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(54) Title: CALCIUM PHOSPHATE COMPOSITION AND METHOD OF PREPARING SAME

(57) Abstract: A method for controlling the formation of a hydroxyapatite bone filler from dry calcium phosphate precursors in an aqueous solution uses coated sodium phosphate powder. The sodium phosphate powder is coated with a water soluble cellulose. Until the calcium dissolves in the aqueous solution setting of the cellulose phosphate cements proceeds slowly but when the exposed sodium phosphate particles start to solubilize in the aqueous solution the setting rate increases.

CALCIUM PHOSPHATE COMPOSITION AND METHOD OF PREPARING SAMEBACKGROUND OF THE INVENTION

The field of this invention is in the preparation of calcium phosphate minerals for bone cement or bone filler applications. More specifically, this invention relates to a calcium phosphate bone cement which consists of a mixture of tetra-calcium phosphate and di-calcium phosphate in an aqueous mixture, which mixture then sets to form a bone cement with a substantial portion of the cement being hydroxyapatite.

Hydroxyapatite is the major natural building block of bone and teeth. It has been found useful in fixing fractures and bone defects to use bone cements which are formed by combining calcium and phosphate precursors in an aqueous solution which initially forms a paste but then hardens into a hydroxyapatite bone cement. Hydroxyapatite has a calcium to phosphorous ratio of approximately 1.67 which is generally the same as the calcium phosphate ratio in natural bone structures. The paste may be placed *in situ* prior to setting in situations where bone has been broken, destroyed, degraded, become too brittle or has been the subject of other deteriorating effects. Numerous calcium phosphate bone cements have been proposed such as those taught by Brown and Chow in U.S. Reissue Patents 33,161 and 33,221 and by Constantz in U.S. Patents 4,880,610 and 5,047,031, the teachings of these patents are incorporated herein by reference.

It has been well known since the initial use of calcium phosphate cements that the addition of sodium phosphate solutions, potassium phosphate solutions or sodium carbonate solutions to the aqueous setting solution of the calcium phosphate precursors can speed setting times. This is documented in the Chow et al. April, 1991 IADR Abstract No.: 2410 and AADR, 1992 Abstract No.: 666 and was well known to those skilled in the art prior to these publications. In addition, such is discussed in Constantz et al. patents 5,336,264, 5,820,632 and 6,002,605.

Controlling bone cement or filler setting time is important because it is necessary for the material to be initially flowable and then moldable but it also must be able to set in place even though quantities of blood may be present. This blood creates problems in providing too much fluid so that the mineral salts are either washed away or dissolved prior to setting up. Obviously, if the material sets up too quickly, the surgeon has difficulty in mixing the calcium phosphate mineral cement and then placing it at the necessary bone site.

It has been found that the addition of sodium phosphate to the aqueous solutions mixed with the dry calcium phosphate precursors speeds the setting of calcium phosphate bone cement, such as those disclosed by Brown and Chow. However, the sodium phosphate may cause the setting to

occur more quickly than desired. Thus, a method for better controlling the setting time of such cements has been found desirable.

The above discussion of background art is included to explain the context of the invention. It is not to be taken as an admission or suggestion that any of the material referred to was published, known or part of the common general knowledge in Australia at the priority date of any of the claims of this specification.

Throughout the description and claims of this specification the word "comprise" and variations of that word such as "comprises" and "comprising" are not intended to exclude other additives, components, integers or steps.

SUMMARY OF THE INVENTION

It is an aspect of the invention to provide a method for better controlling the setting times of calcium phosphate cements.

It is an additional aspect of the invention to provide a method for coating dry highly water soluble sodium phosphate particles with a water soluble material, the thickness of which can be varied to control the time in which the sodium phosphate particle solubilize in the aqueous setting solution. The dry coated sodium phosphate particles are pre-mixed with the dry powdered calcium phosphate precursors to thereby better control setting times.

