

# NFATc1 in mice represses osteoprotegerin during osteoclastogenesis and dissociates systemic osteopenia from inflammation in cherubism

Antonios O. Aliprantis,<sup>1,2</sup> Yasuyoshi Ueki,<sup>3</sup> Rosalyn Sulyanto,<sup>1,4</sup> Arnold Park,<sup>1</sup> Kirsten S. Sigrist,<sup>1</sup> Sudarshana M. Sharma,<sup>5</sup> Michael C. Ostrowski,<sup>5</sup> Bjorn R. Olsen,<sup>3</sup> and Laurie H. Glimcher<sup>1,2</sup>

<sup>1</sup>Department of Infectious Diseases and Immunology, Harvard School of Public Health, Boston, Massachusetts, USA.

<sup>2</sup>Division of Rheumatology, Allergy and Immunology, Department of Medicine, Brigham and Women's Hospital, Harvard Medical School, Boston, Massachusetts, USA. <sup>3</sup>Department of Developmental Biology and <sup>4</sup>Harvard School of Dental Medicine, Boston, Massachusetts, USA.

<sup>5</sup>Department of Molecular and Cellular Biochemistry and Comprehensive Cancer Center, The Ohio State University, Columbus, Ohio, USA.

**Osteoporosis results from an imbalance in skeletal remodeling that favors bone resorption over bone formation. Bone matrix is degraded by osteoclasts, which differentiate from myeloid precursors in response to the cytokine RANKL. To gain insight into the transcriptional regulation of bone resorption during growth and disease, we generated a conditional knockout of the transcription factor nuclear factor of activated T cells c1 (*Nfatc1*). Deletion of *Nfatc1* in young mice resulted in osteopetrosis and inhibition of osteoclastogenesis in vivo and in vitro. Transcriptional profiling revealed NFATc1 as a master regulator of the osteoclast transcriptome, promoting the expression of numerous genes needed for bone resorption. In addition, NFATc1 directly repressed osteoclast progenitor expression of osteoprotegerin, a decoy receptor for RANKL previously thought to be an osteoblast-derived inhibitor of bone resorption. “Cherubism mice”, which carry a gain-of-function mutation in SH3-domain binding protein 2 (*Sh3bp2*), develop osteoporosis and widespread inflammation dependent on the proinflammatory cytokine, TNF- $\alpha$ . Interestingly, deletion of *Nfatc1* protected cherubism mice from systemic bone loss but did not inhibit inflammation. Taken together, our study demonstrates that NFATc1 is required for remodeling of the growing and adult skeleton and suggests that NFATc1 may be an effective therapeutic target for osteoporosis associated with inflammatory states.**

## Introduction

Osteoporosis, the most common pathologic condition resulting from excessive bone destruction, afflicts 10 million Americans over the age of 50 and results in 1.5 million fractures annually (1, 2). Congenital and acquired inflammatory conditions, such as cherubism and RA, cause regional bone loss leading to skeletal deformities (3, 4). A scarcity of molecular targets in bone hinders the development of therapeutics for these diseases. Moreover, an inducible genetic system to terminate bone resorption in mice would facilitate the analysis of preclinical models of bone loss.

Osteoclasts are multinucleated, bone resorbing cells that differentiate from myeloid precursors in response to M-CSF and RANKL. Mice deficient in *Csf1* (M-CSF), *Tnfrsf11* (RANKL), or either of their receptors (*Csf1r* [c-FMS] or *Tnfrsf11a* [RANK]), have increased bone density (osteopetrosis) with an absence of osteoclasts (5, 6). A decoy receptor for RANKL, TNFRSF11B (also known as osteoprotegerin [OPG]), balances osteoclastogenesis in vivo (7, 8). Although multiple tissues express OPG,

most research has focused on osteoblasts as the physiologic source of OPG in the skeletal system (9).

Nuclear factor of activated T cell (NFAT) proteins are a family of 5 inducible transcription factors (NFATc1–4 and NFAT5) that control numerous cellular differentiation processes, including immune cell activation, heart valve formation, and the growth of cartilage and stem cells (10–12). NFATc1 is a target of RANK signaling. During osteoclastogenesis, NFATc1 expression is induced and this transcription factor can be identified at the promoters of osteoclast specific genes (13–19). Studies interrogating NFATc1 in the skeletal system have been limited since *Nfatc1*-deficient mice die at E13.5 of cardiac valve defects (20, 21). The importance of NFATc1 in osteoclastogenesis in vivo was confirmed through fetal liver complementation experiments (22) and by rescuing embryonic lethality by targeting expression of NFATc1 to the developing heart (23). However, a tractable system to explore the role of NFATc1-mediated osteoclastogenesis in pathologic bone destruction was lacking. Moreover, a source of NFATc1-deficient osteoclast progenitors was not available, hindering discovery of the universe of NFATc1-regulated genes.

Cherubism is a rare, autosomal dominant, pediatric disorder characterized by fibroinflammatory infiltrates in the face, leading to disfiguring swelling and cystic changes in the jaw. Large numbers of multinucleated giant cells that express osteoclast markers are found within the lesion (24). Mutations leading to cherubism were mapped to a signaling adapter molecule, SH3-domain binding protein 2 (*SH3BP2*) (25, 26), which may promote NFAT activation (27).

**Nonstandard abbreviations used:** BMOcP, BM-derived CD11b<sup>lo</sup>/CD3e<sup>+</sup>B220<sup>+</sup>c-kit<sup>+</sup>c-fms<sup>+</sup> osteoclast precursor; FACS, fluorescence-activated cell sorting; *Ki*, knock-in; *LysM-Cre*, lysozyme-M-Cre; MROcP, M-CSF- and RANKL-stimulated BMOcP; NFAT, nuclear factor of activated T cells; OPG, osteoprotegerin; poly I:C, polyinosinic-polycytidylic acid; Sh3bp2, SH3-domain binding protein 2; TRAP, tartrate-resistant acid phosphatase.

**Conflict of interest:** Laurie H. Glimcher has equity in and is on the corporate board of directors of the Bristol-Myers Squibb Company.

**Citation for this article:** *J. Clin. Invest.* 118:3775–3789 (2008). doi:10.1172/JCI35711.



Recently, a mouse model of cherubism was developed (3). These knock-in (*KI*) mice contain a missense mutation in *Sh3bp2*, resulting in a proline to arginine substitution (P416R), the most commonly identified mutation in cherubism (P418R in humans). Homozygous *KI/KI* mice exhibit multiorgan inflammation and severe systemic bone loss that is dependent on TNF- $\alpha$  derived from myeloid cells. In mouse models and humans, TNF- $\alpha$  has been implicated in the systemic bone loss associated with inflammatory conditions like RA and postmenopausal estrogen withdrawal states (28). Mutant SH3BP2 enhances TNF- $\alpha$  production in macrophages and promotes osteoclastogenesis via ERK and spleen tyrosine kinase pathways, respectively (3). Since NFATs can promote both osteoclastogenesis (22) and TNF- $\alpha$  production (29, 30), here we have asked which aspects of the cherubism phenotype depend on NFATc1?

Using a conditional knockout strategy, we show that genetic ablation of *Nfatc1* results in osteoclast-poor osteopetrosis. This result complements previous studies (22, 23) and extends the requirement of NFATc1 to bone remodeling in the growing and adult skeleton. Molecular profiling of highly purified osteoclast precursors reveals 2 families of NFATc1-regulated genes in osteoclasts: those that are augmented by NFATc1 and those that absolutely depend on this transcription factor. Remarkably, in the absence of NFATc1, RANKL stimulates the expression of OPG, identifying a previously unrecognized function for NFATc1 in this pathway. Last, we show that NFATc1 functions downstream of mutant SH3BP2 and prevents the enhanced osteoclastogenesis and systemic bone loss characteristic of “cherubism mice”, which carry a gain-of-function mutation in *Sh3bp2* (3). Notably, cherubism mice deficient in NFATc1 still display multiorgan inflammation. Thus, NFATc1 uncouples systemic inflammation from osteopenia in this unique model and may represent a therapeutic target for TNF- $\alpha$ -dependent bone loss.

## Results

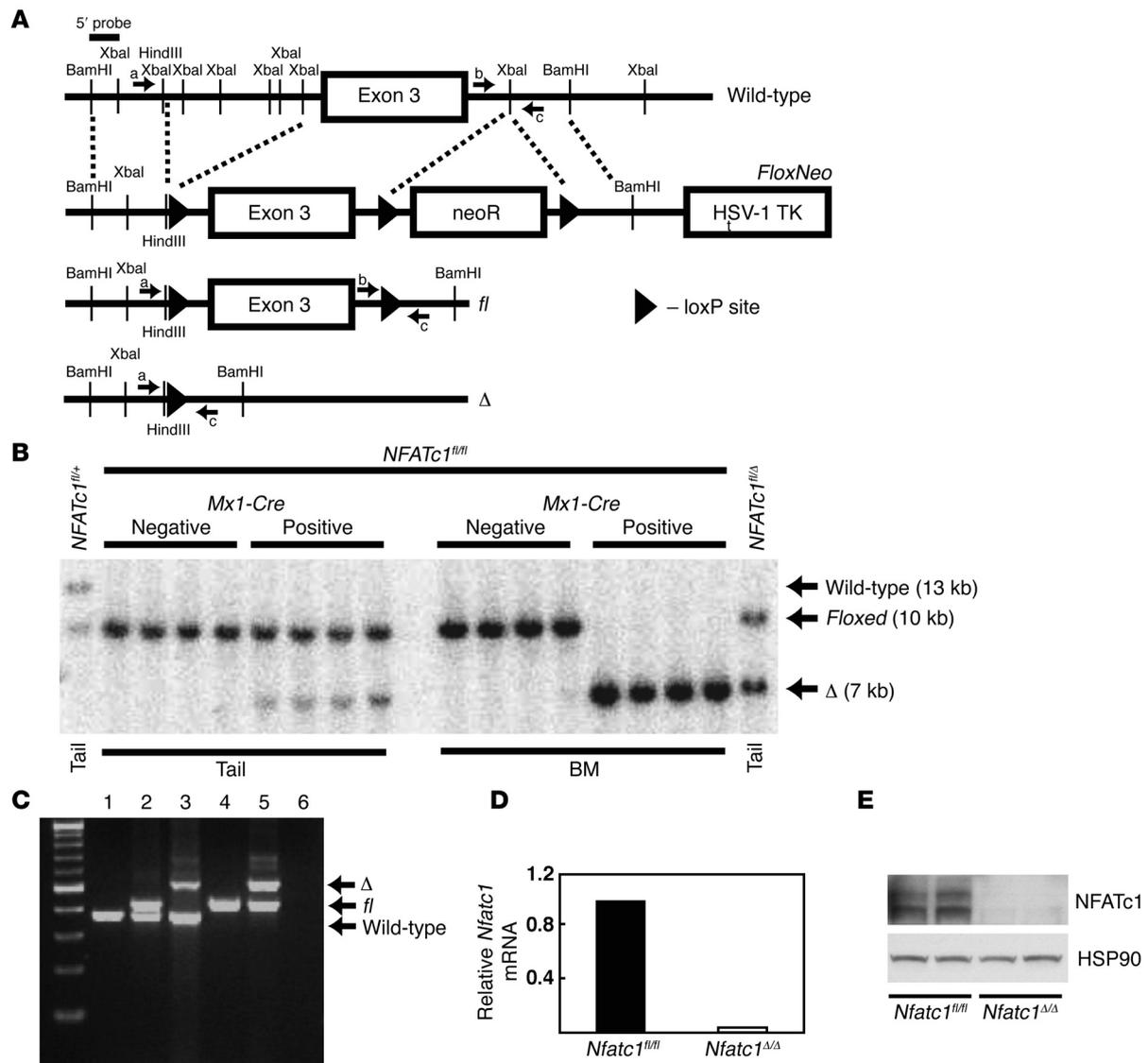
**Creation of an *Nfatc1* conditional knockout mouse.** To circumvent embryonic lethality (20), the exon encoding the regulatory domain of *Nfatc1* (exon 3) was flanked with loxP sites (*Nfatc1<sup>fl/fl</sup>*) (Figure 1A) (31). *Nfatc1*-deficient mice (*Nfatc1<sup>Δ/Δ</sup>*) were generated by crossing *Nfatc1<sup>fl/fl</sup>* mice with a transgenic line expressing the Cre recombinase from a type I interferon inducible promoter (*Mx1-Cre*) (32). Activation of *Mx1-Cre* by i.p. injection of polyinosinic-polycytidylic acid (poly I:C) resulted in complete deletion of exon 3 in BM DNA (Figure 1B), a greater than 95% reduction in *Nfatc1* mRNA levels in BM (Figure 1D), and markedly diminished NFATc1 protein levels in splenic lysates (Figure 1E).

