Tracing the skeletal progenitor transition during postnatal bone formation

Graphical Abstract

Highlights
- Bone formation before and after adolescence is controlled by distinct progenitors
- Chondrocytes and \textit{Lepr}^+ BMSCs mediate bone lengthening and thickening, respectively
- The adult skeletal progenitors derive mainly from developmental skeletal progenitors
- Running enhances osteogenesis by developmental but not adult skeletal progenitors

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In brief
Shu et al. discovered that bone formation before and after adolescence is dominated sequentially by chondrocytes and \textit{Lepr}^+ BMSCs. They regulate bone lengthening and thickening, respectively. This transition explains, from a stem cell perspective, how mammalian limb bones transition from rapid longitudinal growth to slower appositional remodeling after adolescence.
Tracing the skeletal progenitor transition during postnatal bone formation

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SUMMARY

Multiple distinct types of skeletal progenitors have been shown to contribute to endochondral bone development and maintenance. However, the division of labor and hierarchical relationship between different progenitor populations remain undetermined. Here we developed dual-recombinase fate-mapping systems to capture the skeletal progenitor transition during postnatal bone formation. We showed that postnatal osteoblasts arose primarily from chondrocytes before adolescence and from Lepr+ bone marrow stromal cells (BMSCs) after adolescence. This transition occurred in the diaphysis during adolescence and progressively spread to the metaphysis. The osteoblast-forming Lepr+ BMSCs derived primarily from fetal Col2+ cells. Conditional deletion of Runx2 from perinatal chondrocytes and adult Lepr+ BMSCs impaired bone lengthening and thickening, respectively. Forced running increased osteoblast formation by perinatal chondrocytes but not by adult Lepr+ BMSCs. Thus, the short-term developmental skeletal progenitors generated the long-term adult skeletal progenitors. They sequentially control the growth and maintenance of endochondral bones.

INTRODUCTION

Development, maintenance, and regeneration of bones require continuous osteoblast formation by skeletal progenitors (Ono et al., 2019). Skeletal progenitors have been identified in multiple cell types of the endochondral bones in the mouse, including various subsets of bone marrow stromal cells (Méndez-Ferrer et al., 2010; Mizoguchi et al., 2014; Park et al., 2012; Pineault et al., 2019; Rux et al., 2016; Seike et al., 2018; Worthley et al., 2015; Zhou et al., 2014a), chondrocytes (Mizuhashi et al., 2018; Ono et al., 2014; Yang et al., 2014b; Zhou et al., 2014b), and periosteal stromal cells (Debnath et al., 2018; Duchamp de Lageneste et al., 2018; Ortniau et al., 2019).

Single-cell RNA sequencing has defined bone marrow stromal cells (BMSCs) as Leptin receptor (LepR)-expressing BMSCs that abundantly expressed hematopoietic niche factors (Baryawno et al., 2019; Tikhonova et al., 2019). LepR+ BMSCs contain virtually all colony-forming unit-fibroblasts (CFU-Fs) in the bone marrow (Zhou et al., 2014a). Fate mapping with Lepr-Cre labeled most osteoblasts and adipocytes in adult bone marrow (Mizoguchi et al., 2014; Zhou et al., 2014a). Thus, LepR+ BMSCs may represent the main skeletal progenitor population in adult bone marrow, but the lack of a Lepr-CreER allele from which fate mapping could be initiated during adulthood raises the possibility that early postnatal progeny of LepR+ cells may be the key source of osteoblasts during adulthood. It is also uncertain whether LepR+ BMSCs are self-renewable or whether they are replenished continuously by LepR+ progenitors during adulthood.

In the endochondral ossification model, cartilage does not form bone but, rather, serves as a template that is replaced by new bone. Recent studies have challenged this dogma by showing that early postnatal cells marked by Col2-CreER, Sox9-CreER, Acan-CreER, or Col10-Cre contribute to endochondral bones that persist throughout adulthood (Ono et al., 2014; Yang et al., 2014a, 2014b; Zhou et al., 2014b). The relative contributions of chondrocytes and BMSCs to bone formation at different stages have not been determined.

BMSCs have been thought to originate from fetal Osterix (Osx)+ perichondral precursors that invade the cartilage template along with the blood vessels (Maes et al., 2010). Recent evidence has suggested that chondrocytes give rise to a significant portion of BMSCs that persist in adulthood (Ono et al., 2014). This raises the possibility that the BMSCs maintaining adult bones arise from chondrocytes. However, this idea is difficult to address using single-recombinase fate-mapping systems, such as Col2-Cre or Acan-CreER, because it is hard...
Figure 1. Chondrocytes form most osteoblasts in early postnatal bones, but their contribution declines by 2 months of age
(A and B) Confocal imaging of femur sections from AcancreER; R26tdTomato; Col1a1-GFP mice that had been tamoxifen treated at P1–P3. Mice were analyzed at 1 day (A) and 1 month after the treatment (B). Arrows indicate Tomato⁺ osteocytes. n = 3 mice per condition from 3 independent experiments.

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to distinguish whether the undifferentiated BMSCs that arise from these cell populations form osteoblasts independent of the osteoblasts formed by the chondrocytes themselves. In this study, we generated a series of single-recombinase and dual-recombinase fate-mapping systems to trace chondrocytes and Lepr+ BMSCs, comparing their contributions to osteogenesis during bone growth and maintenance and elaborating their relationship.

