

Biomaterial

The present invention relates to the field of synthetic bone, dental materials and regeneration scaffolds for
5 biomedical applications and, in particular, to synthetic bone, dental materials and regeneration scaffolds and their precursors comprising collagen, a calcium phosphate material and one or more glycosaminoglycans.

10 Natural bone is a biocomposite of collagen, non-collagenous organic phases including glycosaminoglycans, and calcium phosphate. Its complex hierarchical structure leads to exceptional mechanical properties including high stiffness, strength, and fracture toughness, which in turn
15 enable bones to withstand the physiological stresses to which they are subjected on a daily basis. The challenge faced by researchers in the field is to make a synthetic material that has a composition and structure that will allow natural bone growth in and around the synthetic
20 material in the human or animal body.

It has been observed that bone will bond directly to calcium phosphates in the human body (a property referred to as bioactivity) through a bone-like apatite layer formed in
25 the body environment. Collagen and copolymers comprising collagen and other bioorganics such as glycosaminoglycans on the other hand, are known to be optimal substrates for the attachment and proliferation of numerous cell types, including those responsible for the production and
30 maintenance of bone in the human body.

- 2 -

Hydroxyapatite is the calcium phosphate most commonly used as constituent in bone substitute materials. It is, however, a relatively insoluble material when compared to other forms of calcium phosphate materials such as brushite, tricalcium phosphate and octacalcium phosphate. The relatively low solubility of apatite can be a disadvantage when producing a biomaterial as the rate of resorption of the material in the body is particularly slow.

Calcium phosphates such as hydroxyapatite are mechanically stiff materials. However, they are relatively brittle when compared to natural bone. Collagen is a mechanically tough material, but has relatively low stiffness when compared to natural bone. Materials comprising copolymers of collagen and glycosaminoglycans are both tougher and stiffer than collagen alone, but still have relatively low stiffness when compared to natural bone."

Previous attempts in the prior art of producing a synthetic bone-substitute material having improved mechanical toughness over hydroxyapatite and improved stiffness over collagen and copolymers of collagen and glycosaminoglycans include combining collagen and apatite by mechanical mixing. Such a mechanical method is described in EP-A-0164 484.

Later developments in the technology include producing a bone-replacement material comprising hydroxyapatite, collagen and chondroitin-4-sulphate by the mechanical mixing of these components. This is described in EP-A-0214070. This document further describes dehydrothermic crosslinking of the chondroitin-4-sulphate to the collagen. Materials

- 3 -

comprising apatite, collagen and chondroitin-4-sulphate have been found to have good biocompatibility. The mechanical mixing of the apatite with the collagen, and optionally chondroitin-4-sulphate, essentially forms collagen /
5 chondroitin-4-sulphate-coated particles of apatite. It has been found that such a material, although biocompatible, produces limited in-growth of natural bone when in the human or animal body and no remodeling of the calcium phosphate phase of the synthetic material.

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The present invention seeks to address at least some of the problems associated with the prior art.

In a first aspect, the present invention provides a
15 process for the production of a composite material comprising collagen, brushite and one or more glycosaminoglycans, said process comprising the steps of providing an acidic aqueous solution comprising collagen, a calcium source and a phosphorous source and one
20 or more glycosaminoglycans, and precipitating the collagen, the brushite and the one or more glycosaminoglycans together from the aqueous solution to form a triple co-precipitate.

25 The term triple co-precipitate encompasses precipitation of the three compounds where the compounds have been precipitated at substantially the same time from the same solution/dispersion. It is to be distinguished from a material formed from the mechanical mixing of the
30 components, particularly where these components have been precipitated separately, for instance in different solutions. The microstructure of a co-precipitate is

- 4 -

substantially different from a material formed from the mechanical mixing of its components.

In the first aspect, the solution preferably has a pH of from 2.5 to 6.5, more preferably from 2.5 to 5.5. More preferably, the solution has a pH of from 3.0 to 4.5. Still more preferably, the solution has a pH of from 3.8 to 4.2. Most preferably, the solution has a pH of around 4.

10 The calcium source is preferably selected from one or more of calcium nitrate, calcium acetate, calcium chloride, calcium carbonate, calcium alkoxide, calcium hydroxide, calcium silicate, calcium sulphate, calcium gluconate and the calcium salt of heparin. A calcium salt of heparin may
15 be derived from the porcine intestinal mucosa. Suitable calcium salts are commercially available from Sigma-Aldrich Inc.

 The phosphorus source is preferably selected from one
20 or more of ammonium-dihydrogen phosphate, diammonium hydrogen phosphate, phosphoric acid, disodium hydrogen orthophosphate 2-hydrate ($\text{Na}_2\text{HPO}_4 \cdot 2\text{H}_2\text{O}$, sometimes termed GPR Sorensen's salt) and trimethyl phosphate, alkali metal salts (e.g Na or K) of phosphate, alkaline earth salts (e.g. Mg
25 or Ca) of phosphate.

 Glycosaminoglycans are a family of macromolecules containing long unbranched polysaccharides containing a repeating disaccharide unit. Preferably, the one or more
30 glycosaminoglycans are selected from chondroitin sulphate, dermatin sulphate, heparin, heparin sulphate, keratin sulphate and hyaluronic acid. Chondroitin sulphate may be

- 5 -

chondroitin-4-sulphate or chondroitin-6-sulphate, both of which are available from Sigma-Aldrich Inc. The chondroitin-6-sulphate may be derived from shark cartilage. Hyaluronic acid may be derived from human umbilical chord.

5 Heparin may be derived from porcine intestinal mucosa.

Preferably, in the precipitation of the triple co-precipitate, the solution has a temperature of from 4.0 to 50°C. More preferably, the solution has a temperature of

10 from 15 to 40°C. The solution may be at room temperature, that is from 20 to 30°C, with a temperature of from 20 to 27°C being preferred. Most preferably, the temperature is around 25°C.

15 The concentration of calcium ions in the aqueous solution is typically from 0.00025 to 1 mol dm⁻³ and preferably from 0.001 to 1 mol dm⁻³. Where the process includes the additional further steps of filtration and/or low temperature drying, the concentration of calcium ions in

20 the aqueous solution is more preferably from 0.05 to 0.5 mol dm⁻³ (for example from 0.08 to 0.25 mol dm⁻³) and most preferably from 0.1 to 0.5 mol dm⁻³. Where the process includes the additional further steps of freeze drying and optionally injection moulding, the concentration of calcium

25 ions in the aqueous solution is more preferably from 0.01 to 0.3 mol dm⁻³ and most preferably from 0.05 to 0.18 mol dm⁻³.

Preferably, the solution comprises phosphate ions and the concentration of phosphate ions in solution is typically

30 from 0.00025 to 1 mol dm⁻³ and preferably from 0.001 to 1 M. Where the process includes the additional further steps of filtration and/or low temperature drying, the concentration

- 6 -

of phosphate ions in solution is more preferably 0.05 to 0.5 moldm⁻³, still more preferably 0.1 to 0.5 M, for example 0.1 to 0.35 moldm⁻³. Where the process includes the additional further steps of freeze drying and optionally injection

5 moulding, the concentration of phosphate ions in solution is more preferably from 0.01 to 0.3 moldm⁻³, still more preferably 0.05 to 0.18 M.