***Nfatc1<sup>Δ/Δ</sup>* mice develop osteoclast-poor osteopetrosis.** *Nfatc1* was ablated in 10-day-old pups to examine its role in the skeleton of growing and adult mice. Compared with littermate controls, *Nfatc1<sup>Δ/Δ</sup>* mice displayed (a) short, club-shaped long bones, with higher radiodensity at the distal femur and proximal tibia (Figure 2, A–C, and Supplemental Figure 1A; supplemental material available online with this article; doi:10.1172/JCI35711DS1); (b) a massive increase in Von Kossa and toluidine blue-positive matrix (calcified cartilage) adjacent to the growth plates (Figure 2, D and E); and (c) growth plate dysplasia, characterized by an increase in thickness, acellular areas, and the retention of hypertrophic chondrocytes beyond the chondro-osseous junction (Figure 2, E and F). Other skeletal sites, such as diaphyseal cortical bone and the epiphysis did not display increased bone density (Supplemental Figure 1B

and A.O. Aliprantis, data not shown). Some *Nfatc1<sup>Δ/Δ</sup>* mice developed upper incisor overgrowth with short or absent lower incisors (Figure 2G). Lateral radiographs of the skull did not show impaction of the lower incisors in the jaw but did reveal widened, hyperdense mandibular condyles (Figure 2H). Histologically, the mandibular condyles of *Nfatc1<sup>fl/fl</sup>* mice showed a thin cap of articular cartilage overlying a mature neck of compact bone, while those from *Nfatc1<sup>Δ/Δ</sup>* mice displayed marked retention of hypertrophic cartilage (Figure 2I). These data suggest that the tooth phenotype observed in *Nfatc1<sup>Δ/Δ</sup>* mice is a consequence of malocclusion from abnormal biomechanical properties at the mandibular condyle, though a selective defect in lower incisor growth cannot be excluded. The increased bone density in *Nfatc1<sup>Δ/Δ</sup>* mice is not due to augmented osteoblast function because (a) these mice showed a trend toward reduced osteoblast numbers (Supplemental Figure 1, C and D) and (b) *Nfatc1<sup>fl/fl</sup>* mice expressing the Cre recombinase from the osteoblast-specific osterix promoter (33) did not display a similar high bone mass phenotype (Supplemental Figure 1E). Taken together, *Nfatc1<sup>Δ/Δ</sup>* mice display features of severe osteopetrosis, with improperly shaped long bones and a failure to resorb primary spongiosa (calcified cartilage), resulting in aberrant endochondral growth and ossification (34, 35).

Osteopetrosis results from either a decrease in osteoclast numbers or a reduction in osteoclast function (6). Therefore osteoclast differentiation was investigated in *Nfatc1<sup>Δ/Δ</sup>* mice in vivo and in vitro. Compared with *Nfatc1<sup>fl/fl</sup>* littermates, *Nfatc1<sup>Δ/Δ</sup>* mice displayed (a) a deficiency in tartrate-resistant acid phosphatase-positive (TRAP-positive) multinucleated cells at the growth plates of the long bones and mandibles (Figure 3A and A.O. Aliprantis, data not shown); (b) reduced osteoclast numbers by histomorphometric analysis (Figure 3B); and (c) decreased TRAP5b levels, a serologic marker of osteoclast function (Figure 3C). Rare TRAP-positive, morphologically normal osteoclasts could be identified in some *Nfatc1<sup>Δ/Δ</sup>* mice (Supplemental Figure 1, F and G), consistent with our observation that serum TRAP5b levels were reduced, but not absent, in these mice. Consistent with markedly reduced osteoclast numbers in vivo, BM cells from *Nfatc1<sup>Δ/Δ</sup>* mice failed to generate osteoclasts in vitro in response to either osteoblasts (Figure 3, D and E) or M-CSF and RANKL (Figure 3F). To establish that the differentiation defect was due to a failure of osteoclast progenitors to differentiate, and not to a lack of progenitors, osteoclast precursors were purified. Within the CD11b<sup>lo</sup>-CD3 $\epsilon$ -B220<sup>-</sup>c-fms<sup>+</sup> population in BM are c-kit<sup>+</sup> and c-kit<sup>-</sup> cells, with the former being the most highly osteoclastogenic (36). Similar numbers of each population were found in *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* BM (Figure 3, G and H). Fluorescence-activated cell sorting-purified (FACS-purified) BM-derived CD11b<sup>lo</sup>-CD3 $\epsilon$ -B220<sup>-</sup>c-kit<sup>+</sup>-c-fms<sup>+</sup> osteoclast precursors (BMOcPs) from *Nfatc1<sup>Δ/Δ</sup>* mice failed to generate osteoclasts (Figure 3, I and J). Importantly, M-CSF induced normal macrophage differentiation in *Nfatc1<sup>Δ/Δ</sup>* BM cells, as determined by CD11b and F4/80 staining (A.O. Aliprantis, data not shown) (37). In summary, *Nfatc1*-deficient mice display a progenitor intrinsic defect in osteoclastogenesis.

***Nfatc1* deletion using lysozyme-M-Cre does not result in osteopetrosis.** Lysozyme-M-Cre (*LysM-Cre*) (38) has been used to interrogate osteoclast function in mice carrying conditional alleles (39, 40). *LysM-Cre* mice express the Cre recombinase from a myeloid specific promoter (38). Surprisingly, no alterations in bone density or osteoclast differentiation were observed in *Nfatc1<sup>fl/fl</sup>*, *LysM-Cre* mice (Supplemental Figure 2, A–C). This result was explained by the fact



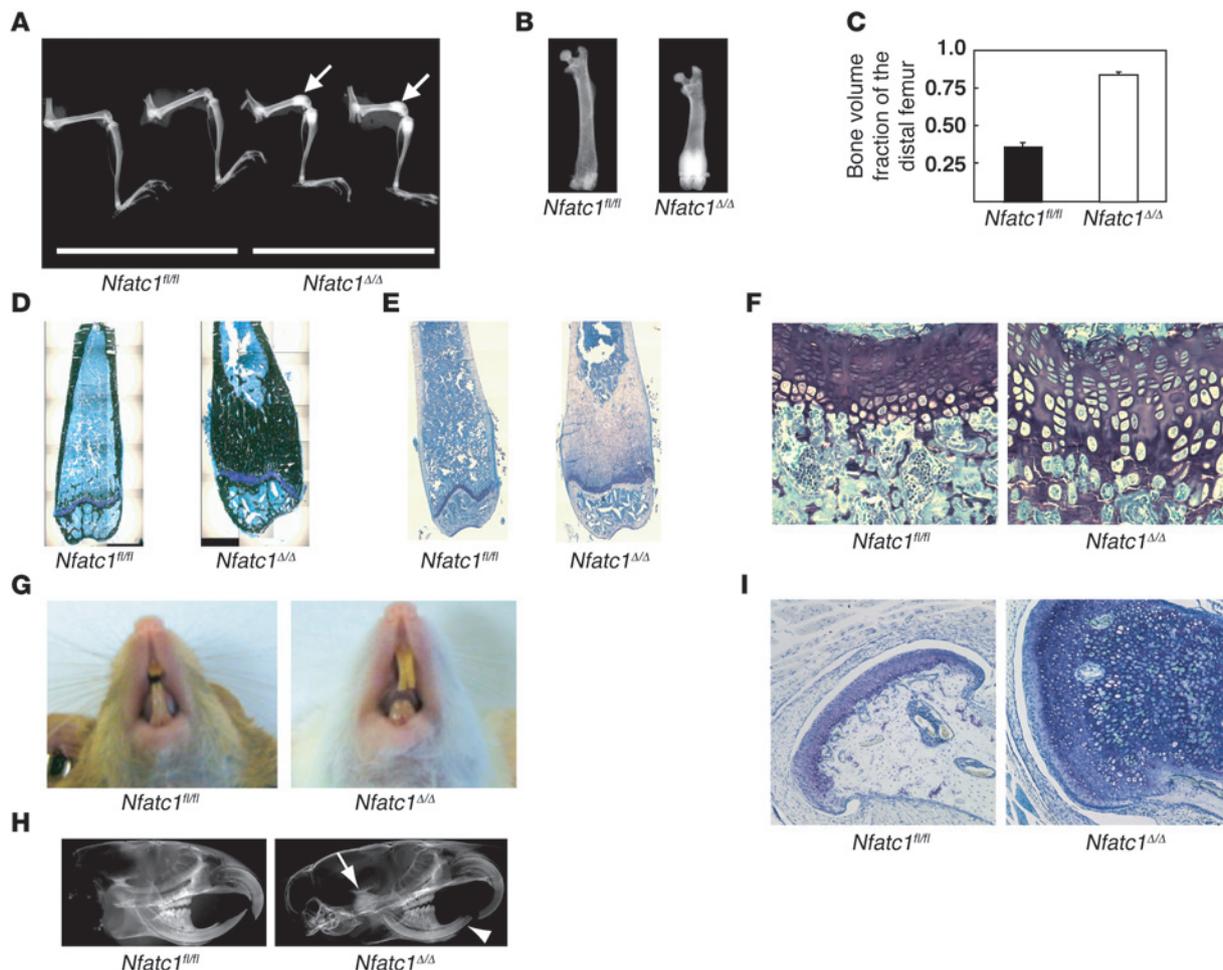
**Figure 1**

Generation of an *Nfatc1* conditional knockout mouse. **(A)** Graphic representation of the *Nfatc1* conditional targeting strategy. Flanking exon 3 of *Nfatc1* with loxP sites generated the floxed allele (*fl*). Cre-mediated recombination of the loxP sites results in the delta allele ( $\Delta$ ). The approximate locations of the 5' probe used for Southern blotting and PCR primers (see Methods) employed for genotyping (a, b, and c) are shown. HSV-1 TK, herpes simplex virus-1 thymidine kinase cassette; neoR, neomycin resistance cassette. **(B)** Southern blot analysis of tail and BM genomic DNA from poly I:C-treated *Nfatc1<sup>fl/fl</sup>* ( $n = 4$ ) and littermate *Nfatc1<sup>fl/fl</sup>*, *Mx1-Cre* ( $n = 4$ ) mice. Genomic tail DNA from *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>fl/Δ</sup>* mice are shown for comparison. **(C)** Three primer genotyping PCR, using primers a, b, and c shown in **A**, of tail genomic DNA from mice of the following genotypes: *Nfatc1<sup>+/+</sup>* (lane 1), *Nfatc1<sup>fl/+</sup>* (lane 2), *Nfatc1<sup>+/Δ</sup>* (lane 3), *Nfatc1<sup>fl/fl</sup>* (lane 4), *Nfatc1<sup>fl/Δ</sup>* (lane 5). A 100-bp ladder and DNA diluent negative control (lane 6) are shown. **(D)** qRT-PCR analysis for *Nfatc1* (using primer set *Nfatc1ex3*; Table 1) in mRNA samples from *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* BM. **(E)** Western blot analysis for NFATc1 and HSP90 (loading control) in protein lysates from *Nfatc1<sup>fl/fl</sup>* ( $n = 2$ ) and *Nfatc1<sup>Δ/Δ</sup>* ( $n = 2$ ) splenocytes.

that *LysM-Cre* failed to efficiently delete the floxed allele in BM macrophage or osteoclast cultures (Supplemental Figure 2D). However, near complete deletion was observed in peritoneal macrophages (Supplemental Figure 2E). These findings are consistent with the original report of this line (38) and indicate that *LysM-Cre* may be of limited value to the study of conditional alleles in osteoclasts.