**RESULTS**

**Perinatal chondrocytes form most of the new osteoblasts by 1 month of age**

To determine the contribution of chondrocytes to postnatal bone formation, we crossed AcancreER knockin mice (Henry et al., 2009) with Rosa26ΔAG-loxP-STOP-loxP-tdTomato (R26tdTomato) knockin mice (Madisen et al., 2010) and Col1a1^{2.3-GFP} (Col1a1-GFP) transgenic mice (Kajacic et al., 2002) to generate AcancreER; R26tdTomato; Col1a1-GFP mice. Acan (Aggrecan) promoter-driven Cre expression did not show any leakiness in the bone marrow without tamoxifen treatment (Figure S1A).

We administered tamoxifen to these mice from post-natal day 1 (P1)–P3. At P2 and P3, Tomato expression was highly restricted to the growth plate (Figures S1B and S1C). At P4, Tomato+ non-chondrocytes appeared underneath the growth plate (Figures 1A and S1D); however, they were negative for aggrecan expression (Figure S1E), suggesting that they are not Acan+ cells but progeny of Acan+ cells. Sox9+ columnar chondrocytes and Col10+ hypertrophic chondrocytes were marked by Tomato (Figures S1F and S1G). Col1a1-GFP+ osteoblasts were uniformly Tomato+ at this stage (Figures 1A and 1F). 1 month after tamoxifen treatment, Tomato+ stromal cells were distributed throughout the bone marrow (Figures 1B and S1N). Osteoblasts, as revealed by Col1a1-GFP (Figure 1B) or anti-Runx2 staining (Figure S1H), were co-labeled extensively by Tomato at the metaphysis and diaphysis. A significant portion of osteocytes, as revealed by anti-sclerostin staining, were also marked by Tomato in 1-month-old mice (Figures 1B, arrows, and S1I). Flow cytometry analysis of enzymatically dissociated femora showed that 55% ± 6.7% of all Col1a1-GFP+ osteoblasts expressed Tomato (Figure 1E), suggesting that perinatal chondrocytes are the major contributors to osteoblast production by 1 month of age.

### The contribution of chondrocytes to osteoblasts decreases by adulthood

2 months after tamoxifen treatment, we still observed many Tomato+ osteoblasts in the metaphysis of AcancreER; R26tdTomato; Col1a1-GFP mice pulsed at P1–P3 (Figure S1M), but there were considerably fewer in the diaphysis (compare Figures 1Bii and S1Mi). By flow cytometry, only 24% ± 6.0% of all Col1a1-GFP+ osteoblasts expressed Tomato in these mice, significantly less than that at 1 month of age (Figure 1F). This value decreased further to 18% ± 4.4% 6 months after tamoxifen treatment (Figure 1F). These data suggested that the contribution of perinatal chondrocytes to osteoblast formation decreased progressively with age.

To determine whether adult chondrocytes form osteoblasts, we administered tamoxifen to 2-month-old AcancreER; R26tdTomato; Col1a1-GFP mice. 1 day after tamoxifen treatment, we observed efficient and specific labeling of chondrocytes in the growth plate (Figures 1C and S1J–S1L). Unlike perinatal chondrocytes, 2-month-old chondrocytes generated few Tomato+ BMSCs (Figures 1D and S1N). Tomato+ osteoblasts were observed but restricted to metaphyseal regions adjacent to the growth plate (Figure 1D). By flow cytometry, only 3.9% ± 2.0% of all Col1a1-GFP+ osteoblasts expressed Tomato in these mice (Figure 1G), 6 months after tamoxifen treatment, we observed even fewer Tomato+ Col1a1-GFP+ osteoblasts (Figure S1O). Thus, adult chondrocytes generate only a few osteoblasts underneath the growth plate.

We determined the contribution of perinatal chondrocytes to LepR+ BMSCs. By flow cytometry, only 4.6% ± 0.54% of all LepR+ BMSCs expressed Tomato in 1-month-old AcancreER; R26tdTomato; Col1a1-GFP mice pulsed at P1–P3 (Figure 1H). This value increased gradually to 20% ± 3.3% by 2 months of age (Figure 1I). Approximately 3% and 8% of all CFU-Fs in the bone marrow expressed Tomato in 1- and 2-month-old mice, respectively (Figure 1K). 0.022% ± 0.016% of all LepR+ BMSCs expressed Tomato in 4-month-old AcancreER; R26tdTomato; Col1a1-GFP mice pulsed at 2 months of age (Figure 1J). These data suggest that perinatal chondrocytes generate a small subset of adult LepR+ BMSCs.

In vertebrae, which are also formed by endochondral ossification, we found that chondrocytes in AcancreER; R26tdTomato; Col1a1-GFP mice formed most osteoblasts before adolescence but few afterward (Figures 1L, S2A–S2C, S2F, and S2G).
Figure 2. Lepr<sup>+</sup> BMSCs form most osteoblasts during adulthood but few during development

(A and B) Confocal imaging of femur sections from Lepr<sup>-creER</sup>; R26<sup>Tomato</sup>; Col1a1-GFP mice that had been tamoxifen treated at P1–P3. Mice were analyzed 1 day (A) or 2 months (B) after the treatment. n = 3 mice per condition from 3 independent experiments.

