

Improved osseointegration of dental titanium implants by TiO₂ nanotube arrays with recombinant human bone morphogenetic protein-2: a pilot in vivo study

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Abstract: TiO₂ nanotube arrays on the surface of dental implants were fabricated by two-step anodic oxidation. Their effects on bone-implant contact were researched by a pilot in vivo study. The implants were classified into four groups. An implant group with TiO₂ nanotube arrays and recombinant human bone morphogenetic protein-2 (rhBMP-2) was compared with various surface implants, including machined surface, sandblasted large-grit and acid-etched surface, and TiO₂ nanotube array surface groups. The diameter of the TiO₂ nanotube window and TiO₂ nanotube were ~70 nm and ~110 nm, respectively. The rhBMP-2 was loaded into TiO₂ nanotube arrays and elution was detected by an interferometric biosensing method. A change in optical thickness of ~75 nm was measured by flow cell testing for 9 days, indicating elution of rhBMP-2 from the TiO₂ nanotube arrays. For the in vivo study, the four groups of implants were placed into the proximal tibia of New Zealand White rabbits. In the implant group with TiO₂ nanotube arrays and rhBMP-2, the bone-to-implant contact ratio was 29.5% and the bone volume ratio was 77.3%. Bone remodeling was observed not only in the periosteum but also in the interface between the bone and implant threads. These values were higher than in the machined surface, sandblasted large-grit and acid-etched surface, and TiO₂ nanotube array surface groups. Our results suggest that TiO₂ nanotube arrays could potentially be used as a reservoir for rhBMP-2 to reinforce osseointegration on the surface of dental implants.

Keywords: TiO₂ nanotube arrays, drug reservoir, dental implant, bone morphogenetic protein

Introduction

Titanium and its alloys are currently used extensively in the field of dental implants because of its good mechanical properties and excellent biocompatibility. However, the native oxide layer of titanium cannot directly bond with bone to promote new bone formation in the early stage of osseointegration. Numerous studies have sought to optimize dental and orthopedic implants by modifying the implant surface chemistry and/or surface topography by methods such as blasting, plasma spraying of hydroxyapatite, sandblasting and etching, and anodic oxidation.¹⁻⁷

TiO₂ nanotube arrays can be formed as “columnar porous” titania layers by anodic oxidation on pure titanium or titanium alloy surfaces, and are of great interest due to their highly ordered nanostructure.⁸⁻¹² The results of recent studies suggest that formation of a titanium implant surface with a nanostructure could reinforce osseointegration because the surface area is markedly increased and the surface topography can be nanomodified to resemble native bone tissue.¹²⁻¹⁴ Recent studies have demonstrated

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that the fabrication of ordered TiO₂ nanotube arrays with controlled empty space diameters can be adjusted by controlling processing factors, such as voltage, current density, and electrolytes.^{8,15,16} TiO₂ nanotube surfaces with the optimal length scale for cell adhesion and differentiation can induce the migration of osteoblasts and mesenchymal stem cells, and hence reinforce interactions between implant surfaces and cells.^{9,10,13,14}

Recent advances in the fabrication, properties, and applications of TiO₂ nanotube arrays have provided new opportunities for research in relation to their use in clinical practice. The more recent studies attempted localized drug delivery via TiO₂ nanotubes.^{11,17} The empty space of TiO₂ nanotubes could be used as a drug reservoir. Antibiotics, anti-inflammatory drugs, and growth factors can be prescribed to be taken orally, intravenously, or intramuscularly. Several drugs, however, are not effective when delivered via these routes. Moreover, systemic delivery of these drugs can produce adverse effects and organ toxicity associated with high concentrations. Therefore, local drug therapy has become an accepted type of treatment.¹¹

Recombinant human bone morphogenetic protein-2 (rhBMP-2) has been shown to improve osteoblast differentiation and bone formation and remodeling.^{18,19} Several studies showed that application of rhBMP-2 at appropriate doses induced bone formation, whereas higher doses were associated with undesirable effects.^{19,20} Moreover, systemic delivery of rhBMP-2 can have adverse effects such as

unwanted ectopic bone formation. Therefore, it is believed that rhBMP-2 should be immobilized on the implant surface to allow sufficient time to promote osseointegration.¹¹

In this study, TiO₂ nanotube arrays were formed on the surface of dental titanium implants by anodic oxidation. TiO₂ nanotube arrays provided empty spaces for drug loading and demonstrated better biocompatibility than a machined surface. We designed a dental implant with TiO₂ nanotube arrays as a suitable structure for inserting drugs such as antibiotics, anti-inflammatory agents, and growth factors. To improve bone-implant contact, the rhBMP-2 was selected and loaded in a storage room of TiO₂ nanotube arrays. The effects of TiO₂ nanotube arrays and rhBMP-2 on bone-implant contact in the dental implants were investigated in vitro and in an in vivo study in rabbits.

