

Alveolar restoration following rapid maxillary expansion with and without corticotomy: A microcomputed tomography study in sheep

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Objective: This study examined bone microstructure restoration after rapid maxillary expansion (RME) with and without corticotomy over multiple retention periods. **Methods:** Eighteen male Dorper sheep were randomly distributed into three groups (n = 6 each group): group 1, RME with corticotomy on the buccal and palatal sides; group 2, conventional RME treatment; and group 3, no treatment. Post-RME, trabecular bone microstructure and new bone formation were evaluated by using microcomputed tomography (microCT) and histomorphometry after a 4- or 12-week retention period. Intergroup differences in bone quality and bone remodeling were analyzed by using two-way analysis of variance with Bonferroni *post-hoc* test. **Results:** The bone volume fraction (bone volume [BV]/total volume [TV]) values relative to the control in groups 1 and 2 were 54.40% to 69.88% after the 4-week retention period and returned to approximately 80% after the 12-week retention period. The pooled BV/TV values of the banded teeth in groups 1 and 2 were significantly lower than those of the control after the 4-week retention period ($p < 0.05$). However, after the 12-week retention period, the pooled BV/TV values in group 2 were significantly lower than those in groups 1 and 3 ($p < 0.05$). Histomorphological analysis showed that the new bone formation area in group 1 was approximately two to three times of those in group 2 and control. **Conclusions:** Corticotomy significantly enhanced the restoration of bone quality after the retention periods for banded teeth. This benefit might result from the increased new bone formation after corticotomy.
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Key words: Corticotomy, Rapid maxillary expansion, Bone microstructure, Microcomputed tomography

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INTRODUCTION

Maintenance of teeth in the corrected position after orthodontic treatment remains a challenge for orthodontists.¹⁻³ Little^{2,3} reported a remarkably high percentage of relapse in all types of treated malocclusions over 10 to 20 years post-retention. Among cases in which either fixed or removable appliances were used to increase the arch length in mixed dentition, only 11% showed a satisfactorily enlarged arch over the follow-up period.^{2,3} In children younger than 13 years, the interpremolar and intermolar width could be expanded up to 7 mm or more, but this dimension reduced by 2 to 3 mm after 2 to 7 years post-retention.⁴ Other authors reported relapse rates of up to 63% for conventional maxillary expansion.⁵ In older patients, surgically assisted rapid maxillary expansion (RME) is more preferable, but the relapse rates were also higher, ranging from 5% to 25%.⁵ Thus, the optimal duration for which patients should wear a retainer to prevent relapse still remains to be identified.⁶

The instability of orthodontic outcomes is believed to originate from physiological processes and orthodontic factors.⁶ As teeth are orthodontically moved into a new position, the balance between teeth and surrounding tissues changes. Instability from orthodontic factors may occur due to occlusal contacts, soft tissue pressure, limits of dentition, and periodontal and gingival fibers.⁶ In addition, bone remodeling might play a crucial role in relative tooth stability in its new position.⁷ After active orthodontic treatment, retainers are used to hold the teeth in position for a certain period to facilitate bone remodeling and establish a new equilibrium between the tooth and its surrounding environment. However, the duration of the remodeling process and its adequacy in retaining teeth in their new position remain to be clarified. Although most protocols have recommended retainer usage for 6 weeks to 8 months,⁸⁻¹¹ there is insufficient information on the time required for complete bone remodeling.

Some authors have hinted that corticotomy might enhance the stability of orthodontic outcomes post-treatment.¹²⁻¹⁴ In 1959, Kole¹² reported that cortical bone interfered with the movement of teeth. By performing corticotomy, the resistance of the cortical bone could be released, the tooth could be moved faster into its new position, and relapse could be prevented.¹² Later, Wilcko et al.¹³ also showed that “periodontal accelerated osteogenic orthodontics” increased tooth stability. Chung et al.¹⁴ stated that when the corticotomized area was ossified, this corticotomized segment was impossible to return to its original position.

Corticotomy has been shown to be useful in assisting maxillary expansion, especially for non-growing patients.¹⁵⁻¹⁷ Lines¹⁵ reported successful outcomes with

a combination of buccal incisions and midpalatal suture splitting. Echchadi et al.¹⁶ showed that piezo-bone perforations on the buccal alveolar bone can correct severe maxillary transverse deficiency. Meanwhile, Le et al.¹⁷ illustrated that an adjunctive buccal and palatal corticotomy is also a potential modality.

Recent advancements in microcomputed tomography (microCT) allow three-dimensional examination of the bone microstructure in detail. Nevertheless, no study has used *in vivo* microCT to investigate bone quality after maxillary expansion. Therefore, this study assessed the bone microstructure changes after RME with and without corticotomy over multiple retention periods.

