Materials Design and Osteoinduction Characteristics of Biomimetic and Functionally Graded Apatites

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Abstract: Bioabsorbable and functionally graded apatites (fg-HAp) were designed using bovine bone by the calcination and partial dissolution-precipitation methods. The fg-HAp ceramics had the specified surface structure and pore size distribution derived from living body. The pore structure of the ceramics with specific surface areas of 30-40 m²g⁻¹ and porosities of 60-80 % was divided into a macro-pore originating from spongy bone and a micro-pore related to body fluid permeation and blood permeability. Recombinant human bone morphogenetic protein-2 (rhBMP-2) solution was added to fg-HAp ceramics to obtain rhBMP-2-loaded fg-HAp (rhBMP-2/fg-HAp) ceramics exhibiting osteoinduction. The fg-HAp and rhBMP-2/fg-HAp ceramics were implanted into the subcutaneous tissue of rats. At 4 weeks after the implantation, for the fg-HAp, body fluid and albumin permeated the bulk region of the ceramics through the micro-pores, and surface- and bulk-degradations of the HAp grains proceeded. At 12 weeks, the fg-HAp ceramics were completely biodegraded and bioabsorbed. For the rhBMP-2/fg-HAp, at 4 weeks after the implantation, some pieces of fg-HAp were incorporated into the induced bone and fatty marrow, suggesting that osteoinduction occurred in conjunction with bone remodeling. The rhBMP-2/fg-HAp ceramics would be newly biomimetic materials with excellent blood permeability and osteoinduction.

Key words: Bioabsorption; Biodegradation; Blood permeability; Functionally graded apatites; Osteoinduction

Introduction

Hydroxyapatite (HAp:Ca₁₀(PO₄)₆(OH)₂) has been widely used for dental and medical implants because of its excellent biocompatibility and osteoconductivity. In recent years, coping with necessity for patients, i.e., implanting location or situation, development and application of tailored HAp ceramics which can take a reasonable balance between absorption rate and mechanical strength in a living body have been expected. HAp ceramics synthesized from reagents show very low bioabsorption characteristics in implanted regions even after several years. Therefore, commercial HAp ceramics cannot always be satisfactory materials for bone regeneration and long-time use. Responding to the background, we focus on the development of biomimetic HAp ceramics having controlled bioabsorption and bone remodeling. HAp ceramics made of natural bone have physical and chemical properties of raw materials such as pore structure and small amounts of metal ions. For instance, bovine bone-originated HAp (designated as b-HAp), which was calcined at 800-1100 °C, can give high porosities and interconnecting pore size as well as appropriate mechanical strength. Surface modification of the calcined HAp ceramics might bring the control of absorption rate of HAp ceramic blocks in a living body.

Bone morphogenetic protein (BMP) is known as one of bone formation factors accelerating osteoinduction, and it has been added to various scaffold materials because of its novel ability to induce bone and cartilage in intramuracular and subcutaneous tissues. Oda et al. and Takahashi et al. demonstrated that interconnected porous HAp and calcium phosphate ceramics would be superior carriers of recombinant human BMP-2 (rhBMP-2) in ectopic and orthotopic sites. Gradations in the grain size and crystallinity of fg-HAp plus rhBMP-2-addition might achieve the desired bio-absorption and osteoinduction characteristics.

The first purpose of this study is to securely design biomimetic and functionally graded HAp (designated as fg-HAp) ceramics using bovine bone by the calcination and partial dissolution-precipitation methods. The second purpose is to design rhBMP-2-loaded fg-HAp (designated as rhBMP-2/fg-HAp) ceramics with an excellent osteoinduction. The third purpose is to histologically and immunohistochemically evaluate the bioabsorption and osteoinduction characteristics of the two ceramics.

