Local Application of Alendronate on $\beta$-Tricalcium Phosphate Accelerated Induction of Osteogenesis with Formation of Giant Osteoclast-Like Cell

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ABSTRACT

Intrinsic osteoinductivity—the ability to induce bone formation in ectopic sites without addition of osteogenic factors—has been reported in various porous materials. Tartrate-resistant acid phosphatase (TRAP)-positive osteoclast-like cells are thought to play an important role in material-induced osteoinduction. To investigate the influence of osteoclastic activity on intrinsic osteoinduction, we loaded alendronate ($10^{-2}$, $10^{-4}$, and $10^{-6}$ M) onto porous $\beta$-tricalcium phosphate ($\beta$-TCP) blocks to inhibit osteoclastic activity, and evaluated osteoinductivity by implantation of the blocks into the dorsal muscles of adult beagle dogs. Alendronate-loaded porous $\beta$-TCP blocks increased both speed and amount of osteoinduction, with the $10^{-4}$ M alendronate-loaded $\beta$-TCP being especially active. This finding indicates that $\beta$-TCP loaded with $10^{-4}$ M alendronate might prove crucial in providing the desirable balance between the degradation rate of bone scaffolds and their osteoinductive replacement. Thus, material-induced osteoinduction may be controlled by local application of alendronate, establishing alendronate loading as a promising therapeutic approach.

Keywords: Osteoinduction; $\beta$-tricalcium Phosphate; Alendronate; Osteoclast; Osteogenesis

1. Introduction

Various porous materials exhibit intrinsic osteoinductivity, which is the ability to induce bone formation at ectopic sites without addition of osteogenic factors, such as bone morphogenetic proteins (BMPs), or cell transplantation [1-10]. The osteoinductivity of materials is affected by their composition [11], macro- and micro-pore size [11-15], as well as the animal models employed in its investigation [4,16,17]. However, the exact mechanism of intrinsic osteoinduction by bioactive materials is yet to be elucidated.

Porous $\beta$-tricalcium phosphate ($\beta$-TCP) with interconnected micro-pores is an osteoinductive material that exhibits intrinsic osteoinductivity in canine dorsal muscles [18]. We previously reported that $\beta$-TCP-induced bone formation at ectopic sites was accelerated by injection of an EP4 agonist (prostaglandin E receptor 4, a prostaglandin E2 receptor subtype). Further, the EP4 agonist enhanced the degradation of porous $\beta$-TCP constructs and stimulated osteoclastogenesis before bone formation [19]. Subsequently, in concert with Kondo et al. [18], we considered the crucial role of tartrate-resistant acid phosphatase (TRAP)-positive osteoclast-like cells in material-induced osteoinduction. Enhanced osteoclastic activity results in an increase of $\beta$-TCP replacement by bone. In addition, we reported that TRAP-positive cells might be an important factor responsible for the cross-species differences in material-induced osteoinduction [17]. While TRAP-positive multinuclear cells were detected and osteoinduction was observed in the dorsal muscles of dogs 4 weeks after the implantation of porous calcium-deficient hydroxyapatite (CDHA), no osteoinduction and only a small number of TRAP-positive cells were observed 6 weeks following the same treatment in rats. Moreover, osteoclast expression of cathepsin K, a cysteine protease that degrades type-I collagen osteoid matrix, was higher in dogs than in rats.

In the present study, to further investigate the influence of osteoclastic activity on intrinsic osteoinduction, we loaded alendronate onto porous $\beta$-TCP blocks. We hypothesized that inhibition of osteoclastic activity by alendronate results in a decrease of $\beta$-TCP replacement by forming bone tissue. Alendronate, a nitrogen-containing bisphosphonate, is one of the most common thera-
H2O and CaCO₃ at a molar ratio of 2:1 were mixed into a slurry with pure water and particles of zirconia in a pot mill for 24 h, and dried at 80°C, leading to formation of calcium-deficient hydroxyapatite. This crystalline solid was converted to β-TCP by calcination at 750°C for 1 h. After sintering of β-TCP powder at 1050°C for 1 h, a porous β-TCP block was obtained. The porosity of the block was 75% and the surface area was 1.4 m²/g. The morphology of microporous/macroporous β-TCP block was examined by scanning electron microscopy (SEM) (S-4700, Hitachi Ltd., Tokyo, Japan). The β-TCP block possessed macropores (100-500 µm) that were interconnected via 100-200 µm pores, and micropores (<5 µm) (Figure 1(b)).