Preferably, the ratio of collagen to the total amount

10 of one or more glycosaminoglycans in the solution prior to precipitation is from 8:1 to 30:1 by weight. More preferably, the ratio of collagen to the total amount of one or more glycosaminoglycans is from 10:1 to 12:1, and most preferably the ratio is from 11:1 to 23:2.

15 Preferably, the ratio of collagen to brushite in the triple co-precipitate is from 10:1 to 1:100 by weight, more preferably from 5:1 to 1:20, still more preferably from 3:2 to 1:10, most preferably from 3:2 to 1:4.

20 The concentration of collagen in the solution prior to precipitation is typically from 1 to 20 g/L, more preferably from 1 to 10g/L. Where the process includes the steps of filtration and/or low temperature drying, the concentration

25 of collagen in the solution is more preferably from 1 to 10 g/L, still more preferably from 1.5 to 2.5 g/L, and most preferably 1.5 to 2.0 g/L. Where the process includes freeze drying and optionally injection moulding, the concentration of collagen in the solution prior to

30 precipitation is preferably from 5 to 20 g/L, more preferably from 5 to 12 g/L, and most preferably from 9 to 10.5 g/L.

- 7 -

The total concentration of the one or more glycosaminoglycans in the solution prior to precipitation is typically from 0.01 to 1.5 g/L, more preferably from 0.01 to 1 g/L. Where the process includes the additional further steps of filtration and/or low temperature drying, the total concentration of the one or more glycosaminoglycans in the solution is more preferably from 0.03 to 1.25 g/L, still more preferably from 0.125 to 0.25 g/L, and most preferably from 0.13 to 0.182 g/L. Where the process includes the additional further steps of freeze drying and optionally injection moulding, the total concentration of the one or more glycosaminoglycans in the solution is more preferably from 0.15 to 1.5 g/L, still more preferably from 0.41 to 1.2 g/L, and most preferably from 0.78 to 0.96 g/L.

Preferably the solution comprises calcium ions and the ratio of collagen to the calcium ions is typically from 1:40 to 500:1 by weight. Where the process includes the additional further steps of filtration and/or low temperature drying, the ratio of collagen to the calcium ions is more preferably from 1:40 to 250:1, still more preferably 1:13 to 5:4, and most preferably 1:13 to 1:2. Where the process includes the additional further steps of freeze drying and optionally injection moulding, the ratio of collagen to the calcium ions is more preferably from 1:8 to 500:1, still more preferably 5:12 to 30:1, and most preferably 5:5 to 5:1.

Precipitation may be effected by combining the collagen, the calcium source, the phosphorous source and one or more glycosaminoglycans in an acidic aqueous solution and

- 8 -

either allowing the solution to stand until precipitation occurs, agitating the solution, titration using basic titrants such as ammonia, addition of a nucleating agent such as pre-fabricated brushite, varying the rate of
5 addition of the calcium source, and any combination of these techniques.

In a second aspect, the present invention provides a process for the production of a composite biomaterial
10 comprising collagen, octacalcium phosphate and one or more glycosaminoglycans, said process comprising the steps of providing a composite material comprising collagen, brushite and one or more glycosaminoglycans, and
converting at least some of the brushite in the
15 composite material to octacalcium phosphate by hydrolysis.

The term biomaterial encompasses a material that is biocompatible with a human or animal body.

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In the second aspect, the composite material preferably comprises or consists essentially of a triple co-precipitate comprising collagen, brushite and one or more glycosaminoglycans. The triple co-precipitate may be formed
25 by a process as herein described in relation to the first aspect of the present invention.

Preferably, the step of hydrolysis (hydrolysis) of brushite to octacalcium phosphate comprises contacting the
30 triple co-precipitate with an aqueous solution, said aqueous solution being at or above the pH at which octacalcium phosphate becomes thermodynamically more stable than

- 9 -

brushite. Preferably, this aqueous solution has a pH of from 6 to 8. More preferably, this aqueous solution has a pH of from 6.3 to 7. Most preferably, this aqueous solution has pH of about 6.65. The aqueous solution may comprise, 5 for example, deionised water whose pH is controlled with a titrant, a buffer solution, a solution saturated with respect to another calcium-containing compound and/or phosphorus-containing compound. A preferred aqueous solution comprises acetic acid titrated to the desired pH 10 using ammonia.

Preferably, the step of hydrolysis of brushite to octacalcium phosphate is performed at a temperature of from 20 to 50°C, more preferably from 30 to 40°C, still more 15 preferably from 36 to 38 °C, most preferably around 37 °C.

Preferably, the step of hydrolysis of brushite to octacalcium phosphate is performed for a time of from 12 to 144 hours, more preferably from 18 to 72 hours, most 20 preferably from 24 to 48 hours.

In a third aspect, the present invention provides a process for the production of a composite biomaterial comprising collagen, apatite and one or more 25 glycosaminoglycans, said process comprising the steps of providing a composite material comprising collagen, brushite and one or more glycosaminoglycans, and converting at least some of the brushite in the composite material to apatite by hydrolysis.

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Apatite is a class of minerals comprising calcium and phosphate and has the general formula: $\text{Ca}_5(\text{PO}_4)_3(\text{X})$, wherein

- 10 -

X may be an ion that is typically OH^- , F^- and Cl^- , as well as other ions known to those skilled in the art. Apatite also includes substituted apatites such as silicon-substituted apatites. Apatite includes hydroxyapatite, which is a
5 specific example of an apatite. The hydroxyapatite may also be substituted with silicon.

In the third aspect, the composite material preferably comprises or consists essentially of a triple co-precipitate
10 comprising collagen, brushite and one or more glycosaminoglycans. The triple co-precipitate may be formed according to the process as herein described in relation to the first aspect of the present invention.

15 Preferably, the step of hydrolysatation (hydrolysis) of brushite to apatite comprises contacting the triple co-precipitate with an aqueous solution, said aqueous solution being at or above the pH at which apatite becomes thermodynamically more stable than brushite. Preferably,
20 for the conversion of brushite to apatite, the aqueous solution has a pH of from 6.65 to 9, more preferably from 7 to 8.5, still more preferably from 7.2 to 8.5. The aqueous solution may comprise, for example, deionised water whose pH is controlled with a titrant, a buffer solution, a solution
25 saturated with respect to another calcium-containing compound and/or phosphorus-containing compound.

Preferably, the step of hydrolysatation of brushite to apatite is performed at a temperature of 20 to 50°C, more
30 preferably from 30 to 40°C, still more preferably from 36 to 38 °C, most preferably around 37 °C.

- 11 -

Preferably, the step of hydrolysis of brushite to apatite is performed for a time of from 12 to 288 hours, more preferably from 18 to 72 hours, most preferably from 24 to 48 hours.

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Methods of increasing the rate of conversion of brushite to octacalcium phosphate and/or apatite include (i) increasing the temperature, (ii) the brushite concentration in solution, and/or (iii) the agitation speed.

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It may be desirable to produce a biomaterial according to the present invention comprising both apatite and octacalcium phosphate. The processes of the second and third aspects of the present invention may be combined to produce a material comprising both octacalcium phosphate and apatite. The brushite in the triple co-precipitate may first be converted to octacalcium phosphate and then the octacalcium phosphate may be partially converted to apatite. Total, or near total (i.e. at least 98%), conversion of brushite or octacalcium phosphate to apatite typically occurs by hydrolysis at a pH of 8.0 or more for a period of about 12 hours. Partial conversion of the brushite and/or apatite in the material may therefore be effected by hydrolysis for a period of less than 12 hours.

