Osteocyte-specific WNT1 regulates osteoblast function during bone homeostasis

Kyu Sang Joeng, …, Catherine Ambrose, Brendan H. Lee


Mutations in *WNT1* cause osteogenesis imperfecta (OI) and early-onset osteoporosis, identifying it as a key Wnt ligand in human bone homeostasis. However, how and where WNT1 acts in bone are unclear. To address this mechanism, we generated late-osteoblast-specific and osteocyte-specific WNT1 loss- and gain-of-function mouse models. Deletion of *Wnt1* in osteocytes resulted in low bone mass with spontaneous fractures similar to that observed in OI patients. Conversely, *Wnt1* overexpression from osteocytes stimulated bone formation by increasing osteoblast number and activity, which was due in part to activation of mTORC1 signaling. While antiresorptive therapy is the mainstay of OI treatment, it has limited efficacy in WNT1-related OI. In this study, anti-sclerostin antibody (Scl-Ab) treatment effectively improved bone mass and dramatically decreased fracture rate in *swaying* mice, a model of global *Wnt1* loss. Collectively, our data suggest that WNT1-related OI and osteoporosis are caused in part by decreased mTORC1-dependent osteoblast function resulting from loss of WNT1 signaling in osteocytes. As such, this work identifies an anabolic function of osteocytes as a source of Wnt in bone development and homoeostasis, complementing their known function as targets of Wnt signaling in regulating osteoclastogenesis. Finally, this study suggests that Scl-Ab is an effective genotype-specific treatment option for WNT1-related OI and osteoporosis.
Osteocyte-specific WNT1 regulates osteoblast function during bone homeostasis

Kyu Sang Joeng,1 Yi-Chien Lee,1 Joohyun Lim,1 Yuqing Chen,1 Ming-Ming Jiang,1 Elda Munivez,1 Catherine Ambrose,2 and Brendan H. Lee1

1Department of Molecular and Human Genetics, Baylor College of Medicine, Houston, Texas, USA. 2Department of Orthopedic Surgery, University of Texas Health Science Center at Houston, Houston, Texas, USA.

Mutations in WNT1 cause osteogenesis imperfecta (OI) and early-onset osteoporosis, identifying it as a key Wnt ligand in human bone homeostasis. However, how and where WNT1 acts in bone are unclear. To address this mechanism, we generated late-osteoblast-specific and osteocyte-specific WNT1 loss- and gain-of-function mouse models. Deletion of Wnt1 in osteocytes resulted in low bone mass with spontaneous fractures similar to that observed in OI patients. Conversely, Wnt1 overexpression from osteocytes stimulated bone formation by increasing osteoblast number and activity, which was due in part to activation of mTORC1 signaling. While antiresorptive therapy is the mainstay of OI treatment, it has limited efficacy in WNT1-related OI. In this study, anti-sclerostin antibody (Scl-Ab) treatment effectively improved bone mass and dramatically decreased fracture rate in swaying mice, a model of global Wnt1 loss. Collectively, our data suggest that WNT1-related OI and osteoporosis are caused in part by decreased mTORC1-dependent osteoblast function resulting from loss of WNT1 signaling in osteocytes. As such, this work identifies an anabolic function of osteocytes as a source of Wnt in bone development and homeostasis, complementing their known function as targets of Wnt signaling in regulating osteoclastogenesis. Finally, this study suggests that Scl-Ab is an effective genotype-specific treatment option for WNT1-related OI and osteoporosis.

Introduction

Wnt signaling is a well-established pathway that regulates skeletal development and homeostasis (1). Genetic studies of LRP5 in human and mouse strongly suggest that canonical Wnt signaling regulates postnatal bone formation (2–5). Further genetic studies with β-catenin and other Wnt ligands using various mouse models provided additional evidence that corroborate the critical function of Wnt signaling in skeletal development and bone homeostasis (6–17). Several recent studies also reported that Wnt signaling directly regulates osteoclast function (18–22). Despite the established function of Wnt signaling in bone, the role of specific Wnt ligands in human bone homeostasis was not clear. Our group and others reported that heterozygous loss-of-function mutations in WNT1 can cause dominantly inherited early-onset osteoporosis, while biallelic mutations resulting in complete loss of function lead to recessively inherited osteogenesis imperfecta (OI) (23–29). The semidominant inheritance of this mutation spectrum underscores the strict temporal, spatial, and dosage requirement of this essential Wnt ligand in human bone homeostasis. Moreover, we have established a swaying mouse model (Wnt1sw/sw mice, which carry a mutation in Wnt1) as a murine model of WNT1-related OI (30). These human and mouse genetic studies strongly suggest that WNT1 is a major Wnt ligand regulating human bone homeostasis; however, the mechanistic basis of WNT1 action in bone homeostasis and its cellular source and targets in bone are unclear.