2.1.2. Preparation of Alendronate-Loaded Porous β-TCP Blocks

Alendronate sodium hydrate (Teiroc® Injection) was supplied by Teijin Pharma Ltd. (Tokyo, Japan). Thirty-six porous β-TCP blocks were used. For histological evaluation, 6 β-TCP blocks were immersed in undiluted alendronate solution (10⁻² M), alendronate solutions diluted with saline (10⁻⁴ and 10⁻⁶ M), and pure saline (control). Additionally, for quantitative real-time polymerase chain reaction (qRT-PCR) analysis of osteoclastic gene expression, we compared the β-TCP blocks pre-loaded with the most dense, undiluted alendronate solution with the control. Each group of 6 β-TCP blocks was immersed in undiluted alendronate solution (10⁻² M) and saline (control), respectively. Each β-TCP block was immersed in 1.5 mL of the corresponding solution. Solutions containing β-TCP blocks were then vacuumized with an oil rotary vacuum pump (G-25SA; ULVAC KIKO Inc., Miyazaki, Japan) for 15 minutes. After vacuumization, the β-TCP blocks were kept immersed in the solutions for 2 days at 4°C. Subsequently, the blocks were removed from solutions to dry at room temperature.

2.1.3. Loading Alendronate on and Release of Alendronate from Porous β-TCP Blocks

Radioactive samples were prepared according to the aforementioned procedure by using ¹⁴C-labeled alendronate from Porous β-TCP Blocks.
nate (Moravek Biochemicals Inc., California, USA). Six β-TCP blocks were immersed in solutions of 14C-labeled alendronate. The loading efficiency of alendronate on β-TCP blocks was calculated as the difference between its initial concentration and its residual concentration in the supernatant. In order to determine the amount of alendronate released from alendronate-loaded β-TCP, 14C-alendronate-loaded β-TCP blocks were immersed in 4 mL of culture medium, and the alendronate concentration was evaluated by liquid scintillation after 1, 4, and 7 days. At the predetermined time intervals, the samples (1 mL) were collected and replaced with equal volumes of fresh medium.

2.2. Canine Model for Osteoinduction
Osteoinductivity was evaluated by implantation of porous β-TCP blocks into the dorsal muscles of adult beagle dogs. Four healthy, 1-year-old, female beagle dogs weighing 10 kg were used for the study. The animals were reared and the experiments were carried out at the Institute of Laboratory Animals, Graduate School of Medicine, Kyoto University, Japan. All procedures were performed according to the Principles of Laboratory Animal Care of the Kyoto University Animal Experiment Committee. The dogs were anesthetized by intramuscular administration of ketamine hydrochloride (7.5 mg/kg) and xylazine (7 mg/kg). The operations were performed under standard sterile conditions. After performing a longitudinal incision in the skin and fascia in the middle of the back, intramuscular pouches were created in the bilateral dorsal muscles. For histological evaluation, 8 β-TCP blocks pre-loaded with alendronate (10^{-2}, 10^{-4}, or 10^{-6} M) or saline (control) were implanted in 8 intramuscular pouches of 3 dogs. For qRT-PCR analysis, 12 β-TCP blocks pre-loaded with alendronate (10^{-2} M) or saline were implanted in 12 intramuscular pouches of the fourth dog. The fascia and skin were sutured in layers after irrigation with saline. The β-TCP samples were completely covered with muscle and dispersed sufficiently not to affect each other.

2.3. Histological Examinations and Histomorphometry
For histological evaluation, appropriate animals were anesthetized and the implants were retrieved after 4 weeks. The samples were fixed in 4% neutral buffered paraformaldehyde (PFA) for 48 h, decalcified with a 20% EDTA solution at 4°C for approximately a week, dehydrated in an ethanol series (70%, 80%, 90%, 99.5%, and 100%), and embedded in paraffin. The samples were sectioned perpendicular to the long axis of the cylinder by using a microtome (SM2000R; Leica Ltd., Tokyo, Japan) to yield 4 pairs of adjacent 5-µm-thick sections. The first pair of sections was approximately 1-mm apart from the end of the cylinder sample, and each subsequent pair of sections was approximately 1-mm apart from the previous one. Half of the section-pairs were stained with hematoxylin and eosin (HE), and the remaining sections were stained with TRAP staining kit (MK300; Takara Bio Inc., Shiga, Japan) to detect osteoclast-like multinuclear cells. Images were acquired with Nikon Eclipse 80i (Tokyo, Japan) light microscope and analyzed using Adobe Photoshop CS5 (Adobe Systems Inc., CA, USA) and ImageJ (National Institutes of Health freeware). The area of newly formed bone, residual β-TCP, and the porous area of β-TCP were calculated in each section stained with HE, and multinuclear TRAP-positive cells (with more than 3 nuclei) were counted in each section stained with TRAP. New bone formation rate (%) was calculated by dividing the area of newly formed bone by the porous area of β-TCP and multiplying this value by 100.

