI and NotI sites
 at the 5' and 3' ends respectively.

(SEQ ID NO: 16)
 3' gtcgctcggttttttcgcgcgtcgataagcccgctgtttcgcgtcgttg
 cggcgtaggagcgcggg 5'

The amino acid sequence of the R5 peptide of Sil1 protein
 from *C. fusiformis* is: SSKKSGSYSGSKGSKRRIL (SEQ ID
 NO: 17)

The amino acid sequence of the R2 peptide of Sil1 protein
 from *C. fusiformis* is: SSKKSGSYSGYSTKKSGSRRIL
 (SEQ ID NO: 18),

The amino acid sequence of the R3 peptide of Sil1 protein
 from *C. fusiformis* is: SSKKSGSYSGYSGKSKRRIL (SEQ
 ID NO: 19),

The amino acid sequence of the R4/R6 peptide of Sil1
 protein from *C. fusiformis* is: SSKKSGSYSGYSGK-
 SKRRNL (SEQ ID NO: 20).

The amino acid sequence of the R1 peptide of Sil1 protein
 from *C. fusiformis* is: SSKKSGSYSGYGTGK (SEQ ID
 NO: 21).

Bone Sialoprotein Precursor [Homo sapiens] Accession
 AAC95490 (GenBank) is:

(SEQ ID NO: 22)
 MKTALILLSI LGMACAFSMK NLHRRVKIED SEENGVFYK
 PRYYLYKHAY FYPHLKRFV QGSSDSSEEN GDDSSSEEEE
 EEETSNEGEN NEESNEDEDS EAENTTLSAT TLGYGEDATP

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GTGYTGLAAI QLPKKAGDIT NKATKEKESD EEEEEEEEGN
 ENEESEAEVD ENEQGINGTS TNSTEAENG GSSGGDNNGEE
 GEEESVTGAN AEGTTETGGQ KGKTSKTTTS PNGGFEPPTP
 PQVYRTTSPF FGKTTTVEYE GEYEYTVNE YDNGYEIYES
 ENGEPRGDNY RAYEDEYSYF KGQGYDGYDG QNYHHQ

Silaffin 1 precursor (natSil-1) [Contains: Silaffin-1B; Silaf-
 fin-1A2; Silaffin-1A1] Accession Q9SE35 (UniProtKB):

(SEQ ID NO: 23)
 MKLTAIFPLL FTAVGYCAAQ SIADLAAANL STEDSKSAQL
 ISADSSDDAS DSSVESVDAA SSDVSGSSVE SVDVSGSSLE
 SVDVSGSSLE SVDDSSSEDSE EEELRILSSK KSGSYSYGT
 KKSGSYSGYS TKKSASRRIL SSKKSGSYSG YSTKKSGSRR
 ILSSKSGSY SGSKGSKRRI LSSKSGSYSG GSKGSKRRNL
 SSKKSGSYSG SKGSKRRILS SKKSGSYSGS KSKRRNLSS
 KKSGSYSGSK GSKRRILSGG LRGS

CRGD15mer-R5: Sequence of fusion protein; fusion of
 CRGD15mer to the amino acid sequence of the R5 peptide of
 Sil1 protein from *C. fusiformis*.

(SEQ ID NO: 24)
 MASMTGGQQM GRGSCRGDTS GRGGLGGQGA GAAAAAGGAG
 QGGYGGGLGSQ GTSGRGGGLG QGAGAAAAAG GAGQGGYGG
 GSQGTSGRGG LGGQGAGAAA AAGGAGQGGY GGLGSQGTSG
 RGGLGGQGAG AAAAAGGAGQ GGYGGLGSQG TSGRGGGLGQ
 GAGAAAAAGG AGQGGYGGGL SQGTSGRGG GQGAGAAAA
 AGGAGQGGYG GLGSQGTSGR GGLGGQGAGA AAAAGGAGQG
 GYGGLGSQGT SGRGGGLGGQ AGAAAAAGGA GQGGYGGLS
 QGTSGRGGLG GQGAGAAAAA GGAGQGGYGG LGSQGTSGRG
 GLGGQGAGAA AAAGGAGQGG YGGLGSQGTSG GRGGLGGQGA
 GAAAAAGGAG QGGYGGGLGSQ GTSGRGGGLG QGAGAAAAAG
 GAGQGGYGG LGSQGTSGRGG LGGQGAGAAA AAGGAGQGGY
 GGLGSQGTSG RGGLGGQGAG AAAAAGGAGQ GGYGGLGSQG
 TSGRGGGLGQ GAGAAAAAGG AGQGGYGGGL SQGTSGRGG
 SEFSSKSGS YSGSGSKRR ILCRHHHHH H

15mer-R5: Sequence of fusion protein; fusion of 15mer to
 the amino acid sequence of the R5 peptide of Sil1 protein
 from *C. fusiformis*.

