Evaluation of Porcine Hybrid Bone Block for Bone Grafting in Dentistry

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Abstract. Background/Aim: The purpose of this study was to develop hybrid bone blocks using porcine-derived collagen and low crystalline porcine-derived hydroxyapatite to overcome the disadvantages of commonly used bone grafts in dentistry. Materials and Methods: Collagen was added to hydroxyapatite particles to increase the spatial integration of particulate bone grafts. Physicochemical examination and in vivo tests were performed to analyze scaffold’s characteristics and evaluate bone regeneration. Results: Porcine hybrid bone block had an irregular and interconnecting macroporous structure that was adequate for bone regeneration and bone ingrowth, and showed a good space-occupying ability to become well positioned. In addition, it showed higher angiogenesis and biodegradability than Bio-Oss Collagen®, a commercialized bone graft used in dental clinics. Conclusion: Our results suggest that improved collagen hybrid bone block can be generated when porcine cancellous bone particles and collagen were reasonably mixed. This hybrid bone block was easy in handling had flexibility, good biodegradability and provided bone regeneration.

Bone graft materials are necessary to fill large bone defects in dentistry. Among the various types of bone grafts, the particle type is commonly used because it enables dense packing in an intraosseous defect with irregular forms (1). However, particulate bone grafts are susceptible to external compressive force and are not easy to be well localized within the defect. Hence, a barrier membrane is required to maintain the grafted particulate bone within the defect and prevent collapse of the site. Because the use of a barrier membrane increases costs and operation time, bone substitutes that can be adjusted in the bone defect without a membrane should be developed (2).

A combination of organic components such as collagen and inorganic particles is a promising strategy for ideal bone grafting because collagen and hydroxyapatite are the main components of bone (3). Collagen is the most abundant protein in mammals, and provides structural and mechanical support to tissues with good biodegradability and biocompatibility (4). Collagen has been widely used as a scaffold in medical applications and tissue engineering, such as for skin and vascular grafts, hemostats, adhesives, and drug delivery (5). Despite its excellent biological properties, collagen has poor mechanical strength and must be cross-linked by physical or chemical treatments, or combined with other polymers or inorganic materials.

Bio-Oss Collagen® (BC) shows enhanced handling characteristics and provides good hemostatic properties by mixing 90% Geistlich Bio-Oss® granules with 10% porcine collagen. Several preclinical and clinical studies have shown that Bio-Oss Collagen® exhibits favorable osteoconductivity and space-making properties. However, Bio-Oss® granules tend to be absorbed very slowly and remain in the new bone tissue. Moreover, the use of bovine-derived materials such as Bio-Oss Collagen® poses a risk of bovine spongiform encephalopathy (BSE).

The aim of the present study was to develop porcine hybrid bone block (PHBB) with characteristics such as easy handling, flexibility, good biodegradability and free from BSE. To confirm the clinical efficacy of the PHBB, we compared bone regeneration and biodegradability of PHBB with those of BC.
Materials and Methods

Preparation of porcine hybrid bone block. Type I collagen was extracted from porcine skin. Briefly, the porcine dermis homogenized with pepsin and 0.5 M acetic acid was incubated for 24 h at 4°C. The insoluble parts were removed and salted out with 5 M NaCl. Pellets were obtained by centrifugation and dialyzed against distilled water for 24 h. The pure collagen was freeze dried for 3 days.

Porcine cancellous bone was trimmed and cut into pieces. Cancellous bones were defatted with toluene and deproteinized with ethylenediamine. The results were subjected to heat treatment at 350°C.

The extracted porcine type I collagen was dissolved in 0.05 M acetic acid and adjusted to pH 5±0.5 with 1 M NaOH. The collagen solution was cross-linked with 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide and its core agent N-hydroxysuccinimide (both Sigma-Aldrich, St. Louis, MO, USA) and then dialyzed against distilled water for 24 h.

The cross-linked collagen solution was mixed to porcine cancellous bone particles (250-850 μm) at a ratio of 1:9 (w/w) and freeze-dried for 2 days (PHBB).

