Osterix regulates corticalization for longitudinal bone growth via integrin β3 expression

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Abstract
Corticalization, coalescence of trabecular bone into the metaphyseal cortex, is important for the longitudinal growth of long bones. However, little is known about the molecular mechanisms controlling corticalization. To understand the molecular mechanisms underlying corticalization, we analyzed osteoblast-specific Osterix-knockout mice (Col-OMT). In control mice, corticalization was initiated after 7 postnatal days, and the number of osteoblasts in the peripheral spongiosa was increased compared to the number in the central spongiosa. In contrast, in Col-OMT mice, corticalization was delayed, and the number of osteoblasts in peripheral zones was unchanged compared to the central zone. Furthermore, femoral length was decreased in Col-OMT mice at 1 month. Because Col-OMT mice exhibited impaired matrix coalescence and osteoblast migration, we evaluated integrin signaling in Col-OMT mice. Osterix bound to the \( \text{Itgb3} \) promoter and increased transcription of the \( \text{Itgb3} \) gene in osteoblast cells. Interestingly, the inner and outer cortical bones were separated in \( \text{Itgb3} \)-null mice at postnatal day 7. In \( \text{Itgb3} \)-null mice, the number of osteoblasts in peripheral zones was not changed, and the femoral length was decreased. Taken together, these results indicate that Osterix regulates corticalization for longitudinal bone growth via the control of integrin β3 expression in osteoblasts. Our findings imply that the ability to control osteoblast function during corticalization may help in the treatment of short stature.

Introduction
The longitudinal growth of mammalian long bones is important in the determination of final body height and normal bone structure. The process of longitudinal bone growth is quite complex and is tightly regulated by several factors1. Following growth plate development, trabecular bone is formed around calcified cartilage in the ossification zone in a process called endochondral bone formation2–5. The newly formed trabecular bone coalesces at the metaphyseal cortex. This event is called "corticalization" and is the process by which longitudinal bone growth is completed6. The corticalization process during early growth is essential for the prediction of trabecular and cortical morphology in adulthood7. Furthermore, impaired corticalization during growth influences the fragility of the distal radius in postmenopausal women8. Thus, corticalization is the last stage of longitudinal growth and plays an important role in completing the structure of adult bone, though the underlying genetic and molecular mechanisms of this process remain unknown.

Longitudinal limb growth depends not only on intracellular chondrocyte signals in the growth plate but also on wide variety of mechanisms involving paracrine factors, multiple hormones, and extracellular matrix (ECM) molecules9. Therefore, normal variation in height cannot be defined solely by growth plate problems but must include many other genetic factors.
Many studies have shown that osteoblasts and osteocytes function, in part, to regulate limb length. Mice lacking or overexpressing major bone integrity proteins exhibit longitudinal limb growth defects when compared to control mice without growth plate anomalies\textsuperscript{10–13}. Moreover, considering that corticalization occurs after growth plate area expansion, osteoblasts and osteocytes can directly affect limb length. However, the function of osteoblasts and osteocytes has not yet been associated with longitudinal growth.

During the process of endochondral bone development, the cells in the inner mesenchymal condensation compartment differentiate into chondrocytes and undergo progressive maturation into hypertrophic chondrocytes\textsuperscript{6}. On the other hand, the outer mesenchymal condensation forms the perichondrium, which directly differentiates into osteoblasts and generates a calcified bone matrix called the “bone collar” that surrounds mature chondrocytes\textsuperscript{14}. At embryonic day 14.5 (E14.5), blood vessels invade the center of hypertrophic chondrocytes, leading to resorption of the cartilage matrix and formation of the trabecular bone matrix, which is secreted by invading osteoblast precursors and chondrocyte-derived osteoblasts\textsuperscript{15–17}. During corticalization, trabecular bone must coalesce into cortical bone; trabecular bone is located in the inner compartment and is separated from the bone collar. After the development period, the trabecular bone will become the cortical bone in the outer compartment (Fig. 1). According to previous studies suggesting the mechanism of corticalization, mechanical stimuli and increased osteoblast surface area are both associated with corticalization\textsuperscript{18,19}. From these proposed mechanisms and the absence of corticalization in the embryonic period, we hypothesized that osteoblast differentiation would be involved in the corticalization mechanism.