Osteocytes are one of the major cell types in bone. They are differentiated from osteoblasts and embedded in bone matrix. It was originally believed that osteocytes are quiescent cells in bone matrix, but numerous studies now show that osteocytes play essential roles in bone homeostasis by regulating osteoblasts and osteoclasts (31). For example, osteocytes can regulate osteoclastogenesis by expressing RANKL and its decoy receptor osteoprotegerin (7, 32–36). Osteocytes can also regulate osteoblast differentiation by secreting sclerostin (SOST), an inhibitor of Wnt signaling (37–39). However, whether osteocytes can directly regulate bone formation as a “Wnt-sending” cell type is not clear.

Mammalian target of rapamycin complex 1 (mTORC1) signaling regulates cell metabolism, growth, proliferation, and survival (40, 41). mTORC1 signaling can be activated by various nutritional and environmental signals, including growth factors and amino acids. The mTORC1 complex is composed of multiple components, including mTOR (a serine/threonine kinase), RAPTOR (regulatory-associated protein of mTOR), mLST8 (an mTOR-associated mammalian homolog of LST8), PRAS40 (proline-rich AKT substrate 40 kDa), and DEPTOR (DEP domain-containing mTOR-interacting protein). The tuberous sclerosis heterodimer (TSC1/2) negatively regulates mTORC1 signaling by inhibiting Rheb (the small GTPase) that is necessary for mTORC1 kinase activity. Rapamycin is a bacterial macrolide that inhibits mTORC1 signaling by forming a complex with FKBP12. Eukaryotic initiation factor 4E–binding protein 1 (4E-BP1) and p70 ribosomal S6 kinase 1 (S6K1) are well-known downstream targets of mTORC1 signaling. Interestingly, recent studies have shown that specific Wnt ligands may regulate osteoblast differentiation, partly through mTORC1

Related Commentary: p. 2539

Authorship note: K.S. Joeng and Y.C. Lee contributed equally to this work.
Conflict of interest: The authors have declared that no conflict of interest exists.
Submitted: January 3, 2017; Accepted: April 27, 2017.
signaling (12, 42–44). However, whether mTORC1 is necessary for WNT1-induced osteoblast differentiation remains unclear.

Current pharmacological treatment options for OI predominantly rely on antiresorptive therapy with bisphosphonates, which inhibits bone resorption by blocking osteoclast differentiation and activity (45, 46). Interestingly, bisphosphonate therapy has been reported to be less effective in patients with WNT1-related OI compared with responses observed in patients with other forms of OI (47). As such, there is an unmet need of new therapeutic approaches for the treatment of WNT1-related OI and osteoporosis. Sclerostin inhibits WNT signaling by binding to and interfering with WNT ligand engagement with LDL receptor–related protein 5/6 (LRP5/6) (37–39), and has been validated as a potent therapeutic target for increasing bone formation in vivo (48–51). In fact, anti-sclerostin antibody (Scl-Ab) is currently in phase III clinical trials as an anabolic therapy for postmenopausal osteoporosis (NCT02016716, ClinicalTrials.gov). But the efficacy of Scl-Ab for treating WNT1-related OI patients who may have significantly reduced endogenous WNT signaling is unknown. At the same time, Scl-Ab may improve bone mass and fracture rates if there is the potential of additional WNT ligands that can work redundantly in bone. Therefore, it is necessary to examine the efficacy of Scl-Ab for treating WNT1-related OI patients as a potential, genotype-specific therapeutic approach.

In this study, we determined the bone-specific function of Wnt1 by generating late-osteoblast/osteocyte-specific Wnt1 loss-and gain-of-function mouse models. Our pharmacological and genetic rescue experiments showed that the function of WNT1 in osteoblasts is partly mediated by mTORC1 signaling. In addition, Scl-Ab treatment improved low bone mass and dramatically decreased fracture rate in a WNT1-related OI mouse model (swaying mice, hereafter referred to as Wnt1(sw/sw)). Overall, our data support the model wherein osteocytes serve as a source of WNT1 and the bone fragility of WNT1-related OI patients is caused in part by decreased osteoblast function due to the loss of WNT1 signaling from osteocytes to osteoblasts. Finally, this study suggests that Scl-Ab may be an effective genotype-specific treatment option for WNT1-related patients with OI and osteoporosis.