Materials and Method

Design of fg-HAp ceramics derived from bovine bone

Spongy and cortical bovine femur bones were used as starting materials, boiled and calcined at 800-1100 °C for 24h in air to obtain crystalline b-HAp ceramics. By the calcination process, all prion proteins and organic residues of bovine bone were completely burned out. The pulverized cortical b-HAp was completely dissolved into a HNO₃ solution, while the spongy b-HAp was partially dissolved into another HNO₃ solution. After the solutions were mixed, NH₄ solution was added and reprecipitated HAp (designated as r-HAp) crystals were carefully precipitated on macro-pores and micro-pores of spongy b-HAp at pH 10.5 and 25 °C. The modified spongy b-HAp was aged for 24h to fabricate a structure of fg-HAp. The fg-HAp ceramics were filtrated and washed with distilled water and dried at 120 °C. To obtain rhBMP-2/fg-HAp ceramics, rhBMP-2(5µg) (supplied by Yamanouchi Pharmaceutical Co Ltd, Tokyo, Japan) solution was added to fg-HAp ceramics blocks (3x3x3mm) in a sterilized dish.
Characterization of biomimetic fg-HAp ceramics

The crystalline phase of the samples was identified by Micro-X-ray diffraction (Micro-XRD) using Cu Kα1 radiation. Scanning electron microscopic (SEM) photographs of the fg-HAp ceramics were taken to observe the morphology and microstructure. The porosities of the ceramics were determined by the water displacement method. The BET-specific surface areas were measured by N2-adsorption at –196 °C. The curves of pore size distribution were measured by the mercury permeation method for macro-pores and the N2-adsorption method for micro-pores.

Histological and immunohistochemical observations of fg-HAp and rhBMP-2/fg-HAp explants

The fg-HAp and rhBMP-2/fg-HAp ceramics blocks were implanted into the subcutaneous tissues of the back region in 4-week-old male Wistar rats. In all procedures, the international guidelines for experiments on animals were followed. At 4, 8, and 12 weeks after implantation, these samples were explanted. The specimens were fixed in neutral buffered formalin, decalcified with formic acid, embedded in paraffin, sectioned and stained with hematoxylin and eosin (HE). They were histologically evaluated using an optical microscope. To investigate the eosinophilic areas in HE sections of the implanted ceramics blocks, immunohistochemical staining was carried out. The specimens were immunostained with polyclonal rabbit antibody against rat albumin (IgG fraction rabbit anti-rat albumin, Inter-Cell Technologies, Inc., USA) using the avidin-biotin complex method. The sections were treated with 3,3′-diaminobenzidine in Tris buffer solution containing hydrogen peroxide for visualization of reaction products. Incubations with normal rabbit IgG or without the primary antibody were used as negative controls.

Results

Characterization of biomimetic fg-HAp ceramics with the specified pore structure

Appearance of biomimetic fg-HAp ceramics blocks (3x3x3mm) is shown in Fig.1. The fg-HAp ceramics blocks had sufficient strength for handling and operation. The Micro-XRD analyses indicated that the degree of crystallinity of the single phase HAp was gradually distributed from the surface layer of the macro-pore wall to the bulk region of the b-HAp body structure. Figure 2 shows SEM photographs of the fg-HAp ceramics prepared by the calcination at 800 °C and partial dissolution-precipitation. Macro-pore sizes of 100-600 μm (Fig.1a) and grain sizes of 0.3-0.5 μm in the surface and the macro-pores (Fig.1b) were observed. When the r-HAp crystals were precipitated on the pore wall surface in the macro-pores and micro-pores of the b-HAp, the microstructures of the fg-HAp ceramics would change to spherical moss-like grains forming needle-like microcrystals of the r-HAp. The specific surface areas, total pore volumes, and porosities of the ceramics were 30-40 m² g⁻¹, 0.379-0.390 cm³ g⁻¹, and 60-80 %, respectively. Comparing with the curves of the pore size distribution for the b-HAp and fg-HAp ceramics, pore volumes in the micro-pore sizes of 10-160 nm were clearly recognized for only the fg-HAp ceramics.

Bioabsorption and osteoinduction of fg-HAp and rhBMP-2/fg-HAp ceramics

For the fg-HAp ceramics blocks, at 4 weeks after the implantation, body fluid extensively permeated into the bulk regions of HAp through the nano-micro-pores of the ceramics. Surface- and bulk-degradations of the HAp proceeded, so that a total size of the HAp block remarkably decreased. The volumetric populations occupied by body fluid were 60 % at 4 weeks and 68 % at 8 weeks in the ceramics explants, indicating drastic bioabsorption. At 12 weeks, the fg-HAp ceramics were completely biodegraded and bioabsorbed. The bioabsorption rate of the fg-HAp ceramics was much faster than that of the commercial porous β-TCP ceramics. No bone or cartilage was seen in any of the specimens. Investigating the eosinophilic region of HE sections marked in the fg-HAp ceramics, the immunohistochemical staining was carried out. The immunostaining of albumin in the fg-HAp explant tissue indicated to be immunopositive.