*Microarray analysis of FACS-sorted osteoclast progenitors reveals 2 families of NFATc1-regulated genes in the developing osteoclast.* NFATc1 is necessary for osteoclast differentiation in vivo and in vitro (Figures 2 and 3) (22). However, the universe of NFATc1-regu-

lated transcripts in osteoclasts is unknown. A protocol to isolate BMOcPs was developed (Supplemental Figure 3, A and B) to facilitate a high fidelity comparison of the mRNA profiles of WT and NFATc1-deficient developing osteoclasts. In response to RANKL, WT BMOcPs (a) displayed an approximately 33-fold higher osteoclast forming capacity on a per cell basis compared with crude BM cells, (b) upregulated *Nfatc1* mRNA and NFATc1 protein, and (c) expressed genes associated with terminal osteoclast differentiation (Supplemental Figure 3, C-F). mRNA profiling, by DNA microarray and qRT-PCR, of *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>*  $\mu$ -CSF-



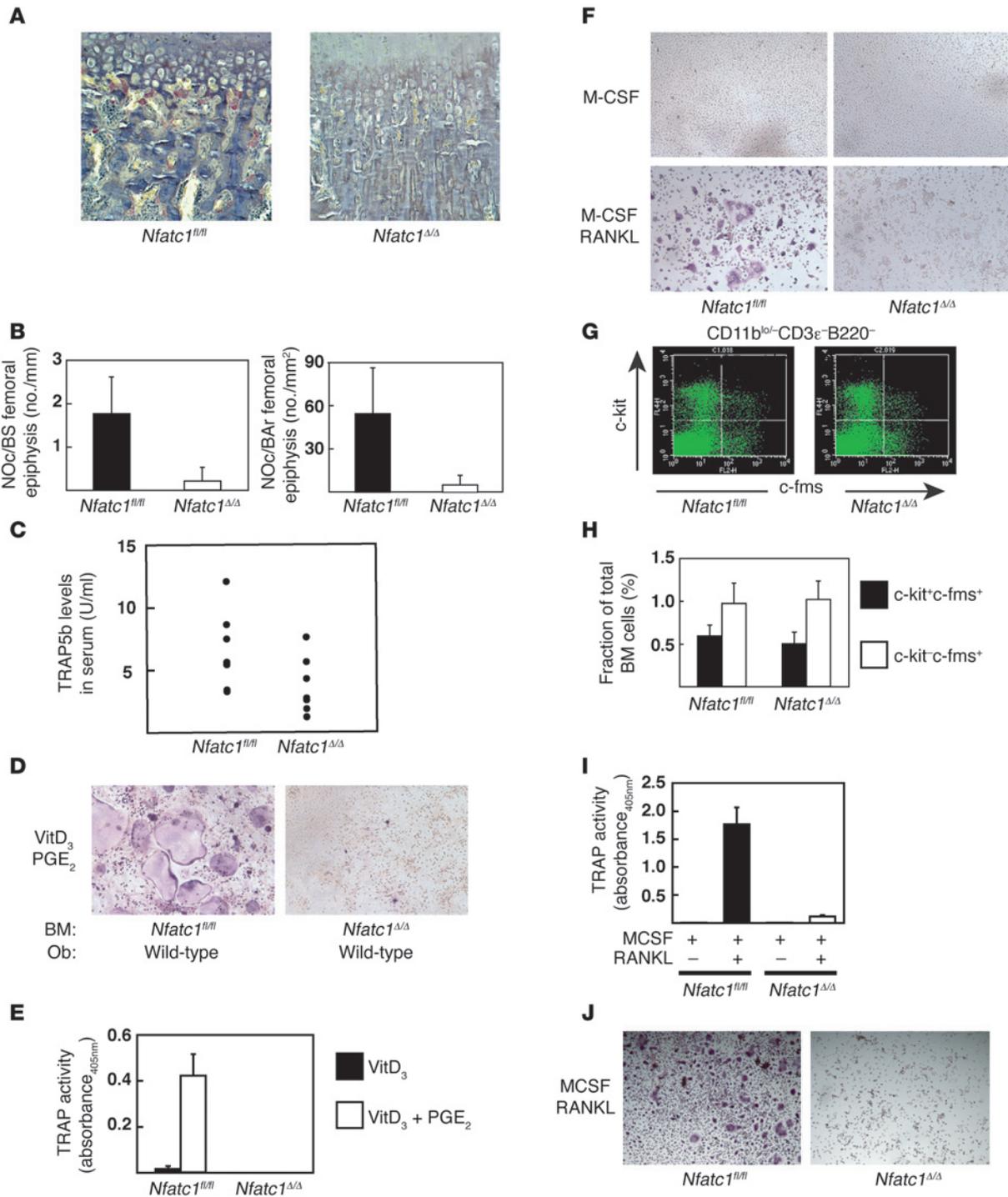
**Figure 2**

*Nfatc1<sup>ΔΔ</sup>* mice develop osteopetrosis. (A) Lateral digital radiograph of the leg of 5-month-old female *Nfatc1<sup>fl/fl</sup>* ( $n = 2$ ) and *Nfatc1<sup>ΔΔ</sup>* ( $n = 2$ ) mice. The white arrows show increased radiodensity at the distal femurs. (B) Anteroposterior digital radiograph of the femur of 5-month-old female *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>ΔΔ</sup>* mice. (C) The bone volume to total volume fraction was determined by microquantitative computed tomography at the distal femur of 6 male *Nfatc1<sup>fl/fl</sup>* (black bar,  $n = 2$  at 2.5 months old and  $n = 4$  at 5 months old) and 4 male *Nfatc1<sup>ΔΔ</sup>* (white bar,  $n = 2$  at 2.5 months old and  $n = 2$  at 5 months old) mice. The data are the mean + SD;  $P < 1 \times 10^{-8}$ . (D) Von Kossa and (E) toluidine blue stains of the distal femur of 5-month-old female *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>ΔΔ</sup>* mice. Pictures are a montage of low-power images. (F) Toluidine blue stain (original magnification,  $\times 400$ ) of the femoral growth plate of 9-week-old male *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>ΔΔ</sup>* mice. (G) Photograph of the snout of 5.5-month-old male *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>ΔΔ</sup>* mice. (H) Lateral radiograph of the skull of 2.5-month-old female *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>ΔΔ</sup>* mice. White arrowhead denotes lack of impaction of the lower incisors. White arrow shows increased radiodensity at the mandibular condyle. (I) Toluidine blue stain (original magnification,  $\times 100$ ) of the mandibular condyle of 3-month-old female *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>ΔΔ</sup>* mice. Radiographs and histology images are representative of at least 8 (long bones) or 3 (mandibles) mice analyzed per genotype.

and RANKL-stimulated BMOcPs (MROcPs) revealed 2 differentially regulated sets of osteoclast genes: (a) NFATc1-dependent genes (Figure 4, A and B) and (b) “NFATc1-augmented” genes (Figure 4, C and D). *Oscar*, *Calcr*, and *Itgb3*, all well-characterized osteoclast genes (6, 19), were absolutely dependent on NFATc1 for expression (Figure 4, A and B). Likewise, *Myo1d* and *Mst1r*, potential regulators of bone resorption (41, 42), depended on NFATc1. Lastly, *Rcan2*, a regulator of the NFAT pathway and homolog of *Rcan1*, an NFATc1 target gene in the heart (43), was only induced in NFATc1-sufficient cells. In contrast, the following genes, whose function in osteoclasts is established (35, 44, 45), could be partially induced by RANKL even in the absence of NFATc1: *Mmp9*, *Acp5*, *Ctsk*, *Car2*, *Mmp14*, and *Cln7* (Figure 4, C and D). However, NFATc1 did augment the expression of all 6

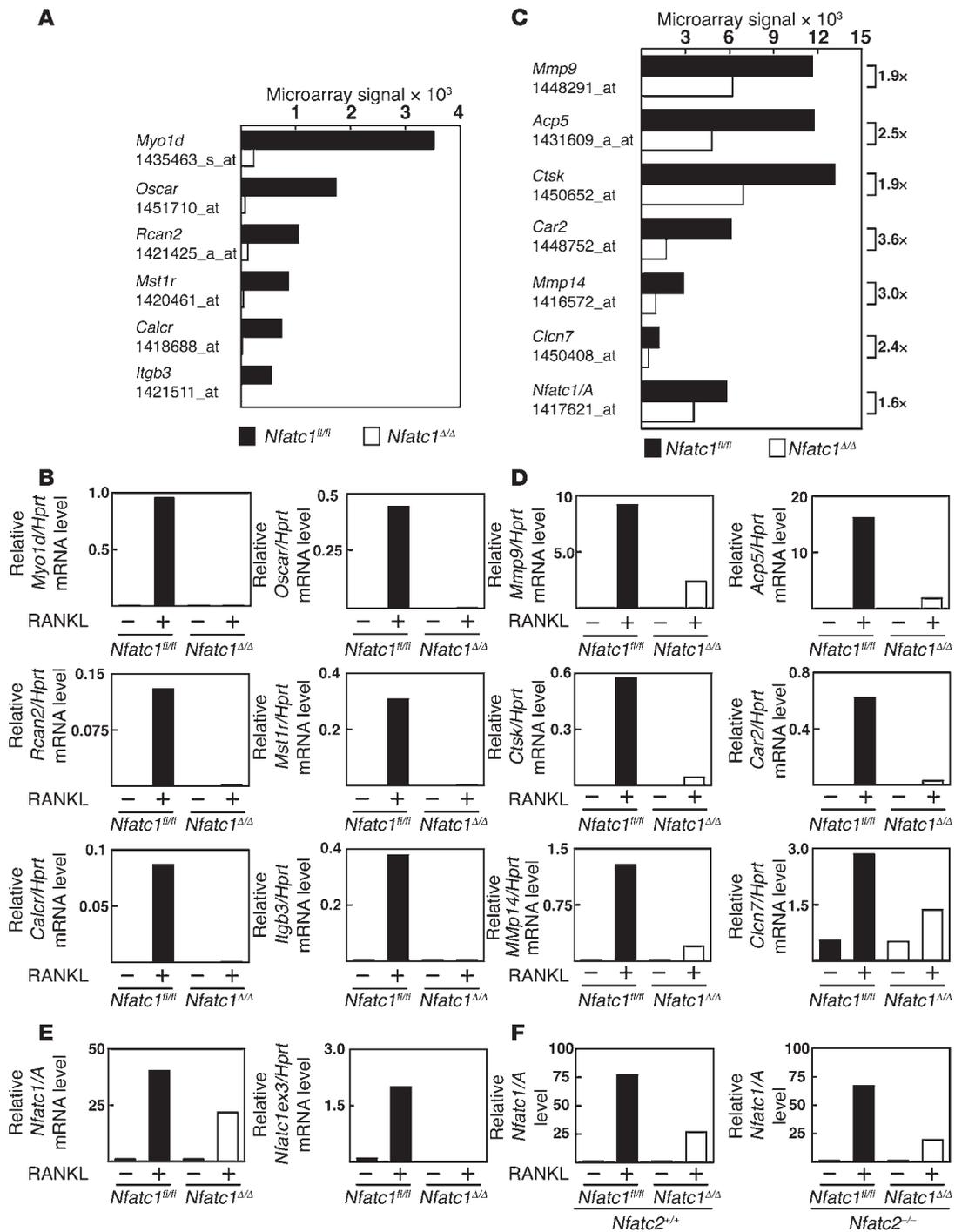
genes. These data support NFATc1 as a broad regulator of osteoclast gene expression but suggest that NFATc1 is more essential to the control of some genetic loci than others.