Results

Specific deletion of Wnt1 in late osteoblasts and osteocytes caused spontaneous fractures and severe loss of bone. To elucidate the bone-specific function of Wnt1, we generated a Wnt1 conditional knockout mouse model (Wnt1fl/fl) by using embryonic stem cells derived from the European Conditional Mouse Mutagenesis Program (EUCOMM) allele (Wnt1EUCOMM; Supplemental Figure 1A; supplemental material available online with this article; https://doi.org/10.1172/JCI92617DS1). We have previously shown that subsets of osteocytes are a potential source of WNT1 in bone via lineage tracing experiments (25); therefore, we deleted Wnt1 in late osteoblasts and osteocytes by generating Wnt1fl/fl Dmp1-Cre mice. Strikingly, Wnt1fl/fl Dmp1-Cre mice showed spontaneous fractures (fracture rate of 67%) and low bone mass in both males and females, without an effect on growth or behavior (Figure 1, A and B, and Supplemental Figure 2, A and B). X-ray images of the hindlimbs showed severe osteopenia in Wnt1fl/fl Dmp1-Cre mice, as
Overexpressing Wnt1 in late osteoblasts and osteocytes showed a high bone mass phenotype. To gain further insight into the therapeutic potential of Wnt1 in bone mass accretion, we generated Dmp1-Cre Rosa26Wnt1/+ mice overexpressing Wnt1 in late osteoblasts and osteocytes (Supplemental Figure 1B). In agreement with the loss-of-function study, both male and female Dmp1-Cre Rosa26Wnt1/+ mice showed a dramatic high bone mass phenotype in femurs and vertebrae (Figure 2, A–C, and Supplemental Figure 2, D–F). X-ray images of the hindlimbs showed increased bone mass in Dmp1-Cre Rosa26Wnt1/+ mice, and μCT analysis revealed an approximately 5-fold increase in trabecular bone mass in mutant mice, which was associated with increased trabecular number and thickness (Figure 2, A and B). Additionally, cortical bone thickness increased about 60% in mutant mice compared with WT controls (Figure 2B).

Histomorphometric analysis showed significant increases in osteoblast number, mineralizing surface, mineral apposition rate, and bone forming rate, whereas osteoclast number per bone surface was unaffected in mutant mice (Figure 2C). Interestingly, serum CTX level is increased in Dmp1-Cre Rosa26Wnt1/+ mice (Supplemental Table 1), consistent with higher total osteoclast number per bone surface resulting from increased total bone surface (data not shown). These data strongly suggest that the high bone mass is caused by increased osteoblast number and activity in Dmp1-Cre Rosa26Wnt1/+ mice in opposite fashion to the conditional loss-of-function model.

Overexpression of Wnt1 increased osteoblast differentiation and mineralization in vitro. To verify the function of Wnt1 in osteoblast differentiation and bone mineralization, we performed in vitro functional studies using a mouse stromal cell line (ST2) transduced with
consistent with enhanced osteoblast differentiation, Wnt1 overexpression markedly enhanced in vitro mineralization (Figure 3B). Based on our mouse genetics and in vitro studies, we conclude that late-osteoblast/osteocyte-derived WNT1 contributes to bone homeostasis by regulating osteoblast function. In addition, these results confirm that impaired osteoblast function due to loss of Wnt1 in osteocytes is a possible pathogenic mechanism of WNT1-related OI.