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Preferably, the step of hydrolysis of octacalcium phosphate to apatite is carried out at a pH of from 6.65 to 10, more preferably from 7.2 to 10, still more preferably from 8 to 9.

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Preferably, the step of hydrolysis of octacalcium phosphate to apatite is performed at a temperature of from

- 12 -

20 to 50°C, more preferably from 30 to 40°C, still more preferably from 36 to 38°C, most preferably around 37°C.

Preferably, the step of hydrolysis of octacalcium
5 phosphate to apatite is performed for a time of from 2 to 144 hours, more preferably from 12 to 96 hours, most preferably from 24 to 72 hours.

In the second and third aspects of the present
10 invention, the conversion of brushite to octacalcium phosphate and/or apatite is preferably conducted at a temperature of from 30 to 40 degrees centigrade. More preferably, the conversion is conducted at a temperature of from 36 to 38 degrees centigrade. Most preferably, the
15 conversion is conducted at a temperature of about 37 degrees centigrade.

Preferably, the processes of the present invention further comprise the step of crosslinking the one or more
20 glycosaminoglycans and the collagen in the triple co-precipitate. By triple co-precipitate this includes the triple co-precipitate comprising collagen, brushite and one or more glycosaminoglycans and derivatives of the co-precipitate. Derivatives include the co-precipitate wherein
25 at least some of the brushite has been converted to octacalcium phosphate and/or apatite, and the co-precipitate that has been shaped or moulded, or subjected to any further chemical or mechanical processing. Crosslinking may be achieved using any of the conventional techniques.

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Preferably, at least some of the brushite is converted to octacalcium phosphate and/or apatite, the

- 13 -

glycosaminoglycan and collagen are crosslinked prior to the conversion of the brushite to octacalcium phosphate and/or apatite. This crosslinking may be effected by subjecting the triple co-precipitate to one or more of gamma radiation, ultraviolet radiation, a dehydrthermal treatment, non-enzymatic glycation with a simple sugar such as glucose, mannose, ribose and sucrose, contacting the triple co-precipitate with one or more of glutaraldehyde, ethyl dimethylaminopropyl carbodiimide and/or nor-dihydroguaric acid, or any combination of these methods. These methods are conventional in the art.

Preferably, if at least some of the brushite is converted to octacalcium phosphate and/or apatite, the glycosaminoglycan and collagen are crosslinked subsequent to the conversion of the brushite to octacalcium phosphate and/or apatite. The crosslinking subsequent to the conversion of the brushite to apatite/octacalcium phosphate may be effected by one or more of the methods mentioned above or a dehydrothermal treatment, or any combination of these methods. A dehydrothermal treatment includes subjecting a substrate to a low pressure atmosphere at a raised temperature. The temperature in the dehydrothermal treatment may be of from 95°C to 135°C. The temperature may preferably be of from 100°C to 110°C, and most preferably of from 105°C to 110°C, if completion of the dehydrothermal treatment is desired in typically 18 to 36 hours. The temperature may preferably be of from 120°C to 135°C, and most preferably of from 125°C to 135°C, if completion of the dehydrothermal treatment is desired in typically 4 to 8 hours.

- 14 -

Preferably, the collagen and the glycosaminoglycan are crosslinked both prior to and subsequent to conversion of the brushite to octacalcium phosphate and/or apatite.

5 The processes of the present invention may comprise the step of shaping the composite biomaterial into a structure suitable for use as a bone or dental substitute. Such a step may occur after formation of the triple co-precipitate, but prior to any conversion of the brushite or crosslinking
10 of the collagen and glycosaminoglycan that may occur.

Alternatively, the step of shaping the biomaterial may occur subsequent to either the conversion of the brushite to apatite and/or octacalcium phosphate or crosslinking of the
15 collagen and the glycosaminoglycan.

Preferably, the composite material is shaped using a technique selected from (i) filtration and/or low temperature drying, (ii) freeze drying, (iii) injection
20 moulding and (iv) cold pressing. Filtration and/or low temperature drying, wherein the temperature is from 15°C to 40°C, most preferably of from 35°C to 40°C, typically results in a dense granular form of material. Freeze drying typically results in an open porous form. Injection
25 moulding results in a wide variety of shapes/morphologies of a material depending on the shape of the dye used. Cold pressing typically results in a dense pellet form.

The present invention further provides a precursor
30 material suitable for transforming into a synthetic biomaterial, said precursor material comprising a composite material comprising collagen, brushite and one or more

- 15 -

glycosaminoglycans. Preferably, the composite material comprises or consists essentially of a triple co-precipitate comprising collagen, brushite and one or more glycosaminoglycans. The triple co-precipitate may be
5 produced according to the process of the first aspect of the present invention.

The present invention also provides a composite biomaterial comprising collagen, brushite and one or more
10 glycosaminoglycans, which biomaterial is obtainable by a process according to the present invention as herein described.

The present invention also provides a composite
15 biomaterial comprising collagen, octacalcium phosphate and one or more glycosaminoglycans, which biomaterial is obtainable by a process according to the second aspect of the present invention.

20 The present invention also provides a composite biomaterial comprising collagen, apatite and one or more glycosaminoglycans, which biomaterial is obtainable by a process according to the third aspect of the present invention.

25

The present invention also provides a composite biomaterial comprising a triple co-precipitate of collagen, glycosaminoglycan and brushite.

30 The present invention also provides a biomaterial comprising particles of one or more calcium phosphate materials, collagen and one or more glycosaminoglycans,

- 16 -

wherein said collagen and said one or more glycosaminoglycans are crosslinked and form a matrix, said particles of calcium phosphate material are dispersed in said matrix, and said calcium phosphate material is selected
5 from one or more of brushite, octacalcium phosphate and/or apatite.

The following description relates to all aspects of the composite biomaterial according to the present invention
10 unless otherwise stated.

The collagen and the one or more glycosaminoglycans have preferably been crosslinked.

15 The collagen is preferably present in the material in an amount of from 5 to 90 (dry) wt%, more preferably from 15 to 60 (dry) wt%, %, more preferably from 20 to 40 (dry) wt%.

20 Preferably, the one or more glycosaminoglycans are present in the material in an amount of from 0.01 to 12 (dry) wt%, more preferably from 1 to 5.5 (dry) wt %, most preferably from 1.8 to 2.3 (dry) wt %.

25 Preferably, if the material comprises brushite, the ratio of collagen to brushite is 10:1 to 1:100 by weight (dry), more preferably 5:1 to 1:20 by weight (dry), most preferably 3:2 to 1:10 by weight (dry), for example 3:2 to 1:4 by weight (dry).

30 Preferably if the material comprises octacalcium phosphate, the ratio of collagen to octacalcium phosphate is 10:1 to 1:100 by weight (dry), more preferably 5:1 to 1:20

- 17 -

by weight (dry), most preferably 3:2 to 1:10 by weight (dry).

Preferably, the ratio of collagen to the total amount
5 of one or more glycosaminoglycans is from 8:1 to 30:1 by weight (dry), more preferably from 10:1 to 30:1 by weight (dry), still more preferably 10:1 to 12:1 by weight (dry), and most preferably 11:1 to 23:2 by weight (dry).

