Mammalian focal adhesion proteins Pinch1 and Pinch2 regulate integrin activation and cell–extracellular matrix adhesion and migration. Here, we show that deleting Pinch1 in osteocytes and mature osteoblasts using the 10-kb mouse Dmp1-Cre and Pinch2 globally (double KO; dKO) results in severe osteopenia throughout life, while ablating either gene does not cause bone loss, suggesting a functional redundancy of both factors in bone. Pinch deletion in osteocytes and mature osteoblasts generates signals that inhibit osteoblast and bone formation. Pinch-deficient osteocytes and conditioned media from dKO bone slice cultures contain abundant sclerostin protein and potently suppress osteoblast differentiation in primary BM stromal cells (BMSC) and calvarial cultures. Pinch deletion increases adiposity in the BM cavity. Primary dKO BMSC cultures display decreased osteoblastic but enhanced adipogenic, differentiation capacity. Pinch loss decreases expression of integrin β3, integrin-linked kinase (ILK), and α-parvin and increases that of active caspase-3 and -8 in osteocytes. Pinch loss increases osteocyte apoptosis in vitro and in bone. Pinch loss upregulates expression of both Rankl and Opg in the cortical bone and does not increase osteoclast formation and bone resorption. Finally, Pinch ablation exacerbates hindlimb unloading–induced bone loss and impairs active ulna loading–stimulated bone formation. Thus, we establish a critical role of Pinch in control of bone homeostasis.
Focal adhesion proteins Pinch1 and Pinch2 regulate bone homeostasis in mice

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Introduction

In vertebrates, adult skeleton is a dynamic tissue that is constantly performing bone remodeling, a process in which the osteoclast-mediated bone resorption is closely coupled with the osteoblast-mediated bone formation. Bone remodeling allows skeleton to adapt its structure to mechanical loading by changing its size, shape, and strength; repairing damaged bone; and maintaining a proper bone mass to support bone homeostasis. Abnormal bone remodeling causes metabolic bone diseases, such as osteoporosis and osteopetrosis. Elucidating mechanisms that regulate bone remodeling is likely to reveal new approaches to the prevention and treatment of these diseases.

Osteocytes originate from osteoprogenitors through osteoblast differentiation and are the most abundant bone cell, comprising 90%–95% of total bone cells. Osteocytes play an important role in regulating bone remodeling under normal physiological and diseased states (1–7). Osteocytes communicate with each other and with osteoclasts and osteoblasts on the bone surfaces through their dendrites. Osteocytes produce several important soluble factors that regulate osteoclast and osteoblast formation and function. Among those are sclerostin, the activator of NF-κB ligand (RANKL), and osteoprotegerin (OPG) (8–11). Osteocytes are the main source of the fibroblast growth factor 23 (FGF23), a major regulator of phosphate homeostasis (12). Mechanical forces are the most potent anabolic stimulus of the skeleton. Mechanical loading from daily movement, exercise, and gravity increases bone mass and strength and improves bone

Mammalian focal adhesion proteins Pinch1 and Pinch2 regulate integrin activation and cell–extracellular matrix adhesion and migration. Here, we show that deleting Pinch1 in osteocytes and mature osteoblasts using the 10-kb mouse Dmp1-Cre and Pinch2 globally (double KO; dKO) results in severe osteopenia throughout life, while ablating either gene does not cause bone loss, suggesting a functional redundancy of both factors in bone. Pinch deletion in osteocytes and mature osteoblasts generates signals that inhibit osteoblast and bone formation. Pinch-deficient osteocytes and conditioned media from dKO bone slice cultures contain abundant sclerostin protein and potently suppress osteoblast differentiation in primary BM stromal cells (BMSC) and calvarial cultures. Pinch deletion increases adiposity in the BM cavity. Primary dKO BMSC cultures display decreased osteoblastic but enhanced adipogenic, differentiation capacity. Pinch loss decreases expression of integrin β3, integrin-linked kinase (ILK), and α-parvin and increases that of active caspase-3 and -8 in osteocytes. Pinch loss increases osteocyte apoptosis in vitro and in bone. Pinch loss upregulates expression of both Rankl and Opg in the cortical bone and does not increase osteoclast formation and bone resorption. Finally, Pinch ablation exacerbates hindlimb unloading–induced bone loss and impairs active ulna loading–stimulated bone formation. Thus, we establish a critical role of Pinch in control of bone homeostasis.
architecture, while skeletal unloading leads to dramatic bone loss. Osteocytes are believed to be a main mechano-sensor of the bone. Osteocytes reside in the lacunae throughout the mineralized bone matrix and are well positioned to sense mechanical signals and transduce them to biochemical signals, which are eventually integrated into cellular responses that regulate bone remodeling (2–6, 13, 14). Osteocytes die through apoptosis (15). However, key signals that control osteocyte function and survival are poorly defined.

Mammalian cells have 2 functional PINCH proteins, PINCH1 (LIMS1) and PINCH2 (LIMS2). They are 5 LIM domain–containing proteins that play important roles in cytoskeletal organization and cell–extracellular matrix adhesion, migration, proliferation, and survival (16–21). Both PINCH1/2 are ubiquitously expressed in most mammalian tissues and organs, including the heart, lung, liver, kidney, and bladder (20). Global deletion of Pinch1 in mice causes lethality (17), while Pinch2–KO mice have no apparent phenotypes (19, 22). Pinch proteins exert their functions in part by forming several distinct functional protein-protein complexes, including the ILK-Pinch-parvin (IPP), Pinch-Nck2, and Pinch-Rsu1 complexes (18, 20, 23–26).