Surprisingly, a microarray probe set (1417621\_at) to *Nfatc1* revealed expression in *Nfatc1<sup>ΔΔ</sup>* MROcPs (Figure 4C). This probe set targets the 3' untranslated region of the *Nfatc1/A* isoform, which is preferentially induced during osteoclastogenesis and suggested to be a target of autoamplification (22). Since neither full-length *Nfatc1* mRNA (Figure 4E, primer set *Nfatc1ex3*, Table 1) nor NFATc1 protein could be detected in *Nfatc1<sup>ΔΔ</sup>* MROcPs (Supplemental Figure 3E), this suggested that an untranslated, mutant *Nfatc1/A* mRNA is induced by RANKL in these cells. qRT-PCR confirmed the expression of *Nfatc1/A* mRNA in *Nfatc1<sup>ΔΔ</sup>* BMOcPs stimulated with RANKL (Figure 4E). Thus, the endogenous



**Figure 3**

*Nfatc1<sup>Δ/Δ</sup>* mice display impaired osteoclast differentiation in vivo and in vitro. **(A)** TRAP stain of the femoral growth plates of 9-week-old male *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* mice. Images are representative of at least 8 mice analyzed per genotype. **(B)** Histomorphometric enumeration of osteoclasts within the femoral epiphysis of 5-month-old female *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* mice ( $n = 4/\text{genotype}$ ). The data is the mean + SD. NOc/BS, number of osteoclasts per mm bone surface; NOc/BAR, number of osteoclasts per mm<sup>2</sup> bone area.  $P < 0.05$  for both panels. **(C)** TRAP5b levels in the serum of 10- to 20-week-old female *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* mice ( $n = 8/\text{genotype}$ );  $P < 0.05$ . **(D)** TRAP stain and **(E)** TRAP assay of *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* M-CSF–primed BM cells cultured with osteoblasts (Obs) and Vitamin D<sub>3</sub> **(E)** or Vitamin D<sub>3</sub> and PGE<sub>2</sub> **(D and E)**. The data in **E** are the mean + SD of triplicate wells and representative of 2 independent experiments. **(F)** TRAP stain of *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* M-CSF–primed BM cells cultured with M-CSF or M-CSF and RANKL. **(G)** c-kit versus c-fms FACS plot of *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* BM cells gated on the CD11b<sup>lo/-</sup>B220<sup>-</sup>CD3ε<sup>-</sup> population. **(H)** Quantification of osteoclast precursors in the BM of *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* mice. The data is the mean + SD ( $n = 5/\text{genotype}$ ). **(I)** TRAP assay and **(J)** TRAP stain of *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* BMOcPs cultured with M-CSF or M-CSF and RANKL. The data in **I** is the mean + SD of triplicate wells and representative of more than 3 independent experiments. Original magnification, ×400 **(A)**; ×100 **(D, F, and J)**.



**Figure 4**

Comparative microarray analysis reveals 2 sets of NFATc1-regulated genes in osteoclasts. (A) Microarray signal intensities for selected NFATc1-dependent mRNAs from *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* MROcPs. (B) qRT-PCR analysis for the expression of the NFATc1-dependent mRNAs identified in A in *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* BMOcPs stimulated with M-CSF or M-CSF and RANKL for 3 days. (C) Microarray signal intensities for selected NFATc1-augmented mRNAs from *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* MROcPs. (D) qRT-PCR analysis for the expression of the NFATc1-augmented mRNAs identified in C in *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* BMOcPs stimulated with M-CSF or M-CSF and RANKL for 3 days. (E) qRT-PCR analysis for the expression of the *Nfatc1* mRNA isoforms in *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* BMOcPs stimulated with M-CSF or M-CSF and RANKL for 3 days. PCR primers targeted the deleted exon (*Nfatc1ex3*) or the *Nfatc1/A* isoform (see Table 1). Note that the qRT-PCR analyses presented in B, D, and E were performed on the same mRNA samples and are representative of at least 2 independent experiments. (F) qRT-PCR analysis for the expression of *Nfatc1/A* mRNA in NFATc2 sufficient (*Nfatc2<sup>+/+</sup>*) and deficient (*Nfatc2<sup>-/-</sup>*) *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>Δ/Δ</sup>* BMOcPs stimulated with M-CSF or M-CSF and RANKL for 4 days. The data in F are representative of at least 2 independent experiments.



**Table 1**  
qRT-PCR primers used in this study

Primer set	Forward primers (5'–3')	Reverse primers (5'–3')
<i>Nfatc1ex3<sup>A</sup></i>	TGCCTTTTGGCAGCAGTATCT	CAGGCAAGGATGGGCTCATAT
<i>Nfatc1/A<sup>B</sup></i>	GGTAACTCTGTCTTTCTAACCTTAAGCTC	GTGATGACCCAGCATGCACCAGTCACAG
<i>Itgb3<sup>C</sup></i>	TCCAGACCTGGGTACCAAG	GCCAATCCGAAGGTTGCTAG
<i>Calcr<sup>C</sup></i>	GCCTCCCATTACATCTGC	CTCCTCGCCTTCGTTGTTG
<i>Myo1d<sup>A</sup></i>	TCATTGGAACACCCCGTACAC	CACGCCACACCTTTTGTAGGA
<i>Oscar<sup>A</sup></i>	TCTGCCCTATGTGCTATCA	AGGAGCCAGAACCTTCGAAAC
<i>Rcan2<sup>A</sup></i>	CCACTCTGGTGGCCTGTGT	CGGAACAGTCCCTCGAATTTTCTTA
<i>Mst1r<sup>A</sup></i>	TCACCACCAAGTCCGATGTGT	AGGTGGAAGGGATCGATATGG
<i>Tnfrsf11b<sup>D</sup></i> (OPG)	TGTCCAGATGGGTTCTTCTCA	CGTTGTCATGTGTTGCATTTCC
<i>Mmp9<sup>E</sup></i>	ACGGACCCGAAGCGGAC	GGGATACCCGTCCTCGTGC
<i>Acp5<sup>E</sup></i>	CAGCAGCCCAAAATGCCT	TTTTGAGCCAGGACAGCTGA
<i>Ctsk<sup>C</sup></i>	GGGCTCAAAGTTCTGCTGC	TGGGTGTCAGCATTTCCTC
<i>Car2<sup>D</sup></i>	TCCCACACTGGGGATACAG	CTCTTGGACGCAGCTTATCATA
<i>Mmp14<sup>D</sup></i>	AGTGACAGGCAAGGCTGATT	AGGGGTGTAATTCGAATGCAG
<i>Cln7<sup>D</sup></i>	CGCCAGTCTCATTCTGCACT	GCTTCTCGTTGTGTGGAATCT
<i>Tnfr<sup>A</sup></i>	CATCTTCTCAAATTCGAGTGACAA	TGGGAGTAGACAAGGTACAACCC

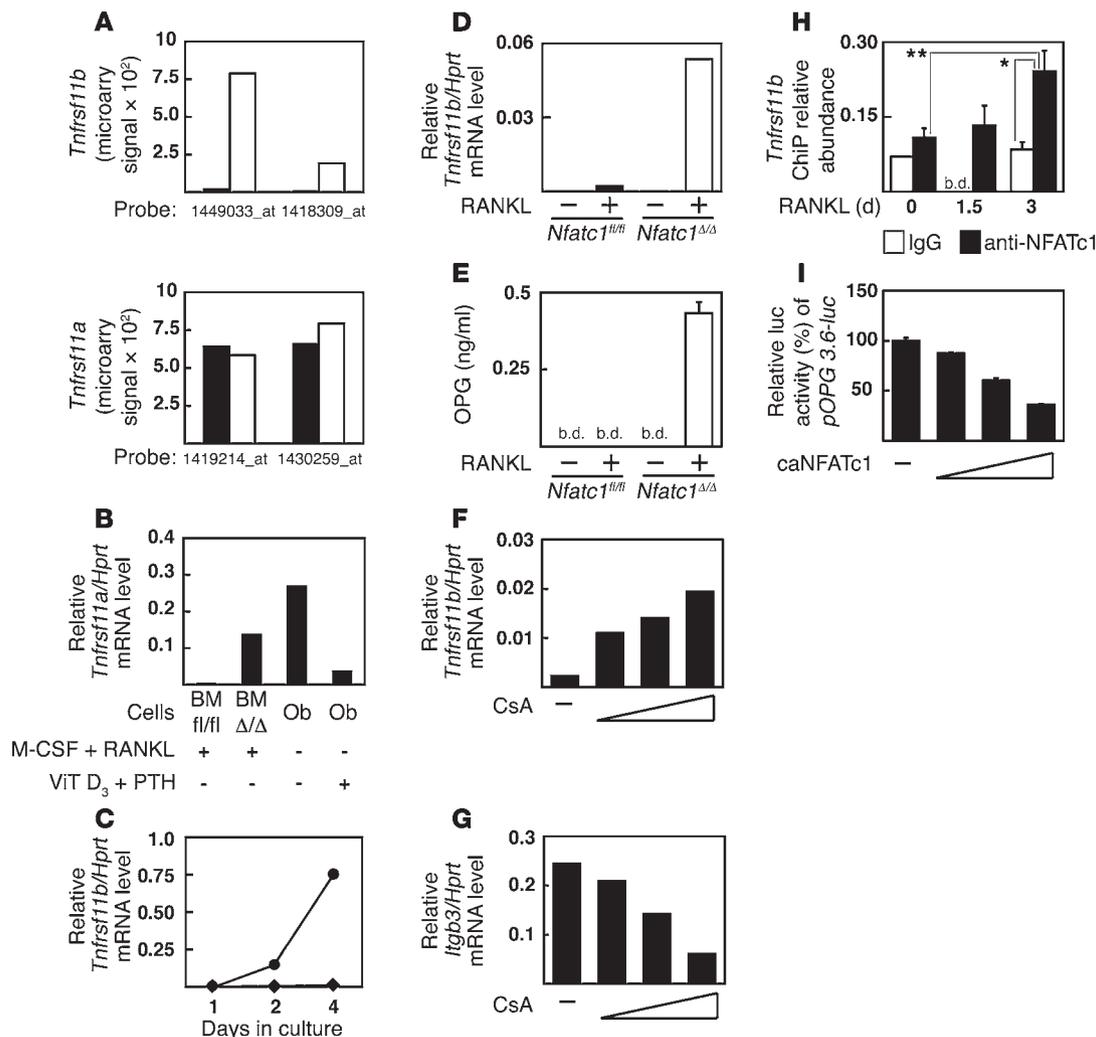
<sup>A</sup>Designed with the use of Primer Express Software (Applied Biosystems). <sup>B</sup>Previously described in ref. 22. <sup>C</sup>Sequences were a kind gift from K. McHugh, Harvard Medical School. <sup>D</sup>The Center for Comparative and Integrative Biology Primer Bank, Harvard Medical School (78). Primerbank IDs 31543882a3 (*Tnfrsf11b*), 31981657a1 (*Car2*), 31982191a2 (*Mmp14*), and 6753436a1 (*Cln7*).