Osterix (Osx) is essential for osteoblast differentiation and bone formation during the embryonic and postnatal periods\textsuperscript{20,21}. Osx also affects osteoblast migration, apoptosis, and death\textsuperscript{22}. Osx deficiency in postnatal mice is associated with reductions in femoral length and thin, porous cortical bone\textsuperscript{11,23}. In particular, trabecular bone is increased without growth plate abnormalities or defects in osteoclastogenesis. These results suggest that Osterix plays an important role in proper bone corticalization.

Corticalization is governed by mechanical stimuli\textsuperscript{19}. Mechanical stimulation triggers signaling mechanisms that promote osteoblast proliferation, differentiation, and survival through integrin signaling\textsuperscript{23–26}. Integrins are transmembrane molecules composed of α and β chains that bind to ECM proteins and play key roles in cell–cell and cell–ECM adhesion as well as in cell migration\textsuperscript{27}. Those roles are analogous to the process of corticalization, in which matrix components and osteoblast in the inner compartment adhere and unify to create the outer compartment.

Here, we show the origin of corticalization over sequential time points in femora. To investigate the role of osteoblasts in longitudinal bone growth and the molecular changes that induce corticalization in osteoblasts at specific stages, we used osteoblast-specific Osx-knockout mice. Our results provide evidence that Osterix regulates corticalization by directly controlling integrin β3 expression.

**Materials and methods**

**Animals**

Osx\textsuperscript{fl/fl} and Col1a1-Cre mouse strains have been previously described\textsuperscript{11}. Itgb3 (B6.129S2-Itgb3\textsuperscript{tm1Hyn/JSemb}) mice were purchased from Jackson Laboratory (Bar Harbor, ME, USA). To generate the Col1a1-Cre;Osx\textsuperscript{fl/fl} (Col-OMT) mice, Col1a1-Cre;Osx\textsuperscript{fl/+} (control) mice were crossed with Osx\textsuperscript{fl/fl} mice as appropriate. Itgb3 heterozygous mice were intercrossed to generate Itgb3-null mice. Mouse offspring were genotyped by polymerase chain reaction (PCR) with previously described primers\textsuperscript{11,28}. Animals were housed in an accredited facility under a 12-h light/dark cycle and were provided water and food ad libitum. The Animal Welfare Committee of Chonbuk National University approved all protocols prior to the start of the studies.

**Histology and immunohistochemistry**

Femora dissected from male mice were fixed in 4% paraformaldehyde at 4°C overnight and decalcified in a 15% ethylenediaminetetraacetic acid solution for 2–3 weeks at 4°C. To observe exactly the same site in each femur, femora were sectioned based on two points. The proximal point was the piriformis insertion site, and the distal point was the anterior or posterior cruciate ligament origin site. Thus, we were able to accurately obtain sections from femoral centers (Figure S1). Embedded tissues were sectioned at a thickness of 5 μm. To evaluate histologic findings, sections were stained with hematoxylin and eosin (H&E), Safranin O or picrosirius red (Sigma-Aldrich, St. Louis, MO, USA). For immunohistochemistry, the EXPOSE Rabbit Specific HRP/DAB Detection IHC Kit (Abcam, Cambridge, MA, USA) was used according to the manufacturer’s instructions. The sections were stained using an antibody against vinculin (1:100; Abcam).

**Femoral length measurement**

To accurately compare femoral lengths, we sectioned the middle part of embedded femoral tissues obtained at each time point. After H&E staining, we measured femoral length from the starting point of the distal
femoral primary spongiosa to the starting point of the proximal femoral primary spongiosa, both of which were identified by stereoscopy.