For example, the IPP complex regulates cell contractility and cytoskeletal dynamics in mice (21, 27–29). Previous studies were primarily focused on roles of Pinch in regulation of multiple properties of cancers, such as growth, survival, apoptosis, progression, invasion, and radio- and chemo-resistance (19, 20, 30–33).

Important roles of focal adhesion proteins in organogenesis have begun to emerge. We demonstrate that the focal adhesion protein Kindlin-2 — which interacts with integrin-linked kinase (ILK), part of the IPP complex — regulates chondrogenesis and early skeletal development by modulating TGF-β signaling and Sox9 expression in chondrocytes and their precursors (34). We further demonstrate that Kindlin-2 directly promotes osteoblastic, but inhibits adipogenic, differentiation of mesenchymal stem cells through control of Yap1/Taz (35). Liang and coworkers reported that Pinch proteins are critical for cardiomyocyte function and heart development in mice. Deletion of both Pinch1/2 in the heart resulted in severe dilated cardiomyopathy and death due to heart failure in mice (36). However, roles of Pinch1/2 in bone have not been defined.

In this study, we deleted Pinch1 expression using the mouse 10-kb Dmp1-Cre transgene, which primarily targets osteocytes and mature osteoblasts, with and without global Pinch2 deletion in mice. Through comprehensive analyses of cells and tissues of the double and single mutant mice, we establish a critical role of Pinch1/2 and a functional redundancy of both factors in control of bone homeostasis through distinct mechanisms.

Results

Pinch1flo/flo; Pinch2+/− (dKO), but not Pinch1flo/flo or Pinch2−/−, mice display a severe osteopenia. To investigate the roles of Pinch1/2 in bone, we generated mice lacking Pinch1 expression in mature osteoblasts and osteocytes by breeding the floxed Pinch1 mice (Pinch1Δflo/flo) with the 10-kb mouse dentin matrix protein 1–Cre (Dmp1-Cre) transgenic mice (Pinch1Δflo/flo), Pinch2-KO (Pinch2−/−), and Pinch1Δflo/flo; Pinch2−/− (double knockout; dKO) (Figure 1, A–D). Results from Western blotting, quantitative PCR (qPCR), and IHC staining analyses demonstrated that Pinch1 expression was significantly reduced in bone and in osteocytes embedded in the cortical bone matrix of dKO mice compared with control mice (Dmp1-Cre). To determine the effects of Pinch deletion on bone mass, we performed micro-computed tomography (μCT) analysis of distal femurs of 3-month-old female control (Dmp1-Cre), Pinch1Δflo/flo, Pinch2−/−, and dKO mice. The bone mineral density (BMD) and bone volume/tissue volume (BV/TV) fraction (BV/TV) were not significantly different among the first 3 groups. However, both parameters were dramatically decreased in dKO mice compared with age- and sex-matched control (Dmp1-Cre) or single mutant mice (Pinch1Δflo/flo or Pinch2−/−) (Figure 1, A–C). The cortical thickness (Ct.Th) was not altered by the double or single mutations (Figure 1D). Specifically, the values of BMD and BV/TV were decreased by 52% and 60%, respectively, in dKO mice compared with those of control mice. Consistent with results from μCT analyses, results from H&E staining of tibial sections of 6-month-old female mice showed much less trabecular bone in dKO mice than in controls (Figure 1E). We performed qPCR analysis using RNA from the osteocyte-enriched femoral drafts (with their BM flushed) from 6-month-old female control and dKO mice and observed that the level of Pinch1 mRNA was dramatically reduced in dKO relative to control femurs (Figure 1F). Similarly, results from Western blotting using protein extracts from femoral drafts revealed that the level of Pinch1 protein was decreased in dKO relative to control bones (Figure 1G). These reductions are specific because expression of Pinch1 in nonbone tissues, such as heart, lung, spleen and kidney, was not reduced in dKO versus control mice (Figure 1, F and G). Results from IHC staining of the longitudinal tibial sections of 6-month-old female control and dKO mice revealed fewer Pinch1-expressing osteocytes embedded in the
bone matrix in dKO bone than in control bones (Figure 1, H and I). The Dmp1-Cre (control), Pinch1Dmp1−/−, Pinch2−/−, and dKO mice were viable, fertile, and born with the expected Mendelian frequency. The body weights were not different between control and dKO mice (Figure 1J).

Pinch deletion causes severe osteopenia in 6- and 14-month-old mice. We next investigated the effects of Pinch deletion on bone homeostasis in greater detail. Because neither Pinch1Dmp1−/− nor Pinch2−/− mice displayed marked skeletal defects, we next focused our study on analyzing the phenotypes of dKO mice in comparison with Dmp1-Cre mice (controls). We harvested femurs from mice of the 2 genotypes of different ages and sexes. μCT analysis of distal femurs of those mice revealed dramatic decreases in BMD and BV/TV in dKO mice compared with control mice at different ages and sexes (Figure 2A). Specifically, BMD was decreased by 47% in 6-month-old and 45% in 14-month-old female dKO mice (Figure 2B), and BV/TV was reduced by 60% at both ages in dKO mice compared with those in sex-matched control littermates (Figure 2C). The trabecular number (Tb.N) was significantly decreased, while the trabecular separation (Tb.Sp) was increased in dKO mice relative to control littermates (Figure 2, D and E). In contrast, the trabecular thickness (Tb.Th) and Ct.Th were not significantly reduced in dKO mice compared with those in control mice (Figure 2, D–G). A similar severe osteopenia was observed in 6-month-old male dKO mice (Figure 2, A–G).