(SEQ ID NO: 25)
 MHHHHHSSG LVPRGSGMKE TAAAKFERQH MDSPDLGTDD
 DDKAMASGRG GLGGQGAGAA AAAGGAGQGG YGGLGSQGTSG
 GRGGLGGQGA GAAAAAGGAG QGGYGGGLGSQ GTSGRGGGLG
 QGAGAAAAAG GAGQGGYGG LGSQGTSGRGG LGGQGAGAAA

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AAGGAGQGGY GGLGSQGTSG RGGLGGQGAG AAAAAGGAGQ
 GGYGGLGSQG TSGRGGGLGGQ GAGAAAAAGG AGQGGYGGLG
 SQGTSGRGLL GGQAGAAAA AGGAGQGGYG GLGSQGTSGR
 GGLGGQGAGA AAAAGGAGQG GYGGGLGSQGT SGRGGGLGGQ
 AGAAAAAGGA QGGYGGLGS QGTSGRGGGL GQGAGAAAA
 GGAGQGGYGG LGSQGTSGR GLGGQGAGAA AAAGGAGQGG
 YGGLGSQGTSG GRGGGLGGQA GAAAAAGGAG QGGYGGLGSQ
 GTSGRGGGLG QGAGAAAAAG GAGQGGYGGL GSQGTSGRGG
 LGGQGAGAAA AAGGAGQGGY GGLGSQGTSG RGGLGGQGAG
 AAAAAGGAGQ GGYGGLGSQG TSSSKKSGSY SGSKGSKRRIL

PCR PRIMER BamH I site, C-DMP1f:
 (SEQ ID NO: 26)
 5'-CAGGATCCAGGGGTGACAACCCAGAT-3'

PCR PRIMER Xho I site, C-DMP1r:
 (SEQ ID NO: 27)
 5' GCCTCGAGGTAGCCATCTTGCAATC-3'.

The references cited herein and throughout the application are incorporated by reference.

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SEQUENCE LISTING

<160> NUMBER OF SEQ ID NOS: 27

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<212> TYPE: PRT

<213> ORGANISM: Nephila clavipes

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Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly
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Thr

<210> SEQ ID NO 2

<211> LENGTH: 4

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic linker peptide

<400> SEQUENCE: 2

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<210> SEQ ID NO 3

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<212> TYPE: PRT

<213> ORGANISM: *Nephila clavipes*

<400> SEQUENCE: 3

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Leu Gly Ser Gln Gly Thr
35

<210> SEQ ID NO 4

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<213> ORGANISM: *Nephila clavipes*

<400> SEQUENCE: 4

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35 40 45

Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala
50 55 60

Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser
65 70 75 80

Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala
85 90 95

Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly
100 105 110

Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly
115 120 125

Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu
130 135 140

Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala
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Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly
165 170 175

Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly
180 185 190

Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly
195 200 205

Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln
210 215 220

Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr
225 230 235 240

Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly
245 250 255

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Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly
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 Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly
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 Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu
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 Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln
 325 330 335
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 Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly
 370 375 380
 Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala
 385 390 395 400
 Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg
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 420 425 430
 Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly
 435 440 445
 Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly
 450 455 460
 Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser
 465 470 475 480
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<210> SEQ ID NO 5
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 35 40 45
 Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly
 50 55 60
 Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu
 65 70 75 80
 Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala
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 Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly
 100 105 110