MATERIALS AND METHODS

Sample selection

Eighteen male Dorper sheep aged 20 to 48 months and weighing 50 to 60 kg were used in this study. The study protocol was approved by Institutional Animal Care and Use Committee (IACUC) at Universiti Putra Malaysia (UPM), Malaysia (No. UPM/IACUC/AUP-R031). The sheep were randomly divided into three groups (n = 6 in each group): group 1, RME treatment with corticotomy; group 2, RME treatment; and group 3, no treatment/the control group. According to resource equation method, if the difference between the total number of animals and the total number of groups is between 10 and 20, the sample size is considered adequate, thus indicating that the sample size in this study is indeed adequate.¹⁸ The sheep selected for this study had good general health with no periodontal disease and showed full eruption of all maxillary premolars.

Hyrax expander fixation and corticotomy were performed under general anesthesia. Sheep were anesthetized with intravenous ketamine 7 mg/kg (Narketan[®] 10; Vetoquinol, Magny-Vernois, France) and diazepam 0.6 mg/kg (Diapine; Atlantic Laboratories Corporation Ltd., Bangkok, Thailand). Endotracheal intubation was performed and anesthesia was maintained with isoflurane at a concentration of 2% to 3% in 100% oxygen.

Corticotomy protocol

In group 1, corticotomy surgery was performed with a 2-mm-diameter round bur—perisegmental from the first to the third premolars on the buccal and palatal aspects in accordance with a previous protocol.¹⁷ The depth and width of the incisions were 2 mm (approximating the thickness of sheep cortical bone). The buccal and palatal mucosa were anesthetized via local infiltration of 2% lidocaine HCl with epinephrine 1:100,000 (Lignospan; Septodont, Saint-Maur-des-Fossés, France). A full-thickness flap was elevated. The cutting lines on the buccal aspect included vertical incisions parallel to the tooth

axis made at the alveoli mesial to the first premolars and distal to the third premolars, which communicated with a horizontal line made above the root apex at approximately 5 mm. On the palatal side, the vertical incisions were similar to those on the buccal side, but the horizontal cut was performed approximately 5 mm from the cemento-enamel junction. Subsequently, the buccal and palatal flaps were closed with absorbable suture material (PDS™ Plus; Ethicon Inc., Somerville, NJ, USA). Hyrax appliances were cemented, and the expander screw was turned by 1 mm immediately after the surgery. After a 5-day lag, the expansion screw was turned at the rate of 0.5 mm/day.

When an incision is placed on the bone, the regional acceleratory phenomenon (RAP) occurs within a few days and accelerates the bone turnover. The RAP is a series of local healing events in response to a noxious stimulus. In this event, normal cellular activities are accelerated to facilitate the healing process. The dominant features of RAP are the burst of anabolic and catabolic events at the injured sites. These activities also lead to demineralization of the local bone.¹⁹ In this study, the expansion screw was only turned 1 mm right after surgery and not turned for 5 days to take the advantage of the demineralization produced by RAP. The bone-healing process after injury has been reported to include two phases, i.e., woven bone formation and transformation of woven bone into lamellar bone. Woven bone started to form within a few days after injury and reached its peak thickness at day 7. After seven days, the woven bone at the cortical area was remodeled to lamellar bone, which increased the bone strength.¹⁹ Thus, ceasing expansion screw turning for 5 days was ideal because it could take the advantage of demineralization produced by RAP, while avoiding the disadvantage of increasing bone strength.

Hyrax appliance design and activation protocol

The Hyrax expander consisted of a 16-mm Super Screw® (Great Lakes Orthodontics, Tonawanda, NY, USA) placed parallel to the midpalatal suture in the region of the second premolars. The extending arms of Super Screw® were soldered to four bands cemented to the first and third premolars with glass ionomer luting cement (Ketac™ Cem Easymix; 3M, St. Paul, MN, USA).

In both groups, the expansion screws were turned by 0.5 mm/day to attain a total of 8 mm of activation in groups 1 and 2. After the active phase, all expanders were kept in position for 4 or 12 weeks. Subsequently, in every group, 3 sheep were sacrificed at 4 or 12 weeks of retention. The upper jaws were separated by a sharp saw and preserved in 4% paraformaldehyde solution for 1 month.

Microcomputed tomography protocol

All maxillary samples of the sheep were covered with Parafilm and scanned using Skyscan 1176 microtomography scanner (Bruker, Kontich, Belgium). MicroCT scanning was performed at a resolution of 18 μm with the following parameters: 90 kV, 278 μA, 0.5-mm aluminum filter, and 360-degree rotation range. All acquired images were reconstructed by using NRecon software (version 1.6.10.4; Bruker) with a Gaussian smoothing of 3, beam hardening correction of 40%, and ring artefact reduction of 12. DataViewer (version 1.5.2.4 64-bit; Bruker) was used to reorient the direction of the reconstructed images to the sagittal plane. Subsequently, all images were imported into CTAn software (version 1.15.4.0+; Bruker) for bone microstructural analysis.