**Figure 3.** mTORC1 signaling mediated enhanced osteoblast differentiation and mineralization by Wnt1 overexpression in vitro. (A) Quantitative reverse transcriptase PCR (RT-PCR) of Wnt1, Runx2, alkaline phosphatase, and osteocalcin in control helper-dependent adenovirus-treated (HDAd-GFP) and WNT1 helper-dependent adenovirus-treated (HDAd-mWnt1) ST2 cells. Results are shown as fold change of the mean of control group ± SD (n = 3 per group). (B) Mineralization assay by alizarin red staining on the seventh day after control virus–treated and HDAd-mWnt1–treated ST2 cells. (C) The representative Western blot analysis showed activated pS6 relative to total S6 and pAkt (Ser473) relative to total Akt of control virus– and HDAd-mWnt1–treated ST2 cells. Results are shown as fold change of the mean of control group ± SD (n = 3 per group). The Western blot represents 3 individual experiments. (D and E) Quantitative RT-PCR of alkaline phosphatase (D) and Lef1 (E) after control treatment (DMSO) or pharmacological inhibition of mTOR signaling by rapamycin in control virus– and HDAd-mWnt1–treated ST2 cells. Results are shown as fold change of the mean of control group ± SD (n = 3 per group). (F) Mineralization assay after control treatment (DMSO) or pharmacological inhibition of mTOR signaling by rapamycin in control virus– and HDAd-mWnt1–treated ST2 cells. The comparison between control and HDAd-mWnt1–treated groups is determined by Student’s t test. *P < 0.05, **P < 0.01, ***P < 0.001.
Phenotypes caused by gain of WNT1 signaling can be reversed by rapamycin. Previous studies by others have shown that Wnt signaling can regulate canonical β-catenin pathways as well as noncanonical targets like the mTOR pathway (12, 42–44). Interestingly, recent studies have demonstrated that WNT ligands may activate mTOR signaling in osteoblasts to affect their differentiation (12, 42). Consistent with these studies, we found that Wnt1 overexpression significantly increased phosphorylation of S6, a major downstream target of mTORC1, but not phosphorylation of Akt, a major downstream target of mTORC2 (Figure 3C). Based on these results, we hypothesized that mTORC1 signaling may mediate WNT1 function in osteoblast differentiation and mineralization. To test this hypothesis, we performed a pharmacological rescue experiment by treating ST2 cells overexpressing Wnt1 with rapamycin, an inhibitor of mTORC1 signaling (52). Rapamycin treatment significantly reduced the expression of alkaline phosphatase, rapamycin treatment significantly reduced Wnt1-induced osteoblast differentiation and in vitro mineralization in a β-catenin–independent manner. To verify these results in vivo, we performed a pharmacological rescue experiment by treating Dmp1-Cre Rosa26 mice with rapamycin. Interestingly, 1 month of rapamycin treatment significantly reversed the high bone mass phenotype of Dmp1-Cre Rosa26 mice (Figure 4A). X-ray radiography revealed a reduction in trabecular bone and cortical bone thickness of rapamycin-treated Dmp1-Cre Rosa26 mice (Figure 4A). μCT parameters showed that rapamycin fully normalized cortical bone thickness and partially reduced trabecular bone volume (Figure 4B), suggesting a differential impact of mTORC1 signaling on WNT1 function in trabecular versus cortical bone compartments. Histomorphometric analysis showed that inhibition of mTORC1 in the WNT1 gain-of-function model was not changes in osteoclast and osteoblast number (Figure 4C). This pharmacological rescue experiment supports that mTORC1 partially mediates the gain of WNT1 signaling in osteoblasts.

Genetic activation of mTORC1 signaling rescued the low bone mass phenotype of the swaying mice. To test the physiological relevance of Wnt1-induced mTORC1 signaling, we determined whether genetic activation of mTORC1 signaling could rescue the low bone mass caused by loss of Wnt1 signaling in bone. To this end, we activated mTORC1 signaling in late osteoblasts/osteocytes by deleting Tsc1, a negative regulator of mTORC1, in Wnt1 mice (Tsc1fl/ fl Wnt1 sw/sw Dmp1-Cre). As expected, Tsc1fl/fl Wnt1 sw/sw Dmp1-
Cre mice no longer suffered spontaneous fractures and had higher trabecular and cortical bone parameters when compared with Wnt1sw/sw mice (Figure 5A). μCT analysis showed partial rescue of trabecular bone and full rescue of cortical bone thickness in Tsc1fl/fl Wnt1sw/sw Dmp1-Cre mice (Figure 5B). Furthermore, histomorphometric analysis revealed that the rescue phenotype was associated with increased bone formation rate and also decreased osteoclast numbers (Figure 5C). Based on our mouse rescue studies and in vitro experiments, we conclude that mTORC1 partially mediates the physiological function of WNT1 signaling in regulating bone formation and mineralization.

Scl-Ab treatment significantly improved the low bone mass phenotype of swaying mice. To test both the therapeutic potential of anti-sclerostin antibody (Scl-Ab) for WNT1-related OI patients and the potential for redundancy of Wnt ligand function in bone, we administered Scl-Ab subcutaneously at a dose of 25 mg/kg to Wnt1sw/sw mice twice a week from 2 to 8 weeks of age. Remarkably, Scl-Ab administration in Wnt1 sw/sw mice caused a significant improvement in fracture rate from 90% to 12.5% (Table 1), as well as increased bone mass (Figure 6A). μCT analysis showed that Scl-Ab-treated Wnt1sw/sw mice exhibited a significant increase in trabecular bone volume and cortical thickness in both femur and lumbar spine (Figure 6B and Supplemental Figure 3). Scl-Ab treatment significantly increased osteoblast activity as indicated by the increased mineralized surface, mineral apposition rate, and bone formation rate (Figure 6C). More importantly, 3-point bending analysis confirmed that the Scl-Ab–treated Wnt1sw/sw mice had significantly increased bone ultimate load, stiffness, and post-yield energy compared with untreated Wnt1sw/sw mice (Figure 6D). Taken together, these results indicate that Scl-Ab treatment significantly improved the bone quantity and quality of Wnt1sw/sw mice.