These aspects are achieved by a method for forming a calcium phosphate bone treatment material comprising mixing at least one dry calcium phosphate mineral power in an aqueous solution causing a reaction which forms a settable material. The rate of reaction is controlled by mixing, as a dry ingredient, a water soluble alkaline/acidic material, such as Na_2HPO_4 or a water soluble NaH_2PO_4 which materials have a water soluble coating, with the dry calcium phosphate mineral precursors prior to or simultaneously with the addition of the aqueous solution. The rate of reaction may be controlled by varying the thickness of the water soluble coating surrounding the water soluble alkaline/acidic material and may be controlled to a lesser degree by the concentration of the sodium phosphate in the mixture. The water soluble coating may be cellulose, such as cellulose acetate, cellulose acetate butyrate, hydroxypropyl cellulose, hydroxymethyl cellulose, carboxymethyl cellulose, hydroxyethyl cellulose, sodium ethyl cellulose sulfate, sodium carboxymethyl cellulose and combinations thereof. The alkaline/acidic material may be any material which is water soluble and forms a solution containing the phosphate ion.

The calcium phosphate mineral may be tetra-calcium phosphate, di-calcium phosphate, tri-calcium phosphate, mono-calcium phosphate and mono-calcium phosphate monohydrate, hydroxyapatite, either alone or in combinations.

5 A further aspect of the present invention provides a method for forming a calcium phosphate cement comprising:

providing a dry powder comprising at least one calcium phosphate mineral at least one soluble phosphate compound encapsulated within a water soluble coating; and

10 mixing the dry calcium phosphate mineral and the encapsulated phosphate compound with an aqueous solution which dissolves the alkaline materials and forms a basic solution which controls the setting time of the calcium phosphate cement.

Yet another aspect of the present invention provides a kit for forming a calcium phosphate cement comprising:

15 a dry powder comprising at least one calcium phosphate mineral and a soluble phosphate encapsulated within a water soluble coating in a first container; and

an aqueous solution in a second container.

DETAILED DESCRIPTION

20 In the preferred embodiment the calcium phosphate dry component is composed of tetra-calcium phosphate and di-calcium phosphate in a 1:1 molar ratio to which about 1% by weight of a powdered and coated sodium phosphate mineral has been added. The preferred sodium phosphate is either Na_2HPO_4 or NaH_2PO_4 which is coated with a cellulose coating by using the method described herein. The amount of sodium phosphate dry particles added, by weight, can be varied to vary the setting time. In the preferred embodiment, from .5% to 1.5% by weight of the coated

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sodium phosphate dry mineral powder is added to the dry calcium phosphate precursors. To ensure complete solubility of the sodium phosphate particles in the aqueous wetting solution, the initial size of the sodium phosphate powder to be coated should be less than 100 μ in size with the coated particles being no more than 100 μ microns in size.

- 5 The amount of coated sodium phosphate, by weight, in the powdered component must be varied depending on the liquid to powder ratio used in the bone cement. For example, if the liquid to powder ratio of aqueous solution to calcium phosphate coated sodium phosphate (by weight) is .25 and approximately 10 grams total of calcium phosphate/coated sodium phosphate dried powder are used (*i.e.* the 10 grams including .75% by weight coated sodium phosphate powder), then if the
10 liquid to powder ratio is raised to .3 (*i.e.* 20% above the .25 ratio) then the sodium phosphate, by weight, must be increased 20% in order to maintain the same setting time (*i.e.* to .9%).

- Furthermore, setting times can be varied by varying the thickness of the coating of the sodium phosphate particles. Having a somewhat thicker coating results in a somewhat longer time for the coating to dissolve thereby releasing the phosphate ions. The increase in coating thickness
15 can be accomplished by increasing the amount of cellulose acetate butyrate in the solution or allowing the solution to sit for a longer period of time (30 to 60 minutes rather than 15 to 30 minutes) after the sodium phosphate particles are added.

 Example 1 details the process for coating the sodium phosphate particles with the results of the calcium phosphate cement setting times for the coated and uncoated particles listed in Table 1.