The furcation areas enclosed by the buccal and lingual roots of premolars were defined as the region of interest (ROI) (Figure 1A). After selection of the ROIs, the automatic (Otsu method) threshold was applied for all images during the analysis. The bone microstructure

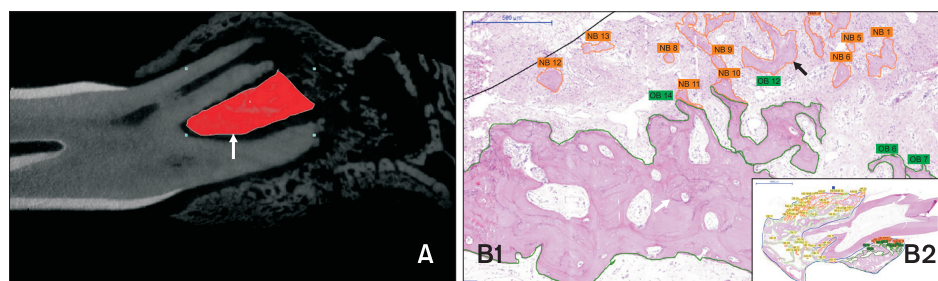


Figure 1. Region of interest (ROI) in quantification of bone microstructure (A) and new bone (NB) and old bone (OB) formation (B1 and B2). A, The furcation enclosed by the buccal and lingual roots of premolars was defined as the ROI (arrow) for measurements of bone microstructure. B, Magnification at hematoxylin and eosin (H&E) stain ($\times 200$) showing NB and OB. NB is defined as the amorphous eosinophilic material with porous bone and pale cement lines (black arrow). OB is the bone with a compact feature and prominent cement lines (white arrow) (B1); the total tissue area (outlined by the blue line) at H&E stain ($\times 20$) (B2).

parameters are presented in Table 1. Since the left and right sides were pooled together, the total number of samples was 36 and every group contained 6 samples taken after 4 and 12 weeks of retention.

Histomorphometric analysis of new bone formation

The obtained maxillary samples (including the second and third premolars) were separated into left and right sides with a sharp saw. A coin was flipped for every sample to choose the left or right side for histological analysis. The total number of samples was 36, in which each group contained 12 samples—6 samples each taken after the 4- and 12-week retention periods. Among these six samples, 3 were the second premolar blocks, while the rest were the third premolar blocks.

The maxillary samples were rinsed with a phosphate-buffered saline solution and decalcified with 10% ethylene diamine tetra-acetic acid (pH 7.4) at 4°C for a month. Following decalcification, the tissue was dehydrated in ethyl alcohol of gradually increasing concentrations from 70% to 100% and embedded in paraffin. All samples were sagittally sectioned at 5- μ m thickness through the second and third premolar roots using a microtome. Two slides were made for each area. Subsequently, all slides were stained with hematoxylin and eosin for histological examination.

The whole-slide imaging data were digitized at 40 \times magnification by using a Panoramic Desk slide scanner (3DHISTECH Ltd., Budapest, Hungary). Quantification of the new bone and old bone areas (as defined in Table

1) was performed by using Panoramic Viewer software (version 1.14.4; 3DHISTECH Ltd.). The ROI referred to the whole area of the tissue surrounding the premolar tooth, as shown on the slide (Figure 1B2). New bone was defined as the amorphous eosinophilic material with porous bone and pale cement lines, while old bone was the bone with a compact feature and prominent cement lines (Figure 1B1).

Statistical analysis

All data were analyzed by using SPSS Statistics version 20.0 (IBM Corp., Armonk, NY, USA). The Shapiro–Wilk test demonstrated that the data were normally distributed ($p > 0.05$). Two-way analysis of variance (ANOVA) with Bonferroni *post-hoc* assessment were employed to compare the intergroup differences. A p -value of 0.05 was considered statistically significant.

RESULTS

Bone microstructure changes after 4- and 12-week retention periods

After the 4-week retention period, bone volume fraction (BV/TV) and Tb.Th values in both intervention groups decreased significantly in comparison with those in the control group ($p < 0.05$; Table 2). The relative values of BV/TV to the control in groups 1 and 2 were 54.40% to 69.88% (Figure 2). After 12 weeks of retention, BV/TV, Tb.Th, and Tb.N values in both intervention groups increased in comparison with those after the

Table 1. Definition of parameters used in the quantification of bone microstructure and new bone formation in the region of interest (ROI)