Both BC and PHBB were prepared in the shape of a disc of diameter 8 mm, weighing 25 mg. The discs were sterilized with ethylene oxide gas and aerated for animal studies.

Characterization of PHBB. BC and PHBB were imaged by low-voltage scanning electron microscopy (LV-SEM, JEOL, Tokyo, Japan) to compare their surface morphology.

Chemical characterization of PHBB was carried out by obtaining infrared spectra of BC, PHBB, porcine cancellous bone, and type I collagen using ATR-FTIR spectroscopy (Bruker TENSOR 37, Bruker AXS, Inc., Germany). The transmission and ATR spectra of each sample were recorded from 4,000 and 600 cm\(^{-1}\) with 64 scans at a resolution of 4 cm\(^{-1}\).

The evaluation of mechanical properties of PHBB was performed at room temperature with a texture analyzer (TMS-Pro, Food Technology Corporation, Sterling, VA, USA). BC and PHBB samples (4×6×7 mm) were immersed in PBS for hydration for 1 h, and then loaded in uniaxial compression at a displacement rate of 10 mm/min to achieve compression of ~50% of the original sample height. Moreover, flexibility of the PHBB was evaluated by checking resilience of the wet sample after the removal of compression.

Animals. Eight-week-old male Sprague-Dawley rats (weighing 250–280 g, SamTako, Osan, Korea) were used in this study. Animals were housed in polycarbonate cages and kept under standard conditions (23±2°C, 50±10% humidity, 12/12 light-dark cycle, with access to water and food ad libitum). The animal care and surgical protocol were approved by the Institutional Animal Care and Use Committee of Chonnam National University and carried out in accordance with the Guidelines for Animal Experiments of Chonnam National University (CNU IACUC-YB-R-2016-42).

Animal surgery. Animals were anesthetized using 10 mg/kg xylazine (Rompun, Bayer, Leverkusen, Germany) and 40 mg/kg ketamine (Ketamine, Yuhan Co., Seoul, Korea) by intraperitoneal injection. The parietal region of rats was shaved and disinfected with 10% povidone-iodine and 70% EtOH, and then the cranium was exposed by incision of the skin and periosteum. An 8-mm critical sized calvarial defect was created using a trephine bur, and then BC or PHBB was grafted into the defect site. The periosteum was closed using 4-0 absorbable sutures (Surgisorb®, Samyang, Seoul, Korea), and then 3-0 non-absorbable sutures (Dafilon®, B.Braun, Melsungen, Germany) were used to close the skin. Five animals in each group were sacrificed by CO\(_2\) at 4 and 8 weeks after operation.

Micro-CT analysis. After harvesting the cranium of rats, micro-computed tomography (CT) scanning was conducted using the SkyScan 1172 Desktop X-ray microtomograph (Bruker, Billerica, MA, USA) at 50 kVp and 200 μA to evaluate bone mineral density and bone volume of the scaffolds before and after implantation.

Histological analysis. Specimens were fixed with 10% buffered formalin and decalcified with Calci-Clear™ Rapid (National Diagnostics, Atlanta, GA, USA). Samples were then dehydrated in an ascending series of alcohol and embedded in paraplast (Sherwood Medical Industries, Lane Cove, Australia). Embedded specimens were cut to 5-μm sections with a microtome (Cambridge Instrument, Somerville, MA, USA). Each slide was stained with hematoxylin and eosin, Masson’s trichrome, and Goldner’s trichrome and observed under a microscope.

Statistical analysis. The experimental data were managed and calculated by using a statistical software (SPSS 21.0, SPSS Inc., Chicago, IL, USA) to compare different groups. Mann–Whitney U-tests were used for statistical analysis.

Results

Analysis of surface morphology. The surfaces of BC and PHBB were scanned by low-voltage scanning electron microscopy (LV-SEM) (Figure 1). LV-SEM images showed that both BC and PHBB have structures with irregular and interconnecting macroporosity. Both BC and PHBB had microstructures in which collagen fibers were embedded with bovine or porcine bone particles.