- 20 Example 1

- Into a round bottom flask were added 150 ml of ethyl acetate and 2.0 g of cellulose acetate butyrate particles (CAB, MW = 200,000, butyrate content = 17%, from Acros/Fisher) stirring continuously. A water bath was placed under the flask for cooling it to below room temperature (25°C). The dry ingredients were allowed to dissolve for 2-3 hours to form a homogenous solution.
25 Then 1.0 g of sodium phosphate particles less than 100 μ in size were added and suspended in the solution for about 15-30 minutes, continuing the stirring. Then 150-200 ml of n-Hexane was added dropwise at the rate of 80-100 drops per minute for 5-10 minutes continuing the stirring. After the addition of the n-Hexane, ice water was added to the bath to help harden the formed microcapsules. After 5-10 minutes, cold (below room temperature) n-Hexane was added to wash loose particles off
30 the microcapsules. The microcapsules are recovered by decantation, washed with cold n-Hexane and air or vacuum dried.

 The process of Example 1 was done for both NaH₂PO₄ and Na₂HPO₄ particles (Na-Phos) less than 100 μ in size. The effect of coated vs. uncoated phosphate powders on setting time of a

tetra-calcium phosphate plus di-calcium phosphate cement as taught in Reissue Patent 33,161 are shown below in Table 1:

Table 1

For NaH_2PO_4 powder:

Liquid/Powder Ratio*	Set Time (Coated Na-Phos Powder)	Set Time (Uncoated Na-Phos Powder)
0.25	40 min.	12 min.
0.30	>60 min.	22 min.

For Na_2HPO_4 powder:

Liquid/Powder Ratio*	Set Time (Coated Powder)	Set Time (Uncoated Powder)
0.25	39 min.	18 min.
0.30	>70 min.	26 min.

*Liquid/"total powder" ratio with the "total powder" consisting of tetra-calcium phosphate, di-calcium phosphate and 0.75% sodium phosphate powder by weight.

As can be seen from Table 1, setting times were dramatically increased by using the sodium phosphate particles coated by the method of the present invention with cellulose acetate butyrate versus uncoated particles. As is well known in the prior art whether the sodium phosphate particles are included in the cement as dried particles or previously solubilized in the aqueous component, it is the concentration of the sodium and phosphate ions in solution which determines the setting times. The use of a coating delays the release of the sodium and phosphate ions thereby retarding the setting time by allowing a slow setting initial phase followed by a faster setting phase (after the coating dissolves and the sodium phosphate solubilizes). This method of controlling setting time is superior to varying the concentration of uncoated sodium phosphate which works to a limited extent but as the concentration of phosphate ions in solution decreases, the affect on setting time becomes erratic and less robust. With the present invention, the concentration of phosphate ions can be kept constant by mixing coated sodium phosphate particles having coatings of varying thicknesses or mixing coated and uncoated particles in the dry calcium phosphate cement precursors.

Although the invention herein has been described with reference to particular embodiments, it is to be understood that these embodiments are merely illustrative of the principles and applications of the present invention. It is therefore to be understood that numerous modifications may be made to the illustrative embodiments and that other arrangements may be devised without departing from the spirit and scope of the present invention as defined by the appended claims.

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A method for forming a calcium phosphate bone treatment material comprising:
 - reacting at least one dry calcium phosphate mineral with an aqueous solution
 - causing a reaction which forms a settable material; and
 - controlling the rate of the reaction by mixing as a dry ingredient a water soluble phosphate compound having a water soluble coating with the calcium phosphate mineral and aqueous solution.
2. The method as set forth in claim 1 where the reaction is controlled by varying the thickness of the coating surrounding the water soluble alkaline material.
3. The method as set forth in claim 2 wherein the water soluble coating is cellulose.
4. The method as set forth in claim 3 having a coating selected from the group consisting of cellulose acetate, cellulose acetate butyrate, hydroxypropyl cellulose, hydroxymethyl cellulose, carboxymethylethyl cellulose, hydroxyethyl cellulose, sodium ethyl cellulose sulfate, sodium carboxymethyl cellulose and a combination thereof.
5. The method as set forth in any one of claims 1 to 4 wherein the coated phosphate compound is selected from the group consisting of potassium phosphates, sodium phosphates and a combination thereof.
6. The method as set forth in claim 5 wherein the at least one calcium phosphate includes tetra-calcium phosphate.
7. The method as set forth in any one of claims 1 to 6 wherein the calcium phosphate mineral includes at least one member selected from the group of tetra-calcium phosphate, di-calcium phosphate, tri-calcium phosphate, mono-calcium phosphate monohydrate, hydroxyapatite and a combination thereof.
8. The method as set forth in any one of claims 1 to 7 wherein the coated water soluble phosphate compound makes up .5 to 1.5% of the combined weight of the dry ingredients.
9. A method for forming a calcium phosphate cement comprising:
 - providing a dry powder comprising at least one calcium phosphate mineral at least one soluble phosphate compound encapsulated within a water soluble coating; and
 - mixing the dry calcium phosphate mineral and the encapsulated phosphate compound with an aqueous solution which dissolves the alkaline materials and forms a basic solution which controls the setting time of the calcium phosphate cement.