Abbreviation	Parameter	Description
Parameters in quantification bone microstructure in the ROI		
BV/TV (%)	Bone volume fraction	Ratio of bone volume to the total volume tissue in the ROI
Tb.Th (μ m)	Trabecular thickness	Average of the thickness of trabecular
Tb.Sp (μ m)	Trabecular separation	Average distance between trabeculae
Tb.N (1/ μ m)	Trabecular number	Average number of trabeculae per unit length in the ROI
Parameters in quantification of new bone formation		
NB (μ m ²)	New bone	The area of new bone in the ROI
OB (μ m ²)	Old bone	The area of old bone in the ROI
TT (μ m ²)	Total tissue	The area of the whole tissue surrounding the tooth.
TB = NB + OB (μ m ²)	Total bone	The area of new and old bone in the ROI
NBF = NB/TB	New bone fraction	The ratio of new bone to total bone area in ROI
OBF = OB/TB	Old bone fraction	The ratio of old bone to total bone area in ROI
TNBF = NB/TT	Total new bone fraction	The ratio of new bone to total tissue area in ROI
TOBF = OB/TT	Total old bone fraction	The ratio of old bone to total tissue area in ROI
TBF = TB/TT	Total bone fraction	The ratio of total bone and total tissue area in ROI

Table 2. Bony microstructural changes in groups 1–3 after the 4- and 12-week retention periods

Bone structural parameter	BV/TV (%)	Tb.Th (µm)	Tb.Sp (µm)	Tb.N (1/µm)
4-week				
G1 (n = 6)				
P1	37.79 ± 6.01	88.11 ± 16.79	99.78 ± 10.43	0.0042 ± 0.0009
P2	33.07 ± 7.91	83.83 ± 13.99	108.92 ± 21.69	0.0039 ± 0.0006
P3	25.04 ± 8.70	74.17 ± 21.04	111.56 ± 30.08	0.0034 ± 0.0008
G2 (n = 6)				
P1	31.46 ± 4.60	72.53 ± 14.50	127.08 ± 19.46	0.0044 ± 0.0080
P2	33.66 ± 4.42	91.64 ± 11.48	146.27 ± 34.26	0.0038 ± 0.0010
P3	27.80 ± 7.51	93.25 ± 33.36	135.48 ± 31.99	0.0032 ± 0.0012
G3 (n = 6)				
P1	46.98 ± 6.39	99.11 ± 15.33	119.03 ± 23.91	0.0053 ± 0.0020
P2	48.17 ± 6.73	113.69 ± 25.21	134.69 ± 28.22	0.0051 ± 0.0029
P3	46.03 ± 8.90	119.39 ± 44.61	121.75 ± 24.71	0.0041 ± 0.0010
12-week				
G1 (n = 6)				
P1	43.95 ± 4.73	92.12 ± 11.18	87.62 ± 21.81	0.0048 ± 0.0006
P2	37.72 ± 9.85	100.98 ± 17.10	130.27 ± 40.33	0.0038 ± 0.0010
P3	37.63 ± 4.25	89.07 ± 18.57	102.32 ± 23.74	0.0043 ± 0.0014
G2 (n = 6)				
P1	38.26 ± 6.17	86.69 ± 11.58	104.63 ± 18.07	0.0045 ± 0.0013
P2	38.74 ± 7.44	113.32 ± 30.61	149.70 ± 26.35	0.0037 ± 0.0011
P3	28.73 ± 4.34	92.49 ± 36.85	156.62 ± 27.87	0.0028 ± 0.0010
G3 (n = 6)				
P1	47.72 ± 6.49	99.11 ± 13.61	116.44 ± 26.28	0.0054 ± 0.0023
P2	48.92 ± 7.06	114.75 ± 27.54	127.76 ± 16.17	0.0052 ± 0.0300
P3	47.32 ± 9.29	119.06 ± 21.47	118.55 ± 28.48	0.0049 ± 0.0022
p-value*				
4-week				
Teeth	0.029 (P1 vs. P3)	0.435	0.326	0.174
Group	0.000 (G1 vs. G3; G2 vs. G3)	0.002 (G1 vs. G3; G2 vs. G3)	0.013 (G1 vs. G2)	0.638
Teeth × Group	0.301	0.431	0.966	0.977
12-week				
Teeth	0.062	0.083	0.004 (P1 vs. P2)	0.353
Group	0.000 (G1 vs. G3; G2 vs. G3)	0.067	0.010 (G1 vs. G2)	0.567
Teeth × Group	0.248	0.570	0.179	0.519

Values are presented as mean ± standard deviation.

G, Group; G1, corticotomy + rapid maxillary expansion (RME); G2, conventional RME; G3, control; P1, first premolar; P2, second premolar; P3, third premolar.

*Two-way analysis of variance with Bonferroni *post-hoc* assessment for comparisons among groups.

See Table 1 for the definitions of each measurement.

4-week retention period, with a higher increment noted in group 1 ($p > 0.05$; Table 2). At this point of time, the BV/TV values in all groups (except for the third premolar region in group 2) returned to approximately 80% relative to those of the control (Figure 2).