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Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg
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Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly
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355						360				365				
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Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Gly
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Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr
		405						410				415		Ser
Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala
		420						425				430		
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450						455				460				
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Ser Gln Ser Thr Glu Glu Gln Gly Asp Ser Glu Ser Asn Glu Ser Leu
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Ser Leu Ser Glu Glu Ser Gln Glu Ser Ala Gln Asp Glu Asp Ser Ser
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Ser Gln Glu Gly Leu Gln Ser Gln Ser Ala Ser Arg Glu Ser Arg Ser
      65             70             75             80
Gln Glu Ser Gln Ser Glu Glu Asp Ser Arg Ser Glu Glu Asn Arg Asp
      85             90             95
Ser Asp Ser Gln Asp Ser Ser Arg Ser Lys Glu Glu Ser Asn Ser Thr
      100            105            110
Gly Ser Thr Ser Ser Ser Glu Glu Asp Asn His Pro Lys Asn Ile Glu
      115            120            125
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Asp His Thr Asp Ser Ser Glu Ser Gly Glu Glu Leu Gly Ser Asp Arg
      35             40             45
Ser Gln Tyr Arg Pro Ala Gly Gly Leu Ser Lys Ser Ala Gly Met Asp
      50             55             60
Ala Asp Lys Glu Glu Asp Glu Asp Asp Ser Gly Asp Asp Thr Phe Gly
      65             70             75             80
Asp Glu Asp Asn Gly Pro Gly Pro Glu Glu Arg Gln Trp Gly Gly Pro
      85             90             95
Ser Arg Leu Asp Ser Asp Glu Asp Ser Ala Asp Thr Thr Gln Ser Ser
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Glu Asp Ser Thr Ser Gln Glu Asn Ser Ala Gln Asp Thr Pro Ser Asp
      115            120            125
Ser Lys Asp His His Ser Asp Glu Ala Asp Ser Arg Pro Glu Ala Gly
      130            135            140
Asp Ser Thr Gln Asp Ser Glu Ser Glu Glu Tyr Arg Val Gly Gly Gly
145            150            155            160
Ser Glu Gly Glu Ser Ser His Gly Asp Gly Ser Glu Phe Asp Asp Glu
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Gly Met Gln Ser Asp Asp Pro Gly Ser Thr Arg Ser Asp Arg Gly His
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Thr Arg Met Ser Ser Ala Asp Ile Ser Ser Glu Glu Ser Lys Gly Asp
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 His Glu Pro Thr Ser Thr Gln Asp Ser Asp Asp Ser Gln Asp Val Glu
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 325 330 335
 Ser Glu Glu Asp Arg Leu Asn Thr Phe Ser Ser Ser Glu Ser Gln Ser
 340 345 350
 Thr Glu Glu Gln Gly Asp Ser Glu Ser Asn Glu Ser Leu Ser Leu Ser
 355 360 365
 Glu Glu Ser Gln Glu Ser Ala Gln Asp Glu Asp Ser Ser Ser Gln Glu
 370 375 380
 Gly Leu Gln Ser Gln Ser Ala Ser Arg Glu Ser Arg Ser Gln Glu Ser
 385 390 395 400
 Gln Ser Glu Glu Asp Ser Arg Ser Glu Glu Asn Arg Asp Ser Asp Ser
 405 410 415
 Gln Asp Ser Ser Arg Ser Lys Glu Glu Ser Asn Ser Thr Gly Ser Thr
 420 425 430
 Ser Ser Ser Glu Glu Asp Asn His Pro Lys Asn Ile Glu Ala Asp Asn
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 <212> TYPE: PRT
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 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Unknown: Unknown bone
 sialoprotein sequence of mammal origin

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<210> SEQ ID NO 9
 <211> LENGTH: 675
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence

-continued

<220> FEATURE:

<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic fusion protein

<400> SEQUENCE: 9

```

Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Cys Arg Gly Asp
 1             5             10             15

Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala
      20             25             30

Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln
      35             40             45

Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala
      50             55             60

Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser
      65             70             75             80

Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala
      85             90             95

Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly
      100            105            110

Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly
      115            120            125

Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu
      130            135            140

Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala
      145            150            155            160

Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly
      165            170            175

Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly
      180            185            190

Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly
      195            200            205

Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln
      210            215            220

Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr
      225            230            235            240

Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly
      245            250            255

Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly
      260            265            270

Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly
      275            280            285

Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly
      290            295            300

Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu
      305            310            315            320

Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln
      325            330            335

Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly
      340            345            350

Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly
      355            360            365

Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly
      370            375            380

Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala
      385            390            395            400

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Met	Ala	Ser	Met	Thr	Gly	Gly	Gln	Gln	Met	Gly	Arg	Cys	Arg	Gly	Asp
1				5					10					15	
Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala
			20					25					30		
Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln
		35					40					45			
Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala
	50					55					60				
Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser
65				70						75					80

Gln	Gly	Thr	Ser	Gly 85	Arg	Gly	Gly	Leu	Gly 90	Gly	Gln	Gly	Ala	Gly 95	Ala
Ala	Ala	Ala	Ala	Gly 100	Gly	Ala	Gly	Gln	Gly 105	Gly	Tyr	Gly	Gly 110	Leu	Gly
Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly 125	Ala	Gly
Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu
Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala
Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly 170	Gln	Gly	Gly	Tyr	Gly 175	Gly
Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly 185	Gly	Gly	Leu	Gly	Gly 190	Gln
Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly
Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln
Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr
Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gln
Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr
Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly
Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly
Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly
Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Ala	Gly	Gln
Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Gly	Arg	Gly	Leu
Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Gly	Gly	Gly	Ala	Gly	Gln
Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly
Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly
Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly
Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala
Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg
Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly
Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly
Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly
Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser
Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala
Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr

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500						505						510					
Ser	Arg	Gly	Asp	Cys	Gly	Ser	Leu	Pro	Val	Ala	Arg	Tyr	Gln	Asn	Thr		
		515					520					525					
Glu	Ser	Glu	Ser	Ser	Glu	Glu	Arg	Thr	Gly	Asn	Leu	Ala	Gln	Ser	Pro		
		530				535					540						
Pro	Pro	Pro	Met	Ala	Asn	Ser	Asp	His	Thr	Asp	Ser	Ser	Glu	Ser	Gly		
545					550					555					560		
Glu	Glu	Leu	Gly	Ser	Asp	Arg	Ser	Gln	Tyr	Arg	Pro	Ala	Gly	Gly	Leu		
				565					570					575			
Ser	Lys	Ser	Ala	Gly	Met	Asp	Ala	Asp	Lys	Glu	Glu	Asp	Glu	Asp	Asp		
			580					585					590				
Ser	Gly	Asp	Asp	Thr	Phe	Gly	Asp	Glu	Asp	Asn	Gly	Pro	Gly	Pro	Glu		
		595					600					605					
Glu	Arg	Gln	Trp	Gly	Gly	Pro	Ser	Arg	Leu	Asp	Ser	Asp	Glu	Asp	Ser		
		610				615					620						
Ala	Asp	Thr	Thr	Gln	Ser	Ser	Glu	Asp	Ser	Thr	Ser	Gln	Glu	Asn	Ser		
625					630					635					640		
Ala	Gln	Asp	Thr	Pro	Ser	Asp	Ser	Lys	Asp	His	His	Ser	Asp	Glu	Ala		
				645					650					655			
Asp	Ser	Arg	Pro	Glu	Ala	Gly	Asp	Ser	Thr	Gln	Asp	Ser	Glu	Ser	Glu		
			660					665					670				
Glu	Tyr	Arg	Val	Gly	Gly	Gly	Ser	Glu	Gly	Glu	Ser	Ser	His	Gly	Asp		
		675					680					685					
Gly	Ser	Glu	Phe	Asp	Asp	Glu	Gly	Met	Gln	Ser	Asp	Asp	Pro	Gly	Ser		
		690				695					700						