**Micro-CT assessment**

The femora were scanned with a micro-CT (1076 SkyScan Micro-CT, Skyscan, Kontich, Belgium) at 8-μm resolution and analyzed with CTAn software (Skyscan). A global thresholding algorithm was applied at a constant threshold for all specimens. The threshold was the intensity (gray value) that corresponded to ~50% of the average intensity of intact cortical bone. Trabecular lesions and metaphyseal cortical bone samples were analyzed in the distal metaphysis extending proximally 1.75 mm from the end of the primary spongiosa.

**Cell culture**

An osteoblastic cell line, MC3T3-E1, was purchased from the American Type Culture Collection (Manassas, VA, USA) and was maintained in α-MEM with 10% FBS and 100 IU/ml penicillin–100 μg/ml streptomycin at 37 °C under 5% CO₂.

**Western blot analysis**

The femora of control and Col-OMT male mice were removed from soft tissue, and the bone marrow was
flushed away. Femora were frozen in liquid nitrogen and then crushed with a mortar and pestle. Then, 10 μg of protein from either bone tissue or cell lysates was separated by 10% SDS-PAGE and transferred to PVDF membranes. After blocking with 5% skim milk, the membrane was probed with a primary antibody against Osx (Santa Cruz Biotechnology), Itgb3 (Cell Signaling Technology, Beverly, MA, USA), Itgb1 (Cell Signaling Technology), p-vinculin (Abcam), GAPDH (Bioworld Technology, St Louis Park, MN, USA), and HSP90 (Enzo Life Sciences, Farmingdale, NY, USA). A horseradish peroxidase-conjugated goat anti-rabbit IgG (Enzo Life Sciences) secondary antibody was used for visualization.

Quantitative real-time PCR with reverse transcription analysis

Total RNA was extracted from dissected bone tissue using Trizol reagent (Invitrogen). RNA was precipitated with isopropanol and dissolved in diethylpyrocarbonate-treated distilled water. First-strand cDNA was generated using the random hexamer primer provided in the first-strand cDNA synthesis kit (Applied Biosystems, Foster City, CA, USA). Specific primers for each gene (Table S1) were designed using qPrimerDepot (http://mouseprimerdepot.nci.nih.gov). Each real-time RT-PCR reaction contained 1 ng of reverse-transcribed total RNA, 2 nM of forward and reverse primers, and PCR master mix and was brought to a final volume of 10 μl. RT-PCR was performed in 384-well plates using the QuantStudio™ 6 Flex Real-Time PCR System instrument (Life Technologies, Foster City, CA, USA).

Transient transfections and reporter gene assays

Transient transfections were performed using Lipo-fectamine 2000 (Invitrogen) according to the manufacturer’s instructions. Briefly, 80% confluent MC3T3-E1 or HEK293 cells were transfected with empty vector (Empty) or Flag-Osx. For the reporter gene assay, we cotransfected either an empty vector or a Flag-Osx overexpression vector with the Renilla luciferase vector phRL-TK and the ITGB3-Prom-Luc reporter construct (SwitchGear Genomics, Menlo Park, CA, USA). After 48 h, cells were harvested in reporter lysis buffer (Promega, Madison, WI, USA). Luciferase activity was determined in whole cell lysates by a luciferase assay kit (Promega, Madison, WI, USA). Luciferase activity was determined in whole cell lysates by a luciferase assay kit (Promega, Madison, WI, USA). Luciferase activity was determined in whole cell lysates by a luciferase assay kit (Promega, Madison, WI, USA). Luciferase activity was determined in whole cell lysates by a luciferase assay kit (Promega, Madison, WI, USA).

ChIP assay

Chromatin immunoprecipitation (ChIP) assays were performed in MC3T3-E1 cells using the SimpleChIP® Enzymatic Chromatin IP kit according to the manufacturer’s instructions. The resulting extract was sonicated and used for immunoprecipitation (IP) with H3, IgG, or Osx rabbit antibodies. ChIP DNA samples were subjected to PCR using primers targeting the Itgb3 promoter region (sense, 5′-CTGCCTCCTTAGGCTTGAAT; antisense, 5′-TAAAATAGGGCAGCGATG) and the RPL30 gene (Cell Signaling Technology).