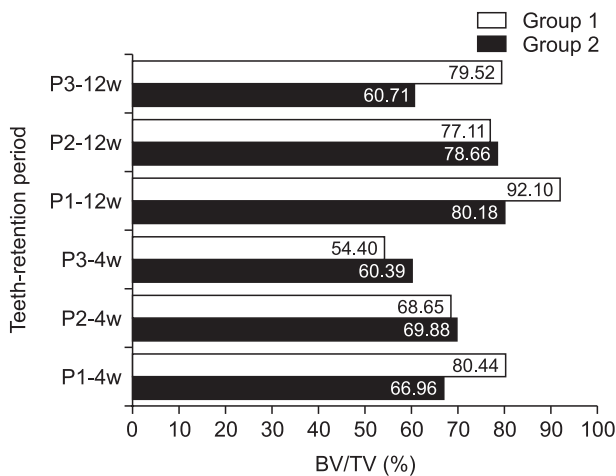


Figure 2. Relative values of bone volume fraction (BV/TV) at the premolar regions compared to the control values in groups 1 (corticotomy + rapid maxillary expansion [RME]) and 2 (conventional RME) after 4 and 12 weeks of retention. P1, First premolar; P2, second premolar; P3, third premolar; 4w, 4-week retention; 12w, 12-week retention.

The bone microstructure data of the banded teeth (first premolar and third premolar) in the three groups were pooled and analyzed as shown in Table 3. After the 4-week retention period, the pooled BV/TV and Tb.Th values in both intervention groups were significantly lower than those in the control group ($p < 0.05$; Table 3). Nevertheless, after the 12-week retention period, the pooled BV/TV value of group 2 was significantly lower than those of groups 1 and 3 ($p < 0.05$; Table 3 and Figure 3). At this point of time, the BV/TV value in group 1 returned to 85.80% relative to that of the control, while it was lower in group 2 (70.47%; Figure 3). The intraclass correlation coefficients (ICCs) were 0.95, 0.96, 0.97, and 0.94 for BV/TV, Tb.Th, Tb.Sp, and Tb.N, respectively.

Histomorphometric analysis of new bone formation after 4- and 12-week retention periods

After the 4-week retention period, the ratio values related to the new bone changes (new bone fraction [NBF] and total new bone fraction [TNBF]) in group 1 were approximately two to four times higher than those obtained for groups 2 and 3 ($p < 0.05$; Table 4). The new bone formation in group 2 was just slightly higher than that in the control group ($p > 0.05$; Table 4). However, the total bone fraction (TBF) values in the two intervention groups were significantly lower than that in the control ($p < 0.05$; Table 4) at this point of time, indicating an overall decrease in the quantity of alveolar bone during the experimental period.

Table 3. Differences in bony microstructural changes of pool banded teeth (P1 + P3) in groups 1–3 after the 4- and 12-week retention periods

Bone structural parameter	BV/TV (%)		Tb.Th (μm)		Tb.Sp (μm)		Tb.N (1/μm)	
	Value	p-value*	Value	p-value*	Value	p-value*	Value	p-value*
4-week		0.000 (G1 vs. G3, G2 vs. G3)		0.024 (G1 vs. G3, G2 vs. G3)		0.053 (G1 vs. G2)		0.862
G1 (n = 12)	31.41 ± 9.76		81.15 ± 19.56		105.67 ± 22.33		0.0038 ± 0.0009	
G2 (n = 12)	29.64 ± 6.24		82.89 ± 26.81		131.28 ± 25.62		0.0038 ± 0.0012	
G3 (n = 12)	46.50 ± 7.40		109.25 ± 25.22		120.39 ± 23.23		0.0047 ± 0.0016	
12-week		0.000 (G1 vs. G2, G2 vs. G3)		0.043 (G2 vs. G3)		0.047 (G1 vs. G2)		0.639
G1 (n = 12)	40.78 ± 5.41		90.60 ± 14.70		94.98 ± 20.98		0.0045 ± 0.0011	
G2 (n = 12)	33.49 ± 7.11		89.59 ± 26.22		130.63 ± 35.19		0.0038 ± 0.0013	
G3 (n = 12)	47.52 ± 7.64		109.08 ± 20.06		117.50 ± 26.15		0.0052 ± 0.0022	

Values are presented as mean ± standard deviation.

G, Group; G1, corticotomy + rapid maxillary expansion (RME); G2, conventional RME; G3, control.

*One-way analysis of variance with Bonferroni *post-hoc* assessment for comparison among groups.

See Table 1 for the definitions of each measurement.