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consistent with our observations in the femur. In calvaria, whose formation involves intramembranous ossification, we detected few Tomato+ stromal cells or osteoblasts in AcancreER; R26tdTomato; Col1a1-GFP mice, regardless of whether tamoxifen was administered at P1–P3 or 2 months of age (Figures 1M, S2D, S2E, S2H, and S2I), suggesting that postnatal chondrocytes are not the origin of osteoblasts in calvaria.

**Perinatal Lepr**+ BMSCs rarely form osteoblasts

To determine the contribution of Lepr+ BMSCs to postnatal bone formation, we generated Lepr-creER BAC (bacterial artificial chromosome)-transgenic mice. These mice were crossed with R26tdTomato and Col1a1-GFP mice to generate Lepr-creER; R26tdTomato; Col1a1-GFP. Lepr promoter-driven Cre expression did not show any leakiness in the bone marrow without tamoxifen treatment (Figure S3A). At 1 day after tamoxifen treatment at P1–P3, a few Tomato+ stromal cells appeared in the bone marrow (Figures 2A and S3B), 2 months after tamoxifen treatment, more Tomato+ stromal cells were detected in the bone marrow (Figure 2B). By flow cytometry, they accounted for 28% ± 8.4% of all Lepr+ BMSCs in 2-month-old bone marrow (Figure 2E). This number decreased to 13% ± 4.2% 4 months after tamoxifen treatment (Figure S3C). On femur sections 2 months after tamoxifen treatment, most of the Col1a1-GFP+ osteoblasts were Tomato negative (Figure 2B). Consistent with this, only 2.2% ± 1.0% of all Col1a1-GFP+ osteoblasts were Tomato+ by flow cytometry (Figure 2H). Thus, perinatal Lepr+ cells make a limited and declining contribution to BMSCs and contribute minimally to bone formation in postnatal bone marrow.

**Adult Lepr**+ BMSCs become the main source of new osteoblasts

We examined the Tomato expression pattern in Lepr-creER; R26tdTomato; Col1a1-GFP mice 1 day after tamoxifen treatment at 2 months of age. Tomato expression was detected in stromal cells throughout the bone marrow (Figure S3F), accounting for 70% ± 5.6% of all Lepr+ BMSCs (Figure 2G). Tomato expression was not detected in Col1a1-GFP+ osteoblasts (Figures 2C and S3D; Video S1), VE-cadherin+ endothelial cells (Figure S3E), aggrecan+ chondrocytes (Figure S3G), and perilipin+ adipocytes (Figure S3H). Thus, Lepr-CreER efficiently and specifically marks Lepr+ BMSCs in 2-month-old mice.

After 4 months of tracing, 91% ± 3.4% of all Lepr+ BMSCs were Tomato+ in the femora of Lepr-creER; R26tdTomato; Col1a1-GFP mice (Figures 2F and 2G). Virtually all Tomato+ stromal cells were Lepr+ (Figures S3I and S3L). Similarly, 85% ± 4.5% of all PDGFRa+ BMSCs were Tomato+ (Figure S3J), and virtually all Tomato+ stromal cells expressed PDGFRa (Figure S3K). Among all CFU-F colonies formed by whole bone marrow cells, 91% ± 6.7% of them expressed Tomato (Figure 2K). Consistent with this, fluorescence-activated cell sorting (FACS) of CD45-Ter119+ CD31 Lepr+ cells recovered ~95% and ~85% of all CFU-Cs from the bone marrow and femur shaft, respectively. The percentage of Lepr+ BMSCs that were labeled by Tomato did not decrease even after tracing for 12 months (Figure 2G). Similarly, when tamoxifen was administered to these mice at 6 months of age, we found that Lepr+ BMSCs essentially overlapped with the Tomato+ BMSCs at 8 months of age (Figures S3O and S3Q). Thus, unlike perinatal Lepr+ cells, adult Lepr+ BMSCs sustain themselves throughout adulthood.

In femur sections from 6-month-old Lepr-creER; R26tdTomato; Col1a1-GFP mice that were treated with tamoxifen at 2 months of age, most Col1a1-GFP+ osteoblasts in the femur, especially those in the diaphysis, were co-labeled by Tomato (Figure 2D). Tomato expression was also detected robustly in osteocytes in the bone (Figures 2D, arrows, and S3N). By flow cytometry, Tomato+ cells accounted for 55% ± 9.0% of all Col1a1-GFP+ osteoblasts in the femur (Figure 2B). This value increased to 69% ± 6.3% 12 months after tamoxifen treatment (Figure 2J). When these mice were treated with tamoxifen at 6 months of age, we detected that 36% ± 5.1% and 47% ± 8.0% of osteoblasts expressed Tomato at 8 and 10 months of age, respectively (Figure S3P). These data suggested that most of the newly formed osteoblasts during adulthood arise from adult Lepr+ BMSCs.

In vertebrae, we found that adult but not perinatal Lepr+ BMSCs from Lepr-creER; R26tdTomato; Col1a1-GFP mice made a major contribution to bone formation (Figures 2L, S4A–S4C, S4F, and S4G), consistent with our observations in...
Figure 3. Adult Lepr+ BMSCs, but not perinatal chondrocytes, contribute to fracture healing

(A) Safranin O staining of femur sections from normal mice or mice 2 or 8 weeks after fracture. n = 3 mice per condition from 3 independent experiments.