Materials and methods

Fabrication of implants with TiO₂ nanotube arrays and in vitro study

Dental implants with four different surface characteristics were used: a machined surface, a sandblasted large-grit and acid-etched SLA implant (TS III™, Osstem Implants, Seoul, South Korea) as a positive control group, TiO₂ nanotube arrays, and TiO₂ nanotube arrays with rhBMP-2 loading were called I1 group, I2 group, I3 group, and I4 group, respectively. All implants were 3.5 mm in diameter and 8.5 mm in length. TiO₂ nanotube arrays on the surface of the implants were fabricated by anodic oxidation in an electrolyte solution

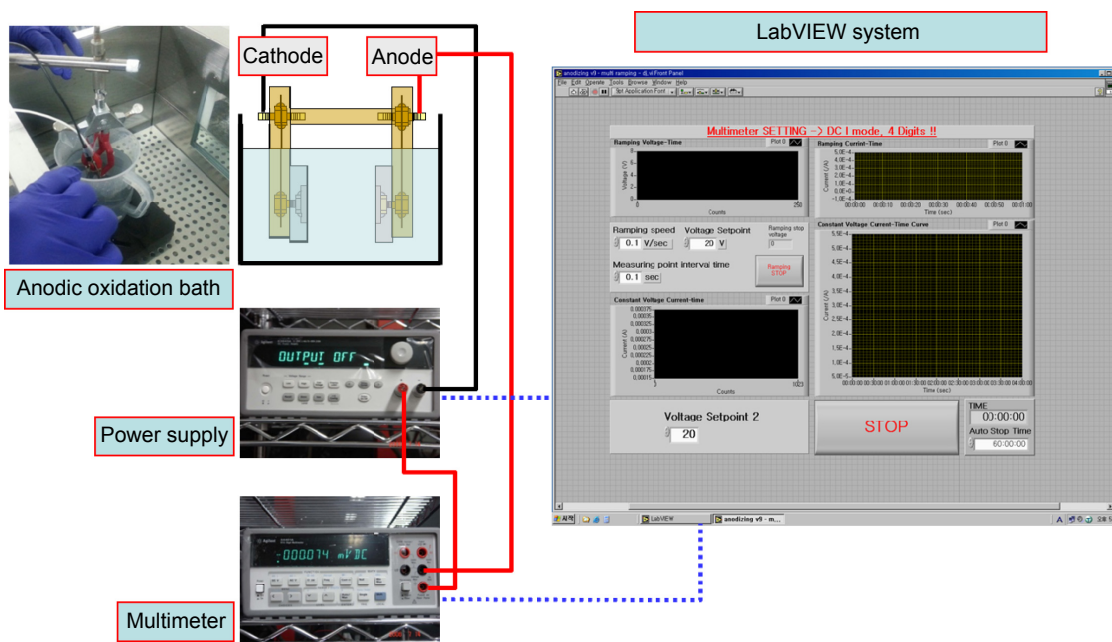


Figure 1 Schematic diagram of anodic oxidation process.

containing ethylene glycol with 0.5 wt% NH₄F.²¹ Figure 1 shows a schematic diagram of the anodic oxidation procedure. To obtain a proper microstructure of TiO₂ nanotube arrays, anodic oxidation voltages and the time provided by a DC power supply were computer-controlled by a LabVIEW program. Two-step anodic oxidation was conducted to obtain the clean surface and open windows of the TiO₂ nanotube arrays. The implant was first anodized at a voltage of 60 V for 60 minutes. The TiO₂ nanotube arrays fabricated by the first anodic oxidation were removed in ultrasonication. TiO₂ nanotube arrays with clean and open windows were then fabricated by the second anodic oxidation. The voltage and duration of the second anodic oxidation were 60 V and 15 minutes, respectively. The thickness and open window size of the TiO₂ nanotube arrays were determined by field emission scanning electron microscopy (Phillips/FEI XL 30SFEG, Eindhoven, the Netherlands) at the Korea Basic Science Institute. For the rhBMP-2-loaded TiO₂ nanotube array implant, rhBMP-2 (Cowellmedi Co Ltd, Busan, South Korea) was loaded into the inner space of the TiO₂ nanotube arrays on the surface of the dental implant by a dip-coating process in a vacuum chamber. The concentration of rhBMP-2 was 1.5 mg/mL, ie, the same concentration as that used by Lee et al¹⁹ and Wikesjo et al.²⁰ Each implant was immersed three times in rhBMP-2 solution for 5 seconds and dried at a maximum temperature of 20°C.

The elution of rhBMP-2 was observed by an interferometric biosensing method.²² Figure 2 shows a schematic diagram of the interferometric biosensing method using TiO₂ nanotube arrays. TiO₂ nanotube arrays fabricated on a flat titanium surface were also used as a sensing porous layer. White light from a tungsten lamp (Ocean Optics, Dunedin, FL, USA) was fed

through one end of a bifurcated fiber optic cable and focused through a lens onto the surface of the TiO₂ nanotube arrays at normal incidence. Reflected light was collected through the same optics, and the distal end of the bifurcated fiber optic cable was input to a S-4000 CCD spectrometer (Ocean Optics).²³ Optical thickness (nL) was determined from the Fabry-Perot relationship ($m\lambda = 2nL$), where m is the fringe order, λ is the wavelength of maximum constructive interference for a spectral fringe of order m , n is the refractive index of the porous layer, and L is the thickness of the porous layer. The change in optical thickness over time was monitored.