2.4. Relative Quantitative Real-time Polymerase Chain Reaction (qRT-PCR) Analyses
To correlate the morphometric evaluation of material-induced osteoinduction with osteoclastic gene expression, the appropriate samples were analyzed for changes in TRAP and cathepsin K mRNA synthesis by qRT-PCR. The 12 β-TCP implants, harvested at 2, 4, and 8 weeks (and immediately frozen in liquid nitrogen), were crushed in a mortar filled with liquid nitrogen. Total RNA was isolated using TRIzol® Reagent (Invitrogen Co., CA, USA) and further purified using RNeasy® Mini kit (Qiagen Inc., CA, USA) according to the manufacturer’s instructions. First-strand cDNA was prepared from 1 µg of total RNA by using Transcriptor First Strand cDNA Synthesis Kit (Roche Diagnostics, Mannheim, Germany), and qRT-PCR was performed using LightCycler® FastStart DNA MasterPLUS SYBR Green I (Roche Diagnostics) according to the manufacturer’s instructions. The sequences of primers used for the amplification of the housekeeping glyceraldehyde 3-phosphate dehydrogenase (GAPDH) gene, as well as the osteoclast marker genes encoding cathepsin K and TRAP, are shown in Table 1. Quantitative data analyses were performed with the LightCycler software 3.5 (Roche Diagnostics). Relative expression levels of TRAP and cathepsin K genes were normalized to GAPDH.

2.5. Statistical Analyses
All values are expressed as mean ± standard deviation. The data concerning new bone formation rate, residual amount of β-TCP, and amount of TRAP-positive mult-
tinuclear cells were analyzed using the Steel-Dwass procedure with a 1-way design for nonparametric all-pairs multiple comparisons. The qRT-PCR data were analyzed using the Mann-Whitney’s U test. The critical level of statistical significance was \( p < 0.05 \).

3. Results

3.1. Amount of Alendronate Loaded on and Released from Porous \( \beta \)-TCP Blocks

The amount of alendronate loaded on and released from the porous \( \beta \)-TCP blocks was evaluated using \(^{14}\)C-labeled alendronate. The results indicated that 67.4% ± 1.6% of the initial amount of alendronate solution was incorporated on porous \( \beta \)-TCP blocks. After 1-, 4-, and 7-day immersion, cumulative alendronate release (%) into the culture medium was 0.21 ± 0.07, 0.39 ± 0.12, and 0.79 ± 0.34, respectively (Figure 2).

3.2. Induction of Bone Formation by Alendronate-Loaded \( \beta \)-TCP Blocks

After 4 weeks, 3 of the 6 implants from the \( 10^{-2} \) M group, 4 of the 6 implants from the \( 10^{-4} \) M group, and 2 of the 6 implants from the \( 10^{-6} \) M group showed osteoinduction, while the process was not observed in the control group (Table 2). Induction of bone formation was more remarkable in the peripheral zone of porous \( \beta \)-TCP than in the center (Figure 3(a)). Newly formed bone tissue appeared to spread not only along the surface of \( \beta \)-TCP, but also within the macroporous spaces, where active angiogenesis was identified (Figure 3(b)). Moreover, cuboidal osteoblast-like cells were lining along induced bones. Osteoclast-like multinuclear cells were also identified on the \( \beta \)-TCP surface in HE stained sections. Again, the formation of osteoblasts and osteoclasts, as well as cap-

dillaries, was more abundant in the peripheral zone than in the central zone. In porous spaces of \( \beta \)-TCP, collagenous condensations of extracellular matrix were observed (Figure 3(c)). New bone formation rate (%) in the \( 10^{-4} \) M group was significantly higher than those in the \( 10^{-2} \) M and \( 10^{-6} \) M groups (Figure 4).