*Nfatc1* locus can still be stimulated by RANKL in the absence of NFATc1. However, NFATc1 was necessary for optimal expression, since higher levels of *Nfatc1/A* mRNA were detected in *Nfatc1<sup>f/f</sup>* BMOcPs stimulated with RANKL compared with *Nfatc1<sup>Δ/Δ</sup>* cells (Figure 4E), consistent with autoamplification (22). Thus, *Nfatc1/A* is an NFATc1 augmented transcript. Since NFAT family members often display redundant functions (46) and NFATc2 is recruited to the *Nfatc1* promoter (22), the induction of *Nfatc1/A* mRNA by RANKL was investigated in NFATc2-deficient and NFATc1/NFATc2 doubly deficient BMOcPs. WT (*Nfatc1<sup>f/f</sup>Nfatc2<sup>+/+</sup>*) and NFATc2-deficient (*Nfatc1<sup>f/f</sup>Nfatc2<sup>-/-</sup>*) BMOcPs strongly upregulated *Nfatc1/A* in response to RANKL treatment for 4 days (Figure 4F). Interestingly, the induction of *Nfatc1/A* mRNA was not further attenuated in doubly deficient (*Nfatc1<sup>Δ/Δ</sup>Nfatc2<sup>-/-</sup>*) BMOcPs compared with those lacking NFATc1 alone (*Nfatc1<sup>Δ/Δ</sup>Nfatc2<sup>+/+</sup>*) (Figure 4F). These data suggest that NFATc2 does not play a dominant or compensatory role in the induction of *Nfatc1/A* mRNA during osteoclastogenesis.

*Tnfrsf11b* is upregulated in osteoclast precursors in the absence of NFATc1. Although the expression and function of OPG by osteoblasts has been well studied (47), OPG expression by osteoclast-like cells has only been reported, to our knowledge, in giant cell tumors (24, 48). Remarkably, 2 probe sets showed expression of *Tnfrsf11b* in *Nfatc1<sup>Δ/Δ</sup>* MROcPs but not in *Nfatc1<sup>f/f</sup>* cells (Figure 5A). In comparison, *Tnfrsf11a* expression levels were similar (Figure 5A). The expression of *Tnfrsf11b* mRNA by *Nfatc1<sup>Δ/Δ</sup>* MROcPs was similar to levels found in WT osteoblasts (Figure 5B), increased over time in culture (Figure 5C), and was dependent on RANKL, as cells cultured in M-CSF alone did not express *Tnfrsf11b* (Figure 5D). In addition, OPG was detected in the culture supernatants of *Nfatc1<sup>Δ/Δ</sup>* MROcPs (Figure 5E). Treatment of *Nfatc1<sup>f/f</sup>* MROcPs with cyclosporine A, an inhibitor of the NFAT activating phosphatase calcineurin (10), reciprocally upregulated *Tnfrsf11b* and downregulated the NFATc1-dependent gene, *Itgb3* (Figure 5, F and G). Despite the dramatic upregulation of *Tnfrsf11b* mRNA

by *Nfatc1<sup>Δ/Δ</sup>* MROcPs, no differences in serum OPG levels were observed between *Nfatc1<sup>f/f</sup>* and *Nfatc1<sup>Δ/Δ</sup>* mice (Supplemental Figure 4). To exclude the possibility that cells of mesenchymal origin (i.e., osteoblasts) were contaminating our BMOcP preparations from *Nfatc1<sup>Δ/Δ</sup>* mice and producing the OPG signal observed in these experiments, osteoclast precursors were derived from spleen cells. Similar to FACS-sorted BMOcPs, *Nfatc1<sup>Δ/Δ</sup>* spleen cells both failed to differentiate into osteoclasts and upregulated *Tnfrsf11b* upon stimulation with RANKL (Supplemental Figure 5, A–D). Interestingly, a putative NFAT binding site was identified in a conserved region of the human TNFRSF11B and mouse *Tnfrsf11b* promoters, located approximately 2 kb and 3 kb from the transcription start site (TSS), respectively (Supplemental Figure 6A). After 3 days of RANKL treatment, NFATc1 was recruited to the *Tnfrsf11b* promoter, as determined by a ChIP assay using primers flanking this conserved sequence (Figure 5H and Supplemental Figure 6A). In contrast, NFATc1 binding to the *Itgb3* promoter peaked at 1.5 days of RANKL treatment and was almost undetectable by 3 days (Supplemental Figure 6B), consistent with previous observations (18). Lastly, a constitutively active version of NFATc1 (caNFATc1) repressed the basal activity of a *Tnfrsf11b* reporter plasmid (*pOPG 3.6-luc*) (Figure 5I). Appropriately, caNFATc1 had little effect on an NF-κB reporter plasmid (*pBII-luc*) and strongly activated an NFAT luciferase reporter construct (*pNFAT-luc*) (Supplemental Figure 6, C and D). Taken together, these results uncover a previously unrecognized regulatory loop downstream of RANKL: calcineurin/NFATc1-mediated repression of OPG expression.

*NFATc1 dissociates systemic bone loss and osteoclastogenesis from inflammation in cherubism.* Our conditional knockout strain represents a unique tool to evaluate the role of NFATc1 in disease. Cherubism is a pediatric, craniofacial, inflammatory disorder, characterized by bone loss in the jaw that is caused by mutations in an adapter molecule *SH3BP2* (3, 25). To clarify the role of NFATc1 in cherubism, *Mx1-Cre* was used to ablate *Nfatc1* in mice carrying 1 (*KI/+*) or 2 (*KI/KI*) cherubism alleles (3). This approach was taken since the cherubism phenotype is intrinsic to myeloid cells (3) and deletion of *Nfatc1<sup>f/f</sup>* by *Mx1-Cre* is complete in the BM (Figure 1, B and D). Deletion of *Nfatc1* reversed the diffuse bone loss and focal cystic changes characteristic of cherubism mice (Figure 6A). In contrast, *Nfatc1<sup>Δ/Δ</sup>KI/KI* mice still developed widespread inflammation in the liver and lungs (Figure 6, B and C) as well as lymph nodes, spleen, and stomach (A.O. Aliprantis and Y. Ueki, data not shown). The systemic bone loss and inflammatory phenotypes of cherubism mice are dependent on TNF-α derived from myeloid cells, which express abnormally high levels of *Tnf* mRNA (3). Consistent with our observation that deletion of *Nfatc1* had no effect on inflammation in vivo, BM macrophage cultures from *KI/+* and *KI/KI* mice