Pinch loss impairs the mechanical properties of dKO femurs. We next performed the 3-point bending tests on femurs of 4-month-old control and dKO female mice to determine whether Pinch loss impairs the mechanical properties of the cortical bones. The results revealed that the values for both stiffness and maximum load were significantly reduced in dKO versus control mice (Figure 2, H and I).

Taken together, these results demonstrate an important role of Pinch1/2 and a functional redundancy of the 2 factors in control of bone mass and function.

Reduced osteoblast formation and function contribute to the osteopenia in dKO mice. To investigate mechanisms through which Pinch loss causes osteopenia, we performed several sets of experiments. We measured the serum level of procollagen type 1 amino-terminal propeptide (P1NP), an in vivo indicator for osteoblast function and bone formation, and found that it was dramatically reduced in 6-month-old dKO mice compared with that of control mice (Figure 3A). We determined the in vivo bone-forming activity by performing a calcein double-labeling experiment and found that the mineralization apposition rate (MAR) and bone formation rate (BFR) in the tibial metaphyseal cancellous bones were significantly reduced in 3-month-old female dKO mice compared with those in control mice (Figure 3, B–D). To determine the effects of Pinch loss on the osteoid production and mineralization, we performed the von Kossa staining of tibia sections of 3-month-old female control and dKO mice and found that the osteoid volume/tissue volume (OV/TV) and mineralized bone volume/tissue volume (mBV/TV) were significantly decreased in dKO mice relative to those in control mice (Figure 3, E–G). Results from Western blotting analyses using tibial protein extracts revealed reduced expression of Osx protein, a marker for osteoblasts and their precursors in dKO relative to control bones (Figure 3, H and I).

Pinch deletion does not increase osteoclast formation and bone resorption. We further investigated whether Pinch loss impacts the osteoclast formation and bone resorption. We first performed the tartrate-resistant acid phosphatase (TRAP) staining of tibial sections for osteoclast formation in bone and found that the osteoclast number/bone perimeter (Oc.Nb/BPm), but not osteoclast surface/bone surface (Oc.S/BS), was increased in both primary and secondary spongiosa bones in dKO mice compared with those of controls; however, the increases did not achieve a statistical significance (Figure 4, A–E). Furthermore, the serum levels of the collagen type I cross-linked C-telopeptide (CTX1), degradation products from type I collagen during in vivo osteoclastic bone resorption, were not significantly altered in 6-month-old female dKO mice relative to control mice, as...
measured by ELISA assay (Figure 4F). Results from Western blotting showed that the level of secreted Rankl was increased in dKO cortical bones relative to that in control cortical bones (Figure 4G). Results from IHC staining revealed increased expression of Rankl protein in dKO osteocytes relative to control osteocytes in bones (Figure 4H). Results from reverse transcription PCR (RT-PCR) analyses showed that the levels of both Rankl and Opg mRNA were similarly upregulated in dKO versus control bones (Figure 4I).

Pinch deletion increases sclerostin expression in osteocytes embedded in the cortical bone matrix. Sclerostin, the Sost gene protein product almost exclusively expressed by osteocytes (8), binds to the Wnt coreceptors Lrp5 and Lrp6 and inhibits the Wnt/β-catenin signaling, osteoblast function, bone formation (37, 38). Interestingly, results from Western blotting revealed that expression of sclerostin was increased in dKO bones compared with that of control bones (Figure 5, A and B). Results from IHC staining confirmed the upregulation of sclerostin in osteocytes embedded in dKO cortical bone matrix (Figure 5C).

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**Table 1**

<table>
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<tr>
<th>Group</th>
<th>BMD(g/cm²)</th>
<th>BV/TV (%)</th>
<th>Tb.N(1/mm)</th>
<th>Tb.Sp(μm)</th>
<th>Tb.Th(μm)</th>
<th>Stiffness(N/mm)</th>
<th>Ultimate Force(N)</th>
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<td>43.02</td>
<td>70.15</td>
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<td>343.7</td>
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**Figure 2. Pinch loss causes severe osteopenia in 6- and 14-month-old male and female mice.**