In vivo study

This study was approved by the ethical committee on animal research at the Institute of Gangneung-Wonju National University (GWNU-2013-3). Three New Zealand White rabbits (weighing 3,060±167 g and aged 13–14 weeks at the start of the experiment) were used in the study. The surgical procedures were performed aseptically under intramuscular anesthesia with a combination of 0.7 mL/kg of ketamine (Yuhan Corporation, Seoul, South Korea) and 0.2 mL/kg of xylazine (Bayer, Leverkusen, Germany). Prior to surgery, the operative sites were shaved and carefully washed with iodine solution and injected with 2% lidocaine containing 1:100,000 epinephrine (Huons, Seoul, South Korea). A skin incision was made along the proximal one third of the tibia using sterile surgical technique. After full-thickness flap reflection, two holes were drilled about 7 mm apart with copious irrigation. The drilling procedures followed the manufacturer's instructions. All 12 implants were surgically placed. Each rabbit received the four different types of implant. Two different types of implant were placed at each

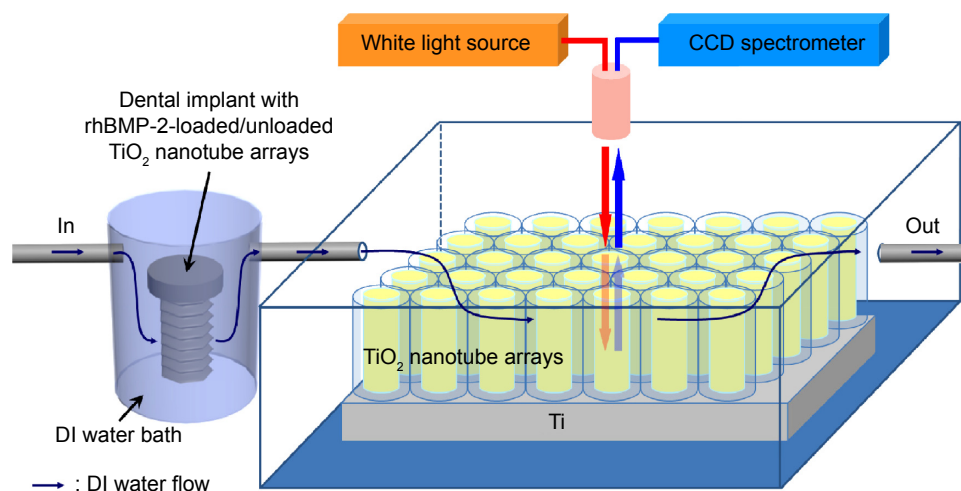


Figure 2 Schematic diagram of interferometric biosensing method using TiO₂ nanotube arrays.

Abbreviations: DI, deionized water; TiO₂, titanium dioxide; rhBMP-2, recombinant human bone morphogenetic protein-2; CCD, charged-coupled device.

tibia in random circulating order into the left and right sides of the tibia to ensure unbiased comparisons. The middle third of each implant was engaged by upper cortical bone only. After surgery, all rabbits received antibiotic cover, ie, 0.15 mL/kg gentamicin sulfate (Dongwha Pharm Co Ltd, Seoul, South Korea) and analgesia, ie, 0.20 mL/kg Sulpyrine (Green Cross Veterinary Products Co Ltd, Yongin, South Korea) by intramuscular injection. The animals were kept in separate cages and allowed full weight bearing after surgery.

For fluorochrome labeling, subcutaneous injections with alizarin red (30 mg/kg; Sigma-Aldrich, St Louis, MO, USA) and calcein green (10 mg/kg; Sigma-Aldrich) were performed 3 and 6 weeks, respectively, after the operation. The rabbits were euthanized in a CO₂ chamber 8 weeks after inserting the implants, and 1 cm × 1 cm bone fragments with dental implants were collected.

After euthanasia, the implant areas were scanned using a 1173 micro-computed tomography scanner (Skyscan, Kontich, Belgium). The voltage and current of the X-ray tube were 130 kV and 60 μA, respectively, with an exposure time of 500 msec. X-ray projections were obtained at 0.3 degree intervals with a scanning angular rotation of 360 degrees. The image pixel size was set at 6.04 μm.

The samples were fixed by immersion in a 10% neutral-buffered formalin solution (Accustain™, Sigma-Aldrich, Steinheim, Germany) for one day, then dehydrated in a graded series of ethanol solutions and embedded in methylmethacrylate resin (Technovit 7200 VLC, Kulzer Co GmbH, Friedrichsdorf, Germany). After dehydration, the specimens were polymerized in a light-based polymerization unit (Exakt Apparatebau, Norderstedt, Germany). The implants were cut mid-axially in a mediolateral plane into 200 μm thick sections using a band saw with a diamond blade (Exakt cutting-grinding system, Exakt Apparatebau). The final section was ground to no thicker than about 20 μm using an Exakt microgrinder and polished to an optical finish utilizing the cutting-grinding technique described by Donath and Breuner.²⁴

All the sections were first examined by immunofluorescence microscopy (Leica Microsystems, Wetzlar, Germany). The specimens were then stained by the Masson-Goldner trichrome method. Images of the specimens were obtained by optical microscopy (BX-50, Olympus America, Melville, NY, USA). One examiner performed the histomorphometric analysis in a blinded manner. The bone-to-implant contact ratio (%) and bone volume ratio (%) was measured in the cortical bone area using Image Pro Plus analysis software (Media Cybernetics Inc, Rockville, MD, USA). The bone-to-implant

contact ratio was defined as the percentage of bone contact measured along the entire surface length of two implant threads. The bone volume ratio was the percentage of total bone tissue within the area of two threads.

Results and discussion

Fabrication of implant with TiO₂ nanotube arrays and in vitro study

Figure 3 shows field emission scanning electron microscopic images of the implants with various surface treatments. Figure 3A and B shows typical images and microstructures of a machined titanium implant. It has mainly unidirectional machined grooves from the manufacturing instrument. The smooth surface by machining with a CNC lathe is also shown in Figure 3B. The surfaces of the SLA implant have very rough surfaces (Figure 3C and D). The rough and irregular sharp surface results from a grit-blasting procedure. The rough morphology enhances osseointegration. The rough surface was additionally obtained by anodic oxidation. Figure 3E and F show the rough surface of TiO₂ nanotube arrays fabricated by anodic oxidation. The surface with the TiO₂ nanotube arrays is less rough than the SLA surface. However, the TiO₂ nanotube surface has nanosized holes for loading drugs such as BMP-2, PEP7, and ibuprofen.