10 The composite biomaterial according to the present invention may be used as a substitute bone or dental material.

The present invention also provides a synthetic bone
15 material, bone implant, bone graft, bone substitute, bone scaffold, filler, coating or cement comprising a composite biomaterial of the present invention. The term coating includes any coating comprising the biomaterial or precursor of the present invention. The coating may be applied to the
20 external or internal surfaces of prosthetic members, bones, or any substrate intended for use in the human or animal body, which includes particulate materials. The composition of the present invention may be used for both in-vivo and ex-vivo repair of both mineralized biological material,
25 including but not limited to bone and dental materials. The biomaterials of the present invention may be used in the growth of allografts and autografts.

The biomaterial according to the present invention
30 comprising octacalcium phosphate may be free or essentially free of any of the precursor brushite phase. This biomaterial may comprise less than 2% by weight of brushite

- 18 -

in total amount of calcium phosphate materials in the biomaterial.

The calcium phosphate material may comprise or consist essentially of phase pure octacalcium phosphate or apatite. By phase pure, this means preferably containing at least 98%, more preferably at least 99%, and most preferably, at least 99.5% of the desired phase (as measured by x-ray diffraction). Alternatively, the biomaterial may comprise a mixture of octacalcium phosphate and apatite, depending on the desired properties of the biomaterial.

The material of the present invention comprising brushite may be used either as a precursor material for making a biomaterial, or may be suitable in itself for use as a biomaterial.

The processes according to the present invention may be preformed using the following sequential method, which may be applied in whole or in part, to produce biocomposites of collagen, one or more glycosaminoglycan and one or more calcium phosphate constituents. The following description is provided by way of example and is applicable to any aspect of the processes according to the present invention.

I: Triple Co-precipitation of Collagen, GAG, and the Calcium Phosphate Brushite at Acidic pH

This step is performed to initiate simultaneous formation, via precipitation from solution, of the three (or more) constituents of the composite, and to control the ratio of the three (or more) respective phases. Control of

- 19 -

the compositional properties of the composite (and in particular the collagen:GAG:CaP ratio) may be achieved by varying one or more of the pH, temperature, ageing time, calcium ion concentration, phosphorous ion concentration, collagen concentration and GAG concentration. The pH may be maintained constant (using, for example, buffers, pH-stat titration or other methods) or be allowed to vary. The possible secondary (contaminant) phases include other acidic calcium phosphates (e.g. monetite, calcium hydrogen phosphate) and complexes including by-products of titration and reactant addition (e.g. ammonium phosphate, ammonium nitrate). Additives to aid crosslinking (e.g. glucose, ribose) or to enhance in-vivo response (e.g. growth factors, gene transcription factors, silicon, natriuretic peptides) may also be added during this step.

II: Net Shape Formation

This step may be performed to produce the desired architecture of the final composite form, with particular emphasis on control of pore architecture. Examples of techniques include filtration and low-temperature drying (resulting in a dense granular form), freeze drying (resulting in an open porous form), injection moulding (resulting in a wide range of shapes depending on the type of dye) and cold pressing (resulting in a dense pellet form).

III: Primary Crosslinking

This step may be performed to preferably ensure that, when placed in a solution of elevated pH, the GAG content of

- 20 -

the composite does not elude rapidly, and, furthermore, to enhance the mechanical and degradation properties of the composite. Examples of techniques include low-temperature physical techniques (e.g. gamma irradiation, ultraviolet
5 radiation, dehydrothermal treatment), chemical techniques (e.g. non-enzymatic glycation with a simple sugar, glutaraldehyde, ethyl dimethylaminopropyl carbodiimide, nordihydroguaric acid), or combination methods (e.g. simultaneous non-enzymatic glycation and gamma-irradiation).
10 In the event that conversion to octacalcium phosphate (i.e. as in step IV) is desirable, primary crosslinking is advantageously performed at a temperature below about 37°C to prevent conversion of the brushite phase to its dehydrated form, monetite, which is a calcium phosphate that
15 does not readily hydrolyse to octacalcium phosphate.

IV: Hydrolysis

This step may be performed to partially or fully
20 hydrolyse the CaP phase from brushite (phase with high solubility at physiological pH) to octacalcium phosphate and/or apatite (phases with lower solubility at physiological pH), and to substantially remove any soluble contaminant phases (e.g. ammonium nitrate, calcium hydrogen
25 phosphate). In the case of hydrolysis to OCP, the selected pH is advantageously maintained constant at about 6.65 (using a buffer, pH stat, or other method), and the temperature at about 37°C for around 24-48 hours. As was the case in Step I, additives to aid in crosslinking (e.g.
30 glucose, ribose) or to enhance in-vivo response (e.g. growth factors, gene transcription factors, silicon, natriuretic

- 21 -

peptides) may also be added during the hydrolysis step (Step IV).

V: Secondary Crosslinking

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This step may be performed to further tailor the mechanical and degradation properties of the composite. Any or all of the crosslinking procedures listed in Step III above may be used to effect secondary crosslinking..

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The following Examples and the accompanying Figures are provided to further assist in the understanding the present invention. The Examples and Figures are not to be considered limiting to the scope of the invention. Any feature described in the Examples or Figures is applicable to any aspect of the foregoing description.

Example 1

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Example 1 is an example of the synthesis method described above, executed via application of steps I through III only. Triple co-precipitation is carried out at room temperature (20-25°C), at a pH of about 3.2 (maintained by titration with ammonia). In this example, co-precipitates are dried at 37°C and crosslinked via a dehydrothermal treatment. Neither hydrolytic conversion of the CaP nor secondary crosslinking is performed in this example.

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Materials

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- 22 -

Collagen: Reconstituted, pepsin-extracted porcine dermal collagen (atelocollagen); 85% Type I, 15% Type III; Japan Meat Packers (Osaka, Japan)

GAG: Chondroitin-6-sulphate from shark cartilage; sodium salt; Sigma-Aldrich Inc (St. Louis, MO, USA)

Calcium Sources: (i) Calcium hydroxide; $\text{Ca}(\text{OH})_2$ Sigma-Aldrich Inc (St. Louis, MO, USA), (ii) Calcium nitrate; $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$; Sigma-Aldrich Inc (St. Louis, MO, USA)

Phosphorous Source: Orthophosphoric acid; H_3PO_4 ; BDH

10 Laboratory Supplies (Poole, United Kingdom)

Titrant: Ammonia; NH_3 ; BDH Laboratory Supplies (Poole, United Kingdom)

Procedure

15

Step I

Solution A:

- $\text{Ca}(\text{OH})_2$ is dissolved in 0.48M H_3PO_4 to a concentration of 0.12M at room temperature, and the resulting solution
20 titrated to pH = 3.2 using ammonia.

Suspension B:

- Chondroitin-6-sulphate is dissolved in dionised water to a concentration of 3.2g/L. Under constant stirring,
Ca(NO₃)₂·4H₂O and Ca(OH)₂ is then added to the chondroitin
25 sulphate solution at a nitrate:hydroxide molar ratio of 1.5, to produce a suspension with a total calcium concentration of 2.4M.

- 0.144g collagen is added to 20mL of Solution A, and blended using a homogeniser until dissolved. 4mL of
30 Suspension B is then added to Solution A under constant stirring.

- 23 -

- Stirring is continued for 60 minutes, and pH monitored to ensure that it remains in the range $3.15 < \text{pH} < 3.30$. The resulting slurry is then allowed to age for 24 hours at room temperature.

5

Step II

- The slurry is allowed to dry at 37°C in air for 5 days, and the remaining triple co-precipitate rinsed with deionised water, and subsequently dried again at 37°C for an additional 24 hours.