Statistical analysis

One-way analysis of variance was used for multiple comparisons, and Student’s t test was used to determine significant differences between two sets of data. Differences were considered significant at P < 0.05. All data were analyzed using SPSS statistical software (version 16.0; SPSS, Chicago, IL, USA).

Results

Corticalization is initiated after birth in mouse long bones

During endochondral bone development, the bone collar, which is formed by intramembranous bone, and trabecular bone, which is formed by endochondral bone, are divided into different compartments. To observe the beginning of corticalization, we first subjected sequential femoral bone samples to Safranin O staining. Safranin O is a red cationic dye that binds to the anionic groups of the glycosaminoglycans (GAGs) that comprise cartilage. At E16.5, the bone collar is completely separated from cartilage-based bone material. In the initial stage after birth, trabecular bone did not coalesce with cortical bone (Fig. 1). However, coalescence of trabecular bone was observed at postnatal day 7 (P7) (Fig. 2a), and trabecular bone had obviously coalesced into the metaphyseal cortex by P10 (Fig. 1).

Disruption of Osx in osteoblasts leads to delayed corticalization, abnormal osteoblast distribution, and decreased limb length

A previous study showed that osteoblast-specific Osx-knockout mice displayed both a reduction in femoral bone length and abnormal corticalization11. To explore the mechanisms responsible for long bone corticalization, we investigated the beginning of corticalization in the femora of osteoblast-specific Osx-knockout (Col-OMT) mice. First, we analyzed Cre activity using X-gal staining in Col1a1-Cre;R26R mice. In the distal femur of reporter mice, most osteoblasts around the trabecular bone were stained at P7, while the outer periosteum and hypertrophic chondrocytes were negative for LacZ at P7 (Figure S2). We also performed immunofluorescence to confirm the expression of Osterix in growing femora. In control mice, Osterix was localized in osteoblast nuclei in trabecular bone lesions at P7. However, the expression of Osterix was completely absent in Col-OMT mice.
Fig. 2 (See legend on next page.)
In P7 control mice, but not in P7 Col-OMT mice, the trabecular bone coalesced with the metaphyseal cortex. Further, trabecular bone volume was increased in P7 Col-OMT mice, though there was no difference in growth plate area (Fig. 2a). Because corticalization is associated with increased osteoblast surface area due to an increased number of osteoblasts in the peripheral spongiosa, we analyzed osteoblast conditions in the peripheral spongiosa of control and Col-OMT mice. At P10, osteoblast clusters were observed in the peripheral spongiosa of control mice but were not observed in Col-OMT mice. At P5 and P6, the number of osteoblasts in peripheral zones tended to be similar in each zone (zones I and IV). Central zones: II and III. Values are presented as the mean ± SEM. *p < 0.05, zone I versus zone II or zone III versus zone IV. **p < 0.01 versus control. Scale bar, 50 µm