Figure 4 shows field emission scanning electron microscopic images of the TiO₂ nanotube arrays and rhBMP-2-loaded TiO₂ nanotube arrays on the surface of the implants. The TiO₂ nanotube arrays were fabricated by two-step anodic oxidation.²⁵ There were traces of the TiO₂ nanotube arrays by first anodic oxidation on the surface of the newly grown TiO₂ nanotube arrays. The diameter of the TiO₂ nanotube window and TiO₂ nanotube were ~70 nm and ~110 nm, respectively. The windows of the TiO₂ nanotubes were clean and open, and these structures were appropriate for loading a drug. The thickness of the TiO₂ nanotube arrays was ~17 μm, as shown in the inset in Figure 4A. The window of the TiO₂ nanotube, as shown in Figure 4D, was slightly clogged with rhBMP-2 loading. The diameter of the TiO₂ nanotube windows was decreased to ~50 nm. The surface-loaded rhBMP-2 was expected to improve the osseointegration.

To observe the elution of rhBMP-2 from the TiO₂ nanotube arrays on the surface of the dental implant, an interferometric biosensing method with a flow cell was used.^{22,26} During this process, deionized water flowed through the dental implant with rhBMP-2-loaded TiO₂ nanotube arrays. The change in optical thickness with rhBMP-2 was monitored in real time. Figure 5 shows the change in optical thickness of the dental implant with

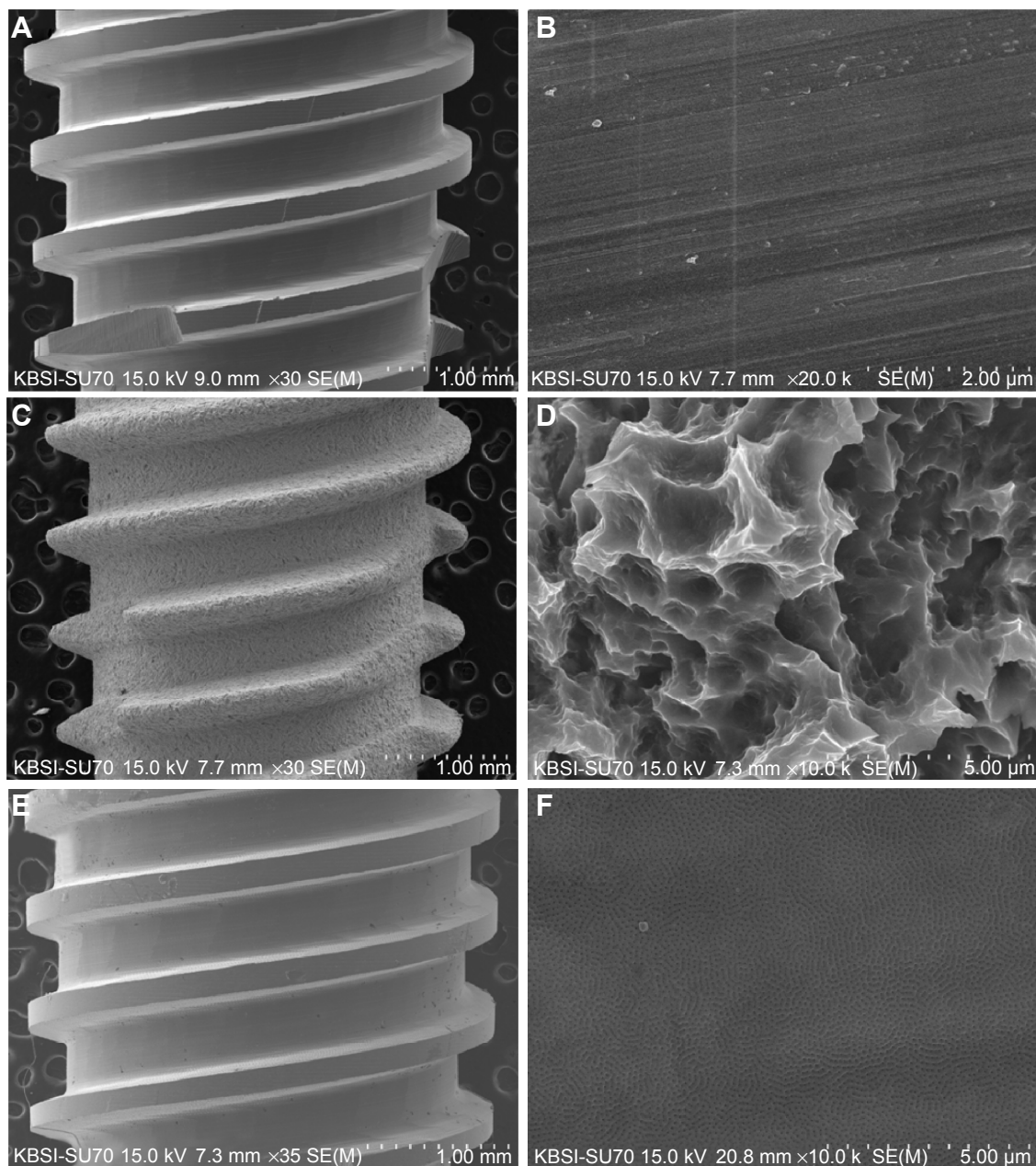


Figure 3 Field emission scanning electron microscopic images of the implants.