(B–D) Confocal imaging of femur sections from AcancreER; R26tdTomato; Col1a1-GFP mice 2 weeks (B and C) or 8 weeks (D) after fracture. Mice were tamoxifen-treated after fracture (B) or at P1–P3 (C and D). n = 3 mice per condition from 3 independent experiments.

(E and F) Confocal imaging of femur sections from Lepr-creER; R26tdTomato; Col1a1-GFP mice 2 weeks (E) or 8 weeks (F) after fracture. Mice were tamoxifen-treated 2 months of age and fractured at 4 months of age. n = 3 mice per condition from 3 independent experiments.

See also Figure S5.
femora. In calvaria, although we detected robust Tomato expression in adult BMSCs, they formed a minor portion of osteoblasts (Figures 2M, S4D, S4E, S4H, and S4I), suggesting that Lepr BMSCs are not the origin of osteoblasts in postnatal calvaria.

Most of the perilipin+ adipocytes expressed Tomato in the bone marrow from 6- and 14-month-old Lepr-creER; R26tdTomato mice pulsed at 2 months of age (Figures S5A and S5C). At 2 weeks after lethal irradiation and bone marrow transplantation, adipocytes were increased markedly in the bone marrow of Lepr-creER; R26tdTomato mice compared with non-irradiated mice (Figure S5B). Over 90% of all perilipin+ bone marrow adipocytes from these mice were Tomato+ (Figures S5B and S5D). Thus, Lepr BMSCs generate most of the adipocytes formed in adult bone marrow.

**Adult Lepr BMSCs, but not perinatal chondrocytes, contribute to fracture healing**

We investigated the contribution of perinatal chondrocytes and adult Lepr BMSCs to bone regeneration. Safranin O staining revealed cartilage formation at the callus of the fractured femur (Figure 3A). Consistent with literature (Zhou et al., 2014b), administering tamoxifen to AcancreER; R26tdTomato; Col1a1-GFP mice during the fracture healing process labeled cartilage and osteoblasts in the callus (Figure 3B). In contrast, few Col1a1-GFP osteoblasts were Tomato+ at the callus from AcancreER; R26tdTomato; Col1a1-GFP mice that had been treated with tamoxifen at P1–P3 (Figure 3C), 8 weeks after bone fracture, when the femur had largely healed (Figure 3A), almost no regenerated osteoblasts expressed Tomato in these mice (Figure 3D). Thus, perinatal chondrocytes do not generate progenitors for fracture healing.

We then performed fractures on 4-month-old Lepr-creER; R26tdTomato; Col1a1-GFP mice that had been tamoxifen treated at 2 months of age. 2 weeks after fracture, Tomato expression was detected in a subset of Col1a1-GFP+ osteoblasts (Figure 3E) and aggrecan+ chondrocytes (Figure S5E). 8 weeks after bone fracture, some regenerated osteoblasts expressed Tomato in Lepr-creER; R26tdTomato; Col1a1-GFP mice (Figure 3F). A few perivascular stromal cells were detected in the periosteum of the metaphysis from Lepr-creER; R26tdTomato; Col1a1-GFP mice pulsed at 2 months of age, but they were not detected at the diaphysis where fracture occurred (Figures S5F and S5G). Adult Lepr BMSCs contribute to osteoblast regeneration upon injury.

**Osteoblast-forming Lepr+ cells derive from Col2+ cells**

Our data above suggested chondrocytes and Lepr+ BMSCs as developmental and adult skeletal progenitors, respectively. To investigate the potential relationship between them, we used Col2-Cre (Hao et al., 2002) and Lepr-DreER, two different recombinases, to simultaneously trace chondrocytes and Lepr+ BMSCs. Dre recombination excises DNA regions flanked by rox recombinase sites (Anastassiadis et al., 2009), activating Tomato expression when crossed with Rosa26AG-AG-lox-STOP-lox-Tomato (R26RSR-ttdTomato) mice. Lepr-DreER did not show leaky recombination (Figure S6A). It recombined efficiently in LeprBMSCs but not in Col21-GFP osteoblasts, perilipin+ adipocytes, or aggrecan+ chondrocytes (Figures S6B–S6E), 3 weeks after tamoxifen treatment, robust labeling of Col1a1-GFP+ osteoblasts by Tomato was detected at diaphysal bones (Figure S6F). In Col2-cre; R26tdTomato; Col1a1-GFP mice, Col2-Cre marked cells included chondrocytes, periosteal cells, BMSCs, and osteoblasts (Figures S7A and S7B), consistent with previous reports (Ono et al., 2014; Szabova et al., 2009).