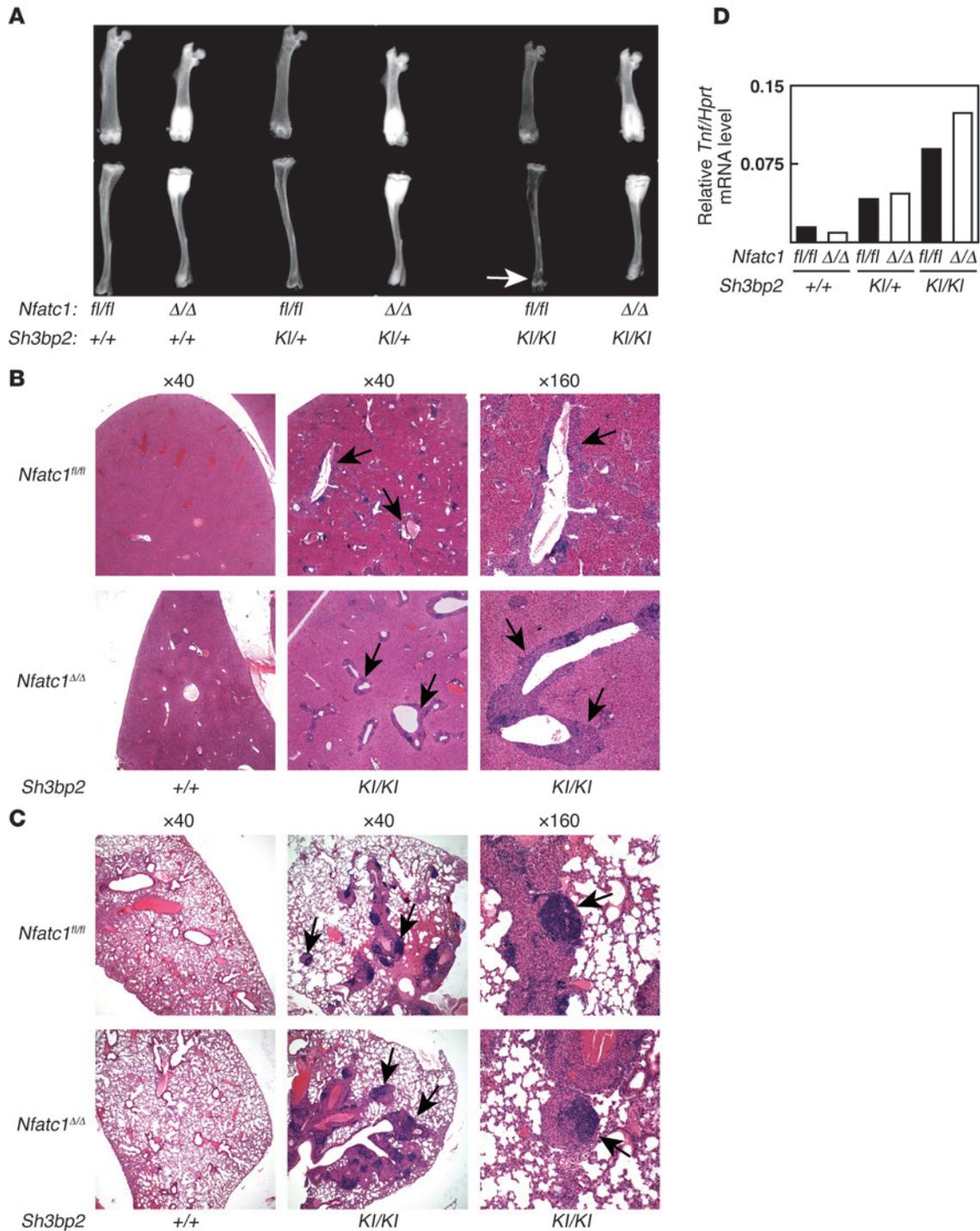


**Figure 5**

RANKL induces *Tnfrsf11b* expression in the absence of NFATc1. (A) Microarray signal intensities for *Tnfrsf11b* (upper panel) and *Tnfrsf11a* (lower panel) mRNAs from *Nfatc1<sup>fl/fl</sup>* (black bars) and *Nfatc1 $\Delta\Delta$*  (white bars) MROcPs. (B–D) qRT-PCR analysis for *Tnfrsf11b* mRNA in (B) *Nfatc1<sup>fl/fl</sup>* and *Nfatc1 $\Delta\Delta$*  MROcPs or WT calvarial osteoblasts with or without Vitamin D<sub>3</sub> and parathyroid hormone (PTH); (C) *Nfatc1<sup>fl/fl</sup>* (diamonds) and *Nfatc1 $\Delta\Delta$*  (circles) BMOCs stimulated with M-CSF and RANKL for 1, 2, or 4 days; and (D) *Nfatc1<sup>fl/fl</sup>* and *Nfatc1 $\Delta\Delta$*  BMOCs stimulated for 3 days with M-CSF or M-CSF and RANKL. (E) ELISA for OPG in the supernatants of *Nfatc1<sup>fl/fl</sup>* and *Nfatc1 $\Delta\Delta$*  BMOCs stimulated for 4 days with M-CSF or M-CSF and RANKL and for 2 days with M-CSF only. b.d., below detection. The data are the mean + SD of 4 independent wells. (F) qRT-PCR analysis for *Tnfrsf11b* and (G) *Itgb3* mRNA in *Nfatc1<sup>fl/fl</sup>* BMOCs stimulated with M-CSF and RANKL in the absence or presence of increasing concentrations of cyclosporine A (CsA) (62.5, 125, or 250 ng/ml CsA). The triangle under the x axis refers to increasing CsA concentrations. (H) NFATc1 ChIP of osteoclast precursors incubated with RANKL for 0, 1.5, or 3 days. Immunoprecipitated chromatin was analyzed by qRT-PCR for *Tnfrsf11b* promoter DNA, which was normalized to input. Data is the mean + SD of 2 independent experiments (\**P* < 0.05, \*\**P* = 0.053). (I) Relative luciferase (luc) activity of 293T cells transfected with *pOPG 3.6-luc* and increasing amounts of *pMSCV-caNfatc1* (0, 8, 40, and 200 ng/transfection). The triangle under the x axis refers to increasing amounts of *pMSCV-caNfatc1* per transfection. Data are the mean + SD of transfections performed in triplicate. The data in C and B, D–G, and I are representative of 2 and 3 similar experiments, respectively.

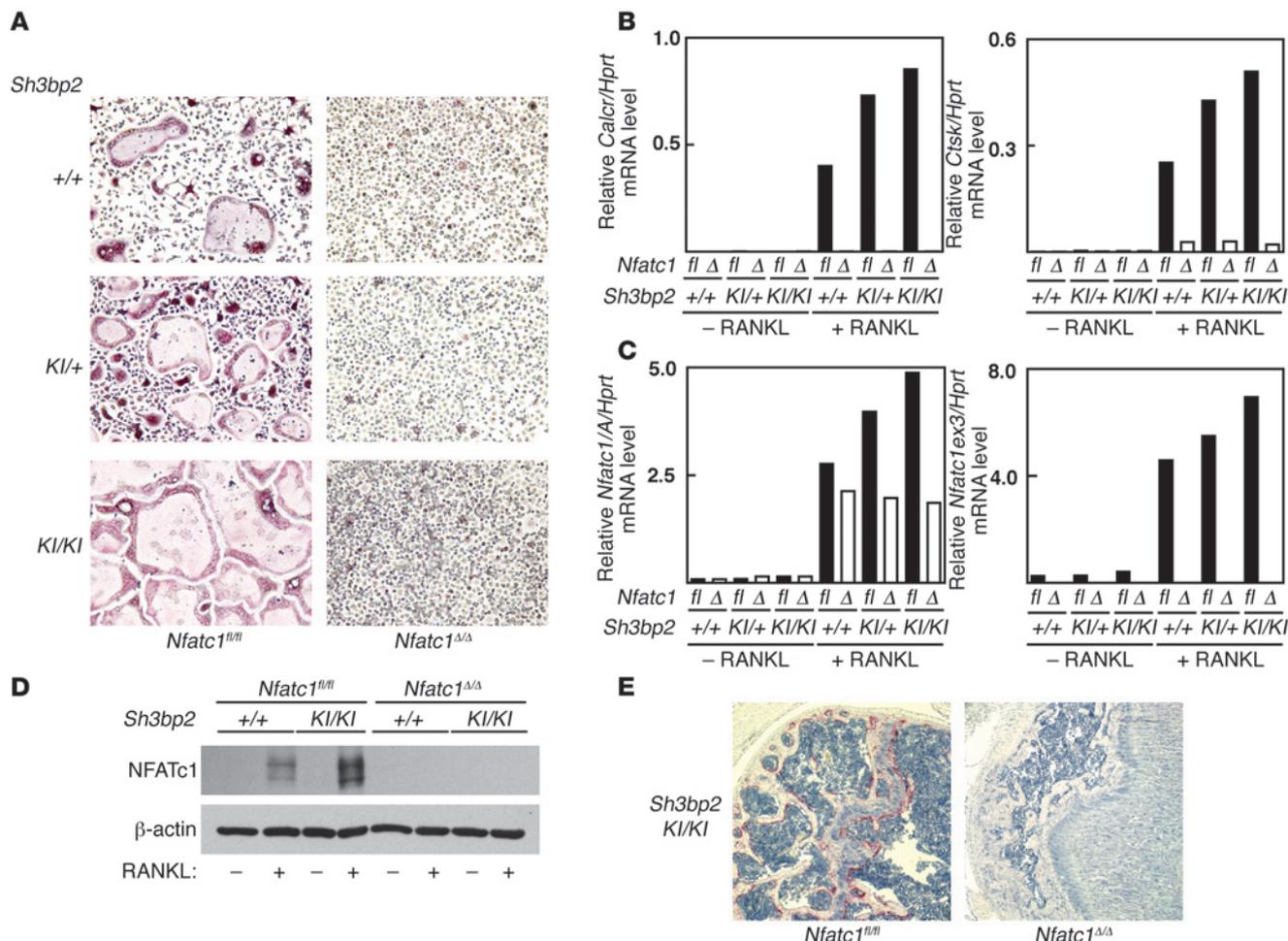
expressed higher levels of *Tnf* mRNA than WT cells irrespective of their *Nfatc1* genotype (Figure 6D). Thus, NFATc1 uncouples diffuse osteopenia from inflammation in this TNF- $\alpha$  driven, clinically relevant mouse model of a human disease. Furthermore, the enhanced osteoclastogenesis conferred by mutant SH3BP2 in vitro, which occurs in a gene dosage-dependent manner, was completely abolished in BM cells lacking NFATc1, as assessed by TRAP staining (Figure 7A) or the expression of NFATc1-regulated genes (Figure 7B). Accordingly, mutant SH3BP2 augmented RANKL induced expression of *Nfatc1/A* in proportion to gene dosage (Fig-

ure 7C) in NFATc1-sufficient cells. Conversely, RANKL did not increase the expression of the mutant *Nfatc1/A* isoform in either *Nfatc1 $\Delta\Delta$ KI/+* or *Nfatc1 $\Delta\Delta$ KI/KI* BM cells, indicating that NFATc1 protein is necessary for SH3BP2 to mediate this effect (Figure 7C). Western blotting confirmed increased NFATc1 expression in *Nfatc1<sup>fl/fl</sup>KI/KI* BM cells treated with RANKL compared with WT cells (Figure 7D). Moreover, NFATc1 was not detected in *Nfatc1 $\Delta\Delta$*  cells, irrespective of their *Sh3bp2* genotype (+/+ or KI/KI) or RANKL treatment (Figure 7D). These observations strongly suggest that SH3BP2 promotes osteoclastogenesis by augment-



**Figure 6**

NFATC1 uncouples bone loss from inflammation in a mouse model of cherubism. **(A)** Anteroposterior digital radiograph of the femurs and tibias of 12-week-old female mice. White arrow denotes cystic changes at the distal tibia of *Nfatc1<sup>fl/fl</sup>KI/KI* mice. Two other *Nfatc1<sup>fl/fl</sup>KI/KI* mice (1 mouse each at 12 and 17 weeks old) had a similar radiographic appearance to the *Nfatc1<sup>Δ/Δ</sup>KI/KI* sample displayed. **(B)** H&E stain of the liver and **(C)** lung of 12-week-old female mice. Black arrows in **B** and **C** indicate examples of inflammatory infiltrates. The original magnifications used to obtain the images are indicated above each column. The organ histology of *Nfatc1<sup>fl/fl</sup>KI/KI* mice was identical to that previously described for mice bearing 2 *KI* alleles (3). A second *Nfatc1<sup>Δ/Δ</sup>KI/KI* mouse had similar histology to the *Nfatc1<sup>Δ/Δ</sup>KI/KI* sample displayed. **(D)** qRT-PCR analysis for the expression of *Tnf* mRNA in BM cells cultured for 5 days with 50 ng/ml M-CSF.



**Figure 7**

NFATc1 is downstream of SH3BP2 in osteoclastogenesis in vitro and in vivo. **(A)** TRAP stain (original magnification, ×100) of BM cells incubated with 50 ng/ml M-CSF and 25 ng/ml RANKL for 4 days. **(B and C)** qRT-PCR analysis for the expression of the indicated genes in BM cells incubated with 50 ng/ml M-CSF and 25 ng/ml RANKL for 3 days. The data in **A–C** is representative of 2 independent experiments. **(D)** Western blot analysis for NFATc1 and β-actin in BM cells incubated with 50 ng/ml M-CSF and 25 ng/ml RANKL for 3 days. **(E)** TRAP stain (original magnification, ×100) of the proximal humerus of 12-week-old female mice. The histology in **E** is representative of at least 3 *Nfatc1<sup>Δ/Δ</sup>KI/KI* mice.

ing NFATc1 autoamplification. Since TNF-α has been reported to induce osteoclast differentiation independent of RANKL (49) and *Sh3bp2* mutant mice have elevated levels of TNF-α (3), osteoclast differentiation was assessed in vivo in cherubism mice lacking NFATc1. Consistent with our observations of increased bone density in *Nfatc1<sup>Δ/Δ</sup>KI/KI* mice (Figure 6A) and failure of BM cells from these mice to differentiate into osteoclasts in vitro (Figure 7, A and B), a dramatic reduction in TRAP-positive cells was seen at the proximal humerus (Figure 7E) and distal femur (A.O. Aliprantis and Y. Ueki, data not shown) of *Nfatc1<sup>Δ/Δ</sup>KI/KI* compared with *Nfatc1<sup>fl/fl</sup>KI/KI* mice. These data imply that TNF-α in this model cannot drive NFATc1-independent osteoclast differentiation.