Discussion

Our current studies significantly advance our understanding of the function of Wnt ligand in bone homeostasis and the pathogenic mechanism of WNT1-related skeletal disorders. Given the CNS phenotype found in WNT1-related OI patients and the Wnt1sw/sw mouse model (23–29, 53), the bone fragility of WNT1-related OI could be caused by loss of WNT1 function in bone as well as in other organs. In addition, the extremely low expression of Wnt1 in bone has also hindered determination of the tissue-specific function of WNT1. Therefore, our mouse genetic studies using a conditional model provide strong evidence that WNT1 specifically functions in bone. We generated a conditional mouse model deleting Wnt1 in late osteoblasts/osteocytes using the Dmp1-Cre transgenic line. Interestingly, Dmp1-Cre–specific deletion of Wnt1 causes spontaneous fractures and low bone mass, recapitulating the phenotypes of Wnt1sw/sw mice. These data strongly suggest that
the bone fragility in Wnt1sw/sw mice and WNT1-related OI patients is mainly caused by loss of WNT1 function in bone.

Our Wnt1 loss- and gain-of-function studies suggest that osteoblasts are the major target cell of WNT1 signaling. The low bone mass phenotype in the loss-of-function study is mainly associated with decreased osteoblast activity, while the high bone mass phenotype in the gain-of-function study is associated with increased osteoblast number and activity. It is worth noting that osteoblast number was only affected in the gain-of-function study, which may be partly because of overexpression of Wnt1. Nonetheless, we conclude that late-osteoblast/osteocyte-derived Wnt1 contributes to bone homeostasis by regulating osteoblast function.

Our study also provides functional evidence that osteocytes are the major source of WNT1. Our previous lineage-tracing experiment using the transgenic Wnt1-Cre transgenic mice supports the hypothesis that osteocytes are a potential source of WNT1 in bone, although we could not rule out nonspecific effects of the transgene in those experiments (25). Our current Dmp1-Cre–specific Wnt1 loss-of-function study provides strong evidence supporting the idea that the major skeletal source of WNT1 is a subset of osteocytes in bone, as suggested by the lineage-tracing experiment. This is also consistent with the extremely low expression of Wnt1 in bone. The specificity of Dmp1-Cre to late osteoblasts and osteocytes does not rule out the possibility that the late osteoblasts may also express Wnt1. But we found only GFP-positive osteocytes and no GFP-positive osteoblasts in our previous lineage-tracing experiment (25). This strongly suggests that osteocytes are a major source of WNT1 in bone rather than osteoblasts. Further investigation of Wnt1 deletion with early-osteoblast Cre lines will further elucidate the function of WNT1 in early osteoblast lineage cells.

Our finding of osteocytes as a source of WNT1 ligand suggests that osteocytes could have a dual role as a sender and receiver of Wnt ligands during bone homeostasis (Figure 6E). As sending cells, osteocytes stimulate osteoblastogenesis by triggering Wnt signaling in osteoblasts. This is consistent with previous studies in which mice with bone-specific Lrp5 and Lrp6 knockout showed osteoblast-specific phenotypes (54, 55). Our study and those of others strongly suggest that the regulation of osteoblastogenesis by osteocytes can be mediated by a β-catenin–independent pathway like mTOR signaling (12, 42). As receiving cells, osteocytes can modulate osteoclastogenesis by triggering cell-autonomous β-catenin–dependent signaling. Consistent with this notion, osteocyte–specific β-catenin knockout mice show predominantly an osteoclast phenotype (33). In addition, previous studies showed that osteocytes regulate osteoclastogenesis by secreting RANKL and osteoprotegerin that are downstream targets of canonical β-catenin–dependent WNT signaling (7, 32–36). Our current study and other previous studies also strongly suggest that there is specificity between WNT ligands and downstream cell signaling targets (10, 12, 14, 42). Hence, the osteocyte can be seen increasingly as the central regulator of bone mass. As a receiver of Wnt signaling, it regulates osteoclastogenesis in a β-catenin–dependent fashion. As a sender cell of Wnt signaling, it controls anabolic aspects of osteoblast function by targeting mTORC1, though we cannot rule out additional roles in early stages of osteoblast differentiation. This last function together with its production of sclerostin shows that it can also serve as a negative and positive regulator of osteoblast differentiation and function, respectively (Figure 6E).