Thr	Arg	Ser	Asp	Arg	Gly	His	Thr	Arg	Met	Ser	Ser	Ala	Asp	Ile	Ser		
705					710					715					720		
Ser	Glu	Glu	Ser	Lys	Gly	Asp	His	Glu	Pro	Thr	Ser	Thr	Gln	Asp	Ser		
				725					730					735			
Asp	Asp	Ser	Gln	Asp	Val	Glu	Phe	Ser	Ser	Arg	Lys	Ser	Phe	Arg	Arg		
			740					745					750				
Ser	Arg	Val	Ser	Glu	Glu	Asp	Asp	Arg	Gly	Glu	Leu	Ala	Asp	Ser	Asn		
		755					760					765					
Ser	Arg	Glu	Thr	Gln	Ser	Val	Ser	Thr	Glu	Asp	Phe	Arg	Ser	Lys	Glu		
		770				775					780						
Glu	Ser	Arg	Ser	Glu	Thr	Gln	Glu	Asp	Thr	Ala	Glu	Thr	Gln	Ser	Gln		
785					790					795					800		
Glu	Asp	Ser	Pro	Glu	Gly	Gln	Asp	Pro	Ser	Ser	Glu	Ser	Ser	Glu	Glu		
				805					810					815			
Ala	Gly	Glu	Pro	Ser	Gln	Glu	Ser	Ser	Ser	Glu	Ser	Gln	Glu	Gly	Val		
			820					825					830				
Ala	Ser	Glu	Ser	Arg	Gly	Asp	Asn	Pro	Asp	Asn	Thr	Ser	Gln	Thr	Gly		
		835					840					845					
Asp	Gln	Arg	Asp	Ser	Glu	Ser	Ser	Glu	Glu	Asp	Arg	Leu	Asn	Thr	Phe		
		850				855					860						
Ser	Ser	Ser	Glu	Ser	Gln	Ser	Thr	Glu	Glu	Gln	Gly	Asp	Ser	Glu	Ser		
865					870					875					880		
Asn	Glu	Ser	Leu	Ser	Leu	Ser	Glu	Glu	Ser	Gln	Glu	Ser	Ala	Gln	Asp		
				885					890					895			
Glu	Asp	Ser	Ser	Ser	Gln	Glu	Gly	Leu	Gln	Ser	Gln	Ser	Ala	Ser	Arg		
			900					905					910				
Glu	Ser	Arg	Ser	Gln	Glu	Ser	Gln	Ser	Glu	Glu	Asp	Ser	Arg	Ser	Glu		
		915					920					925					

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Glu Asn Arg Asp Ser Asp Ser Gln Asp Ser Ser Arg Ser Lys Glu Glu
 930 935 940

Ser Asn Ser Thr Gly Ser Thr Ser Ser Ser Glu Glu Asp Asn His Pro
 945 950 955 960

Lys Asn Ile Glu Ala Asp Asn Arg Lys Leu Ile Val Asp Ala Tyr His
 965 970 975

Asn Lys Pro Ile Gly Asp Gln Asp Asp Asn Asp Cys Gln Asp Gly Tyr
 980 985 990

<210> SEQ ID NO 11
 <211> LENGTH: 568
 <212> TYPE: PRT
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 fusion protein

<400> SEQUENCE: 11

Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser Ala Met
 1 5 10 15

Ala Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala
 20 25 30

Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln
 35 40 45

Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala
 50 55 60

Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser
 65 70 75 80

Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala
 85 90 95

Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly
 100 105 110

Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly
 115 120 125

Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu
 130 135 140

Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala
 145 150 155 160

Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly
 165 170 175

Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly
 180 185 190

Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly
 195 200 205

Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln
 210 215 220

Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr
 225 230 235 240

Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly
 245 250 255

Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly
 260 265 270

Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly
 275 280 285

Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly
 290 295 300

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Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu
 305 310 315 320
 Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Gly Gly Ala Gly Gln
 325 330 335
 Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly
 340 345 350
 Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Gly Gly Ala Gly
 355 360 365
 Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly
 370 375 380
 Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Gly Gly Ala
 385 390 395 400
 Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg
 405 410 415
 Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Gly Gly
 420 425 430
 Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly
 435 440 445
 Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Gly
 450 455 460
 Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser
 465 470 475 480
 Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala
 485 490 495
 Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr
 500 505 510
 Ser Glu Phe Pro Val Gln Ser Ser Ser Asp Ser Ser Glu Glu Asn Gly
 515 520 525
 Asn Gly Asp Ser Ser Glu Glu Glu Glu Glu Glu Glu Asn Ser Asn
 530 535 540
 Glu Glu Glu Asn Asn Glu Glu Asn Glu Asp Ser Asp Gly Asn Glu Asp
 545 550 555 560
 Lys Leu His His His His His His
 565

<210> SEQ ID NO 12

<211> LENGTH: 578

<212> TYPE: PRT

<213> ORGANISM: Artificial Sequence

<220> FEATURE:

<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic fusion protein

<400> SEQUENCE: 12

Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser Cys Arg
 1 5 10 15
 Gly Asp Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly Ala
 20 25 30
 Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly
 35 40 45
 Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly
 50 55 60
 Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu
 65 70 75 80
 Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala
 85 90 95

Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly
			100					105					110		
Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly
			115				120				125				
Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly
			130			135					140				
Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln
			145		150				155						160
Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr
			165					170						175	
Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly
			180				185						190		
Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly
			195			200					205				
Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly
			210			215					220				
Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly
			225		230				235						240
Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu
			245					250					255		
Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln
			260				265						270		
Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly
			275			280						285			
Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly
			290			295					300				
Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly
			305		310					315					320
Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala
			325					330					335		
Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg
			340				345						350		
Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly
			355			360					365				
Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly
			370			375				380					
Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly
			385		390				395						400
Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser
			405					410						415	
Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly
			355			360					365				
Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly
			370			375				380					
Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly
			385		3										

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Ser Ser Ser Asp Ser Ser Glu Glu Asn Gly Asn Gly Asp Ser Ser Glu
 530 535 540

Glu Glu Glu Glu Glu Glu Glu Asn Ser Asn Glu Glu Glu Asn Asn Glu
 545 550 555 560

Glu Asn Glu Asp Ser Asp Gly Asn Glu Asp Lys Leu His His His His
 565 570 575

His His

<210> SEQ ID NO 13
 <211> LENGTH: 37
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 oligonucleotide

<400> SEQUENCE: 13

ggatcctgtc gcggtgacac tagtcgcggt gactgtg 37

<210> SEQ ID NO 14
 <211> LENGTH: 37
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 oligonucleotide

<400> SEQUENCE: 14

ggatccacag tcaccgcgac tagtgtcacc gcgacag 37

<210> SEQ ID NO 15
 <211> LENGTH: 64
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 oligonucleotide

<400> SEQUENCE: 15

aattcagcag caaaaaaagc ggcagctatt cgggcagcaa aggcagcaaa cgccgcatcc 60

tcgc 64

<210> SEQ ID NO 16
 <211> LENGTH: 64
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 oligonucleotide

<400> SEQUENCE: 16

gtcgtcgttt ttttcgccgt cgataagccc gtcgtttccg tcgtttgcgg cgtaggagcg 60

cggg 64

<210> SEQ ID NO 17
 <211> LENGTH: 19
 <212> TYPE: PRT
 <213> ORGANISM: *Cylindrotheca fusiformis*

<400> SEQUENCE: 17

Ser Ser Lys Lys Ser Gly Ser Tyr Ser Gly Ser Lys Gly Ser Lys Arg
 1 5 10 15

Arg Ile Leu

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<210> SEQ ID NO 18
 <211> LENGTH: 22
 <212> TYPE: PRT
 <213> ORGANISM: *Cylindrotheca fusiformis*

<400> SEQUENCE: 18

Ser Ser Lys Lys Ser Gly Ser Tyr Ser Gly Tyr Ser Thr Lys Lys Ser
 1 5 10 15

Gly Ser Arg Arg Ile Leu
 20

<210> SEQ ID NO 19
 <211> LENGTH: 20
 <212> TYPE: PRT
 <213> ORGANISM: *Cylindrotheca fusiformis*

<400> SEQUENCE: 19

Ser Ser Lys Lys Ser Gly Ser Tyr Ser Gly Tyr Ser Lys Gly Ser Lys
 1 5 10 15

Arg Arg Ile Leu
 20

<210> SEQ ID NO 20
 <211> LENGTH: 20
 <212> TYPE: PRT
 <213> ORGANISM: *Cylindrotheca fusiformis*

<400> SEQUENCE: 20

Ser Ser Lys Lys Ser Gly Ser Tyr Ser Gly Tyr Ser Lys Gly Ser Lys
 1 5 10 15

Arg Arg Asn Leu
 20

<210> SEQ ID NO 21
 <211> LENGTH: 15
 <212> TYPE: PRT
 <213> ORGANISM: *Cylindrotheca fusiformis*

<400> SEQUENCE: 21

Ser Ser Lys Lys Ser Gly Ser Tyr Tyr Ser Tyr Gly Thr Lys Lys
 1 5 10 15

<210> SEQ ID NO 22
 <211> LENGTH: 317
 <212> TYPE: PRT
 <213> ORGANISM: *Homo sapiens*

<400> SEQUENCE: 22

Met Lys Thr Ala Leu Ile Leu Leu Ser Ile Leu Gly Met Ala Cys Ala
 1 5 10 15

Phe Ser Met Lys Asn Leu His Arg Arg Val Lys Ile Glu Asp Ser Glu
 20 25 30

Glu Asn Gly Val Phe Lys Tyr Arg Pro Arg Tyr Tyr Leu Tyr Lys His
 35 40 45

Ala Tyr Phe Tyr Pro His Leu Lys Arg Phe Pro Val Gln Gly Ser Ser
 50 55 60

Asp Ser Ser Glu Glu Asn Gly Asp Asp Ser Ser Glu Glu Glu Glu Glu
 65 70 75 80

Glu Glu Glu Thr Ser Asn Glu Gly Glu Asn Asn Glu Glu Ser Asn Glu
 85 90 95

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Asp	Glu	Asp	Ser	Glu	Ala	Glu	Asn	Thr	Thr	Leu	Ser	Ala	Thr	Thr	Leu
			100					105					110		
Gly	Tyr	Gly	Glu	Asp	Ala	Thr	Pro	Gly	Thr	Gly	Tyr	Thr	Gly	Leu	Ala
		115					120					125			
Ala	Ile	Gln	Leu	Pro	Lys	Lys	Ala	Gly	Asp	Ile	Thr	Asn	Lys	Ala	Thr
	130					135					140				
Lys	Glu	Lys	Glu	Ser	Asp	Glu	Glu	Glu	Glu	Glu	Glu	Glu	Gly	Asn	
145					150					155				160	
Glu	Asn	Glu	Glu	Ser	Glu	Ala	Glu	Val	Asp	Glu	Asn	Glu	Gln	Gly	Ile
				165					170					175	
Asn	Gly	Thr	Ser	Thr	Asn	Ser	Thr	Glu	Ala	Glu	Asn	Gly	Asn	Gly	Ser
			180					185					190		
Ser	Gly	Gly	Asp	Asn	Gly	Glu	Glu	Gly	Glu	Glu	Glu	Ser	Val	Thr	Gly
		195					200					205			
Ala	Asn	Ala	Glu	Gly	Thr	Thr	Glu	Thr	Gly	Gly	Gln	Gly	Lys	Gly	Thr
	210					215					220				
Ser	Lys	Thr	Thr	Thr	Ser	Pro	Asn	Gly	Gly	Phe	Glu	Pro	Thr	Thr	Pro
225					230					235					240
Pro	Gln	Val	Tyr	Arg	Thr	Thr	Ser	Pro	Pro	Phe	Gly	Lys	Thr	Thr	Thr
				245					250					255	
Val	Glu	Tyr	Glu	Gly	Glu	Tyr	Glu	Tyr	Thr	Gly	Val	Asn	Glu	Tyr	Asp
			260					265					270		
Asn	Gly	Tyr	Glu	Ile	Tyr	Glu	Ser	Glu	Asn	Gly	Glu	Pro	Arg	Gly	Asp
		275					280					285			
Asn	Tyr	Arg	Ala	Tyr	Glu	Asp	Glu	Tyr	Ser	Tyr	Phe	Lys	Gly	Gln	Gly
	290					295					300				
Tyr	Asp	Gly	Tyr	Asp	Gly	Gln	Asn	Tyr	Tyr	His	His	Gln			
305					310					315					

<210> SEQ ID NO 23

<211> LENGTH: 265

<212> TYPE: PRT

<213> ORGANISM: *Cylindrotheca fusiformis*

<400> SEQUENCE: 23

Met	Lys	Leu	Thr	Ala	Ile	Phe	Pro	Leu	Leu	Phe	Thr	Ala	Val	Gly	Tyr
1				5					10					15	
Cys	Ala	Ala	Gln	Ser	Ile	Ala	Asp	Leu	Ala	Ala	Ala	Asn	Leu	Ser	Thr
			20					25					30		
Glu	Asp	Ser	Lys	Ser	Ala	Gln	Leu	Ile	Ser	Ala	Asp	Ser	Ser	Asp	Asp
		35					40					45			
Ala	Ser	Asp	Ser	Ser	Val	Glu	Ser	Val	Asp	Ala	Ala	Ser	Ser	Asp	Val
	50					55				60					
Ser	Gly	Ser	Ser	Val	Glu	Ser	Val	Asp	Val	Ser	Gly	Ser	Ser	Leu	Glu
	65				70				75					80	
Ser	Val	Asp	Val	Ser	Gly	Ser	Ser	Leu	Glu	Ser	Val	Asp	Asp	Ser	Ser
			85					90					95		
Glu	Asp	Ser	Glu	Glu	Glu	Glu	Leu	Arg	Ile	Leu	Ser	Ser	Lys	Lys	Ser
			100				105						110		
Gly	Ser	Tyr	Tyr	Ser	Tyr	Gly	Thr	Lys	Lys	Ser	Gly	Ser	Tyr	Ser	Gly
	115					120						125			
Tyr	Ser	Thr	Lys	Lys	Ser	Ala	Ser	Arg	Arg	Ile	Leu	Ser	Ser	Lys	Lys
	130					135					140				
Ser	Gly	Ser	Tyr	Ser	Gly	Tyr	Ser	Thr	Lys	Lys	Ser	Gly	Ser	Arg	Arg
145					150					155					160

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Ile Leu Ser Ser Lys Lys Ser Gly Ser Tyr Ser Gly Ser Lys Gly Ser
      165                      170                      175