10. The method as set forth in claim 9 where the reaction is controlled by varying the thickness of the coating surrounding the water soluble phosphate compound.

11. The method as set forth in claim 9 wherein the water soluble coating is cellulose.

5 12. The method as set forth in claim 11 having a coating selected from the group consisting of cellulose acetate, cellulose acetate butyrate, hydroxypropyl cellulose, hydroxymethyl cellulose, carboxymethylethyl cellulose, hydroxyethyl cellulose, sodium ethyl cellulose sulfate, sodium carboxymethyl cellulose and a combination thereof.

10 13. The method as set forth in any one of claims 9 to 12 wherein the coated phosphate compound is selected from the group consisting of potassium phosphates, sodium phosphates and a combination thereof.

14. The method as set forth in claim 13 wherein the at least one calcium phosphate includes tetra-calcium phosphate.

15 15. The method as set forth in any one of claims 9 to 14 wherein the calcium phosphate mineral includes a member selected from the group of tetra-calcium phosphate, di-calcium phosphate, tri-calcium phosphate, mono-calcium phosphate monohydrate, hydroxyapatite and a combination thereof.

20 16. The method as set forth in any one of claims 9 to 15 wherein the coated water soluble phosphate compound makes up .5 to 1.5% of the combined weight of the dry ingredients.

17. A kit for forming a calcium phosphate cement comprising:
a dry powder comprising at least one calcium phosphate mineral and a soluble phosphate encapsulated within a water soluble coating in a first container; and
an aqueous solution in a second container.

25 18. The kit as set forth in claim 17 where the reaction is controlled by varying the thickness of the coating surrounding the water soluble alkaline material.

19. The kit as set forth in claim 17 wherein the water soluble coating is cellulose.

30 20. The kit as set forth in claim 17 or claim 19 having a coating selected from the group consisting of cellulose acetate, cellulose acetate butyrate, hydroxypropyl cellulose, hydroxymethyl cellulose, carboxymethylethyl cellulose, hydroxyethyl cellulose, sodium ethyl cellulose sulfate, sodium carboxymethyl cellulose and a combination thereof.

21. The kit as set forth in any one of claims 17 to 20 wherein the phosphate material is selected from the group consisting of potassium phosphates, sodium phosphates and a combination thereof.

22. The kit as set forth in claim 19 wherein the at least one calcium phosphate includes tetra-calcium phosphate.
23. The kit as set forth in any one of claims 17 to 21 wherein the calcium phosphate mineral includes a member selected from the group of tetra-calcium phosphate, di-calcium phosphate, tri-calcium phosphate, mono-calcium phosphate monohydrate, hydroxyapatite and a combination thereof.
24. The method as set forth in any one of claims 17 to 23, wherein the coated water soluble phosphate compound make up .5 to 1.5% of the combined weight of the dry ingredients.
25. A calcium phosphate bone treatment material prepared by the method of claim 1.
26. A calcium phosphate cement prepared by the method of claim 9.
27. A method for forming a calcium phosphate bone treatment material according to claim 1 substantially as hereinbefore described.
28. A method for forming a calcium phosphate cement according to claim 9 substantially as hereinbefore described.
29. A kit for forming a calcium phosphate cement according to claim 17 substantially as hereinbefore described.

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