10

- The x-ray diffraction pattern of the resultant triple coprecipitate is shown in Figure 1 (Cu-K(alpha) radiation) and an SEM image is shown in Figure 2.

15 Step III

Triple co-precipitates are crosslinked via dehydrothermal treatment (DHT) at 105°C, under a vacuum of 50 mTorr, for 48 hours. A TEM image of the triple co-precipitate following DHT is shown in Figure 3. Figure 4 shows the x-ray

20 diffraction pattern of the triple co-precipitate following DHT and indicates that the brushite phase has converted to its dehydrated form monetite.

Example 2

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Example 2 is an example of the synthesis method described above, executed via application of steps I through IV only. Triple co-precipitation is carried out at room temperature, and a pH of 4.0. In this example, pH control

30 is effected by careful control of the calcium hydroxide and calcium nitrate concentrations - an approach that also enables control of the mass ratio of brushite to collagen

- 24 -

plus GAG in the triple coprecipitate. The resulting triple co-precipitates are then frozen to -20°C , placed under vacuum and then heated to induce sublimation of unbound water (i.e. ice). Primary crosslinking is performed using a 1-ethyl 3-(3-dimethyl aminopropyl) carbodiimide treatment. The resulting dried triple coprecipitate is then converted to octacalcium phosphate via hydrolysis at a pH of 6.67 at about 37°C . In this example, secondary crosslinking is not performed.

Materials

Type I: Acid solubilised from bovine tendon
Integra Life Sciences Plainsboro, NJ, USA

GAG: Chondroitin-6-sulphate from shark cartilage; sodium salt; Sigma-Aldrich Inc (St. Louis, MO, USA)

Calcium Sources: (i) Calcium hydroxide; $\text{Ca}(\text{OH})_2$ Sigma-Aldrich Inc (St. Louis, MO, USA), and (ii) Calcium nitrate; $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$; Sigma-Aldrich Inc (St. Louis, MO, USA)

Phosphorous Source: Orthophosphoric acid; H_3PO_4 ; BDH Laboratory Supplies (Poole, United Kingdom)

Titrant: None

Crosslinking agents: (i) 1-Ethyl-3-(3-Dimethylaminopropyl) Carbodiimide (=EDAC); Sigma-Aldrich Inc (St. Louis, MO, USA), and (ii) N-Hydroxysuccinimide (=NHS); Sigma-Aldrich Inc (St. Louis, MO, USA)

Procedure

Step I

- A target mass ratio of brushite to collagen plus glycosaminoglycan of 1:1 is selected.

- 25 -

- The concentration of collagen plus GAG in a total reaction volume of 200mL is set at 21mg/mL.
- Using an empirical, 3-dimensional map of pH variation (produced at a constant $[\text{Ca}^{2+}]$ to $[\text{P}]$ reactant ion ratio of 1.0) with differing (i) ionic concentrations (i.e. $[\text{Ca}^{2+}] = [\text{H}_3\text{PO}_4]$) and (ii) ratios of calcium nitrate:calcium hydroxide, a locus of points over which pH remained constant at 4.0 is identified. This is shown in Figure 5 (sets of combinations of ionic concentration and calcium nitrate:calcium hydroxide ratio for maintaining pH = 4.0).
- Superimposing this locus of points onto a map of brushite mass yield with identical axes, and identification of its intersection with the 21 mg/mL contour allows the set of reactant concentrations for which a triple coprecipitate slurry containing a 1:1 mass ratio of calcium phosphate (21mg/mL) to collagen plus GAG (21mg/mL) can be produced at pH 4.0 ($[\text{Ca}^{2+}] = [\text{H}_3\text{PO}_4] = 0.1383\text{M}$; $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O} : \text{Ca}(\text{OH})_2 = 0.1356$) See Figure 6: identification of conditions for pH 4.0 synthesis of a triple coprecipitate slurry containing a 1:1 mass ratio of calcium phosphate to collagen plus GAG.
- 3.8644g collagen is dispersed in 171.4mL of 0.1383M H_3PO_4 cooled in an ice bath, by blending over 90 minutes at 15,000rpm, using a homogeniser equipped with a stator 19mm in diameter, to create a highly viscous collagen dispersion.
- 0.3436g chondroitin-6-sulphate (GAG) is allowed to dissolve in 14.3mL of 0.1383M at room temperature, by shaking periodically to disperse dissolving GAG, producing a GAG solution.
- After 90 minutes, the 14.3mL of GAG solution is added to the mixing collagen dispersion at a rate of approximately 0.5mL/min, under continuous homogenisation at 15,000rpm, and

- 26 -

the resulting highly-viscous collagen/GAG dispersion blended for a total of 90 minutes

- After 90 minutes of mixing, 1.804g Ca(OH)_2 and 0.780g $\text{Ca(NO}_3)_2 \cdot 4\text{H}_2\text{O}$ are added to the highly-viscous collagen/GAG

5 dispersion over 30 minutes under constant blending at 15,000rpm, creating a collagen/GAG/CaP triple coprecipitate slurry, after which time an additional 14.3mL of 0.1383M H_3PO_4 is blended into the slurry

- The pH of the triple coprecipitate slurry is approximately
10 4.0

- The triple coprecipitate slurry is allowed to remain at 25°C for a period of 48 hours.

Step II

15 - The triple coprecipitate slurry is placed in a freezer at -20°C and allowed to solidify overnight.

- The frozen slurry is then removed from the freezer, placed in a vacuum of approximately 80mTorr, and the temperature allowed to rise to room temperature, thus inducing
20 sublimation of ice from the slurry, which is allowed to proceed over 48 hours.

- The x-ray diffraction pattern of the collagen/GAG/brushite triple co-precipitate following removal of unbound water (Cu-K (alpha) radiation) is shown in Figure 7, and an SEM
25 image of the surface of a co-precipitate is shown in Figure 8 (secondary (SE) and backscattered electron (BSE) images of surface of triple co-precipitate with CaP: collagen + GAG = 1:1).

30 Step III

- 27 -

- After complete removal of unbound water, 1.25g of the resulting dry triple coprecipitate is hydrated in 40mL deionised water for 20 minutes.

5 - 20mL of a solution of 0.035M EDAC and 0.014M NHS is added to the container containing the triple coprecipitates and deionised water, and the triple coprecipitates allowed to crosslink for 2 hours at room temperature under gentle agitation.

10 - The EDAC solution is removed, and the triple coprecipitates rinsed with phosphate buffer solution (PBS) and allowed to incubate at 37°C for 2 hours in fresh PBS under mild agitation.

15 - After two hours in PBS, the triple coprecipitates are rinsed with deionised water, and allowed to incubate for two 10-minute intervals at 37°C under mild agitation.

- The triple coprecipitates are then dried at 37°C for 72 hours. Figure 9 shows an x-ray diffraction pattern of the collagen/GAG/brushite triple coprecipitate following EDAC crosslinking (Cu-K (alpha) radiation).

20

Step IV

- Crosslinked triple coprecipitate granules are placed in 50mL deionised water at 37°C, and the pH of the solution adjusted to 6.67 using ammonia.

25 - Temperature and pH are maintained constant for 48 hours, after which time the co-precipitates are filtered, rinsed in deionised water, and dried at 37°C in air.

- An x-ray diffraction pattern of the coprecipitates following conversion to OCP is shown in Figure 10 (EDAC-crosslinked collagen/GAG/CaP triple co-precipitate following
30 conversion at 37°C to OCP over 72 hours at pH 6.67, to form a collagen/GAG/OCP biocomposite, Cu-K (alpha) radiation).