In Col2-cre; LeprdreER; R26SL-ZsGreen; R26RSR-ttdTomato mice, Lepr+ derived cells and Col2- derived cells were labeled by Tomato and ZsGreen, respectively (Figure 4A). Osteoblasts were marked by anti-Osx staining (Figures 4B and S5H–S5K). Mice were treated with tamoxifen 4 and 7 weeks after birth and analyzed at 10 weeks. Confocal imaging and flow cytometry analysis revealed significant overlap of most Tomato+ (Lepr-derived) and ZsGreen+ (Col2-derived) BMSCs (Figures 4B and 4C). Moreover, Tomato (Lepr derived) and ZsGreen (Col2 derived) also displayed extensive overlap in Osx+ osteoblasts, especially at the diaphysis (Figure 4B). Similar results were obtained when Acan+ progenitors and Lepr BMSCs were traced simultaneously by Acan-Dre and Lepr-CreER, respectively (Figures S6G–S6I). Thus, Col2 progenitors, Lepr BMSCs, and their osteolineages are genetically related.

The simultaneous tracing experiments could not tell which population is upstream of the other. To address this issue, we developed a mutually exclusive tracing system using the R26AGAG-loxp-STOP-loxp-ZsGreen-AG-AG-lox-STOP-Tomato (R26RSR-ttdTomato) interlaced reporter (IR) (He et al., 2017). In Col2dre; Leprcre, IR mice, Lepr-Cre- and Col2-Dre-mediated recombination was mutually exclusive in the same cell and its progeny. Whichever appeared first would prevent recombination by the other. This allowed us to determine which one appears first (Figure 4D). As expected, we never detected ZsGreen+Tomato+ cells in Col2dre; Leprcre; IR mice (Figures 4G and 4J).
Figure 5. Chondrocytes and Lepr+ BMSCs generate spatially separated subsets of osteoblasts during adolescence

(A) Confocal imaging of femur sections from 2-month-old AcancreER; LeprdreER; R26LSL-ZsGreen; R26RSR-tdTomato mice that had been tamoxifen-treated at P1–P3. n = 3 mice from 3 independent experiments.
Flow cytometry analysis showed that the frequencies of Tomato+ cells in the bone marrow of 5-month-old Col2dre; IR and Col2dre; Leprcre; IR mice were comparable (Figures 4E and 4G). In contrast, the frequency of ZsGreen+ cells in the bone marrow of Col2dre; Leprcre; IR mice was only about 11% of that in Leprcre; IR mice (Figures 4F and 4G), suggesting that Col2-Dre-mediated recombination prevents Lepr-Cre-mediated recombination in most BMSCs. Consistent with this, femur sections of 5-month-old Col2dre; Leprcre; IR mice largely resembled those of Col2dre; IR mice, with few ZsGreen+ BMSCs (Figures 4H–4J). Most Osx+ osteoblasts were ZsGreen‘Tomato+’ (Figure 4J), suggesting that most Lepr-derived osteoblasts originate from Col2+ cells. Thus, the Col2 lineage is upstream of Lepr-derived BMSCs and osteoblasts.

Perinatal chondrocytes and Lepr+ BMSCs form spatially separated subsets of osteoblasts during adolescence

To investigate the relationship between perinatal chondrocytes and Lepr+ BMSCs, we generated AcancreER; LeprdreER; R26LSL-ZsGreen; R26RSR-tdTomato mice. Femur sections from 2-month-old mice that had been tamoxifen treated at P1–P3 showed extensive labeling of BMSCs and bone-lining osteoblasts by ZsGreen but few by Tomato (Figure 5A), consistent with our finding that perinatal chondrocytes, but not Lepr+ BMSCs, make the major contribution to osteoblast formation during childhood.

We next treated AcancreER; LeprdreER; R26LSL-ZsGreen; R26RSR-tdTomato mice with tamoxifen 1 day, 4 weeks, and 7 weeks after birth. At 10 weeks of age, we observed a clear gradient formed by ZsGreen and Tomato fluorescence along the endostem (Figure 5B). Osx+ osteoblasts in the metaphysis were mostly ZsGreen‘Tomato−’ (Acan derived), whereas those in the diaphysis were mostly ZsGreen‘Tomato+’ (Lepr derived; Figures 5B and 5C), suggesting that perinatal chondrocytes and Lepr+ BMSCs generate spatially separated subsets of osteoblasts in 10-week-old mice.

By flow cytometry, only a small subset of Tomato+ (Lepr-derived) BMSCs were ZsGreen+ (Acan derived) in AcancreER; LeprdreER; R26LSL-ZsGreen; R26RSR-tdTomato mice (Figure 5D), although most of the ZsGreen+ (Acan derived) BMSCs were Tomato+ (Lepr derived; Figure 5E). Consistent with Figures 1H–1J, these data suggested that perinatal chondrocytes give rise to a small subset of Lepr+ BMSCs.

Runx2 is required by Lepr+ BMSCs for bone maintenance in adult mice

We used a non-inducible Lepr-Cre allele to delete Runx2 (DeFalco et al., 2001). Two-month-old Leprcre; Runx2fl/fl mice showed normal body size, normal femur length, and normal trabecular and cortical bone parameters (Figures 6M–6X), consistent with the limited contribution of Lepr+ cells to osteoblasts during development. Four-month-old Leprcre; Runx2fl/fl mice also had normal body size, normal femur length, and normal trabecular and cortical bone parameters but reduced cortical area and thickness (Figures 6Y–6AJ). Seven-month-old Leprcre; Runx2fl/fl mice had more severe defects in bone formation because the defects appeared in trabecular and cortical parameters (Figures 6J–6T). Reduced bone thickness was also observed in 4-month-old AcancreER; Runx2fl/fl mice that were treated with tamoxifen at 2 months of age (Figures 6S–6T). These data suggested that conditional deletion of Runx2 from Lepr+ BMSCs impairs bone thickening in adult mice.