Figure 3 shows photomicrographs of histological sections at 4 weeks after the implantation of the rhBMP-2/fg-HAp ceramics. The HAp fragmented by body fluid permeation and new bone formation from surface and bulk regions of the ceramics were observed. The HAp, which had been cut into small pieces, was incorporated into the induced bone with fatty marrow, suggesting that osteoinduction occurred with bone remodeling. The morphometric analysis conducted after the implantation indicated that the volumetric populations were 27% for HAp, 59% for bone and marrow, and 14% for connective tissue.

Discussion

The calcination and partial dissolution-precipitation method proposed in this study is expected to produce a structure of fg-HAp, in which the composition ratios of r-HAp/b-HAp decrease from the surface layer of pore wall to a bulk region of the fg-HAp body in the micropore matrix, forming gradations in both the crystallinity and the grain size of HAp. The fg-HAp ceramics with activated pore surfaces had a significantly higher specific surface area than the b-HAp ceramics. In the curves of the pore size distribution for the fg-HAp ceramics, pore volumes in the micro-pore sizes of 10-160 nm were surely recognized. The specified pore structure may be constructed in the macro-pores. Based on these results, it was found that the pore structure of the fg-HAp was classified into a macro-pore (100-600 μm) originating from spongy bone and a micro-pore (10-160 nm) related to body fluid permeation and blood permeability. The structure of fg-HAp ceramics can permit body fluid to easily permeate the parts of a living body where it is used because of the number of graded micro-pores with nano-order. The easy permeation of body fluid into the fg-HAp structure implies better blood permeability, together with albumin included in blood. Microcracks in bulk regions of the ceramics formed by partial dissolution with HNO₃ will be related to auto-degradation, body fluid permeation, and cellular invasion and proliferation. Consequently, the bioabsorbable and biomimetic fg-HAp ceramics may improve the blood permeability and bioabsorbability of bioceramics to advance the permeation rate of serum proteins in blood.

Conclusions

The absorbable fg-HAp ceramics with the specified pore structures and gradations in both crystallinity and grain size were newly developed by the calcination and partial dissolution-precipitation methods. After implantation into the subcutaneous tissue of the back region of 4-week-old Wistar rats, the fg-HAp ceramics gave excellent bioabsorption characteristics because of the surface- and bulk-degradations of HAp grains, which were successively followed by body fluid permeation, blood permeability and albumin-adsorption. Concerning the rhBMP-2/fg-HAp ceramics, permeation and diffusion of body fluid and active bone formation were observed at significant levels within 4 weeks of implantation. Accordingly, the rhBMP-2/fg-HAp ceramics can be a new biomimetic implant material exhibiting cellular absorption by giant cells or release characteristics of BMP, since the gradation conditions in crystallinity and grain size, accompanied by the needle-like pore wall of the fg-HAp would
effectively control the bioabsorption rate and the permeation rate of body fluid.

Reference

Fig. 1. Appearance of fg-HAp ceramics (3mm~3mm~3mm).

Fig. 2. SEM photographs of fg-HAp ceramics prepared by the calcination at 800•• for 24h and partial dissolution-precipitation. (a) Macro-pores with 100-600 µm were observed. (b) Spherical moss-like grains with 0.3-0.5 µm forming needle-like microcrystals were seen in the surface and the macro-pores.

Fig. 3. Photomicrographs of HE sections at 4 weeks after the implantation of rh BMP-2/fg-HAp ceramics. B: bone, H: HAp containing body fluid, F: fatty marrow, : fragmentized HAp, : osteoinduction with bone remodeling. HAp fragmentized by body fluid permeation and new bone formation from surface and bulk regions of the ceramics were observed, while HAp incorporated by the induced bone with fatty marrow were found.