Recent studies also showed that Wnt signaling could directly regulate osteoclastogenesis (18–22). Further studies will be required to elucidate the direct regulation of osteoclasts by osteocyte-derived Wnt signaling. The code for the underlying ligand–cell specificity likely integrates both temporal and spatial requirements that help to explain the complexity of both cellular (osteoblastic versus osteoclastic) and tissue compartment (trabecular versus cortical, and appendicular versus axial) phenotypes observed in different genetic models of Wnt signaling components. With no doubt, additional study and improved genetic reagents will be required to resolve this Wnt signaling code in the skeleton.

Scl-Ab treatment significantly improved the bone quantity and quality of Wnt1sw/sw mice. The rescued bone phenotypes of Wnt1sw/sw mice strongly suggest that other WNT ligands, including excellent candidates such as WNT16, WNT10b, and WNT7b, can compensate for loss of Wnt1 in Wnt1sw/sw mice (10–17). Despite the potential redundancy of multiple WNT ligands that affect bone formation, the incomplete restoration of bone volume by Scl-Ab treatment in Wnt1sw/sw mice suggests that WNT1 clearly has a critical function in bone homeostasis. Overall, this study supports the potential utility of Scl-Ab in the management of OI patients with WNT1 mutations. Further investigation to determine the efficacy of Scl-Ab in human WNT1-related OI patients and early-onset osteoporosis will be required for clinical application.

Methods

Animals. Embryonic stem cells carrying knockout-first Wnt1 allele (Wnt1EUCOMM) were obtained from the European Conditional Mouse Mutagenesis Program (EUCOMM) and injected into embryos in house. Wnt1EUCOMM mice were crossed with Rosa26-Flipase (Flip) mice to delete β-gal and the neo-cassette in order to generate the conditional knockout allele (Wnt1f/F). Rosa26Wnt1+/+ mice were provided by Thomas Carroll and Andrew McMahon (56), and Dmp1-Cre transgenic mice were provided by Jian Feng (57). The swaying (Wnt1sw/sw) mouse (58, 59) and Tsc1f/f mouse (60) models were obtained from the Jackson Laboratory. Ramapycin was purchased from LC Laboratories, and subcutaneous rapamycin (4 mg/kg) was administered daily to female Rosa26Wnt1+/+ mice for a month. The sclerostin-neutralizing antibody (Sc-Al) was provided by Amgen Inc. and UCB Pharma. Sc-Al was diluted 1:10 in PBS to a concentration of 3.6 mg/ml for administration. Female Wnt1sw/sw and WT mice were randomly assigned and treated with subcutaneous Sc-Al (25 mg/kg) or the control vehicle (PBS) twice per week, starting at 3 weeks old, for 6 weeks and were eutha-
Figure 6. Wnt\textsuperscript{floxed} mice showed phenotypic corrections in femurs after treatment with Scl-Ab. (A) X-ray radiograph of femurs of 2-month-old female WT and Wnt\textsuperscript{floxed} mice treated with vehicle control or sclerostin-neutralizing antibody (Scl-Ab). (B) \(\mu\)CT analysis of femoral trabecular bone for bone volume/total volume (BV/TV) and of cortical bone for cortical thickness (Cort.Th) in WT and Wnt\textsuperscript{floxed} mice treated with vehicle control or Scl-Ab. Results are shown as means ± SD (\(n=7\) for WT, \(n=3\) for WT with Scl-Ab, \(n=8\) for Wnt\textsuperscript{floxed}, \(n=7\) for Wnt\textsuperscript{floxed} with Scl-Ab). (C) Histomorphometric analysis of L4 vertebrae for mineral surface per bone surface (MS/BS), mineral apposition rate (MAR), and bone formation rate (BFR) in WT and Wnt\textsuperscript{floxed} mice treated with vehicle control or Scl-Ab. Results are shown as means ± SD (\(n=6\) for WT, \(n=4\) for WT with Scl-Ab, \(n=9\) for Wnt\textsuperscript{floxed}, \(n=7\) for Wnt\textsuperscript{floxed} with Scl-Ab). (D) Biomechanical testing results by 3-point bending assay for maximum load, stiffness, and post-yield energy of femurs in WT and Wnt\textsuperscript{floxed} mice treated with vehicle control or Scl-Ab. Results are shown as means ± SD (\(n=7\) for WT, \(n=4\) for WT with Scl-Ab, \(n=8\) for Wnt\textsuperscript{floxed}, \(n=8\) for Wnt\textsuperscript{floxed} with Scl-Ab). The comparisons of WT mice with vehicle versus Scl-Ab treatment and Wnt\textsuperscript{floxed} mice with vehicle versus Scl-Ab treatment are determined by Mann-Whitney U test. *\(P<0.05\), **\(P<0.01\), ***\(P<0.001\). (E) Schematic model of osteocyte function as a sender and receiver of Wnt signaling. OPG, osteoprotegerin.
nized at 8 weeks old. Fracture rate was defined by callus formation in tibia during final dissection of mouse at 8 weeks old. Either unilateral or bilateral fracture in a mouse was counted as 1 incident fracture. No blinding was possible during treatment because Scl-Ab or control vehicle was injected according to group allocation. In all subsequent analyses, the investigators were blinded to genotype and treatment group. Calcine (250 μg i.p.) was injected first and followed by alizarin red (800 μg i.p.) injections with a 5-day interval before euthanasia to assess the parameters of dynamic bone histomorphometry. Spines and right femurs were collected and fixed in 10% formalin for 48 hours for subsequent radiography and μCT imaging. After imaging, spines were embedded in plastic for histomorphometry. Left femurs were wrapped in saline-soaked gauze and stored at −20°C for biomechanical testing.