(A) Three-dimensional (3-D) reconstruction from micro-computerized tomography (μCT) scans of distal femurs from 6- and 14-month-old female and 6-month-old male control and dKO mice. Scale bar: 100 μm. (B–G) Quantitative analyses of bone mineral density (BMD), bone volume/tissue volume (BV/TV), trabecular number (Tb.N), trabecular thickness (Tb.Th), trabecular separation (Tb.Sp), and cortical thickness (Ct.Th) of distal femurs from A. n = 6 mice per group. *P < 0.05, **P < 0.01, ***P < 0.001 vs. controls. Unpaired Student’s t test. Results are expressed as mean ± SD. (H and I) Three-point bending tests. Femurs isolated from 4-month-old control and dKO female mice were subjected to the 3-point bending test as described in Materials. Stiffness (H) and maximum load (I). n = 8 mice per group. **P < 0.01, ***P < 0.001 vs. controls. Unpaired Student’s t test. Results are expressed as mean ± SD.
Conditioned media from dKO bone slice cultures contain abundant sclerostin protein and suppresses osteoblast differentiation in primary BMSC and calvarial osteoblast cultures. Although above results showed increased expression of sclerostin in osteocytes of dKO bones, we did not observe a significant increase in serum sclerostin in dKO mice compared with control mice (Figure 5D). We next prepared the conditioned media (CM) from ex vivo–cultured middiaphyseal femoral and tibial slices from 6-month-old control and dKO mice and observed a dramatic increase in sclerostin in dKO CM compared with control CM (Figure 5E). Primary BM stromal cells (BMSCs) were isolated from normal C57BL/6 mice and treated with the CM under osteoblast differentiation condition for 7 days. The results showed that...
Figure 4. Pinch loss does not impair osteoclast formation and bone resorption. (A–E) Tartrate-resistant acid phosphatase (TRAP) staining. Tibial sections of 6-month-old female control and dKO mice were used for TRAP staining (A). Osteoclast surface/bone surface (Oc.S/BS) (B and D) and osteoclast number/bone perimeter (Oc.N/BPm) (C and E) of primary and secondary spongiosa bones of the 2 genotypes were measured using Image-Pro Plus 7.0. Scale bar: 50 μm. n = 8 mice per group. Unpaired Student's t test. Results are expressed as mean ± SD. (F) Serum level of collagen type I cross-linked C-telopeptide (CTX1). Sera were collected from 6-month-old female control and dKO mice and subjected to ELISA assay for CTX1. n = 8 mice per group. Unpaired Student's t test. Results are expressed as mean ± SD. (G) Western blot analysis. Protein extracts were isolated from osteocyte-enriched middiaphyseal femoral shafts (with their BM flushed) of 6-month-old female control and dKO mice and subjected to Western blotting using an anti-Rankl antibody. Western blotting was repeated 3 times. (H) IHC staining of tibial sections of 6-month-old female control and dKO mice with an anti-Rankl antibody. Scale bar: 50 μm. (I) qPCR analyses. Total RNAs were isolated from femurs of 6-month-old female control and dKO mice and used for qPCR analysis for expression of Rankl and Opg genes, which were normalized to Gapdh mRNA. n = 3 mice per group. *P < 0.05 vs. controls. Unpaired Student’s t test. Results are expressed as mean ± SD.
Figure 5. Pinch loss increases sclerostin expression in osteocytes and indirectly suppresses osteoblast differentiation. (A) Western blot analyses. Protein extracts were isolated from femurs of 6-month-old female control and dKO mice and subjected to Western blotting using an antibody against sclerostin. (B) Quantitative data. n = 3 mice per group. Unpaired Student’s t test. Results are expressed as mean ± SD. (C) IHC staining of tibial sections of 6-month-old female control and dKO mice with an antibody against sclerostin. Scale bar: 50 μm. (D) Serum level of sclerostin. Sera were obtained from 6-month-old female control and dKO mice and subjected to ELISA assay for sclerostin. n = 8 mice per group. Unpaired Student’s t test. Results are...
expressed as mean ± SD. (E) Western blot analysis. Mid diaphyseal femoral shafts (with their BM flushed) of 6-month-old female control and dKO mice were cut into small pieces and cultured in serum-free α-MEM supplemented with 1% penicillin/streptomycin for 48 hours and conditioned media (CM) for 48 hours, followed by Western blotting with an antibody against sclerostin. (F–J) Primary BM stromal cells (BMSCs) were isolated from normal C57BL/6 mice and treated with control and mutant CM from 3-day-old normal C57BL/6 mice and treated with control and mutant CM from E in osteoblast differentiation media (α-MEM containing 10% FBS, 1% penicillin/streptomycin, 50 μg/mL ascorbic acid, and 5 mM β-glycerophosphate) for 7 days, followed by qPCR analyses for expression of indicated mRNAs, which were normalized to Gapdh mRNA (F) or Western blotting using indicated antibodies (G and I). Quantitative data for G (H) and I (J). (K–N) Primary calvarial osteoblasts were isolated from 3-day-old normal C57BL/6 mice and treated with control and mutant CM in osteoblast differentiation media for 48 hours, followed by Western blotting using indicated antibodies (K) or alkaline phosphatase (ALP) staining (M and N). Quantitative data of K (L). Scale bar: 100 μm. *P < 0.05, **P < 0.01, ***P < 0.001 vs. controls. Unpaired Student’s t test. Results are expressed as mean ± SD. Western blotting analyses in this figure were repeated 3 times.

To investigate whether Pinch loss in osteocytes and mature osteoblasts impacts the BM cells, we prepared fresh BM cells of the 2 genotypes and performed colony forming unit-fibroblast (CFU-F) assays and observed reduced number of CFU-F colony in dKO relative to the control group (Figure 6A). Furthermore, Pinch loss also drastically reduced the number of CFU-osteoblast (CFU-OB) assays (Figure 6B). We determined the in vitro proliferation capacity of primary BMSCs from control and dKO mice by performing the CCK8 assay and observed a reduced proliferation rate in dKO versus control cells (Figure 6C), which may contribute to the reduced formation of both CFU-F and CFU-OB in dKO mice.