Integrin β3 signaling is regulated by osterix in osteoblasts

Because osteoblasts with deleted osterix could not migrate to peripheral areas and could not cluster in peripheral zones, we evaluated integrin signaling in Col-OMT mice. Integrins bind to the ECM cytoskeletal assembly and associate with signaling proteins to form focal adhesions (FAs). Integrin-mediated FAs composed of talin, vinculin, filamin, kindlin, tensin, and paxillin mediate cell adhesion, migration, and signaling. In particular, vinculin plays a major role in stabilizing and strengthening FAs. The phosphorylation of vinculin is required for its activation and talin-binding capacity. Thus, we analyzed p-vinculin in osteoblasts in the peripheral spongiosa at P6. Immunohistochemical staining revealed that p-vinculin was expressed in osteoblasts, which were located in the peripheral spongiosa zone of control mice. In contrast, reduced levels of p-vinculin expression were observed in osteoblasts in the same lesion in Col-OMT mice (Fig. 3a, b). To identify the cause of reduced integrin signaling in Col-OMT mice, we analyzed the mRNA levels of integrins related to osteoblast differentiation and bone formation in P6 bone samples. Itgb3 mRNA expression levels were significantly decreased in Col-OMT mice, while no significant differences were observed in the levels of other genes (Fig. 3c). Furthermore, the protein levels of integrin β3 and p-vinculin were decreased in Col-OMT mice (Fig. 3d). These data suggest that Osx regulates integrin β3 signaling at the beginning of corticalization. To confirm if Osx could positively regulate Itgb3 expression, MC3T3-E1 cells (osteoblast precursor cells from mouse calvaria) were transfected with an empty vector or an Osx overexpression construct. Osx overexpression was confirmed by western blotting. Itgb3 protein levels were increased in Osx-overexpressing MC3T3-E1 cells (Fig. 3e). To investigate whether Osx transcriptionally regulates the Itgb3 promoter, we performed cotransfection experiments in HEK-293 cells (which do not express endogenous Osx) by the use of an Osx expression plasmid and a luciferase reporter driven by the 11 kb native Itgb3 promoter. Increasing amounts of Osx plasmid correlated with increased activity of the Itgb3 promoter reporter (Fig. 3f), suggesting that Osx is a transcriptional activator of the Itgb3 gene. We next performed ChIP assays to...
Fig. 3 Direct regulation of integrin \( \beta 3 \) by osterix.  

**a** Peripheral spongiosa lesions in the distal femora of P6 mice were immunostained with an anti-p-vinculin antibody.  

**b** Quantification of p-vinculin-positive cells per unit area. Values are presented as the mean ± SEM (\( n = 5 \)). **p < 0.01 versus control.**  

**c, d** Total RNA and protein was isolated from the bones of each littermate male mouse at P6. mRNA and protein levels of the indicated genes were determined by real-time RT-PCR and western blotting, respectively. Values are presented as the mean ± SEM (\( n = 5 \)). *p < 0.05 versus controls.**  

**e** MC3T3-E1 cells were transfected with either an empty vector or an Osx overexpression vector, and protein levels were analyzed by western blotting.  

**f** HEK293 cells were transiently transfected with an \( \text{Itgb3} \) luciferase reporter with or without increasing amounts of Osx, as indicated. Values are presented as the mean ± SEM (\( n = 6 \)). *p < 0.05, **p < 0.01 versus control (no Osx).**  

**g** ChIP assays were carried out in MC3T3-E1 cells. An anti-Osx antibody was used for ChIP analysis, while H3 and IgG antibodies were used as positive and negative controls, respectively. Immunoprecipitated genomic DNA was analyzed using PCR to assess protein binding to the \( \text{Itgb3} \) promoter or a control region (RPL30). Scale bar, 25 µm.
determine whether the Osx protein binds to the Itgb3 promoter endogenously. Cross-linked and sonicated chromatin from MC3T3-E1 cells was immunoprecipitated with antibodies specific to Osx, H3 (positive control), or immunoglobulin G (negative control). The binding of immunoprecipitated proteins to the Itgb3 promoter DNA or the control RPL30 region was determined by PCR. ChIP demonstrated that Osx specifically binds to the endogenous Itgb3 promoter. By contrast, Osx did not bind to the endogenous control region within MC3T3-E1 chromatin (Fig. 3g). These results indicate that the Itgb3 gene is a direct transcriptional target of Osx.

Integrin β3 regulates corticalization

To investigate whether integrin β3, which is modulated by Osx, affects corticalization, we analyzed femora from Itgb3-null mice. Interestingly, in the femora of Itgb3-null mice, the inner and the outer cortical bone was separated, and the trabecular bone volume of the central spongiosa was increased at P7 with no observable growth plate differences (Fig. 4a, Fig. S4). Furthermore, these bone phenotypes were also observed in tibia. However, the detachment of bone from cortical bone vanished gradually with age (Fig. 4c). To determine whether the bone separation was a result of a reduction in bone resorption in the bone collar (intramembranous ossification) or detachment of the trabecular bone (endochondral ossification), we performed Safranin O staining. Separated bone was stained red (Fig. 4a). This showed that bone formed by endochondral ossification did not coalesce with the bone collar and was separated in Itgb3-null mice.