Notes: (A) Low magnification (30 \times) of machined implant, (B) high magnification (20,000 \times) of machined implant, (C) low magnification (30 \times) of SLA implant, (D) high magnification (10,000 \times) of SLA implant, (E) low magnification (35 \times) of anodized implant, and (F) high magnification (10,000 \times) of anodized implant.

Abbreviation: SLA, sandblasted large-grit and acid-etched.

rhBMP-2-unloaded TiO₂ nanotube arrays (I3 group) and rhBMP-2-loaded TiO₂ nanotube arrays (I4 group) for 10 days. A baseline was established using the optical thickness of deionized water for ~20 hours. The solution was subsequently introduced by deionized water passed through the dental implant with TiO₂ nanotube arrays. The optical thickness remained constant for 10 days in I3 group. However, in the I4 group, optical thickness was dramatically increased by elution of rhBMP-2 from the

TiO₂ nanotube arrays and slowly increased for 9 days. A change in optical thickness of ~75 nm confirms the elution of rhBMP-2 from TiO₂ nanotube arrays.

In vivo study

This study was designed to evaluate the possibility of using TiO₂ nanotube arrays as a drug reservoir, and rhBMP-2 was selected as a drug that might improve the induction of new bone formation and osseointegration around an implant

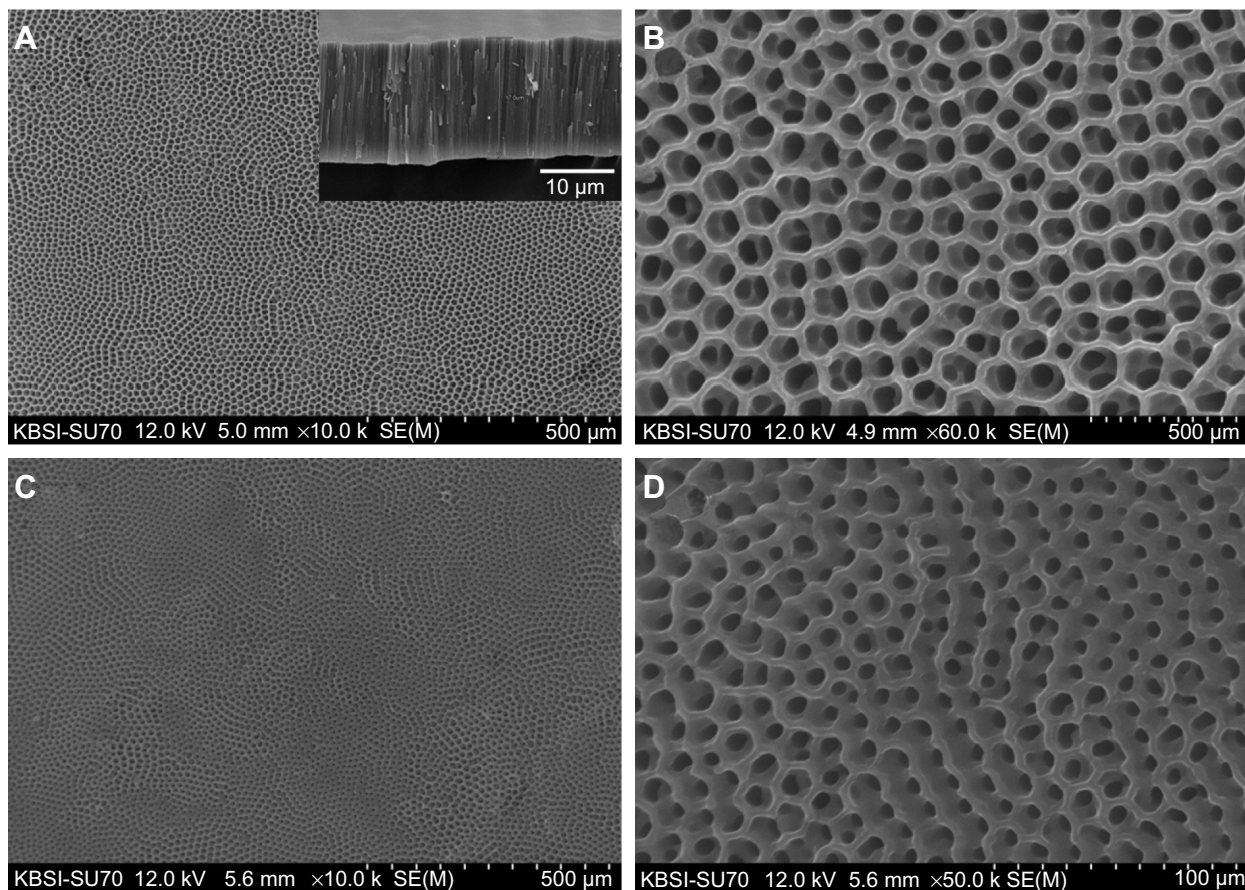


Figure 4 Field emission scanning electron microscopic images of TiO_2 nanotube arrays on the implant surface.

Notes: (A) Low magnification (10,000 \times) of TiO_2 nanotube arrays, (B) high magnification (60,000 \times) of TiO_2 nanotube arrays, (C) low magnification (10,000 \times) of rhBMP-2-loaded TiO_2 nanotube arrays, and (D) high magnification (50,000 \times) of rhBMP-2-loaded TiO_2 nanotube arrays. Note the open pore size was decreased by loading rhBMP-2 on nanotubes.