Pinch deletion increases BM adiposity, and primary BMSCs from dKO mice display decreased osteoblastic, but increased adipogenic differentiation capacity. Interestingly, we found that primary BMSCs from dKO mice displayed reduced expression of osteoblast differentiation genes, including Runx2, Osterix, Col1a1, and ALP activity (Figure 6, D–F). In contrast, dKO BMSCs exhibited increased expression of adipogenic differentiation genes — including those encoding peroxisome proliferator-activated receptor γ (Ppara), adipocyte Protein 2 (aP2), and C/EBPα, but not preadipocyte factor-1 (Pref-1) or adiponectin (Figure 6, G and H) — and enhanced adipogenic differentiation, as evidenced by Oil Red O staining (Figure 6I). Results from H&E staining of tibial sections of 6-month-old female control and dKO mice showed dramatically increased adiposity in dKO BM (Figure 6J). Finally, the expression levels of Yap1 and Taz proteins and mRNAs, which were reported to play a role in controlling the MSC fate (35), were drastically decreased in dKO versus control BMSC cultures (Figure 6C, K–M).

Pinch ablation decreases integrin β3, ILK, and α-parvin; increases active caspase-3 and -8 expression; and promotes osteocyte apoptosis. We further performed the TUNEL staining of tibial cross sections of 6-month-old female control and dKO mice and found that apoptotic osteocytes were scarcely detected in control cortical bones; the number of apoptotic osteocytes was significantly increased in dKO cortical bones (Figure 7, A–C). We isolated primary osteocytes for flow cytometry analysis to further assess cell apoptosis. To confirm that the isolated cells from cortical bone were Dmp1+ osteocytes, we performed immunofluorescence staining using an anti-Dmp1 antibody. The result showed that almost all cells were Dmp1+ cells (Figure 7D). Results from flow cytometry analysis of those cells revealed increased apoptosis in dKO cultures compared with that of control cultures (Figure 7E). Consistently, results from Western blotting revealed that the levels of active caspase-8 and caspase-3 proteins, but not their total protein levels, were increased in dKO relative to control osteocytes (Figure 7, F and G). The levels of several Pinch-related proteins, including the ILK, integrin β3, and α-parvin, were decreased in dKO bones compared with those of control bones (Figure 7H).

Pinch deficiency exacerbates hindlimb unloading–induced bone loss and impairs active ultra loading–induced bone formation. Skeleton throughout life is constantly loaded by mechanical forces generated by daily physical activities and gravity. Osteocytes are believed to be the main mechano-sensors of bone. Whereas increased mechanical forces promote bone formation, decreased loading causes bone loss and skeletal fragility. The severe osteopenia of dKO mice prompted us to investigate whether Pinch expression in Dmp-1–expressing cells is involved in mediation of bone mechanotransduction. We first utilized a well-established tail suspension mouse model to create a condition of hindlimb unloading (HLU) as previously described by Robling et al. (39).
Three-month-old female control and dKO mice were subjected to HLU for 21 days, followed by µCT analysis of distal femur. The results showed that HLU induced significant bone loss in both control and dKO mice, as demonstrated by significant decreases in BV/TV, Tb.N, and Ct.Th and an increase in Tb.Sp induced by HLU in both control and dKO mice (Figure 8, A–F). However, the magnitudes in the HLU-induced decreases in BV/TV and Tb.N were increased in dKO mice compared with those in control mice (P < 0.05, Con-HLU/Con versus dKO-HLU/dKO) (Figure 8, B and C). These results suggest that Pinch is probably involved in mechanical unloading–induced bone loss. Next, we utilized an active ulna loading model to determine whether Pinch is required for mechanical loading. In this experiment, 3-month-old female control and dKO mice were subjected to active mechanical loading as previously described (40). µCT analysis of ulnas of control and dKO mice with and without loading revealed that the Ct.Th was increased by loading in control ulna; this increase was lost in dKO ulna (P < 0.05, Con-load/Con versus dKO-load/dKO) (Figure 8G). Results from the calcine double-labeling experiments showed that, although both control and dKO ulnas responded to loading by increasing MAR and BFR, the magnitudes stimulated by loading were decreased by Pinch loss (P < 0.05, Con-load/Con versus dKO-load/dKO) (Figure 8, H–J).

Discussion

In this study, we establish a critical role of Pinch proteins, through expression in mature osteoblasts and osteocytes, in regulation of bone mass and mechanotransduction in mice. We also demonstrate a functional redundancy of Pinch1/2 in control of bone mass. We define distinct cellular and molecular mechanisms through which Pinch regulates bone homeostasis. The findings of this study may provide a potential therapeutic target for the prevention and treatment of metabolic bone diseases, such as osteoporosis.