As osteoblasts did not migrate to the peripheral spongiosa in Col-OMT mice, the number of osteoblasts in peripheral zones was not changed compared to the central zone in Itgb3-null mice at P7 (Fig. 4b). At 1 month of age, femoral length decreased significantly in Itgb3-null mice compared to Itgb3+/− mice (Fig. 4c). In the μCT analysis of trabecular bone at 1 month of age, the trabecular number and the bone volume to total tissue volume ratio (BV/TV) were higher in Itgb3-null mice than in Itgb3+/− mice. Trabecular separation (Tb.Sp) was significantly lower in Itgb3-null mice. On the other hand, BV/TV and cortical thickness were significantly decreased in the metaphyseal cortical bone of the distal femur in Itgb3-null mice (Fig. 4c). As in Col-OMT mice, Itgb3-null mice exhibited an abnormal distribution of osteoblasts at the beginning of corticalization and decreased limb length and delayed corticalization at 1 month. However, unlike in Col-OMT mice, cortical BMD was not changed in Itgb3-null mice compared to Itgb3+/− mice. These results suggest that Osx regulates bone mineralization and corticalization, whereas integrin β3 controls corticalization without affecting bone mineralization.

Discussion

This study showed that corticalization affecting longitudinal bone growth takes place at postnatal day 7 with the migration of osteoblasts to the peripheral spongiosa. Additionally, the regulation of integrin β3 by osterix in osteoblasts plays an important role in corticalization (Fig. 5).

Longitudinal bone growth is largely determined by the rate of growth plate chondrogenesis. However, taking into consideration that corticalization (the process by which the bone formed in the growth plate is fused to the metaphyseal cortical bone) is the last step in the longitudinal bone growth process and that corticalization is modulated by Osx in osteoblasts (osteoblast differentiation), osteoblasts play a role in limb growth (Fig. 5). A large meta-analysis genome-wide association study showed that genes in loci associated with the regulation of normal variation in adult human height tend to be highly expressed in osteoblast-related tissues as well as in chondrocytes. Furthermore, many studies in knockout mouse models have shown that osteoblasts affect growth retardation. Fibrillins, the major structural components of extracellular microfibrils, were observed in the endosteal surface of trabecular bone and are secreted from osteoblasts. Fibrillin-1 and fibrillin-2 knockout mice exhibited differences in bone length compared to controls without growth plate anomalies. Bone sialoprotein (BSP), a SIBLING protein (small, integrin-binding ligand N-linked glycoprotein), is a component of mineralized tissues such as bone. Femoral length was decreased in BSP knockout mice compared to control mice with no growth plate differences or nutritional defects. Similar to Itgb3-null mice, BSP knockout mice displayed a high trabecular bone mass. In particular, osteocyte-specific knock-in mice expressing a dominant-active β-catenin in osteocytes failed to show longitudinal limb growth despite a marked increase in trabecular bone. These mice also did not display differences in growth plate thickness or serum phosphate levels. These results support the finding that osteoblasts and osteocytes regulate longitudinal limb growth without affecting the growth plate.

In this study, osteoblasts in the primary spongiosa migrated to the peripheral spongiosa and formed clusters (cell–cell) for coalescence in cortical bone (cell–matrix) by postnatal (mechanical stimulation) day 10 (Figs. 1 and 2). These changes indicate a role for integrin signaling. Therefore, we analyzed integrin signaling in Col-OMT mice, which exhibited delayed corticalization.