<p>| Table 1. The sequences of the primers. |</p>
<table>
<thead>
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<th>Gene</th>
<th>Sequence (5' - 3')</th>
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<td>GAPDH F: AACATCATCCCTGCTTCCAC R: GACCACCTGGTCCTCAGTGT</td>
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<tr>
<td>Cathepsin K F: AAGAACCAGGGTCAGTGTGG R: GAGGTCAGGCTTGCATCAAT</td>
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<tr>
<td>TRAP F: GATGCCAACGACAAGGTT R: CTTTGAAGCGTAGCCGGTAG</td>
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<p>| Table 2. Number of samples that showed induction of bone formation. |</p>
<table>
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<th>Group</th>
<th>( 10^{-2} ) M</th>
<th>( 10^{-4} ) M</th>
<th>( 10^{-6} ) M</th>
<th>Control</th>
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<tbody>
<tr>
<td>Number of samples</td>
<td>3/6</td>
<td>4/6</td>
<td>2/6</td>
<td>0/6</td>
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Figure 2. Cumulative alendronate release (%) of alendronate-loaded porous \( \beta \)-TCP blocks in the culture medium.

Figure 3. Osteoinduction of porous \( \beta \)-TCP blocks loaded with alendronate (\( 10^{-4} \) M) 4 weeks after implantation. (a) Induction of bone (black arrows) was found in a peripheral area of materials more than a central area. Bar = 1000 \( \mu \)m. *, \( \beta \)-TCP; (b) High magnification image of (a) showed that osteoinduction (black arrows) extended along the surface of \( \beta \)-TCP (*) and also within the macroporous spaces where angiogenesis (black arrow heads) was active. Osteoclast-like multinuclear cells (white arrows) attached to a surface of \( \beta \)-TCP. Osteoblast-like cells (white arrow heads) were lining along newly formed bone. Bar = 100 \( \mu \)m; (c) High magnification image of another specimen showed induced bones (black arrows) and collagenous condensations of extracellular matrix (black arrow heads). Bar = 100 \( \mu \)m. *, \( \beta \)-TCP.
3.3. Residual Amount of Porous β-TCP

After 4 weeks, the residual β-TCP area in the control group was the smallest and significantly different from those in the alendronate-loaded groups (Figure 5). The residual β-TCP area in the alendronate-loaded groups increased with the density of alendronate; however, there were no significant differences between the alendronate-loaded groups (Figure 5).

3.4. Histomorphometric Features and Amount of TRAP-Positive Multinuclear Cells

In the alendronate-loaded groups, giant TRAP-positive osteoclast-like cells with more than 10 nuclei were found (Figures 6(a) and (b)). In addition, giant osteoclast-like cells often detached from β-TCP surface, and other mononuclear cells, interposed between the β-TCP surface and the giant osteoclast-like cells, were observed; this finding proves that the detachment was not due to tissue shrinkage during histologic preparation (Figure 6(a)). On the other hand, no giant osteoclast-like cells were found, and the TRAP-positive multinuclear cells were always tightly adherent to the β-TCP surface in the control group (Figure 6(c)). After 4 weeks, the amount of TRAP-positive multinuclear cells divided by the remaining amount of β-TCP in the alendronate-loaded groups was smaller than in the control; however, no significant differences between the groups were observed (Figure 7). Among the alendronate-loaded groups, the amount of TRAP-positive multinuclear cells was lowest in the 10^{-2} M group. However, the obvious tendency to decrease the number of TRAP-positive multinuclear cells by increasing the amount of loaded alendronate was not seen.

3.5. Relative Quantitative Real-Time Polymerase Chain Reaction (qRT-PCR) Analyses

qRT-PCR analysis showed that the expression of TRAP was decreased in the β-TCP specimens loaded with alendronate (Figure 8). Expression of cathepsin K was also slightly decreased in the alendronate-loaded specimens. There were no significant differences between the control and the 10^{-2} M group in the same harvest period. In both groups, the expression of TRAP and cathepsin K increased with time, and the expression of TRAP was

Figure 4. New bone formation rate (%) of porous β-TCP blocks loaded with alendronate (10^{-2} M, 10^{-4} M and 10^{-6} M) and saline (control) 4 weeks after implantation. There was no new bone formation at the control group. *Significant compared with the 10^{-2} M and 10^{-6} M groups (p < 0.05).