In this study, we establish that Pinch expression in osteocytes and mature osteoblasts plays a critical role in control of osteoblast function and bone formation. This notion is supported by the following lines of experimental evidence: (a) Pinch loss reduces the serum level of P1NP, which is a bona fide bone formation marker; (b) results from our calcine double-labeling experiments provide direct evidence that the osteoblast-mediated bone forming activity is impaired by Pinch loss, as demonstrated by dramatic decreases in MAR and BFR in dKO versus control mice; (c) reduced osteoid production and deposition and mineralization further point to an impairment in osteoblast number and/or function in dKO mice; and (d) Pinch loss decreases expression of Osx, a marker for preosteoblasts and osteoblasts, in bones. It should be noted that Pinch loss does not markedly impact the Rankl/Opg ratio, osteoclast formation, and bone resorption. Thus, impaired osteoblast formation and function primarily contributes to the osteopenia in mutant mice. We provide convincing evidence that Pinch loss greatly increases expression of sclerostin in osteocytes, which contributes, at least in part, to the impaired osteoblast formation and function in mutant mice. Indeed, Pinch loss largely increases sclerostin expression in osteocytes embedded in the bone matrix. Of particular interest, CM from ex vivo cultured mutant bones contain abundant sclerostin and decrease the expression of β-catenin, Runx2, Osx, Colla1, and Alp in primary BMSCs and calvarial osteoblasts. It is interesting to note that the level of sclerostin in circulation is not increased in mutant mice; this suggests that Pinch regulates the level of sclerostin primarily in the local bone microenvironment and controls osteoblast function and bone formation. Given the important role of sclerostin in control of bone mass and the fact that osteocytes are a main source of sclerostin (8, 41), results of the present study demonstrating that...
Pinch proteins function as a critical inhibitory signal for sclerostin expression in osteocytes represents an important finding in osteocyte biology.

Pinch loss in mature osteoblasts and osteocytes indirectly impacts the formation of OFU-F and OFU-OB, cell proliferation, osteoblast gene expression, and differentiation in primary BMSC cultures. These results suggest that the BMSCs are influenced before in vitro experiments in the bone microenvironment by signals initiated by Pinch loss in osteocytes and mature osteoblasts. In addition to reduced osteogenesis, BM and BMSCs of mutant mice display increased adipogenesis. Since there is a discrepancy in literature regarding the role of Yap1/Taz in regulation of MSC differentiation (35, 42), whether or not the reduced Yap1/Taz expression in dKO BMSCs plays a role for the altered BMSC differentiation capacity of those cells remains to be determined.

We utilize 2 unique mouse models to demonstrate an involvement of Pinch in mediation of bone mechanotransduction. We show that Pinch loss exacerbates the HLU-induced bone loss. We further use an active ulna loading mouse model and demonstrate a requirement of Pinch in active ulna loading–induced bone formation in vivo. Batra and coworkers showed that (a) mechanical stress-activated integrin α5β1 induces opening of connexin 43 (Cx43) hemichannels (43); (b) AKT activation by fluid flow shear stress directly phosphorylates Cx43 and integrin α5; and (c) Ser-373 of Cx43 plays a predominant role in mediating the interaction between these 2 proteins and Cx43 hemichannel opening, a crucial step to mediate the anabolic function of mechanical loading in bone (44). A more recent study showed that IGF1 assembles the formation of a complex between IGF1R and the integrin β3 subunit to mediate mechanical stimuli in osteoblastic cells (45). Our results of the present study reveal that Pinch loss reduces expression of integrin β3 and 2 integrin-related proteins, ILK and α-parvin, in osteocytes. Thus, it is likely that Pinch mediates bone mechanotransduction by modulating, in part, the integrin signaling in osteocytes. Future study will explore this possibility.

Our results suggest that Pinch proteins protect osteocytes from apoptosis and Pinch loss promotes osteocyte apoptosis associated with increased expression of cleaved caspase-3 and -8.

Liang and coworkers demonstrate that ablation of either Pinch1 or Pinch2 in cardiomyocytes in mice does not cause obvious defects in the heart structure and function at the basal level. However, loss of either of the 2 proteins deteriorates myocardial infarction induced by coronary artery ligation. Deletion of both Pinch1/2 in the heart in mice result in severe dilated cardiomyopathy and death due to heart failure (36). In the present study, we demonstrate a functional redundancy of Pinch1/2 in control of bone mass.

In summary, we establish critical roles for Pinch1/2 in regulation of osteocyte survival and function, bone homeostasis, and bone mechanotransduction, and findings of the present study emphasize the importance of investigating the role of Pinch proteins in human metabolic bone diseases.

Methods

Animal studies. Dmp1-Cre mice, in which the 10-kb mouse dentin matrix protein 1 (Dmp1) gene promoter drives Cre expression in mature osteoblasts and osteocytes, were previously described (46). Pinch1fl/fl and Pinch2–/– mice were previously described (36). To obtain the double mutant mice, Pinch1fl/fl and Pinch2–/– mice were first crossed with Pinch2–/– mice to generate Pinch2–/– mice. Pinch1fl/fl; Pinch2–/– mice were then crossed to each other and generate the Pinch1fl/fl Pinch2–/– mice. Hemizygous Dmp1-Cre transgenic mice were crossed with Pinch1fl/fl Pinch2–/– mice to generate Dmp1-Cre, Pinch1fl/fl; Pinch2–/– mice. Finally, Dmp1-Cre, Pinch2–/–, Pinch2–/– mice were bred with Pinch1fl/fl, Pinch2–/– mice to generate Dmp1-Cre, Pinch1fl/fl, Pinch2–/– and other genotypes. All mice used in this study have been crossed with normal C57BL/6 mice for at least 10 generations. All animal experiments were conducted in the specific pathogen free (SPF) Experimental Animal Center of SUSTech. Animals were housed 4/cage at 20°C–24°C, exposed to a 12-hour/12-hour light-dark period, and given water and rodent chow ad libitum. Mice were monitored and weighed daily. μCT analysis,
HLU, and ulna loading are described in detail below. Mouse age and sex and numbers of mice used for each experiment are specified in the figure legends.