Abbreviations: TiO_2 , titanium dioxide; rhBMP-2, recombinant human bone morphogenetic protein-2.

surface. Dental implants were classified into four groups: machined surface implants, SLA surface implants, TiO_2 nanotube array surface implants, and TiO_2 nanotube array surface implant with rhBMP-2 (groups I1, I2, I3, and I4, respectively). Four group implants were placed into the proximal tibia of

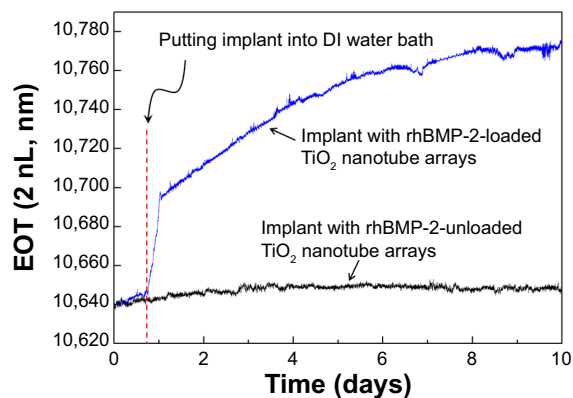


Figure 5 Change in optical thickness of dental implant with rhBMP-2-unloaded TiO_2 nanotube arrays and rhBMP-2-loaded TiO_2 nanotube arrays over 10 days.

Abbreviations: DI, deionized water; EOT, effective optical thickness; TiO_2 , titanium dioxide; rhBMP-2, recombinant human bone morphogenetic protein-2.

rabbits for 8 weeks. Clinical healing was uneventful, with no cases of implant exposure or loss in any of the rabbits. After 8 weeks, all the implants were histologically in direct contact with the surrounding bones along the threads.

Figure 6 and Table 1 show the results of the histomorphometric analysis. The highest mean bone-to-implant contact ratio was found in the I4 group (29.5%), followed by the I3 group, I2 group, and I1 group (16.3%, 14.7%, and 11.1%, respectively). The bone volume ratios were measured around the implant threads. The highest bone volume ratio (77.3%) was also found in the I4 group, with values of 67.2%, 53.7%, and 66.9% in the I3 group, I2 group, and I1 group, respectively. These results suggest that the TiO_2 nanotube array surface had more of an osseointegration effect than the machined surface or the SLA surface implant, and that rhBMP-2 loaded into TiO_2 nanotube arrays had the biochemical effect of bone induction.

Our findings concerning the effect of TiO_2 nanotube arrays are similar to those of previous studies reporting an

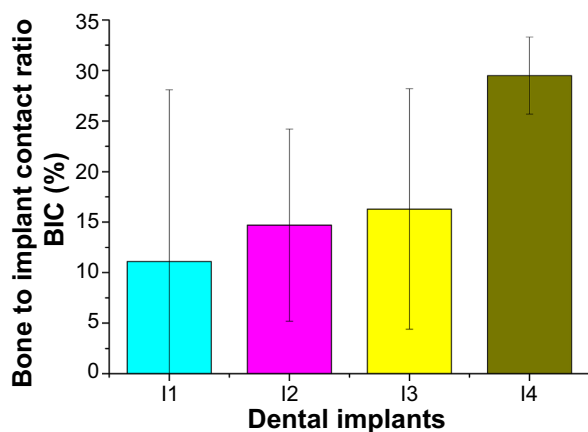


Figure 6 Bone-to-implant contact ratios of machined surface, SLA surface, TiO₂ nanotube array surface, and TiO₂ nanotube array surface with rhBMP-2.

Notes: (I1) machined surface implants, (I2) SLA surface implants, (I3) TiO₂ nanotube array surface implants, and (I4) TiO₂ nanotube array surface implant with rhBMP-2. **Abbreviations:** BIC, bone-to-implant contact ratio; SLA, sandblasted large-grit and acid-etched.

enhanced bone formation and cell adhesion effect with the TiO₂ nanotube array implant.^{9,12,14} Bjursten et al¹² reported that TiO₂ nanotubes significantly improved bone bonding strength, bone-implant contact, and new bone formation when compared with titanium grit-blasted surfaces. Because TiO₂ nanotubes have a good oxide microstructure for growing new bone, TiO₂ nanotube arrays are able to influence protein interactions and components of bone for rapid and permanent bone bonding.

A number of previous studies have reported the osteoinduction effect of the rhBMP-2-coated implant.^{18–20,27} However, conventional rhBMP-2 coating methods cannot provide a long-lasting osteoinduction effect by rhBMP-2. As suggested by Çalışkan et al²⁸ TiO₂ nanotube arrays enhance water wettability, and can therefore act as a drug reservoir. Recently, a TiO₂ nanotube implant combined with rhBMP-2 was reported to improve bone formation.²⁹ In this study, the I4 group in which rhBMP-2 was loaded into TiO₂ nanotube arrays showed the greatest bone-to-implant contact ratio and bone volume ratio 8 weeks after implantation. This can be taken as evidence of the long-lasting effect of rhBMP-2, because bone remodeling, bone formation, and

Table I Histomorphometric analysis

	I1 group	I2 group	I3 group	I4 group
Bone to implant contact (%)	11.1±17.0	14.7±9.5	16.3±11.9	29.5±3.8
Bone volume ratio (%)	66.9±6.7	53.7±11.5	67.2±7.6	77.3±8.8

Note: The data are shown as the mean ± standard deviation.