Running differentially regulates osteoblast formation by perinatal and adult skeletal progenitors

Finally we investigated how perinatal and adult skeletal progenitors respond to physiological stimuli, such as running. One-month-old AcancreER; R26tdTomato; Col1a1-GFP mice that had been tamoxifen treated at P1–P3 were subjected to forced running for 2 h once every day at a speed of 14 m/min (Figure 7A). Two months after running, the Col1a1-GFP osteoblast pool contained significantly more Tomato+ cells than controls by flow cytometry (Figures 7B–7D). Consistent with this, we observed more Tomato+ osteoblasts in the diaphysis of AcancreER; R26tdTomato; Col1a1-GFP mice after running compared with controls (Figures 7E and 7F). These data suggest...

(B and C) Confocal imaging of femur sections from 10-week-old AcancreER; LeprdreER; R26LSL-ZsGreen; R26RSR-tdTomato mice that had been tamoxifen treated at P1–P3 and 4 and 7 weeks after birth. The white dashed line artificially separated the metaphysis from the diaphysis (B). Acan- and Lepr-derived cells were marked by ZsGreen and Tomato, respectively. Osteoblasts were marked by anti-Osx staining. The ratios of Tomato+ZsGreen+ osteoblast numbers were quantified (C). n = 3 mice from 3 independent experiments.

(D and E) Flow cytometry analysis of enzymatically dissociated bone marrow cells from 10-week-old AcancreER; LeprdreER; R26LSL-ZsGreen; R26RSR-tdTomato mice that had been tamoxifen treated at P1–P3, and 4 and 7 weeks after birth. n = 3 mice from 3 independent experiments.

(F) Schematic depicting the progenitor transition during postnatal bone growth. The contributions of perinatal chondrocytes and Lepr+ BMSCs to osteoblast formation are separated temporally and spatially.
Figure 6. Deletion of Runx2 from perinatal chondrocytes and from adult Lepr⁺ BMSCs impairs bone lengthening and thickening, respectively. Different sexes were examined in different experiments so that most data reflected male and female mice. Two-tailed Student’s t tests were used to assess the statistical significance of differences between sex-matched littermates (*p < 0.05, **p < 0.01, ***p < 0.001).
that running augments osteoblast formation by chondrocytes in juvenile mice.

In contrast, when 3-month-old Lepr-creER; R26rtomato; Col1a1-creFOP mice that had been tamoxifen treated at 2 months of age were subjected to running for 2 months (Figure 7G), we did not detect significant changes in the frequency of Tomato+ osteoblasts of these mice by flow cytometry (Figures 7H–7J) or by confocal imaging (Figures 7K and 7L). These data suggest that running does not affect osteoblast formation by adult Lepr+ BMSCs.

**DISCUSSION**

Our work has established that postnatal osteoblast production by chondrocytes and BMSCs is separated temporally and spatially. This skeletal progenitor transition occurs during adolescence, which explains, from a stem cell perspective, why mouse endochondral bones transition from rapid longitudinal growth to slower appositional remodeling after adolescence. By inducible lineage tracing, we provided direct evidence that adult Lepr+ BMSCs are self-renewable. Interestingly, a recent study identified a subpopulation of Lepr+ BMSCs that expressed osteolectin (Shen et al., 2021). The Lepr+osteolectin+ cells were short-lived osteogenic progenitors. Combined with our data, it revealed a hierarchy in adult skeletal stem cells: the self-renewable skeletal stem cells enriched in Lepr+osteolectin+ BMSCs give rise to the short-lived Lepr+osteolectin+ osteogenic progenitors. These cells actively form osteoblasts and are replaced continuously by new Lepr+ osteolectin+ cells during aging.

Col2-Cre-marked cells gave rise to ~70%, not all, of Lepr+ BMSCs (Figure 4C). One possibility is that Col2-Cre did not recombine in all chondrocytes. If Col2-Cre worked as expected, then this would raise the possibility that the remaining 30% of unlabeled Lepr+ BMSCs are formed by cells other than Col2+ cells. One candidate is perichondral Osx+ cells (Maes et al., 2010; Matsushita et al., 2020; Mizoguchi et al., 2014). Some may be generated through endothelial-to-mesenchymal transition (Kenswil et al., 2021). Future studies need to determine the other cells of origin of adult skeletal progenitors and osteoblasts.

**LIMITATIONS OF THE STUDY**

Unlike Acan-creER, Col2-Cre also marked periosteal cells (compare Figures S1C and S7A; Szabova et al., 2009). Acan-Cre might have the same pattern. It is currently difficult to distinguish the fates of chondrocytes and periosteal cells marked by Col2-Cre. Nevertheless, we found that Col10-Cre-marked cells, which excluded periosteal cells (Figure S7C), gave rise to ~58% of all Lepr+ BMSCs (Figures S7D and S7E), whereas Col2-Cre-marked cells give rise to ~70% (Figure 4C). Thus, there were ~12% of Lepr+ BMSCs that might derive from Col2+Col10+ periosteal cells. On one hand, it suggested that we underestimated the contribution of fetal chondrocytes to adult skeletal progenitors; on the other hand, it implied that some adult skeletal progenitors derive from the periosteum. Because the periosteal collar is made by an intramembranous process, this suggests that intramembranous and endochondral bone formation are actually one spectrum. Moreover, only a minor fraction of chondrocytes and osteoblasts at the callus derived from Lepr+ BMSCs during fracture healing (Figures S5E and S5F), supporting the idea that periosteal cells are also involved in fracture healing (Debnath et al., 2018; Ortinau et al., 2019). It will be particularly interesting and important to generate tools to distinguish the fates of progenitors from the growth plate, bone marrow, and periosteum and to dissect their division of labor during bone growth, remodeling, and regeneration.