Both male and female mice were characterized for Wnt1 gain- and loss-of-function studies. Male mouse data are presented in Supplemental Figure 2. For the mTORC1 genetic rescue experiment, female mice were mainly characterized at 2 months of age. We also characterized male mice using μCT (n = 3, data not shown). We performed the rapamycin rescue experiments to be consistent with the mTORC1 genetic rescue experiment. Therefore, female mice were characterized at 2 months of age. For the Scl-Ab treatment experiment, female mice were used to prevent artificial fractures in the swaying mouse model, which occurred more frequently in male mice, likely due to their propensity for physical movement. The bone phenotypes for the Scl-Ab rescue experiment were characterized in 2-month-old mice.

**ST2 cell culture.** The bone marrow–derived stromal cell line ST2 (61) was provided by the laboratory of Fanxin Long (Department of Orthopedic Surgery, Washington University School of Medicine, St. Louis, Missouri, USA) (the cells were not recently further profiled or tested for mycoplasma contamination). ST2 cells were cultured in growth medium (α-MEM HyClone with 10% FBS, 100 U/ml penicillin, and 100 μg/ml streptomycin). GFP and WNT1 helper-dependent adenoviruses were generated as previously described (62). The viruses were added to the medium (500 viral particles per cell) for 24 hours. After virus transfection, the medium was replaced with mineralization medium (growth medium with 500 μM ascorbic acid and 10 mM β-glycerophosphate) (day 0). The mineralization medium was changed every 2 days. Rapamycin in DMSO (20 nM) was added to the mineralization medium and fed to the virus-infected ST2 cells from day 0. ST2 cells were collected for real-time PCR on day 4. Alizarin red staining was performed on day 10.

**Quantitative real-time reverse transcriptase PCR.** Total RNA from virus-transfected ST2 cells was extracted with Trizol reagent (Invitrogen). The Superscript III reverse transcriptase PCR (RT-PCR) system (Invitrogen) was used to synthesize cDNA. We performed quantitative RT-PCR on a LightCycler 96 (Roche) using gene-specific primers and β-mercaptoethanol. The pS6 (2211, Cell Signaling), pAkt (Ser473; 4060, Cell Signaling), S6 (2217, Cell Signaling), and Akt (4691, Cell Signaling) monoclonal antibodies were used with a 1:1,000 dilution in Tris-buffered saline and Tween 20 containing 5% BSA. Quantification of the density of each band was performed by ImageJ software (NIH).

**Immunoblotting and antibodies.** Protein from virus-transfected ST2 cells was extracted with Laemmli buffer containing 5% β-mercaptoethanol. The pS6 (2211, Cell Signaling), pAkt (Ser473; 4060, Cell Signaling), S6 (2217, Cell Signaling), and Akt (4691, Cell Signaling) monoclonal antibodies were used with a 1:1,000 dilution in Tris-buffered saline and Tween 20 containing 5% BSA. Quantification of the density of each band was performed by ImageJ software (NIH).

**Radiography, μCT imaging, and bone histomorphometry.** Radiography was performed with the Xpert 80 (Kubtec) system, and μCT was conducted with the Scanco μCT-40 system (55-peak kilovoltage and 145-μA x-ray source). A standardized region of the distal femoral metaphysis and the L4 vertebrae were scanned at 16-μm resolutions. Mouse spine samples were then embedded in plastic and sectioned with tungsten carbide blades. Trichrome staining and tartrate-resistant acid phosphatase staining were performed for visualizing osteoblasts and osteoclasts, respectively. Calcine and alizarin red double labeling was used to study dynamic histomorphometry. Results were quantified with the Bioquant Osteo 2014 Image Analysis System for evaluation of bone formation and resorption parameters.