Abbreviations: I1, machined surface implants; I2, SLA surface implants; I3, TiO₂ nanotube array surface implants; I4, TiO₂ nanotube array surface implant with recombinant human bone morphogenetic protein-2.

bone reduction occurs very slowly. The slow elution of rhBMP-2 as shown in Figure 5 indicates that rhBMP-2 has a long-lasting effect.

Figure 7 shows the micro-computed tomographic images for the four implant groups. Vertical bone ingrowth in I4 group was increased, and the I3 and I4 groups showed more bone deposition around the implant threads than the I1 and I2 groups.

Figure 8 shows the fluorescence and histologic staining images for the four groups as observed by optical microscopy. The fluorescence and histologic staining images in Figure 8A–D are shown on the right and left sides, respectively. The fluorescence images were formed by fluorochrome labeling with alizarin red and calcein green. Different patterns of bone formation and bone remodeling were observed between the four groups. The red and green areas represent new bone formation at 3 weeks and 6 weeks after implant installation, respectively. Bone formation and bone remodeling, as indicated by white arrows, were mainly observed near the periosteum in the I1, I2, and I3 groups. However, the bone formation and remodeling in the I4 group were observed not only near the periosteum but also all around the implant threads.

The bone formation and bone remodeling in the I1, I2, and I3 groups can be explained by an osteoblast-rich periosteum. In the I4 group, the active bone formation and bone remodeling at the implant thread site are due to the osteogenic effect of rhBMP-2 eluted from the TiO₂ nanotube arrays. The I4 group also showed stronger fluorochrome labeling. Fluorochrome labeling near the periosteum indicates bone formation, whereas fluorochrome labeling around the implant thread is thought to indicate bone remodeling and improving osseointegration. The results suggest that TiO₂ nanotube arrays with rhBMP-2 can enhance bone formation and bone remodeling around an implant, and thus reinforce osseointegration on the surface of dental implants.

The aim of the present study was to fabricate thicker TiO₂ nanotubes on the surface of dental implants for slower release of rhBMP-2 than that in a previous report.²⁹ Slower release of rhBMP-2 from the nanotubes was observed in our in vitro study, and a higher bone-to-implant contact ratio was observed in the I4 group in our in vivo study. However, a weak point in our study was the small number of samples which precluded a statistical analysis. Further, the mechanical analysis like a removal torque and its evaluation on the different time points which could not be not performed due to same reason. A further study with a larger number of samples and various evaluation methods is necessary to