- 28 -

Example 3

Example 3 is an example of the synthesis method described above, executed via application of steps I through V inclusive. Triple co-precipitation is carried out at room temperature, and a pH of about 4.5. As in example 2, pH control is effected by careful control of the calcium hydroxide and calcium nitrate concentrations, without the use of titrants. The resulting co-precipitates are then frozen to -20°C , placed under vacuum and then heated to induce sublimation of unbound water (i.e. ice). Primary crosslinking is performed using a 1-ethyl 3-(3-dimethyl aminopropyl) carbodiimide treatment. The resulting dried coprecipitate is then converted to apatite at pH 8.50, at 37°C . Secondary crosslinking performed using gamma irradiation.

Materials

20

Type I: Acid solubilised from bovine tendon

Integra Life Sciences Plainsboro, NJ, USA

GAG: Chondroitin-6-sulphate from shark cartilage; sodium salt; Sigma-Aldrich Inc (St. Louis, MO, USA)

25 Calcium Sources: (i) Calcium hydroxide; $\text{Ca}(\text{OH})_2$ Sigma-Aldrich Inc (St. Louis, MO, USA), and (ii) Calcium nitrate; $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$; Sigma-Aldrich Inc (St. Louis, MO, USA)

Phosphorous Source: Orthophosphoric acid; H_3PO_4 ; BDH

Laboratory Supplies (Poole, United Kingdom)

30 Titrant: None

Crosslinking agents: (i) 1-Ethyl-3-(3-Dimethylaminopropyl) Carbodiimide (=EDAC); Sigma-Aldrich Inc (St. Louis, MO, USA)

- 29 -

and (ii) N-Hydroxysuccinimide (=NHS); Sigma-Aldrich Inc (St. Louis, MO, USA)

Procedure

5

Step I

- A target mass ratio of brushite to collagen plus glycosaminoglycan of 3:1 is selected.
- The concentration of collagen plus GAG in a total reaction
10 volume of 200mL is set at 10mg/mL.
- Using an empirical, 3-dimentional map of pH variation (at a constant $[Ca^{2+}]$ to $[P]$ reactant ion ratio of 1.0) with differing i) ionic concentrations (i.e. $[Ca^{2+}] = [H_3PO_4]$) and ii) ratios of calcium nitrate: calcium hydroxide, a locus of
15 points over which pH remained constant at 4.5 is identified. This is shown in Figure 11 (set of combinations of ionic concentration and calcium nitrate: calcium hydroxide ratio for maintaining pH = 4.5).
- Superimposing this locus of points onto a map of brushite
20 mass yield (with identical axes), and identification of its intersection with the 30mg/mL (i.e. 3 times the concentration of collagen plus GAG) contour allows the set of reactant concentrations for which a triple coprecipitate slurry containing a 3:1 mass ratio of calcium phosphate
25 (30mg/mL) to collagen plus GAG (10mg/mL) can be produced at a pH of 4.5 ($[Ca^{2+}] = [H_3PO_4] = 0.1768M$; $Ca(NO_3)_2 \cdot 4H_2O$: $Ca(OH)_2 = 0.049$). This is show in Figure 12:
identification of conditions for pH 4.5 synthesis of a triple coprecipitate slurry containing a 3:1 mass ratio of
30 calcium phosphate to collagen plus GAG.
- 1.837g collagen is dispersed in 171.4mL of 0.1768M H_3PO_4 cooled in an ice bath, by blending over 90 minutes at

- 30 -

15,000rpm, using a homogeniser equipped with a stator 19mm in diameter, to create a collagen dispersion.

- 0.163g chondroitin-6-sulphate (GAG) is allowed to dissolve in 14.3mL of 0.1768M at room temperature, by shaking

5 periodically to disperse dissolving GAG, to produce a GAG solution.

- After 90 minutes, the 14.3mL of GAG solution is added to the mixing collagen dispersion at a rate of approximately 0.5mL/min, under continuous homogenisation at 15,000rpm, and
10 the resulting collagen/GAG dispersion blended for a total of 90 minutes.

- After 90 minutes of mixing, 2.498g Ca(OH)_2 and 0.380g $\text{Ca(NO}_3)_2 \cdot 4\text{H}_2\text{O}$ are added to the collagen/GAG dispersion over 30 minutes under constant blending at 15,000rpm, creating a
15 collagen/GAG/CaP triple coprecipitate slurry, after which time an additional 14.3mL of 0.1768M H_3PO_4 were added to the mixing slurry.

- The pH of the triple coprecipitate slurry is approximately 4.5.

20 - The triple coprecipitate slurry is allowed to remain at 25°C for a period of 48 hours.

Step II

- The triple coprecipitate slurry is placed in a freezer at
25 -20°C and allowed to freeze overnight.

- The frozen slurry is then removed from the freezer, placed in a vacuum of approximately 80mTorr, and the temperature allowed to rise to room temperature, thus inducing sublimation of the ice from the slurry, which is allowed to
30 proceed over 48 hours. The x-ray diffraction trace of the collagen/GAG/brushite triple co-precipitate following

- 31 -

removal of unbound water (Cu-K(alpha) radiation) is shown in Figure 13.

Step III

- 5 - After complete removal of unbound water, 1.25g of the resulting dry triple coprecipitate is hydrated in 40mL deionised water for 20 minutes.
- 20mL of a solution of 0.018M EDAC and 0.007M NHS is added to the container containing the triple coprecipitates and
10 deionised water, and the triple coprecipitates allowed to crosslink for 2 hours at room temperature, under gentle agitation.
- The EDAC solution is removed, and the triple coprecipitates are rinsed with phosphate buffer solution
15 (PBS) and allowed to incubate at 37°C for 2 hours in fresh PBS under mild agitation.
- After two hours in PBS, the triple coprecipitates are rinsed with deionised water, and allowed to incubate for two 10-minute intervals at 37°C under mild agitation.
- 20 - The triple coprecipitates are then dried at 37°C for 72 hours. The x-ray diffraction pattern of collagen/GAG/brushite triple coprecipitate following EDAC crosslinking (Cu-K(alpha) radiation) is shown in Figure 14.

25 Step IV

- Crosslinked triple coprecipitate granules are placed in 50mL deionised water pre-saturated with respect to brushite at 37°C, and the pH of the solution adjusted to 8.50 using ammonia.
- 30 - The temperature and pH are maintained constant for 72 hours, after which time the co-precipitates are filtered, rinsed in deionised water, and dried at 37°C in air. An x-

- 32 -

ray diffraction pattern of the co-precipitates following conversion to apatite is shown in Figure 15 (EDAC-crosslinked collagen/GAG/CaP triple co-precipitate following conversion at 37°C to apatite over 72 hours at pH 8.50, to form a collagen/GAG/apatite biocomposite (Cu-K(alpha) radiation)).

Step V

- The dried collagen/GAG/Ap triple coprecipitates are subjected to a 32.1kGy dose of gamma irradiation. Figure 16 shows the x-ray diffraction pattern following gamma irradiation (EDAC-crosslinked collagen/GAG/Ap triple coprecipitates after secondary crosslinking via gamma irradiation).