ATR-FTIR spectroscopy. Chemical characteristics of BC, PHBB, porcine cancellous bone, and type I collagen were evaluated by using ATR-FTIR spectroscopy. Porcine cancellous bone exhibited distinctive peaks at frequencies of 1,454, 1,444, 1,020, and 600-550 cm\(^{-1}\). Porcine type I collagen showed five characteristic absorption bands at the frequencies of 3,296, 2,920, 1,647, 1,544, and 1,236 cm\(^{-1}\) (Figure 2). In case of BC, two absorption band peaks were observed at 1020 and 600-550 cm\(^{-1}\). PHBB exhibited additional peaks between 1,714 and 1,433 cm\(^{-1}\) as well as distinctive peaks at 1,020, and 600-550 cm\(^{-1}\).

Mechanical properties. Uniaxial external forces were applied to evaluate the compressive strength of dry/wet BC and PHBB. Both BC and PHBB showed significant differences in mechanical properties between dry and wet samples at room temperature. The dry compressive strength of the BC
was 53.49±11.59 N, whereas the wet strength was 4.86±0.88 N. In comparison, the PHBB exhibited dry compressive strength of 11.22±1.53 N and wet compressive strength of 2.33±0.37 N (Figure 3). The resilience of wet PHBB after the removal of compression is shown in Figure 4.

**Micro-CT analysis.** Both BC and PHBB were well positioned within the calvarial defects (Figure 5). New bone formation was greater in BC and PHBB groups than in the critical defect (CD) group.

In micro-CT analysis, bone mineral density (BMD) and bone volume (BV) were increased over time in all groups (Figure 6A and B). Both BMD and BV were significantly higher in defects containing BC or PHBB than untreated defects at 4 and 8 weeks (p<0.01). In addition, BMD and BV of the BC group were higher than those of the PHBB group (p<0.05). However, the BC-PHBB ratio of BMD and BV were decreased at 8 weeks after implantation from 1.91 to 1.23 (BMD) and from 2.00 to 1.22 (BV) compared to those of scaffolds before implantation.

**Histological findings.** In both BC and PHBB groups, the bone graft space was maintained well without collapse unlike

![Figure 1. LV-SEM images of bone block surface. (A) Bio-Oss Collagen® (BC) (×200), (B) porcine hybrid bone block (PHBB) (×200). Cross-linked collagen and macroporous structures of the surface were observed in both bone grafts.](image1)

![Figure 2. ATR-FTIR spectra of BC, PHBB, porcine cancellous bone, and collagen.](image2)

![Figure 3. Compressive strength of PHBB and BC. Data are expressed as means±SEM (n=5 per group).](image3)
in the CD group (Figure 7A, E and I). At week 4, bone-like material and new bone formation were observed along the surface of the graft materials in both BC and PHBB groups. However, the PHBB group showed higher angiogenesis and less residual grafts than the BC group (Figure 7).

At week 8, the defect site was mostly filled with fibrous connective tissue in the CD group (Figure 8). Relatively, the defect sites of BC and PHBB groups were filled with residual grafts, bone-like material, and newly formed bone in addition to fibrous connective tissue (Figure 8E and I). Moreover, new bone formation above the dura mater appeared characteristically in both BC and PHBB groups, which was not observed at 4 weeks after bone grafting. The newly formed bone tended to become more mature in both
Bone grafts play a pivotal role in filling extensive bone defects in dentistry. Particulate bone grafts are favored because they fill sufficiently irregular intraosseous defects during dental treatment and maxillofacial surgery (1). However, particulate bone grafts may escape from the grafted site, and thus additional materials and methods such as a barrier membrane are needed to place the grafted materials in the proper position and avoid collapse of the grafted site. The use of a barrier membrane involves additional expenses and prolonged surgical time, exposing patients to greater discomfort. Thus, bone substitutes with a good space maintenance capacity without any additional materials are necessary for better clinical outcomes (2).