**Discussion**

Evidence for the role of NFATc1 in osteoclastogenesis in vivo has been provided by 2 other studies. The first showed that *Nfatc1*-deficient fetal liver or embryonic stem cells do not rescue the osteopetrotic phenotype of *c-fos*<sup>-/-</sup> mice (22). The second study targeted NFATc1 expression to the developing heart to rescue

embryonic lethality. *Nfatc1*<sup>-/-</sup>, *Tie2-Nfatc1* mice showed osteopetrosis with absent osteoclasts (23). These studies examined mice at 6 and 3 weeks, respectively, and provided important evidence for the function of NFATc1 in specifying osteoclast differentiation during development. However, the requirement for NFATc1 in adult skeletal remodeling remained unresolved. Here *Mx1-Cre* was used to delete *Nfatc1* in mice at 10 days of age as was previously done to investigate the role of IκB kinase β in osteoclast survival (50). *Nfatc1<sup>Δ/Δ</sup>* mice developed osteoclast-poor osteopetrosis that was most prominent in the long bones and mandibles (Figures 2 and 3). Our data complement these previous studies (22, 23) and extend the requirement for NFATc1 to remodeling of the growing and adult skeleton. Moreover, unlike osteopetrotic, M-CSF-deficient *op/op* mice, which resolve their phenotype with age (51), *Nfatc1<sup>Δ/Δ</sup>* mice up to the age of 5 months still displayed osteopetrosis. Interestingly, TRAP5b levels were reduced but not absent in *Nfatc1<sup>Δ/Δ</sup>* mice (Figure 3C). This observation may be due to variability in *Nfatc1<sup>fl/fl</sup>* deletion by *Mx1-cre* at anatomic sites not examined in the current study (e.g., the vertebral skeleton). In our



experience, within the bone marrow of long bones, *Mx1-cre*-mediated deletion of *Nfatc1<sup>fl/fl</sup>* was nearly complete at the level of DNA recombination or mRNA expression (Figure 1, B and D).

NFATc1 binds the promoters of many osteoclast-specific genes, including *Nfatc1* itself (13–19, 22). However, a source of NFATc1-deficient osteoclast progenitors from mouse BM was not available prior to this study. Therefore, the relative contribution of NFATc1 to the expression of the osteoclast transcriptome could not be determined using loss of function experiments. To facilitate this analysis, a strategy to isolate a pure population of BMOcPs was devised (Supplemental Figure 3). Transcriptional profiling revealed NFATc1 as a broad regulator of osteoclast gene expression (Figure 4). Two sets of NFATc1-regulated genes were identified: those that are augmented by NFATc1 (Figure 4, C and D) but still induced to some extent by RANKL in NFATc1-deficient cells and those that absolutely depend on NFATc1 for expression (Figure 4, A and B). The temporal recruitment of NFATc1 to different genetic loci may underlie this observation. Recently, ChIP analyses have demonstrated that NFATc1 binds the *Itgb3*, *Oscar*, *Acp5*, and *Ctsk* promoters during osteoclastogenesis (16–18, 52). Interestingly, whereas NFATc1 was recruited to the *Itgb3* and *Oscar* promoters (NFATc1-dependent genes) within 12–48 hours of RANKL stimulation (17, 18, 52), NFATc1 was only detected at the *Acp5* and *Ctsk* promoters (NFATc1-augmented genes) at day 5 after RANKL stimulation (16). The fact that we see differences in expression of *Acp5* and *Ctsk* at day 3 (Figure 4D) between NFATc1-sufficient and -deficient cells is likely due to our use of highly osteoclastogenic BMOcPs. We hypothesize that NFATc1-dependent genes need early NFATc1 recruitment to initiate transcription, whereas NFATc1-augmented genes employ NFATc1 at later time points to effect optimal expression. Other possible reasons these 2 sets of genes differentially require NFATc1 include compensatory use of other osteoclast transcription factors, like AP-1, c-Fos, Pu.1, and Mitf (19, 53), and indirect effects of NFATc1 deficiency.

The most intriguing finding from our profiling experiments was the dramatic upregulation of *Tnfrsf11b* mRNA and protein by RANKL in NFATc1-deficient BMOcPs (Figure 5). OPG, a decoy receptor for RANKL, is a potent negative regulator of osteoclastogenesis. Hence, OPG-deficient mice are osteoporotic (54) while OPG transgenic mice, like RANK knockout mice, display increased bone mass (7, 55). Osteoblasts are thought to be the physiologic source of OPG in the skeleton (47). However, contaminating osteoblasts are very unlikely to be the source of OPG in our experiments, since a pure population of BMOcPs, isolated by flow cytometry, was used. Moreover, osteoclast precursors derived from *Nfatc1<sup>Δ/Δ</sup>* spleen cells also upregulated *Tnfrsf11b* mRNA in response to RANKL. Although, the physiologic significance of our observation is presently unknown, we suggest that NFATc1 dampens the induction of OPG that occurs as a consequence of signaling pathways shared by RANK and other TNF receptor family members. The RANKL/RANK/OPG axis has a well-described role outside of the skeleton in regulating adaptive immune responses (9). In human B cells and dendritic cells, CD40 ligation induces OPG expression (56, 57). CD40, like RANK, is a member of the TNF receptor superfamily. Both receptors utilize TNF receptor-associated factor 6 as a proximal signaling component (58, 59). However, RANK is unique in its ability to promote calcium oscillations and upregulate NFATc1 (59). Consistent with these observations, WT BMOcPs stimulated with RANKL in the presence of cyclosporine A, which blocks calcium-mediated activation of the calcineurin/

NFAT pathway, express more *Tnfrsf11b* mRNA (Figure 5F). Furthermore, it is likely that NFATc1 directly, negatively regulates the *Tnfrsf11b* promoter (Figure 5, H and I), which is similar to the observation in previous reports that NFAT family members repress *Cdk4* expression (12, 60). Taken together, these observations suggest that suppression of OPG expression downstream of RANKL is a previously unrecognized function of NFATc1 in osteoclastogenesis. Interestingly, giant cell tumors of bone, which contain multinucleated giant cells with features of osteoclasts, have been shown to express OPG (24, 48). This intriguing clinical correlate suggests that dysregulation of OPG expression by cells of the osteoclast lineage may occur in pathologic conditions.

Our *Nfatc1* conditional knockout mouse provides a genetic system to study the role of this transcription factor in disease. Here, a mouse model of the human disorder, cherubism, was investigated. Cherubism mice carry a gain-of-function mutation in *Sh3bp2* and develop systemic inflammation and bone loss due to overproduction of TNF- $\alpha$  by myeloid cells (3). NFATc1-deficient mice homozygous for the cherubism mutation still developed inflammation in multiple organs (Figure 6, B and C). In addition, macrophages grown from the BM of *Nfatc1<sup>Δ/Δ</sup>* mice had similarly elevated levels of *Tnf* transcripts compared with their NFATc1-sufficient counterparts (Figure 6D). These results indicate that NFATc1 is not downstream of SH3BP2 for TNF- $\alpha$  production and are consistent with a recent report that NFATc1 does not promote TNF- $\alpha$  expression in T cells (30). However, the inflammatory phenotype of cherubism may depend on other NFAT family members, like NFATc2 (30, 61, 62). Genetic ablation of *Nfatc1* did protect cherubism mice from systemic bone loss (Figure 6A) and completely abrogated the enhanced osteoclastogenesis observed in vitro in BM cells carrying mutant alleles of *Sh3bp2* (Figure 7, A and B). Mutant *Sh3bp2* cells also displayed increased induction of the *Nfatc1/A* isoform by RANKL in a manner proportional to gene dosage. This effect was not observed in NFATc1-deficient cells (Figure 7C). Taken together, these observations situate SH3BP2 upstream of NFATc1 in osteoclastogenesis, likely within the NFATc1 autoamplification loop. Cherubism mice also develop TNF- $\alpha$ -dependent arthritis with cortical erosions (3). Whether NFATc1 deficiency will protect from local bone loss around inflamed joints, as has been described for osteoclast-deficient mice in other arthritis models (63, 64), is under investigation.

In conclusion, this study reveals that NFATc1 is an important regulator of osteoclastogenesis in growing and adult mice consistent with previous reports exploring its function in embryonic and early postnatal development (22, 23). NFATc1 deficiency also protects from systemic bone loss in a mouse model of the human disease, cherubism, without inhibiting the inflammatory aspects of the disease. A gene profiling approach revealed that most genes classically associated with terminal osteoclast differentiation require NFATc1 for expression. Intriguingly, we found that RANKL promoted a dramatic induction of OPG in NFATc1-deficient cells. We propose that pharmacologic perturbation of the NFAT pathway in osteoclast precursors may transform these cells into potent OPG producers, which could inhibit resorption in the bone microenvironment and lead to novel treatments for osteolysis.

## Methods

**Creation of an *Nfatc1* conditional knockout allele.** A targeting vector containing exon 3 of *Nfatc1* and a neomycin resistance cassette, both flanked by loxP sites, was generated (Figure 1A) using lambda phage murine *Nfatc1*



DNA kindly provided by Edgar Serfling and Eriks Jankevics of the University of Wuerzburg, Wuerzburg, Germany. As indicated in Figure 1, 3 kb of DNA were deleted upstream of exon 3 in crafting the targeting construct. C57BL/6 ES cells were transfected with the linearized targeting construct (clone 3D5) and antibiotic selected to obtain stable clones, whose integration site was confirmed by Southern blotting. Targeted ES cells were injected into blastocysts to achieve initial germ-line transmission (*Nfatc1<sup>loxNeo</sup>*). *Nfatc1<sup>loxNeo/+</sup>* mice were crossed onto transgenic *EIIA-Cre* mice (65) (The Jackson Laboratory), and progeny were selected by Southern blotting for selective loss of the neomycin resistance cassette to finally generate *Nfatc1<sup>fl/+</sup>* mice, which were intercrossed with other lines used in this study.

**Mice.** The following genetically engineered mice have been previously described: *Nfatc2<sup>-/-</sup>* (gift of Anjana Rao, Harvard University) (66), *Mx1-Cre* (The Jackson Laboratory) (32), *LysM-Cre* (gift of Ira Tabas, Columbia University, New York, New York, USA) (38), and *Osterix-Cre* (gift of Andrew McMahon, Harvard University) (33). Cherubism mice bearing a mutation in *Sh3bp2* were described (3). In all experiments where histology or imaging is reported, *Nfatc1<sup>fl/fl</sup>*, *Mx1-Cre* mice were injected i.p. with 0.25 ml of 1 mg/ml poly I:C (Amersham Biosciences) in PBS every other day starting at postnatal day 10 for a total of 3 doses to generate *Nfatc1<sup>fl/fl</sup>* mice. Littermates without the *Mx1-Cre* transgene were identically treated with poly I:C and served as NFATc1-sufficient (*Nfatc1<sup>fl/fl</sup>*) controls. In some experiments, in which BMOcPs were isolated for cell culture, mice were treated with poly I:C between 6 and 8 weeks of age. In all experiments, littermates were used as controls except in the cherubism studies where mice had been backcrossed 6 generations onto the C57BL/6 background. This allowed us to generate mice with the 6 relevant genotypes from parallel breeding pairs. A pathogen-free facility at the Harvard School of Public Health housed all the mice in this report. Experimental protocols were reviewed and approved by the Standing Committee on Animals at the Harvard Medical School and were designed in accordance with institutional and NIH guidelines for the humane use of animals.