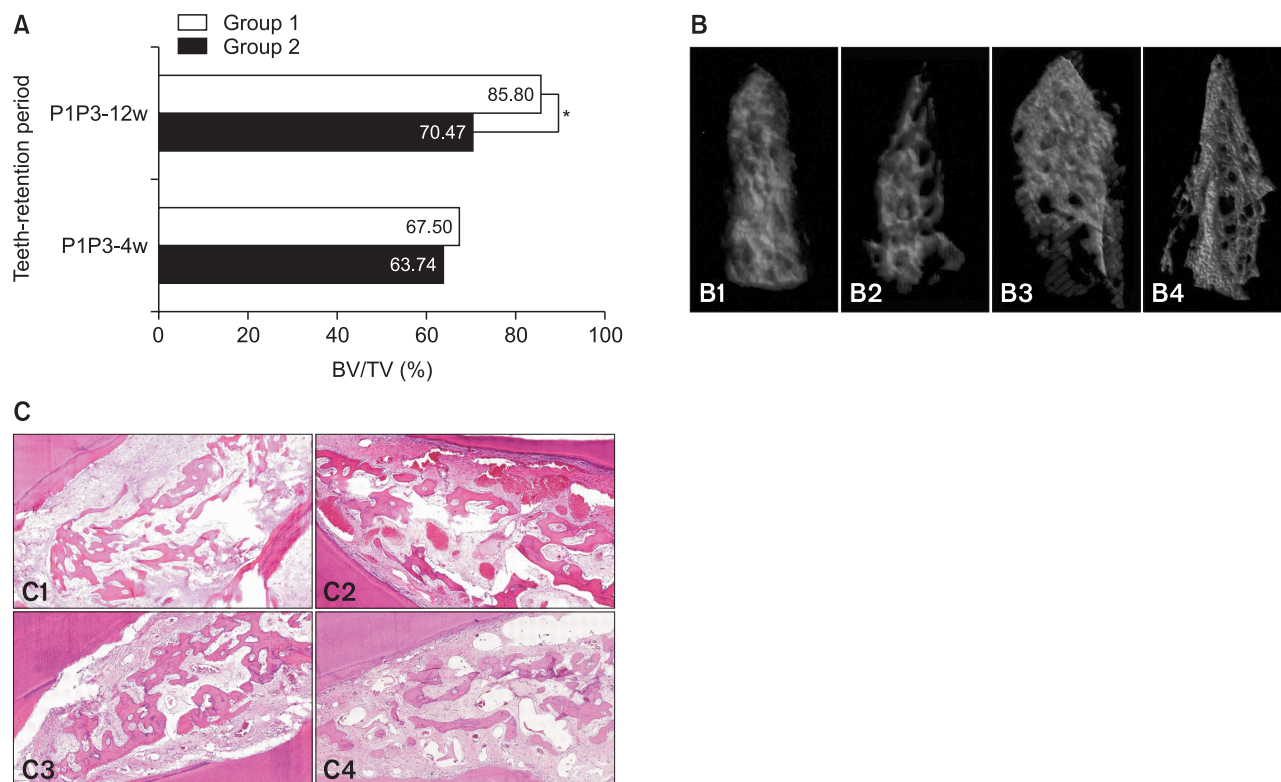


Figure 3. **A**, Relative values of bone volume fraction (BV/TV) values compared to the control values for the banded teeth (the first and third premolars) in groups 1 (corticotomy + rapid maxillary expansion [RME]) and 2 (conventional RME) after 4 and 12 weeks of retention. **B**, Alveolar trabecular bone microstructural images of group 1 after 4 (B1) and 12 weeks of retention (B3) and of group 2 after 4 (B2) and 12 weeks of retention (B4). **C**, Histological images at H&E stain (×100) showed that the bone volume in group 1 after 4 and 12 weeks of retention (C1 and C3) were higher than those in group 2 after 4 and 12 weeks of retention (C2 and C4) (H&E stain, ×100).

P1, First premolar; P3, third premolar; 4w, 4-week retention; 12w, 12-week retention.

* $p < 0.05$.

After the 12-week retention period, NBF value in group 1 slightly decreased, but they were still significantly higher (two to three times) than those in groups 2 and 3 ($p < 0.05$; Table 4). Interestingly, the high levels of new bone formation activity may be responsible for the fact that the TBF in group 1 was not significantly lower than that in the control group ($p > 0.05$), while this value in group 2 was still rather low ($p < 0.05$). The ICC results were 0.93, 0.96, and 0.97 for new bone, old bone, and total tissue, respectively.

DISCUSSION

Adequate bone remodeling around orthodontically moved roots should play an important role in limiting the tooth from turning back to its previous position. Recently, corticotomy has emerged as a promising intervention for this purpose. One of the suggested advantages of corticotomy is that it can enhance the stability

of orthodontic results,¹²⁻¹⁴ but its impact on tooth stability has still not been elucidated. Therefore, the benefits of corticotomy in rehabilitating bone quality during retention periods should be elaborated in comparison with conventional RME.