**STAR METHODS**

Detailed methods are provided in the online version of this paper and include the following:

- **KEY RESOURCES TABLE**
- **RESOURCE AVAILABILITY**
  - Lead contact
  - Materials availability
  - Data and code availability
- **EXPERIMENTAL MODEL AND SUBJECT DETAILS**
  - Animal models
- **METHOD DETAILS**

See also Figure S7.
**Figure 7.** Forced running significantly increases osteoblast formation by perinatal chondrocytes but not by adult Lepr⁺ BMSCs.

(A) Schematic showing the experimental timeline for AcancreER; R26tdTomato; Col1a1-GFP mice.

(B–D) Flow cytometry analyses of femora from AcancreER; R26tdTomato; Col1a1-GFP mice with (C) or without running (B) and quantification of percentages of Col1a1-GFP⁺ osteoblasts that were Tomato⁺ in the femora from the control and the running group (D). Data represent mean ± SD of 5 mice per condition from 5 independent experiments. Two-tailed Student’s t tests were used to assess the statistical significance of differences (**p < 0.01).
SUPPLEMENTAL INFORMATION

Supplemental information can be found online at https://doi.org/10.1016/j.stem.2021.08.010.

ACKNOWLEDGMENTS

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AUTHOR CONTRIBUTIONS

H.S.S. and B.O.Z. initiated the project. H.S.S. performed most of the imaging and flow cytometry experiments. Y.L.L. performed the running exercise experiments and was involved in micro-CT analysis. X.T.T. and X.S.Z. participated in some of the confocal imaging experiments. B.O.Z. provided the R26<sup>tm1.2<sup>WLA</sup>/Rosa26<sup>Tm</sup>cre</sup> and IR mice. W.Z. maintained the AcancreER<sup>+</sup> mice. B.O.Z. and H.S.S. wrote the manuscript and interpreted the data.

DECLARATION OF INTERESTS

The authors declare no competing interests.

REFERENCES


## KEY RESOURCES TABLE

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### Experimental models: Organisms/strains

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| Mouse: R26<sup>Cre-STOP-loxP-loxP-STOP-loxP-Zsgreen</sup> | The Jackson Laboratory | JAX:007906 |
| Mouse: AcancreER | The Jackson Laboratory | JAX:019148 |
| Mouse: Lepr<sup>cre</sup> | The Jackson Laboratory | JAX:008320 |
| Mouse: ACTB-cre | The Jackson Laboratory | JAX:019099 |
| Mouse: R26<sup>Cre-STOP-loxP-loxP-Zsgreen-loxP-tdTomato</sup> | Dr. Bin Zhou | He et al., 2017 |
| Mouse: Col2-cre | Dr. Xiao Yang | Hao et al., 2002 |

(Continued on next page)
RESOURCE AVAILABILITY

Lead contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Bo O. Zhou (bo.zhou@sibcb.ac.cn).

Materials availability
New mouse lines generated in this study will be provided by the lead contact upon request.

Data and code availability
This study did not generate large datasets.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Animal models
All mice were maintained under C57BL/6 background in the Animal Facility at Shanghai Institute of Biochemistry and Cell Biology (SIBCB), Chinese Academy of Sciences. R26R<creR26stop-loxP-loxP-tdTomato> mice were generated by crossing Actb-cre with R26R<creR26stop-loxP-loxP-tdTomato> mice (Madisen et al., 2015) to excise the second loxP-flanked stop cassette. Lepr-creER transgenic mouse line was generated by injecting the BAC clone RP23-184F13, in which the Lepr gene locus was inserted with the creER sequence, into the fertilized eggs from C57BL/6 mice. LeprdreER knock-in mouse line was generated by knocking the dreER sequence into the endogenous Lepr gene locus. Col2dre knock-in mouse line was generated by knocking the dre sequence into the endogenous Col2 gene locus. Dre recombinase excises DNA regions flanked by rox recombination sites, together with Cre, allowing simultaneous tracing of two independent cell population. In the interleaved reporter (IR) mice, recombination by one recombinase should inherently remove one recombinase-recognition site of the other system to render its reporter inert to subsequent recombination (for example, recombination using Dre–rox should remove one loxP site and prevent subsequent Cre–loxP-mediated recombination). All mouse procedures were approved by the Institutional Animal Care and Use Committees of SIBCB.

METHOD DETAILS

Tamoxifen treatment
For perinatal induction of CreER activity, mice were intraperitoneally injected with 30 μg 4-OH tamoxifen at postnatal days 1 to 3. Otherwise, mice were intraperitoneally injected with 100 μL tamoxifen (10 mg/ml) for 5 days. For induction of DreER activity in juvenile mice, mice were intraperitoneally injected with 100 μL tamoxifen for 3 days at 4 and 7 weeks after birth, respectively.