Figure 5. Residual amount of porous β-TCP blocks with alendronate (10^{-2} M, 10^{-4} M and 10^{-6} M) and saline (control) 4 weeks after implantation. *Significant compared with the control group (p < 0.05).

Figure 6. TRAP staining sections at 4 weeks. (a) The 10^{-2} M group; giant TRAP-positive multi-nuclear cells with more than 10 nuclei (black wide arrows) were found. Some of them detached from the β-TCP surface and there were mononuclear cells interposed between the β-TCP surface and the giant osteoclast-like cell (black narrow arrows). Bar = 100 µm; (b) The 10^{-4} M group; Giant and hypernucleated TRAP-positive multi-nuclear cells were found (black arrows). Bar = 100 µm; (c) The control group; TRAP-positive multi-nuclear cells were tightly adherent to the β-TCP surface (black arrows). Bar = 100 µm.
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Figure 7. The amount of TRAP-positive multinuclear cells revised by remaining amount of β-TCP in porous β-TCP blocks pre-loaded with alendronate (10^{-2} M, 10^{-4} M and 10^{-6} M) and saline (control) harvested at 4 weeks.

Figure 8. Effect of alendronate on osteoclastic gene expression using quantitative real-time polymerase chain reaction (qRT-PCR). TRAP (a) and Catepsin K (b) mRNA expression in the control group and the alendronate (10^{-2} M) loaded group retrieved at 2, 4 and 8 weeks after implantation.

greater than that of cathepsin K, except in the control group after 2 weeks.

4. Discussion

Although we hypothesized that osteoinduction by alendronate-loaded materials decreases because of the inhibited osteoclastic activity, our study disclosed that local application of alendronate enhances material-induced osteoinduction by porous β-TCP. Bone formation induced by porous β-TCP without any additional osteogenic factors was usually observed 6-8 weeks after implantation in heterotrophic sites [18,19]. However, as reported in the present study, porous β-TCP blocks pre-loaded with alendronate (10^{-2} M, 10^{-4} M, and 10^{-6} M) induced osteogenesis as early as 4 weeks after implantation in dorsal muscles of canines, while no osteoinduction was observed at that time-point in the control group.

Ripamonti et al. [26] reported that local application of zoledronate onto coral-derived hydroxyapatite/calcium carbonate macroporous constructs (7% HA/CC) inhibited osteoinduction 2 months after implantation in heterotopic sites of Papio ursinus. Conversely, the results of this study indicated that local application of alendronate on β-TCP blocks enhanced osteoinduction after 4 weeks much earlier than in the aforementioned report. We contend that this difference in effect of bisphosphonate on osteoinduction depends mainly on the material characteristics. Ripamonti et al. used macroporous materials without microporous structures, and indicated that osteoinduction was inhibited by zoledronate as the osteoclastic post-implantation modifications of the materials decreased, which was followed by the reduction of the macro- and micro-patterned topographies, important for the differentiation of resident stem cells into osteoblast-like cells. We, however, used β-TCP blocks possessing microporous structures on the macropore walls. Consequently, this original topography proved essential for increased osteoinduction observed in this study.

Induction of bone formation was more remarkable in the peripheral zone of the porous β-TCP, where the formation of osteoblasts, osteoclasts, and capillaries was more abundant than in the center. In the macroporous area, mesenchymal collagenous condensations of extracellular matrix were also observed. They are considered the source of tissue patterning and morphogenesis, ultimately promoting osteogenesis within the macroporous materials [27]. These findings suggest that the capillaries generated in the peripheral zone of the porous β-TCP contribute to bone induction through advancement of osteoblast and osteoclast formation and tissue morphogenesis. In addition, we observed newly formed bone tissue spread from pore surface to the central part of macroporous spaces in the alendronate groups, as shown in Figure 3(b). While different from the results obtained by Kondo et al. [18], who indicated that a newly forming bone induced by a single β-TCP implant was always directly attached to the material surface, our finding is consistent with that of Nasu’s report [19], which suggests enhanced osteoinduction by EP4 agonist spread within macroporous spaces of β-TCP. Cumulatively, all aforementioned data indicate that osteoinduction by β-TCP initially occurs on the pore surface and subsequently extends to the central zone of porous spaces, as the process accelerates.