This study involved adult sheep, whose growth had ceased. In addition, this animal had a larger dental arch than rodents, and the shapes of sheep teeth were convenient for retention of tooth-borne expanders, which facilitated the use of maxillary expansion procedures similar to those used in humans.

After the 4-week retention period, a significant decrease in BV/TV and Tb.Th was observed in the two intervention groups in comparison with the control. These results imply that RME influenced the trabecular thickness and BV/TV more than other bone quality indices. After 12 weeks of retention, increments were observed in the BV/TV, Tb.Th, and Tb.N values for the two intervention groups. The BV/TV values relative to the control

Table 4. New and old bone components in supporting tissue in groups 1–3 after 4 and 12 weeks of retention

Ratio of bone parameter	NBF	OBF	TNBF	TOBF	TBF
4-week					
G1 (n = 3)					
P2	0.199 ± 0.075	0.801 ± 0.075	0.003 ± 0.0002	0.013 ± 0.007	0.015 ± 0.006
P3	0.165 ± 0.059	0.835 ± 0.059	0.002 ± 0.0006	0.010 ± 0.003	0.012 ± 0.004
G2 (n = 3)					
P2	0.055 ± 0.030	0.945 ± 0.030	0.001 ± 0.0003	0.018 ± 0.005	0.019 ± 0.005
P3	0.064 ± 0.032	0.936 ± 0.032	0.001 ± 0.0002	0.014 ± 0.011	0.015 ± 0.011
G3 (n = 3)					
P2	0.035 ± 0.011	0.965 ± 0.011	0.001 ± 0.0002	0.031 ± 0.009	0.032 ± 0.009
P3	0.041 ± 0.018	0.959 ± 0.018	0.001 ± 0.0002	0.033 ± 0.016	0.034 ± 0.016
12-week					
G1 (n = 3)					
P2	0.116 ± 0.046	0.912 ± 0.036	0.003 ± 0.0008	0.023 ± 0.005	0.025 ± 0.006
P3	0.125 ± 0.045	0.875 ± 0.045	0.002 ± 0.0007	0.016 ± 0.001	0.019 ± 0.001
G2 (n = 3)					
P2	0.081 ± 0.054	0.919 ± 0.055	0.001 ± 0.0008	0.014 ± 0.002	0.015 ± 0.001
P3	0.056 ± 0.026	0.944 ± 0.026	0.001 ± 0.0003	0.015 ± 0.003	0.017 ± 0.003
G3 (n = 3)					
P2	0.031 ± 0.016	0.971 ± 0.018	0.001 ± 0.0009	0.031 ± 0.011	0.032 ± 0.012
P3	0.032 ± 0.010	0.968 ± 0.010	0.001 ± 0.0002	0.030 ± 0.010	0.031 ± 0.010
p-value*					
4-week					
Teeth	0.777	0.777	0.043 (P2 vs. P3)	0.753	0.709
Group	0.000 (G1 vs. G2; G1 vs. G3)	0.000 (G1 vs. G2; G1 vs. G3)	0.000 (G1 vs. G2; G1 vs. G3)	0.009 (G1 vs. G3; G2 vs. G3)	0.011 (G1 vs. G3, G2 vs. G3)
Teeth × Group	0.650	0.447	0.152	0.883	0.859
12-week					
Teeth	0.78	0.766	0.356	0.523	0.517
Group	0.004 (G1 vs. G3)	0.009 (G1 vs. G3)	0.003 (G1 vs. G2; G1 vs. G3)	0.004 (G1 vs. G3; G2 vs. G3)	0.006 (G2 vs. G3)
Teeth × Group	0.371	0.337	0.954	0.552	0.605

Values are presented as mean ± standard deviation.

G, Group; G1, corticotomy + rapid maxillary expansion (RME); G2, conventional RME; G3, control; P2, second premolar; P3, third premolar.

*Two-way analysis of variance with Bonferroni *post-hoc* assessment for comparison among groups.

See Table 1 for the definitions of each measurement.

recovered to more than 77% in most of the regions, except for the third premolar region in group 2 (60.71%). Interestingly, the pooled BV/TV value of banded teeth in group 1 was significantly higher than that in group 2 after the 12-week retention period. Therefore, corticotomy surgery improved the recovery of bone quality of banded teeth in group 1 in comparison with that in

group 2.