15

Example 4

Materials

20 Collagen: reconstituted, pepsin-extracted porcine dermal collagen (atelocollagen); 85% by weight of Type I, 15% by weight of Type III; Japan Meat Packers (Osaka, Japan)
GAG: Chondroitin-6-sulphate from shark cartilage; sodium salt; Sigma-Aldrich Inc (St. Louis, MO, USA)
25 Calcium Sources: (i) Calcium hydroxide; $\text{Ca}(\text{OH})_2$ Sigma-Aldrich Inc (St. Louis, MO, USA), and (ii) Calcium nitrate; $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$; Sigma-Aldrich Inc (St. Louis, MO, USA)
Phosphorous Source: Orthophosphoric acid; H_3PO_4 ; BDH Laboratory Supplies (Poole, United Kingdom)
30 Titrant: Ammonia; NH_3 ; BDH Laboratory Supplies (Poole, United Kingdom)

- 33 -

Procedure

Step I

- Solution A was prepared by dissolving Ca(OH)_2 in 0.48M
5 H_3PO_4 to a concentration of 0.12M at room temperature, and
the resulting solution titrated to pH of 3.2.

- Suspension B was prepared by dissolving Chondroitin-6-
sulphate in deionised water to a concentration of 3.2g/L.
Under constant stirring, $\text{Ca(NO}_3)_2 \cdot 4\text{H}_2\text{O}$ and Ca(OH)_2 then added
10 to chondroitin sulphate solution at a nitrate:hydroxide
molar ratio of 1.5, to produce a suspension with a total
calcium concentration of 2.4M.

- 0.144g collagen were added to 20mL of Solution A, and
blended using a homogeniser until dissolved. 4mL of
15 Suspension B was then added to Solution A under constant
stirring. Stirring was continued for 60 minutes, and pH
monitored to ensure that it remained in the range
3.15<pH<3.30. The resulting slurry was then allowed to age
for 24 hours at room temperature.

20

Step II

- The slurry was allowed to dry at 37°C in air for 5 days,
and the remaining triple co-precipitate rinsed with
deionised water, and subsequently dried again at 37°C for an
25 additional 24 hours.

Step III

- Co-precipitates were placed in dilute acetic acid (pH =
3.2), and irradiated with a gamma irradiation dose of 30kGy.
30 The crosslinked precipitates were then removed from
solution, rinsed, and dried at 37°C in air.

- 34 -

Step IV

- Crosslinked, co-precipitate granules were placed in 50mL deionised water at 37°C, and the pH of the solution adjusted to 6.65 using ammonia. Temperature and pH were maintained constant for 48 hours, after which the co-precipitates were filtered, rinsed in deionised water, and dried at 37°C in air.

Step V

- Crosslinked, hydrolysed, co-precipitate granules were placed in a vacuum oven at room temperature, and a vacuum of 50mTorr applied, after which the temperature was then increased to 105°C. After 24 hours, the temperature was reduced to room temperature and the vacuum released.

Figure 17 shows the x-ray diffraction pattern of the composite immediately following triple co-precipitation and drying (Steps I and II). This pattern confirms the major phase present to be brushite.

Figure 18 shows an SEM micrograph of the structure of co-precipitate granules following primary crosslinking (Step III). It is worthy to note the microstructurally homogeneous nature of the granules.

The progression of hydrolysis to octacalcium phosphate (Step IV) is illustrated in the XRD Pattern of Figure 19.

Progressive decreases in the intensity of the brushite peak at 12.5°, and increases of the major octacalcium

phosphate(OCP) peak at 4.5° indicate the conversion of the inorganic phase to OCP over a period of 48 hours.

- 35 -

A TEM image of the composite is shown in Figure 20. A random distribution of 10-20nm low aspect-ratio calcium phosphate crystals dispersed in a collagen/GAG matrix is evident.

5

The composite biomaterials of the present invention may be used as a bioresorbable material. Following implantation, it is expected that a device fabricated from the material would resorb completely, leaving behind only healthy,

10

regenerated tissue, with no remaining trace of the implant itself.

Claims:

1. A process for the production of a composite material comprising collagen, brushite and one or more
5 glycosaminoglycans, said process comprising the steps of providing an acidic aqueous solution comprising collagen, a calcium source and a phosphorous source and one or more glycosaminoglycans, and
precipitating the collagen, the brushite and the one or
10 more glycosaminoglycans together from the aqueous solution to form a triple co-precipitate.
2. A process as claimed in claim 1, wherein the solution has a pH of from 2.5 to 6.5, more preferably from 2.5 to
15 5.5.
3. A process as claimed in claim 2, wherein the solution has a pH of from 3 to 4.5.
- 20 4. A process as claimed in claim 3, wherein the solution has a pH of from 3.8 to 4.2.
5. A process as claimed in any one of the preceding claims, wherein the calcium source is selected from one or
25 more of calcium nitrate, calcium acetate, calcium chloride, calcium carbonate and calcium alkoxide, calcium hydroxide, calcium silicate, calcium sulphate, calcium gluconate and the calcium salt of heparin.
- 30 6. A process as claimed in any one of the preceding claims, wherein the phosphorus source is selected from one or more of ammonium-dihydrogen phosphate, diammonium

- 37 -

hydrogen phosphate, phosphoric acid, disodium hydrogen orthophosphate 2-hydrate and trimethyl phosphate.

7. A process as claimed in any one of the preceding
5 claims, wherein the one or more glycosaminoglycans are selected from chondroitin sulphate, dermatin sulphate, heparin, heparin sulphate, keratin sulphate and hyaluronic acid.

10 8. A process as claimed in any one of the preceding claims, wherein the solution has a temperature of from 4 to 50°C.

9. A process as claimed in any one of the preceding
15 claims, wherein the solution has a temperature of from 15 to 40°C.

10. A process as claimed in any one of the preceding claims wherein the ratio of collagen to the total amount of one or
20 more glycosaminoglycans in the solution is from 8:1 to 30:1 by weight.

11. A process as claimed in any one of the preceding claims, wherein the solution comprises calcium ions and the
25 ratio of collagen to the calcium ions is from 1:40 to 500:1 by weight, preferably from 1:40 to 250:1 by weight, more preferably from 1:13 to 5:4.

12. A process as claimed in any one of the preceding
30 claims, wherein ratio of collagen to brushite in the co-precipitate is from 10:1 to 1:100 by weight, preferably from 5:1 to 1:20.

- 38 -

13. A process as claimed in any one of the preceding claims wherein the solution comprises calcium ions and the concentration of calcium ions in solution is from 0.00025 to
5 1 M, preferably from 0.001 to 1 M.

14. A process as claimed in any one of the preceding claims wherein the solution comprises phosphate ions and the concentration of phosphate ions in solution is from is from
10 0.00025 to 1 M, preferably from 0.001 to 1 M.

15. A process as claimed in any one of the preceding claims wherein the concentration of collagen in the solution is from 1.0 to 20 g/L, more preferably 1.0 to 10 g/L.

16. A process as claimed in any one of the preceding claims wherein the total concentration of the one or more glycosaminoglycans in the solution is from 0.01 to 1.5 g/L, more preferably from 0.01 to 1 g/L.

20 17. A process for the production of a composite biomaterial comprising collagen, octacalcium phosphate and one or more glycosaminoglycans , said process comprising the steps of
providing a composite material comprising collagen,
25 brushite and one or more glycosaminoglycans, and
converting at least some of the brushite in the composite material to octacalcium phosphate by hydrolysis.