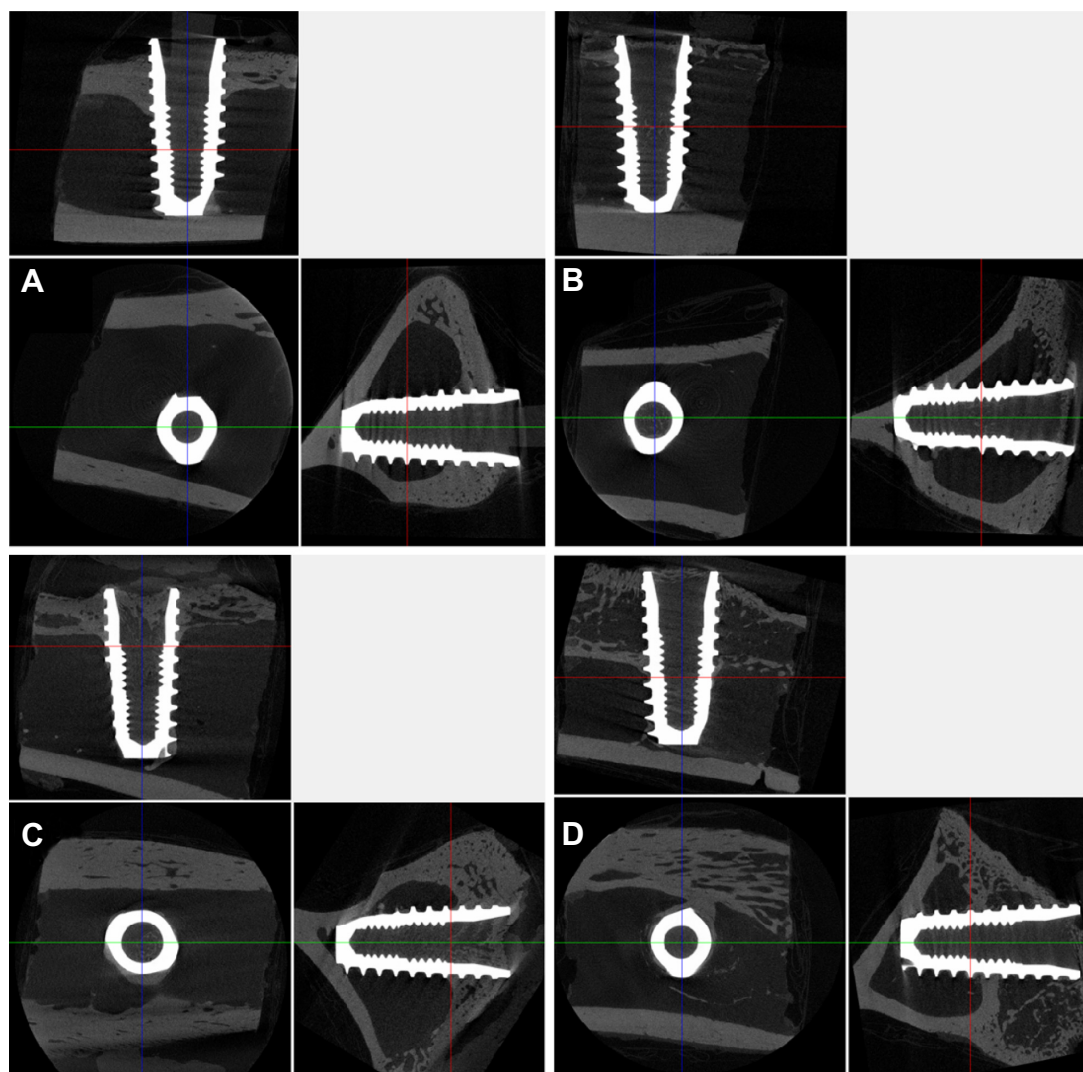


Figure 7 Micro-computed tomographic images 8 weeks after implant installation. (A) Machined surface group, (B) SLA surface group, (C) TiO₂ nanotube array surface group, and (D) TiO₂ nanotube array surface with rhBMP-2 group.

Abbreviations: TiO₂, titanium dioxide; rhBMP-2, recombinant human bone morphogenetic protein-2; SLA, sandblasted large-grit and acid-etched.

clarify the effects of TiO₂ nanotube arrays on the surface of dental implants with rhBMP-2.

Conclusion

TiO₂ nanotube arrays on the surface of titanium dental implants were fabricated by anodic oxidation. To load rhBMP-2 into TiO₂ nanotube arrays, clean and open windows of TiO₂ nanotubes were obtained by two-step anodic oxidation. Elution of rhBMP-2 from the TiO₂ nanotube arrays was measured by an interferometric biosensing method with a flow cell. An optical thickness change of ~75 over 9 days indicated slow release of rhBMP-2. Implants with a TiO₂ nanotube array surface containing rhBMP-2 had the highest bone-implant contact and enhanced bone remodeling in the in vivo study. TiO₂ nanotube arrays on the surface of the

implant demonstrated the feasibility of the drug reservoir and were helpful for bone formation and cell adhesion. This structure could be very useful as a drug reservoir for various treatments, such as bone growth factors, anti-inflammatory drugs, and antibacterial agents.

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Disclosure

The authors report no conflicts of interest in this work.

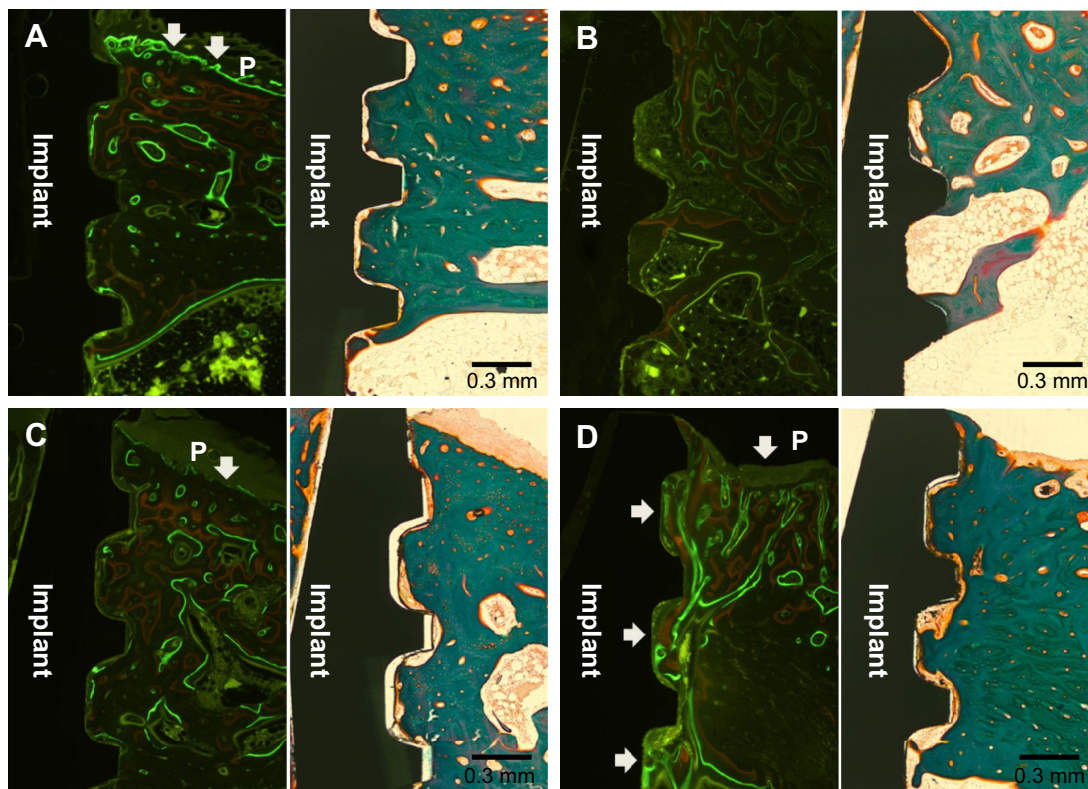


Figure 8 Fluorescence and histologic staining images 8 weeks after implant installation.

Note: (A) Machined surface group, (B) SLA surface group, (C) TiO₂ nanotube array surface group, and (D) TiO₂ nanotube array surface with rhBMP-2 group. P indicates a periosteum area. The white arrow indicates a region of bone formation and bone remodeling.

Abbreviations: TiO₂, titanium dioxide; rhBMP-2, recombinant human bone morphogenetic protein-2; SLA, sandblasted large-grit and acid-etched.

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