Some studies have quantified the alveolar bone density after orthodontic tooth movement.²⁰⁻²² Verna et al.²⁰ reported that the mesial tooth movement in rats caused a noticeable reduction in the BV/TV. This decrease occurred at the alveolar bone adjacent to the displaced teeth and also extended to the bone surrounding its

neighboring teeth. Recent studies in humans have shown that the alveolar density around the roots reduced by 20% to 29%. When the tooth was moved more than 0.5 mm, the density reduction increased up to 59.00% to 69.10%.^{21,22} However, these studies did not reveal the bone quality recovery during the retention period. In addition, the values of bone microstructure parameters evaluated via cone-beam computed tomography images were less accurate when compared to those obtained from microCT.²³ Although our study did not evaluate the bone quality changes immediately after the expansion, the BV/TV values after the 4-week retention period were around 60% than of those of the control group. This result implies that the bone quality would have decreased more than 60%, which is in agreement with past studies.²⁰⁻²² Another advantage of this study refers to the bone microstructure parameters, which were measured by using microCT to ascertain measurement accuracy.

A previous study reported that the rate of tooth relapse and bone quality changes after tooth movement in 10 days.²⁴ Franzen et al.²⁴ showed that the molars relapsed very quickly from the first day and the amount of relapse was up to 73% of the achieved tooth movement. The relapse rate started to stabilize after 7 days, and the total relapse after 12 days was very high at 93%. The BV/TV increased from 17% to 25% within 21 days of follow-up.²⁴ In our study, we did not evaluate the bone volume changes right after the activation phase since the high tendency of tooth relapse at this point of time might affect the results for bone quality. After the 4-week retention, bone microstructure recovered to a certain level to prevent the tooth from immediately returning back to its previous position.

Studies on the effect of retention after long-term orthodontic treatment are scarce. van Leeuwen et al.⁷ reported that teeth in the group without retention showed relapse immediately after orthodontic appliance removal. Conversely, teeth in the 3-month retention group showed a lower degree of relapse and at a slower rate. These authors also suggested that retention was crucial when the tooth was moved greater than 4 mm.⁷ As the bone microstructural value after the 12-week retention period was higher than that after 4 weeks, this study showed that the retention period did influence the recovery of bone quality after orthodontic treatment. However, at the 12-week retention period, the bone quality only returned to approximately 80% of the baseline. Therefore, retention periods exceeding 3 months are crucial to maintain the achieved treatment outcomes since a 3-month retention time is inadequate to remodel the alveolar bone for complete rehabilitation.

The role of corticotomy in bone quality rehabilitation is more obvious for the banded teeth. After 12 weeks of retention, the pooled BV/TV value in group 1 was sig-

nificantly higher than that in group 2. Banded and wire-supported teeth would bear different loads; therefore, their mechanical and biological responses would differ.²⁵ In addition, some authors confirmed that the accumulated load from the expander caused root resorption even after the activation period.²⁶ These findings might explain the slow recovery of bone microstructure around the banded teeth in group 2. Noticeably, the relative value of BV/TV to the control in the third premolar region in group 2 increased slightly (from 60.39% to 60.71%) after the 12-week retention period, which may signify an overloaded expansion force to this tooth. However, decortication was believed to release the resistance of cortical bone.¹² Chung et al.¹⁴ showed that the disruption of cortical bone facilitates the displacement of the bone segment. Corticotomy resulted in discontinuation of the cortical bone and may help reduce the accumulative stress. Therefore, the bone quality recovered better in group 1 than in group 2. In another aspect, the benefit of corticotomy was not expressed clearly on wire-supported teeth, which could be due to the fewer incisions (only horizontal incisions at the apex) around the second premolars.

The benefits of corticotomy may also be derived from RAP. Sebaoun et al.²⁷ claimed that the anabolic turnover increased by two- to three-fold after decortication in rats. The hike in anabolic activity induces greater bone apposition in the trabecular bone around the molar roots.²⁷ The findings reported by Sebaoun et al.²⁷ are in agreement with other studies, in which anabolic activity and bone surface area were shown to dramatically increase some weeks after corticotomy.^{28,29} Parallel to previous studies, this study reveals that the rate of new bone formation in group 1 was three to four times greater than that in group 2 over the retention periods. This implies that the bone remodeling activities were very active in the corticotomy group; therefore, the rehabilitation of the alveolar bone was higher in group 1 than in group 2.

To date, the relative bone quality changes during the retention period, especially after maxillary expansion, have not been well-documented. This study revealed that bone quality in terms of bone microstructure increased during the retention period. However, a 3-month retention period is insufficient for complete rehabilitation of bone microstructure, which explains the possibility of relapse after retention. In addition, corticotomy remarkably enhanced the restoration of bone quality and therefore had the potential to enhance the stability of orthodontic outcomes.

CONCLUSION

Corticotomy significantly enhanced the restoration of

bone microstructure of banded teeth during the retention periods and might have the potential to increase the stability of orthodontic outcomes. The rate of bone remodeling was very active in the corticotomy group. New bone formation in group 1 was two to three times higher of those in groups 2 and 3.

CONFLICTS OF INTEREST

No potential conflict of interest relevant to this article was reported.

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