30 18. A process as claimed in claim 17, wherein the composite material comprises or consists essentially of a triple co-

- 39 -

precipitate comprising collagen, brushite and one or more glycosaminoglycans.

19. A process as claimed in claim 18, wherein the triple
5 co-precipitate is formed according to a process as defined in any one of claims 1 to 16.

20. A process as claimed in any one of claims 17 to 19,
wherein the step of hydrolysis of brushite to octacalcium
10 phosphate comprises contacting the composite material with an aqueous solution, said aqueous solution being at or above the pH at which octacalcium phosphate becomes thermodynamically more stable than brushite.

15 21. A process as claimed in claim 20, wherein said aqueous solution has a pH of from 6 to 8.

22. A process as claimed in claim 21, wherein said aqueous
20 solution has a pH of from 6.3 to 7.

23. A process as claimed in claim 22, wherein said aqueous
solution has pH of about 6.65.

24. A process for the production of a composite biomaterial
25 comprising collagen, apatite and one or more glycosaminoglycans, said process comprising the steps of
providing a composite material comprising collagen,
brushite and one or more glycosaminoglycans, and
converting at least some of the brushite in the
30 composite material to apatite by hydrolysis.

- 40 -

25. A process as claimed in claim 24, wherein the composite material comprises or consists essentially of a triple co-precipitate comprising collagen, brushite and one or more glycosaminoglycans.

5

26. A process as claimed in claim 25, wherein the triple co-precipitate is formed according to a process as defined in any one of claims 1 to 16.

10 27. A process as claimed in any one of claims 24 to 26, wherein the step of hydrolysis of brushite to apatite comprises contacting the composite material with an aqueous solution, said aqueous solution being at or above the pH at which apatite becomes thermodynamically more stable than
15 brushite.

28. A process as claimed in claim 27, wherein said aqueous solution has a pH of from 6.65 to 9, preferably from 7 to 8.5.

20

29. A process as claimed in any one of claims 17 to 28, wherein the conversion of brushite to octacalcium phosphate and/or apatite is carried out at a temperature of from 20 to 50°C, preferably from 30 to 40°C.

25

30. A process as claimed in claim 29, wherein said temperature is from 36 to 38°C.

31. A process as claimed in claim 30, wherein said
30 temperature is about 37°C.

- 41 -

32. A process as claimed in any one of the preceding claims further comprising the steps of crosslinking the collagen and the one or more glycosaminoglycans in the composite material or triple co-precipitate.

5

33. A process as claimed in claim 32, wherein, if at least some of the brushite is converted to octacalcium phosphate and/or apatite, the glycosaminoglycan is crosslinked prior to the conversion of the brushite to octacalcium phosphate and/or apatite.

10

34. A process as claimed in claim 33, wherein the crosslinking is effected by one or more of subjecting the triple co-precipitate to gamma radiation and/or ultraviolet radiation, non-enzymatic glycation with a simple sugar such as glucose, mannose, ribose or sucrose, contacting the triple co-precipitate with glutaraldehyde, ethyl dimethylaminopropyl carbodiimide and/or nor-dihydroguariaretic acid, and subjecting the triple co-precipitate to a dehydrothermal treatment.

15

20

35. A process as claimed in any one of claims 32 to 34, wherein, if at least some of the brushite is converted to octacalcium phosphate and/or apatite, the collagen and one or more of glycosaminoglycans are crosslinked subsequent to the conversion of the brushite to octacalcium phosphate and/or apatite.

25

36. A process as claimed in claim 35, wherein the crosslinking is effected by one or more of: subjecting the triple co-precipitate to gamma radiation, ultraviolet radiation or dehydrothermic treatment, non-enzymatic

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- 42 -

glycation with a simple sugar such as glucose, mannose, ribose and sucrose, contacting the triple co-precipitate with one or more of glutaraldehyde ethyl dimethylaminopropyl carbodiimide and/or nor-dihydroguaric acid.

5

37. A process as claimed in any one of claims 32 to 36, wherein the collagen and the one or more glycosaminoglycan are crosslinked both prior to and subsequent to conversion of the brushite to octacalcium phosphate and/or apatite.

10

38. A process as claimed in any one of the preceding claims, further comprising the step of shaping the composite biomaterial into a structure suitable for use as a bone or dental substitute.

15

39. A process as claimed in any claim 38, wherein the composite material is shaped using a technique selected from filtration and low temperature drying, freeze drying, injection moulding and cold pressing.

20

40. A precursor material for transforming into a synthetic biomaterial, said precursor material comprising a composite material comprising collagen, brushite and one or more glycosaminoglycans.

25

41. A precursor material as claimed in claim 40, wherein the composite material comprises or consists essentially of a triple co-precipitate comprising collagen, brushite and one or more glycosaminoglycans.

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- 43 -

42. A precursor material as claimed in claim 41, wherein said triple co-precipitate is produced according to a process as defined in any one of claims 1 to 16.

5 43. A composite biomaterial comprising collagen, brushite and one or more glycosaminoglycans, which biomaterial is obtainable by a process as defined in any one of claims 1 to 39.

10 44. A composite biomaterial comprising collagen, octacalcium phosphate and one or more glycosaminoglycans, which biomaterial is obtainable by a process as defined in any one of claims 17 to 39.

15 45. A composite biomaterial comprising collagen, apatite and one or more glycosaminoglycans, which biomaterial is obtainable by a process as defined in any one of claims 24 to 39.

20 46. A composite biomaterial comprising collagen, brushite and one or more glycosaminoglycans.

47. A composite biomaterial comprising collagen, octacalcium phosphate and one or more glycosaminoglycans.

25

48. A composite biomaterial as claimed in any one of claims 43 to 47 for use as a substitute bone or dental material.

49. A composite biomaterial comprising a triple co-precipitate of collagen, brushite and one or more glycosaminoglycans.

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- 44 -

50. A material as claimed in any one of claims 40 to 49, wherein the collagen and the one or more glycosaminoglycans have been crosslinked.

5 51. A material as claimed in any one of claims 40 to 50, wherein the collagen is present in the material in an amount of from 5 to 90 wt%, preferably from 15 to 60 wt%.

10 52. A material as claimed in any one of claims 40 to 51, wherein the one or more glycosaminoglycans are present in the material in an amount of from 0.01 to 12 wt%, preferably from 1 to 5.5 wt%.

15 53. A material as claimed in any one of claims 40 to 52, wherein, if the material comprises brushite, the ratio of collagen to brushite is 10:1 to 1:100 by weight, preferably from 5:1 to 1:20 by weight.

20 54. A material as claimed in any one of claims 40 to 52, wherein, if the material comprises octacalcium phosphate, the ratio of collagen to octacalcium phosphate is 10:1 to 1:100 by weight, preferably from 5:1 to 1:20 by weight.

25 55. A material as claimed in any one of claims 40 to 54, wherein the ratio of collagen to the total amount of one or more glycosaminoglycans is from 8:1 to 30:1 by weight.

30 56. A biomaterial comprising particles of one or more calcium phosphate materials, collagen and one or more glycosaminoglycans, wherein said collagen and said one or more glycosaminoglycans are crosslinked and form a matrix,

- 45 -

said particles of calcium phosphate material are dispersed in said matrix, and

said calcium phosphate material is selected from one or more of brushite, octacalcium phosphate and/or apatite.

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57. A synthetic bone material, bone implant, bone graft, bone substitute, bone scaffold, filler, coating or cement comprising a material as defined in any one of claims 43 to 56.

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