We found that the amount of alendronate released from β-TCP blocks was low. This result indicates that...
most of the loaded alendronate was combined with the blocks after implantation, and that alendronate probably acts on osteoclast-like cells attached to the surface of β-TCP. Moreover, giant, hypernucleated, and detached osteoclast-like cells were found in the sections of the alendronate-loaded implants. The histomorphometric features of osteoclast-like cells reported in this study are the same as those found in the study of bone-biopsy specimens obtained from patients treated with orally administered alendronate, in which the giant, hypernucleated, and detached osteoclasts were observed to undergo protracted apoptosis [23]. Osteoclasts are produced by the fusion of mononuclear precursors and usually acquire up to 8 nuclei before dying by apoptosis, probably as a result of exposure to the high extracellular concentration of released calcium occurring during bone resorption [28, 29]. Inhibition of β-TCP resorption by alendronate would decrease the signal for apoptosis of osteoclast-like cells resulting from calcium release, consequently prolonging the duration of apoptosis and extending the time for fusion of osteoclast-like cells with additional mononuclear progenitors [23].

Duque et al. reported that alendronate exerted an anabolic effect through the stimulation of osteogenic differentiation of human mesenchymal stem cells (MSCs) [30]. What is more, while the inhibitive activity of nitrogen-containing bisphosphonates on osteoblastogenesis using mature calvarial osteoblasts was reported [31,32], a more recent report has shown that alendronate potentiated the effect of geranylgeranyltransferase inhibitor on geranylgeranylation, as well as reverted the effect of farnesylation inhibitor on protein farnesylation, both in favor of osteoblastogenesis [33].

In light of the stimulating effect of alendronate on osteogenic differentiation of MSCs, loading β-TCP blocks with alendronate may contribute to the enhancement of ectopic osteogenesis in the porous β-TCP through activation of the MSCs in muscles around implants.

A particularly interesting finding of this study indicated that pre-loading the porous β-TCP blocks with 10^{-4} M alendronate solution most significantly enhanced osteoinduction. We believe that the 10^{-4} M alendronate-loaded β-TCP proved crucial in providing the desirable balance between the degradation rate of bone scaffolds and their osteoinductive replacement, under the conditions of our study. Whereas high biodegradability and solubility are characteristics important for β-TCP to induce angiogenesis [34], osteoblastogenesis [35], and bone formation, the prolonged exposure to high concentration of Ca^{2+} and phosphate ions released from β-TCP is detrimental to cells, including osteoblasts, osteoclasts, and osteocytes, in vivo [36]. Therefore, the concentration of Ca^{2+} and phosphate ions released from the 10^{-4} M alendronate-loaded blocks seems to be advantageous for both induction of osteogenesis and cell survival.

Finally, the role of osteoclastic activity in material-induced osteoinduction in vivo, as well as the entire mechanism of material-induced osteoinduction remains to be elucidated. Lack of significant suppression of osteoclastic activity after local application of alendronate, even at its highest undiluted density (10^{-2} M), as indicated by qRT-PCR analysis results (Figure 8), constitutes a serious limitation of our study. Significant inhibition of osteoclastogenesis at an early stage, e.g., by means of RANKL (receptor activator of nuclear factor κ-B ligand) antibodies, could prove an interesting approach to further investigation of the influence of osteoclastic activity on intrinsic osteoinduction in vivo.

5. Conclusion

This study showed that loading alendronate on porous β-TCP blocks increased both speed and amount of material-induced osteoinduction of β-TCP after implantation in heterotopic dorsal muscles of canines. The 10^{-4} M alendronate-loaded β-TCP displayed the highest activity, inducing a significantly large amount of new bone formation. We contend that β-TCP loaded with 10^{-4} M alendronate proved crucial in providing the desirable balance between the degradation rate of bone scaffolds and their osteoinductive replacement in this study. Thus, ectopic material-induced osteoinduction may be controlled by local application of alendronate, establishing alendronate loading as a promising therapeutic approach.

6. Disclosures

Olympus Terumo Biomaterials Corp. provided porous β-TCP blocks samples and Teijin Pharma Ltd. provided alendronate. However, there is no conflict of interest related to the work.

7. Acknowledgements

We thank Olympus Terumo Biomaterials Corp. and Teijin Pharma Ltd. for supplying β-TCP and alendronate, respectively. We greatly appreciate the technical support of Haruyasu Kohda and Keiko Furuta for the SEM studies and Hideo Miyatake for the radioisotope studies.

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