In this study, we added porcine-derived type I collagen to porcine-derived low crystalline hydroxyapatite to achieve properties such as easy handling, flexibility, good biodegradability and elimination of the risk for BSE. To evaluate the biodegradability and bone regeneration of

![Figure 6. Micro CT measurement of bone formation in rat calvarial defects after implantation. Bone mineral density (A) and bone volume (B) of raw and implanted scaffolds in the defect sites. *p<0.05, **p<0.01. Data are expressed as means±SEM (n=5 per group).](image_url)
developed the bone substitute (PHBB), we used a rat critical-sized calvarial defect model and compared the characteristics of PHBB with those of BC.

Examination of the surface structures of BC and PHBB by LV-SEM revealed irregularly and fully interconnecting macropores. These macropores generally enhance resorption and facilitate the movement of Ca and P ions into the intercellular medium beyond the bone graft material, improving osteoconductivity (10-12). Thus, both BC and PHBB were expected to show good osteoconductivity in vivo.

The chemical properties of the BC and the PHBB were assessed using ATR-FTIR spectroscopy. The FTIR spectra of porcine cancellous bone, one of the constituents of the PHBB, showed the characteristic peaks of PO43- (1,020 and 600-550 cm–1) and carbonate (1,454 and 1,444 cm–1), which are typical hydroxyapatite peaks (13). Porcine type I collagen, another constituent of PHBB, showed five typical bands of collagen at the frequencies of 3,296, 2,920, 1,647, 1,544, and 1,236 cm–1 corresponding to amide A, B, I, II, and III, respectively (14). The spectrum of PHBB was characterized by absorption bands arising from hydroxyapatite (1,444, 1,020, and 600-550 cm–1) and collagen (1,680 and 1,647 cm–1).

The compressive strength of BC decreased dramatically from 53.49±11.59 N to 4.86±0.88 N when it was wet. In comparison, the difference in compressive strength between dry and wet PHBB was smaller than that of BC (11.22±1.53 vs. 2.33±0.37 N, respectively). PHBB exhibited approximately one half of the compressive strength of BC under wet condition. However, the compressive strength of dental bone graft is not a very critical factor once the bone graft can maintain the space and act well as a bone void filler.
The flexibility of PHBB was assessed by checking resilience of the wet sample after the removal of compression. PHBB showed good recovery to its original shape when the compression was removed. This property can be attributed to type I collagen that holds hydroxyapatite granules together and enhances handling characteristics of PHBB.

In the animal study, well-localized materials at the grafted site were confirmed by micro-CT analysis. Both BC and PHBB groups showed significant differences in bone mineral density and bone volume compared to the CD group at 4 and 8 weeks after bone graft. And interestingly, the BC-PHBB ratio of BMD and BV were decreased over time after implantation indicating that new bone formation occurs faster in the PHBB group than in the BC group.

In histological analysis, the defect site of the CD group was collapsed with a small amount of connective tissue. In contrast, the defect sites of the BC and PHBB groups were well maintained with residual grafts, bone-like material, and newly-formed bone in addition to fibrous connective tissue. At week 4, the PHBB group showed more angiogenesis than the BC group. This higher angiogenesis may cause mesenchymal cells to differentiate into osteoblast progenitors, particularly during intramembranous ossification (15). Moreover, blood vessels in the bone graft site supply various growth factors including vitamins and parathyroid hormones (PTHs) which facilitate the healing process, particularly at an early stage of bone healing (16). At week 8, replacement of the grafted materials with bony tissue was prominent in the PHBB group. This indicates that the bone particles of PHBB degrade faster than the Bio-OSS® granules, providing more space for new bone formation. These results are consistent with those of other studies describing the fast resorption rate.

Figure 8. Histological findings of rat calvarial defects at 8 weeks after implantation (A-D) the CD group, (E-H) the BC group, and (I-L) the PHBB group. Samples (A, B, E, F, I, and J) were stained with H&E. Samples (C, G, and K) were stained with Masson’s Trichrome. Samples (D, H, and L) were stained with Goldner’s Trichrome. *Residual grafts, BV: blood vessel, NB: newly-formed bone. [(A, E, I: original magnification ×40), (B, C, D, F, G, H, J, K, L: original magnification ×200)].
of porcine bone grafts (16, 17). Typically, faster resorption of grafted materials is expected to induce earlier substitution with new bone (15), and thus a similar result was predicted for the PHBB grafted site (18).

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