HLU experiment. Three-month-old female control and dKO mice were treated with or without tail suspension for 21 days as previously described (39). Mice were euthanized at day 21 by CO₂ asphyxiation. Femurs from each group were subjected to μCT analysis as described below.

Ulna loading model. In vivo ulna mechanical loading was performed under anesthesia using a Biomaterial mechanical system Electroforce 3230 (TA Instrument) as described (40). Three-month-old female control and dKO mice commenced unilateral cyclic axial compression of the left ulna at day 1, day 4, day 7, day 10, and day 13. Briefly, 900 cycles were applied at a rate of 4 Hz with a peak force of 2.3 N for 3.75 minutes. Right ulnas were used as contralateral nonloaded controls. Calcein was injected at day 3 and day 10. Mice were euthanized at day 14 by CO₂ asphyxiation.

Three-point bending tests. Femurs isolated from 4-month-old control and dKO female mice were placed in a fixture attached to a materials testing machine and loaded until broken as described (47). This assay primarily tests the mechanical properties of the mid-diaphysis, which is typically all cortical bone.

Histological evaluation and bone histomorphometry. For histology, bone tissues were fixed in 10% formalin at 4°C for 24 hours, decalcified in 10% EDTA (pH 7.4) for 10–14 days, and embedded in paraffin.
Bone sections were used for H&E staining using our standard protocol. Fixed nondecalcified bone slices were used for μCT analysis in the Department of Biology of SUSTech using a Bruker CT (SkyScan 1172 Micro-CT, Bruker MicroCT) following the standards of techniques and terminology recommended by the American Society for Bone and Mineral Research (ASBMR) (48). For trabecular bone parameters, transverse slices were obtained in the region of interest in the axial direction from the trabecular bone 0.5 mm below the growth plate (bottom of the primary spongiosa). Contours were defined and drawn close to the cortical bone. The trabecular bone was then removed and analyzed separately. A 3-dimensional analysis was performed on 200 trabecular bone slices. A 0.5-mm section was used to obtain midlateral cortical bone thickness. The analysis of the specimens involved the following bone measurements: BMD, BV/TV, Tb.N, Tb.Th, Tb.Sp, and Ct.Th.

**Calcein double labeling, MAR, and BFR.** Calcein double labeling, MAR, and BFR were measured as we previously described (49). Briefly, mice were injected i.p. with calcein (20 mg/kg) at days 6 and 2 before sacrifice. Nondecalcified bone slices were imaged using an Envision+System-HRP (DAB) kit (Dako North America Inc.) according to the manufacturer’s instruction. For IF, sections were stained with antibodies or control IgG using the EnVision+System-HRP (DAB) kit (Dako North America Inc.) according to the manufacturer’s instruction. Calcein labeling was visualized using EVOS fl microscopy (Invitrogen). Metaphyseal trabecular bone sections were then incubated into the BM area and diaphyseal cortical bones were evaluated, and the distance between the double-labeled bands was measured using Image Pro Plus 7.0 software. The MAR was calculated as the mean distance between the double labels divided by the number of the days between the calcein injections. The mineralizing surface per bone surface (MS/BS) was calculated as the percentage of bone surface exhibiting mineralizing activity. MS/BS = (dLS + sLS/2)/BS, where dLS is the number of double-labeled surfaces and sLS is the number of single-labeled surfaces. The BFR per bone surface (BFR/BS or BFR) is the volume of mineralized bone formed per unit time and per unit bone surface. BFR/BS is calculated as BFR/BS = MAR × (MS/BS).

**IHC and immunofluorescence (IF).** IHC staining was performed as we previously described (34). Briefly, 5-μm sections were stained with antibodies or control IgG using the EnVision+System-HRP (DAB) kit (Dako North America Inc.) according to the manufacturer’s instruction. For IF, cells were seeded and cultured on sterile glass coverslips in 6-well plates. Twenty-four hours later, cells were fixed in 10% formaldehyde for 30 minutes at 37°C. Cells were permeabilized with 0.2% Triton X-100 containing DAPI for 5 minutes and blocked with 2% BSA (MilliporeSigma, A1933) for 1 hour. The cells were then stained with a primary antibody overnight at 4°C. After washing, cells were incubated with anti-mouse Alexa Fluor 594 (Invitrogen) secondary antibodies (1:300) for 1 hour at room temperature. Cells were then imaged using a confocal microscope (SP2-AOBS Leica Microsystems).

**TUNEL staining.** Cell survival was evaluated using the ApopTag Peroxidase In Situ Apoptosis Detection Kit according to the manufacturer’s instruction (EMD Millipore, S7100) as we previously described (34).