Lys Arg Arg Ile Leu Ser Ser Lys Lys Ser Gly Ser Tyr Ser Gly Ser
      180                      185                      190

Lys Gly Ser Lys Arg Arg Asn Leu Ser Ser Lys Lys Ser Gly Ser Tyr
      195                      200                      205

Ser Gly Ser Lys Gly Ser Lys Arg Arg Ile Leu Ser Ser Lys Lys Ser
      210                      215                      220

Gly Ser Tyr Ser Gly Ser Lys Gly Ser Lys Arg Arg Asn Leu Ser Ser
      225                      230                      235                      240

Lys Lys Ser Gly Ser Tyr Ser Gly Ser Lys Gly Ser Lys Arg Arg Ile
      245                      250                      255

Leu Ser Gly Gly Leu Arg Gly Ser Met
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<210> SEQ ID NO 24
<211> LENGTH: 551
<212> TYPE: PRT
<213> ORGANISM: Cyindrotheca fusiformis

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<400> SEQUENCE: 24

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      20                      25                      30

Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu Gly
      35                      40                      45

Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala Gly
      50                      55                      60

Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly Leu
      65                      70                      75                      80

Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly Ala
      85                      90                      95

Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly Gly
      100                      105                      110

Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln Gly
      115                      120                      125

Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr Gly
      130                      135                      140

Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly Gln
      145                      150                      155                      160

Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly Tyr
      165                      170                      175

Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly Gly
      180                      185                      190

Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly Gly
      195                      200                      205

Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu Gly
      210                      215                      220

Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln Gly
      225                      230                      235                      240

Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Gly Arg Gly Gly Leu
      245                      250                      255

Gly Gly Gln Gly Ala Gly Ala Ala Ala Ala Ala Gly Gly Ala Gly Gln
      260                      265                      270

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Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly
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	290					295						300			
Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly
	305				310					315					320
Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala
				325					330					335	
Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg
			340					345					350		
Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly
		355					360					365			
Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly
	370					375					380				
Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly
	385				390				395						400
Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser
			405					410						415	
Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala
			420				425					430			
Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr
		435				440						445			
Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala
	450				455					460					
Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly
	465				470				475						480
Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala
			485					490					495		
Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln
			500				505						510		
Gly	Thr	Ser	Arg	Gly	Asp	Cys	Gly	Ser	Glu	Phe	Ser	Ser	Lys	Lys	Ser
		515				520						525			
Gly	Ser	Tyr	Ser	Gly	Ser	Lys	Gly	Ser	Lys	Arg	Arg	Ile	Leu	Cys	Gly
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Arg	His	His	His	His	His	His									
	545				550										

<210> SEQ ID NO 25

<211> LENGTH: 561

<212> TYPE: PRT

<213> ORGANISM: *Cylindrotheca fusiformis*

<400> SEQUENCE: 25

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		20						25					30		
Ser	Pro	Asp	Leu	Gly	Thr	Asp	Asp	Asp	Lys	Ala	Met	Ala	Ser	Gly	
		35				40					45				
Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly
	50					55				60					
Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser
	65				70				75					80	
Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala
			85					90						95	

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Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr
			100					105					110		
Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala
		115					120					125			
Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly
	130					135					140				
Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala
145					150					155					160
Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln
				165					170					175	
Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala
			180					185					190		
Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser
		195					200					205			
Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala
	210					215					220				
Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly
225					230					235					240
Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly
				245					250					255	
Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly	Leu
			260					265					270		
Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly	Ala
		275					280					285			
Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly	Gly
	290					295					300				
Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln	Gly
305					310					315					320
Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr	Gly
				325					330					335	
Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly	Gln
			340					345					350		
Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly	Tyr
		355					360					365			
Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly	Gly
	370					375					380				
Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly	Gly
385					390					395					400
Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu	Gly
				405					410					415	
Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln	Gly
			420					425					430		
Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly	Leu
		435					440					445			
Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly	Gln
	450					455					460				
Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly	Gly
465					470					475					480
Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala	Gly
				485					490					495	
Gln	Gly	Gly	Tyr	Gly	Gly	Leu	Gly	Ser	Gln	Gly	Thr	Ser	Gly	Arg	Gly
			500					505					510		
Gly	Leu	Gly	Gly	Gln	Gly	Ala	Gly	Ala	Ala	Ala	Ala	Ala	Gly	Gly	Ala
		515					520						525		

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Gly Gln Gly Gly Tyr Gly Gly Leu Gly Ser Gln Gly Thr Ser Ser Ser
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Lys Lys Ser Gly Ser Tyr Ser Gly Ser Lys Gly Ser Lys Arg Arg Ile
 545 550 555 560

Leu

<210> SEQ ID NO 26
 <211> LENGTH: 26
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 primer

<400> SEQUENCE: 26

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26

<210> SEQ ID NO 27
 <211> LENGTH: 26
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
 primer

<400> SEQUENCE: 27

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26

What is claimed is:

1. A method for forming a fibrous protein inorganic-composite comprising:

(a) contacting a fusion protein with an inorganic material capable of mineralizing for a sufficient period of time to allow mineralization of the inorganic material, wherein the fusion protein comprises a fibrous protein domain and a mineralizing domain, wherein the fibrous protein domain is obtained from silk and comprises a repeat unit as set forth in SEQ ID NO: 1 or SEQ ID NO: 3, and wherein the mineralizing domain is capable of inducing mineralization and wherein the mineralizing domain is obtained from bone sialoprotein (BSP), silaffin-1 (Sil1) protein, or functional fragments thereof.

2. The method of claim 1, wherein the fibrous protein domain is from the silk protein Spidroin 1.

3. The method of claim 1, wherein the fibrous protein domain comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 4 and 5.

4. The method of claim 1, wherein the mineralizing domain induces the formation of hydroxyapatite, silica, cadmium sulfide or magnetite.

5. The method of claim 1, wherein the mineralization domain comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 8 and 17-23.

6. The method of claim 1, wherein the fusion protein comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 11, 12, 24, and 25.

7. The method of claim 1, wherein the inorganic material forms hydroxyapatite or silica.

8. The method of claim 1, wherein the fusion protein comprises a fiber, a film, or a sponge.

9. The method of claim 8, wherein the fiber, film, or sponge further comprises an agent.

10. The method of claim 9, wherein the agent is selected from the group consisting of a protein, peptide, nucleic acid, PNA, aptamer, antibody and a small molecule.

11. A fusion polypeptide comprising a fibrous protein domain and a mineralizing domain, wherein the fibrous protein domain is obtained from silk and comprises a repeat unit as set forth in SEQ ID NO: 1 or SEQ ID NO: 3, and wherein the mineralizing domain is capable of inducing mineralization and wherein the mineralizing domain is obtained from bone sialoprotein (BSP), silaffin-1 (Sil1) protein and functional fragments thereof.

12. The fusion polypeptide of claim 11, wherein the fibrous protein domain is from the silk protein Spidroin 1.

13. The fusion polypeptide of claim 11, wherein the fibrous protein domain comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 4 and 5.

14. The fusion polypeptide of claim 11, wherein the mineralizing domain induces the formation of hydroxyapatite, silica, cadmium sulfide or magnetite.

15. The fusion polypeptide of claim 11, wherein the mineralization domain comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 8 and 17-23.

16. The fusion polypeptide of claim 11, wherein the fusion protein comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 11, 12, 24, and 25.

17. The fusion polypeptide of claim 11, wherein the fusion polypeptide comprises a fiber, a film, or a sponge.

18. The fusion polypeptide of claim 17, wherein the fiber, film, or sponge further comprises an agent.

19. The fusion polypeptide of claim 18, wherein the agent is selected from the group consisting of a protein, peptide, nucleic acid, PNA, aptamer, antibody and a small molecule.

* * * * *

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 8,501,437 B2
APPLICATION NO. : 13/347801
DATED : August 6, 2013
INVENTOR(S) : David L. Kaplan et al.

Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

In the Specification

At Column 1, Line numbers 20-25, delete:

“GOVERNMENT SUPPORT

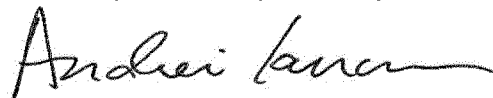
This invention was supported by FA9550041-0363 awarded by the U.S. Air Force and EB003210-01 awarded by the National Institutes of Health. The government has certain rights in the invention.”

And replace it with:

--GOVERNMENT SUPPORT

This invention was made with government support under grants EB003210 and DE011657 awarded by the National Institutes of Health and by grant FA9550-04-1-0363 awarded by the United States Air Force. The government has certain rights in the invention.--

Signed and Sealed this
Twenty-first Day of May, 2019



Andrei Iancu
Director of the United States Patent and Trademark Office