Flow cytometry of whole bone marrow cells
Bone marrow was flushed from femur with 1 mL HBSS digestion buffer containing 5 mg Collagenase I, 5 mg Collagenase IV and 0.1 mg Dnase I and incubated in a shaking bath at 37°C for 30 minutes. Digested cells were centrifuged in 4°C centrifuge at 1500 rpm for 5 minutes. Supernatant was discarded, cells were suspended with 100 μL HBSS buffer and incubated for 30 minutes on ice with the following antibodies: CD45-FITC, Ter119-FITC, PDGFRα-biotin, Lepr-biotin. Streptavidin-APC or Streptavidin-brilliant violet 421 were used as secondary antibodies. Flow cytometry was performed on Attune NxT flow cytometer (Thermo).
Flow cytometry of femur shafts
Marrow-depleted femurs were first cut to small pieces by scissors. Fragments were digested in 37°C shaking bath in 1 mL HBSS digestion buffer (same as above). 30 minutes later, HBSS was transferred into a new 5 mL FACS tube. 1 mL pre-warmed HBSS digestion buffer was added to fragments for another 30 minutes digestion. Digested cells were centrifuged in 4°C centrifuge at 1500 rpm for 5 minutes. Supernatant was discarded and cells were suspended with 100 μL HBSS buffer and incubated for 30 minutes on ice with the following antibodies: CD45-APC, CD31-APC, Ter119-APC. Flow cytometry was performed on Attune NxT flow cytometer (Thermo).

Immunofluorescence and confocal imaging
Femurs were fixed in 4% PFA overnight at 4°C and followed by 5-day decalcification in 20% EDTA. OCT-embedded femurs were sectioned using the CryoJane tape-transfer system (Leica Biosystems). Sections were blocked in PBS with 5% donkey serum and 0.1% triton for 1 hour and then stained overnight with primary antibodies. Primary antibodies used in this study were listed as above. Non-immune immunoglobulins of the same isotype as the primary antibodies were used as negative controls. Images were acquired by Leica TCS SP8 WLL or Leica TCS SP8 STED confocal microscope.

Safranin O Staining
Decalcified bones were dehydrated in ethanol step by step (75%, 85%, 95%, 100%). Subsequently, the specimens were embedded in paraffin and sectioned at 10 μm by Leica manual rotary microtome (Leica RM2235). Prepared paraffin sections went through dewaxing, hydration, and stained with safranin O and fast green. Finally, Slides were mounted with neutral mounting medium and images were acquired by BX51 (Olympus).

Running experiments
Forced running were performed according to literature (Marędziak et al., 2015). Mice were forced to run by putting them on the SA101 treadmill (SANS Biotechnology, China) for 2 hours per day for 2 months. Adaption training started at low speed (7 m/min) to ultimate speed (14 m/min) were performed for one week before mice were given high speed running experiments (14 m/min). Electric shock was on during the whole experiments. Mice were allowed to rest for 4 minutes in the middle of the experiments.

Micro-CT analysis
Dissected femurs in 70% ethanol were scanned by SKYSCAN 1272 3D X-ray microscope (Bruker). The femurs were scanned at the isotropic voxel sizes of 7 μm. The voltage and current were: 60 kV, 166 μA and 80 kV, 125 μA for 2- and 4-month-old mice, respectively. We set the bottom of the distal growth plate as start point and continued for 200 slices to measure trabecular bone parameters. 100 slices were chosen in the middle of diaphysis to measure cortical bone parameters. Bone structure parameters were interpreted according to previous guidance (Bouxsein et al., 2010).

CFU-F forming efficiency assay
Whole bone marrow cells were cultured in 6-well plates at densities ensuring that colonies would form at clonal density. The cultures were incubated at 37°C in a gas-tight chamber (Billups-Rothenberg) with 5% O2 and 10% CO2 for 7 days.

Bone fracture
Mice were anaesthetized by administered with 150 μL 7% chloral hydrate. One of the legs was shaved and scrubbed with 75% ethanol. One 25G needle was used to punch a hole into the marrow through the knee. A stainless steel wire (diameter less than 0.2 mm) was then inserted into intramedullary canal of the femur to stabilize the impending fracture. Three-point bending was performed to make the fracture. Buprenorphine was then used. Steel wire was taken out at 1 week after fracture. Mice were euthanized for analysis at 2 weeks or 8 weeks after fracture.

Lethal irradiation and bone marrow transplantation
Mice were given two doses of 540 rads by RS 2000 X-ray irradiator (Rad Source). At least 2 hours were needed between exposures. One million bone marrow cells were transplanted into irradiated mice in 24 hours to avoid hematopoietic failure. Mice were feed with antibiotic water (neomycin sulfate 1.11 g/l and polymixin B 0.121 g/l) for 14 days after transplantation.

QUANTIFICATION AND STATISTICAL ANALYSIS
Standard two-tailed Student’s t tests and one-way ANOVA were used to analyze statistically significant values. Experiments were performed on at least 3 independent samples. P value less than 0.05 was considered as significant difference. Unless specifically pointed out, data were represented by mean ± SD.