We studied integrin signaling in Col-OMT mice for an additional reason. SIBLINGs are a family of five integrin-binding proteins (osteopontin (OPN), bone sialoprotein (BSP), dentin matrix protein 1 (DMP1), dentin sialophosphoprotein (DSPP), and matrix extracellular phosphoglycoprotein (MEPE)) that modulate cell adhesion by interacting with integrins. Notably, DMP1 binds to the
Fig. 4 Corticalization is controlled by integrin β3. a Picrosirius red staining and Safranin O staining of femora of 7-day-old male littermates. Yellow arrowheads indicate the cartilaginous portion of the trabecular bone. b The number of osteoblasts according to zone at P5, P6, and P7. Values are presented as the mean ± SEM (n = 5). *p < 0.05, zone I versus zone II or zone III versus zone IV. c Morphometric indices in trabecular bone or metaphyseal cortex of the distal femur at 1 month of age. Femur length was measured from the proximal end of the femur to the distal end using micro-CT. Values are presented as the mean ± SEM (n = 5). *p < 0.05, **p < 0.01 versus controls. Scale bar, 50 µm
Because DMP1 is significantly reduced in Osx-knockout mice21,41, we investigated whether disruption of Osx affected integrin signaling. In control mice, DMP1 expression was increased in trabecular bone at P6 compared to P4. In contrast, DMP1 expression did not change in Col-OMT mice (data not shown). The expression of an Osx-modulated ligand (DMP1) also partially affected the induction of integrin signaling.

In the present study, we showed that integrin β3 signaling is involved in the migration of osteoblasts to the peripheral zone at the beginning of corticalization. Considering the mechanism by which integrin β3 signaling leads to the migration of osteoblasts to the peripheral zone, we speculated that mechanical forces or osteoblast differentiation stimuli would be increased in the peripheral zone compared to the central zone. Furthermore, osteoblasts could be recruited and forced to cluster in the peripheral zone due to mechanical loading-induced secretion of cytokines or growth factors from osteocytes in cortical bone42,43. Second, ECMs (SIBLINGs) in the metaphyseal cortex, which is formed through intramembranous ossification in the bone collar, are thought to trigger the migration of osteoblasts to the peripheral zone through integrin β3.

The increase in trabecular bone volume could be affected by osteoclast function in Itgb3-null mice. β3−/− osteoclasts cultured from bone marrow macrophages were defective in their resorption capacity and showed abnormal cytoskeletal morphology36. However, abnormal distribution of osteoblasts and the detachment of trabecular bone was not modulated by osteoclast function. Furthermore, because osteoblasts in the primary spongiosa expressed integrin β3 signaling at P6, integrin β3 signaling might conceivably play a role in osteoblast functioning (Fig. 3). Early studies indicated that several integrins interact with ECM components, control osteoblast functions and are involved in bone formation26,44,45. In addition, some clinical studies have suggested that delayed corticalization can bring about an increase in trabecular bone volume. Cortical thickness and cortical volumetric BMD were significantly lower in the distal radius and distal tibia of males compared to females, despite increased trabecular BV/TV and trabecular thickness during puberty4. Reports have suggested that this transitory deficit in cortical bone formation results from a delay in trabecular corticalization. Based on these reports, it has been suggested that the increase in trabecular bone volume is partially influenced by integrin β3 signaling in osteoblasts due to delayed corticalization.

Glanzmann thrombasthenia (GT), a rare hereditary autosomal-recessive bleeding disorder, is associated with integrin β327. Although GT bone phenotypes are reported less frequently than the blood disorders associated with this disease (mainly focused on platelet function), several reports showed that patients with GT exhibited short stature (below the tenth percentile)46,47. In particular, patients with αIIbβ3 integrin activation failure due to a point mutation in KINDLIN3 displayed heights within only the third percentile. Further, osteopetrosis was observed in bone proximal to the growth plate at 7 months of age48. Malinin et al. suggested that integrin activation may be involved in osteogenesis as well as in bone resorption, as subjects’ mesenchymal stem cells produced substantially higher amounts of bone. Furthermore, allogenic bone marrow transplantation (BMT) resolved the macroscopic bone density defects, and bone production by the mesenchymal stem cells also reverted to control levels after BMT. Our results and those of previous reports suggest that integrin β3 plays a role in longitudinal bone growth.

In conclusion, osteoblasts contribute to longitudinal limb growth through corticalization after birth. Corticalization is mediated by integrin β3 signaling in differentiated osteoblasts. Because differences in osteoblast function can affect normal variations in height, the control of osteoblast function during corticalization may help in the treatment of short stature.

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