**Primary BMSC culture and CFU-F and CFU-OB assays.** Primary BMSC were isolated from tibiae and femurs of 6-month-old female mice as previously described (49). CFU-F and CFU-OB assays were performed as we previously described (50). Briefly, for the CFU-F assays, BM nucleated cells from 5-month-old female control and dKO mice were seeded in a 6-well plate with a cell density of 2 × 10^6 per well and cultured using the Mouse MesenCult proliferation kit (CFU-F assay) for 14 days, followed by Giemsa staining. For the CFU-OB assays, BM nucleated cells from 5-month-old female control and dKO mice were seeded in a 6-well plate at a density of 4 × 10^4 per well and cultured in osteoblast differentiation media (complete α-MEM containing 50 μg/mL L-ascorbic acid and 2.0 mM β-glycerophosphate) for 21 days (media were changed every 48 hours), followed by alizarin red staining.

**Primary calvarial osteoblasts and osteocyte cultures.** Primary calvarial osteoblasts were isolated from 3-day-old WT C57 mice. Briefly, calvariae were isolated, and soft tissues were removed as much as possible. Calvariae were then rinsed with cold 1× PBS 3 times, cut into small pieces, and digested with trypsin for 37°C for 30 minutes. The calvariae were rinsed and further digested with 1 mg/mL of collagenase IA for 60 minutes. Cells were harvested by centrifugation, resuspended, and filtered through a Mesh Of 70 Mesh Filters (Corning, 431751). Cells were cultured in low-glucose DMEM with 10% FBS (Thermo Fisher Scientific, 16000044) and 1% penicillin/streptomycin (HyClone, SV30010). For isolation of primary osteocytes from cortical bones, we utilized a protocol that was provided by Lynda Bonewald of the Indiana University School of Medicine (Indianapolis, Indiana, USA). Briefly, middiaphyseal femurs and tibiae of 9-month-old control and dKO mice were cut into small pieces and digested with collagenase IA at 37°C for 30 minutes 3 times and then alternately digested with 5 mM EDTA and collagenase IA at 37°C for 30 minutes 6 times. Cells from the last digestion were collected...
for experiments. Cells were then cultured in α-MEM (HyClone, SH30265.01) supplemented with 10% FBS (Thermo Fisher Scientific, 16000044) and 1% penicillin/streptomycin (HyClone, SV30010).

Osteoclast formation in bone. The TRAP staining of bone sections was performed as we previously described (51). The Oc.S/BS and osteoclast numbers/bone perimeter (Oc.N/BPm) in both primary and secondary spongiosa of tibiae were measured using Image Pro Plus 7.0 software as we previously described (50).

Preparation of conditional media from ex vivo bone tissue cultures. Tibiae and femurs were isolated from 6-month-old control and dKO mice, and soft tissues were removed. Both ends were cut, and BM was flushed. The remaining cortical bones were cut into small pieces and were cultured in serum-free α-MEM media for 48 hours, and CM were collected.

ELISA assays. Serum levels of P1NP were measured using the RatLaps EIA Kit (Immunodiagnostic Systems Limited, catalog AC-33F1) according to the manufacturer’s instruction. Serum levels of CTX-1, degradation products from type I collagen during osteoclastic bone resorption, were measured using the RatLaps EIA Kit (Immunodiagnostic Systems Limited, catalog AC-06F1) as previously described (51).

qPCR and Western blotting analyses. Total RNAs were isolated using the RNeasy Mini Kit (QIAGEN, catalog 74104) according to the manufacturer’s instruction. RT was performed using 1 μg denatured RNA and 100 pmol of random hexamers (Applied Biosystems) in a total volume of 25 μL containing 12.5 U MultiScribe reverse transcriptase (Applied Biosystems) as previously described (52). qPCR analysis was performed to measure the relative mRNA levels using the SYBR Green kit (Bio-Rad) as previously described (52). Samples were normalized to Gapdh expression. The DNA sequences of primers used for qPCR are summarized in Supplemental Table 1 (supplemental material available online with this article; https://doi.org/10.1172/jci.insight.131692DS1). Western blot analysis was performed as previously described (53). Briefly, protein extracts were fractionated on a 10% SDS-PAGE gel and transferred onto nitrocellulose membranes (Schleicher & Schuell). The membrane was blocked in 5% nonfat milk in Tris-buffered saline/Tween 20 buffer; probed with primary antibodies, followed by incubation with secondary antibodies conjugated with horseradish peroxidase; and visualized using a Western Blotting Detection Kit (GE Healthcare, catalog RPN2106). Antibodies used in this study are listed in Supplemental Table 2.

Statistics. The sample size for each experiment was determined based on our previous experience. Animals used in experiments of this study were randomly grouped. IHC, IF, and histology were performed and analyzed in a double-blinded manner. Statistical analyses were completed using the Prism GraphPad. Unpaired 2-tailed Student’s t test (2 groups) and 2-way ANOVA (multiple groups) were used. P < 0.05 was considered statistically significant.

Study approval. All animal studies were approved by the IACUC of SUSTech.

Author contributions
GX, HC, and YW designed the study. YW, QY, YZ, XL, SL, PZ, LM, and YL conducted the study and collected data. YW, QY, and HC analyzed data. GX, HC, XB, CW, YW, CL, JQF, and DC interpreted data. GX, HC, and YW drafted the manuscript. GX, HC, and YW take the responsibility for the integrity of the data analysis.

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