**Biomechanical testing by 3-point bending.** Left femurs were tested by 3-point bending, with a span of 6 mm, using an Instron 5848 device (Instron Inc.). All the femurs were tested wet at room temperature. They were preloaded to 1N at a rate of 0.05 N/s for 5 seconds. Following preloading, the femurs were compressed to failure at a rate of 0.1 mm/s. Load and displacement data were captured at a rate of 40 Hz using Bluehill Software (Instron). Maximum load was determined by finding the highest load values recorded before the specimen failed. Stiffness was determined by the steepest slope of the linear portion of the load and displacement curve. The yield point was calculated using a 0.2% offset from the steepest slope. Using the trapezoidal numerical integration method, energy to failure was calculated as the area under the load-displacement curve and was divided into the elastic energy (up to the yield point) and plastic energy (from the yield point until the failure point).

**Serum CTX-1 level by ELISA.** Blood samples were collected from 2-month-old mice by retro-orbital bleeding under anesthesia before euthanasia. Serum was separated by centrifuge in the tube with serum separator (BD Microtainer) and stored at −80°C. The bone resorption marker C-terminal telopeptide of type I collagen (CTX-1) was measured in serum. The assay was performed using a CTX-1 ELISA kit (RatLaps, Immunodiagnostic Systems) according to the manufacturer’s instructions. Data were analyzed with SigmaPlot 11.0 (Systat Software Inc.).

**Statistics.** For comparisons between 2 groups following normal distribution, we performed 2-tailed Student’s t tests with a significance level of 0.05. For comparisons among 4 groups, and for those observations that did not follow normal distribution, we performed the Kruskal-Wallis test to detect heterogeneity among groups and applied the Mann-Whitney U test for multiple comparisons between groups. Bonferroni adjustment was applied for multiple comparisons. Each P value was compared with the critical P value divided by the total number of comparisons made to determine whether differences between groups were significant. To determine the initial sample size per group of mice, we performed power analysis, which showed that a group size of 6 mice is required to detect a minimal difference of 20% in bone mass (BV/TV) by μCT between Dmp1-Cre Rosa26+/- mice and WT mice. We used the statistical software R (3.2.4) for all statistical analyses.

**Study approval.** The animal care and use for this study have been approved by the Institutional Animal Care and Use Committee of Baylor College of Medicine (Houston, Texas, USA).

**Author contributions**

KSJ, YCL, and BHL designed the research studies. KSJ, YCL, JL, YC, MMJ, and EM conducted the experiments. KSJ and YCL analyzed the data. CA conducted the biomechanical studies. KSJ, YCL, and BHL wrote the manuscript.
Acknowledgments

The authors thank Brian Dawson and Carrie Jiang for technical assistance. We also appreciate Thomas Carroll, Andrew McMahan, and Jian Feng for providing Rosa26-Wnt1 and Dmp1-Cre mouse lines. This work was supported by the Baylor College of Medicine (BCM) Intellectual and Developmental Disabilities Research Center (HD024064) of the Eunice Kennedy Shriver National Institute of Child Health and Human Development, the BCM Advanced Technology Cores with funding from the NIH (AI036211, CA125123, and RR024574), the Rolanette and Berdon Lawrence Bone Disease Program of Texas, and the BCM Center for Skeletal Medicine and Biology. Research funding to authors includes NIH grants PO1 HD22657 (to BHL), PO1 HD070394 (to BHL), K01 AR069002 (to KS), and the Rising Star Award of American Society for Bone and Mineral Research (to KS).

Address correspondence to: Brendan H. Lee, Baylor College of Medicine, One Baylor Plaza, Houston, Texas 77030, USA. Phone: 713.798.8835; Email: blee@bcm.edu.

6. Day TF, Guo X, Garrett-Beal L, Yang Y. Wnt/β-catenin signaling in mesenchymal progenitors controls osteoblast and chondrocyte differentiation during vertebrate skeletogenesis. Dev Cell. 2005;8(5):739–750.
by erythropoietin depends on Wnt1, the PI 3-K/mTOR pathway, Bad, and Bcl-xL. Aging (Albany NY). 2012;4(3):187–201.


58. Lane PW. Mouse news lett. 1967;36:40.


61. Tong J, Kishi H, Matsuda T, Muraguchi A. A bone marrow-derived stroma cell line, ST2, can support the differentiation of fetal thymocytes from the CD4+ CD8+ double negative to the CD4+ CD8+ double positive differentiation stage in vitro. Immunology. 1999;97(4):672–678.