**Southern blotting of genomic DNA and PCR genotyping of *Nfatc1* conditional knockout mice.** Five micrograms of genomic DNA was digested with BamHI (NE Biolabs), resolved by agarose gel electrophoresis, transferred to a Gene Screen membrane (Perkin-Elmer), and crosslinked and hybridized with a radiolabeled BamHI/XbaI (NE Biolabs) 5' probe. Mice were genotyped using the following 3 PCR primers, which discriminate the wild-type, floxed, and  $\Delta$  *Nfatc1* alleles (Figure 1, A and C): INT2-S5, 5'-AAGGAATTACTGGGAAGCCTGGCA-3' (primer a, Figure 1A); INT3-S2, 5'-AGGGACTATCATTTGGCAGGGACA-3' (primer b, Figure 1A); and INT3-AS2, 5'-ACAGGAAACAGCTCTGTTCACAC-3' (primer c, Figure 1A). An annealing temperature of 66°C was used.

**SDS-PAGE and Western blotting.** A single-cell suspension was prepared from spleens incubated for 1 hour with collagenase as described previously (67). Splenocytes were washed with PBS and lysed with RIPA buffer (20 mM Tris, pH 8.0, 150 mM NaCl, 0.1% SDS, 0.5% NaDeoxycholate, 1% Triton X-100) plus protease (Roche) and phosphatase inhibitors. Protein concentrations were determined by the DC protein assay (Bio-Rad). Fifty micrograms of protein per sample were resolved by SDS-PAGE and analyzed by Western blotting using anti-NFATc1 (mAb 7A6; BD Biosciences – Pharmingen) and rabbit anti-mouse HSP90 (Santa Cruz Biotechnology Inc.) antibodies along with the appropriate HRP-conjugated secondary reagents.

**RNA extraction and real-time PCR.** Total RNA was extracted using either RNeasy kits (Qiagen) or TRIzol (Invitrogen) and solubilized in RNAase-free water. cDNA was synthesized using either kits from Stratagene or Bio-Rad. qRT-PCR reactions were performed using the SYBR Green PCR Master Mix (Applied Biosystems) or the Brilliant II SYBR Green QPCR Master Mix (Stratagene) in a PRISM 7700 system (Applied Biosystems) or Mx3005P QPCR System (Stratagene). Ct values for duplicate samples

were averaged and the amounts of mRNA relative to *Hprt* were calculated using the  $\Delta$ Ct method. qRT-PCR primers used in this study are reported in Table 1. All qRT-PCR reactions yielded products with single peak dissociation curves.

**Histology and histomorphometry.** The Musculoskeletal Biology group at the Indiana University School of Medicine performed all histology except for the mandible and cherubism studies. Briefly, formalin-fixed tissues were embedded in methyl methacrylate without decalcification, sectioned with a tungsten-carbide knife, and stained with either (a) Von Kossa/MacNeal tetrachrome, (b) a TRAP protocol counterstained with Gill's Hematoxylin no. 3 (68), or (c) toluidine-blue. Cellular histomorphometry measurements were made in the femoral epiphysis using a  $\times 40$  objective, within 0.25 mm of the cortical surface and growth plate throughout, using a semiautomatic analysis system (Bioquant OSTEO 7.20.10; Bioquant Image Analysis Co.) attached to a Nikon Optiphot 2 microscope (Nikon). The epiphysis was chosen as the site to report cellular histomorphometry, since the parameters to which cell numbers are normalized to were similar in the epiphysis of *Nfatc1<sup>fl/fl</sup>* and *Nfatc1<sup>fl/fl</sup>* mice. A trend toward reduced osteoclast numbers was also observed in the metaphysis of *Nfatc1<sup>fl/fl</sup>* mice. For the analysis of mandibles, formaldehyde-fixed skulls were decalcified for 4.5 days in 10% EDTA, embedded in paraffin, and sectioned starting behind the ear at an angle parallel to the mandibular ramus. Sections were stained with toluidine-blue or for TRAP (69). Histologic analyses for the cherubism studies were performed as described (3).

**Radiography and microquantitative computed tomography.** Plain radiographs were taken on a HP Faxitron 43855A (skulls) or a Faxitron Mx-20 (long bones). Distal femoral bone volume fraction was assessed with microquantitative computed tomography. Femora were immersed in saline and scanned ( $\mu$ CT-40; Scanco Medical) with the following settings 70 kV, 114  $\mu$ A, and 0.012 mm isotropic voxels. After reconstruction, the bone volume fraction was determined by calculating the ratio of bone voxels to total voxels in the volume of interest, which included both cortical and trabecular bone compartments, since the distal femur of *NFATc1<sup>fl/fl</sup>* mice was too dense to separate these compartments (70).

**Purification of highly enriched BMOcPs.** A scheme of the protocol is shown in Supplemental Figure 3A. Briefly, a single-cell suspension of mouse BM cells was incubated for 20 minutes at 5  $\times 10^7$  per ml with anti-CD117 (c-kit)-APC (1  $\mu$ l per 10<sup>7</sup> cells) at 4°C in  $\alpha$ MEM-10 ( $\alpha$ MEM [Cellgro] supplemented with 10% FCS, 1 $\times$  penicillin/streptomycin). Cells were washed twice with ice-cold PBS plus 0.5% BSA and 2 mM EDTA (MACS buffer) and incubated with anti-APC microbeads (Miltenyi Biotec) (2  $\mu$ l per 10<sup>7</sup> cells) for 20 minutes at 4°C. After 2 washes with ice-cold MACS buffer, CD117<sup>+</sup> cells were positively selected on a MACS LS column (Miltenyi Biotec). The CD117 enriched fraction was stained with labeled antibodies (anti-B220-FITC, anti-CD3 $\epsilon$ -Alexa Fluor 488, anti-CD115(c-fms)-biotin/streptavidin-PE, anti-CD11b-PerCP-Cy5.5, and anti-CD117-APC). Cells were sorted at the Harvard Medical School Center for Neurologic Diseases Cell Sorting Facility on a FACS Aria II sorter. All conjugated antibodies were from Becton Dickinson except anti-CD115-biotin, which was from eBioscience. For the quantification of CD11b<sup>lo/-</sup>CD3 $\epsilon$ B220<sup>+</sup>c-fms<sup>+</sup>c-kit<sup>+</sup> and CD11b<sup>lo/-</sup>CD3 $\epsilon$ B220<sup>+</sup>c-fms<sup>+</sup>c-kit<sup>+</sup> osteoclast progenitors, BM cells were stained with an identical cocktail of antibodies and analyzed by flow cytometry.

**Macrophage, osteoclast, and osteoblast cultures.** Cell culture experiments were performed in  $\alpha$ MEM-10 (Cellgro) at 37°C with 5% CO<sub>2</sub>. Four populations of osteoclast progenitors were used: (a) BM cells cultured overnight with M-CSF and purified over a Histopaque 1083 (Sigma-Aldrich) gradient (71), used at an initial density of 1  $\times 10^5$  cells per well in 96-well plates; (b) highly enriched BMOcPs used at an initial density of 2,500 cells per well in 96-well plates for TRAP assays or at 20,000 cells per well in 24-well plates for RNA isolation; (c) crude BM cells for the cherubism studies as described (3); or (d)



spleen cells incubated with M-CSF for 3 days. M-CSF (R&D Systems) and RANKL (RANKL/TRANCE was a kind gift from Y. Choi, University of Pennsylvania, Philadelphia, Pennsylvania, USA) were used at final concentrations of 20 ng/ml and 200–300 ng/ml, respectively. For the cherubism studies, M-CSF and RANKL were purchased from Peprotech and were used at the concentrations indicated in the figure legends. For coculture experiments, neonatal calvarial osteoblasts were purified from BALB/c mice as described (72) and plated at a density of 6,000 cells per well in 96-well plates. After an overnight incubation,  $1 \times 10^5$  M-CSF-primed BM cells, 10 nM Vitamin D<sub>3</sub> (Sigma-Aldrich), and 10  $\mu$ M PGE<sub>2</sub> (Calbiochem) were added. Biochemical assays for TRAP secreted into the culture supernatant (73) or staining of cellular TRAP (using either a commercially available kit [Sigma-Aldrich] or as previously described in ref. 69) were performed after 4 or 5 days (recombinant cytokines) or after 8 days (coculture). Where indicated, osteoblasts were treated for 2 hours with 10 nM Vitamin D<sub>3</sub> (Sigma-Aldrich) and 10 nM parathyroid hormone (Sigma-Aldrich). To generate samples for Southern blot, peritoneal macrophages were collected 3–4 days after i.p. injection of 40  $\mu$ g of Concanavalin A (Sigma-Aldrich) as described previously (74).

**DNA microarray analysis of MROcPs.** Independently prepared duplicate samples containing pooled cells from multiple mice for each genotype were analyzed. Briefly, CD11b<sup>lo</sup>/CD3 $\epsilon$ <sup>+</sup>B220<sup>-</sup>c-kit<sup>+</sup>c-fms<sup>+</sup> BMOcPs from 8- to 13-week-old *Nfatc1*<sup>ΔΔ</sup> and *Nfatc1*<sup>β/β</sup> littermate controls ( $n = 4$ /genotype, replicate 1 and  $n = 3$ /genotype, replicate 2; respectively) were incubated at an initial density of 20,000 per well in 24-well plates, with 1.2 ml  $\alpha$ MEM-10 supplemented with M-CSF and RANKL. After 3 days of culture, mRNA was purified (RNeasy; Qiagen) and RNA profiling was performed by the microarray facility at the Harvard Medical School and Partners Center for Genetics and Genomics using the Affymetrix mouse 430A 2.0 array, which contains probes for approximately 14,000 well-characterized mouse genes. Data for all 4 samples were scaled to 500 and replicate samples were averaged for reporting in Figure 4. A gene was classified as “NFATc1 augmented” if (a) the expression level of that gene was higher in *Nfatc1*<sup>β/β</sup> samples compared with *Nfatc1*<sup>ΔΔ</sup> samples and (b) a “present” detection call was designated in both samples for the probe set recognizing that gene. “NFATc1 dependent” genes had an “absent” detection call for either one or both of the *Nfatc1*<sup>ΔΔ</sup> samples and were “present” at higher expression levels in *Nfatc1*<sup>β/β</sup> samples. A similar experimental design was used to generate independent RNA samples for validation of the microarray results by qRT-PCR.

**In silico analysis of the *Tnfrsf11b* promoter and NFATc1 ChIP assay.** The mouse and human genomic sequences 3.6 kb upstream of the *Tnfrsf11b* gene were extracted from Ensembl database ([www.ensembl.org](http://www.ensembl.org)). Z picture and rVista options in ECR browser (75) were used to identify and align regions of high conservation and putative NFATc1 binding sites in this region, respectively. ChIP assays were performed as described previously (16). Briefly, WT M-CSF-primed spleen cells were incubated with RANKL and immunoprecipitation was carried out on precleared, sheared soluble chromatin with anti-NFATc1 (Santa Cruz Biotechnology Inc.) or nonspecific IgG overnight at 4°C. Immune complexes were isolated using Protein G agarose, washed, decrosslinked, and purified as described previously (16). ChIP samples were analyzed by qRT-PCR for the *Tnfrsf11b* promoter using the FastStart TaqMan Master kit (Roche) and the universal probe library (Roche Diagnostics) with the following primers: 5'-TTAGGGAATACCTCAGGAAATACA-3' (forward) and 5'-TTGTAGGAGCACGAGGTGAA-3' (reverse). Analysis for the *Itgb3* promoter was performed as described (18). The signal for the promoter being studied was adjusted using input threshold values as a reference and is reported as relative enrichment.

**Promoter luciferase assays.** A firefly luciferase reporter plasmid containing 3.6 kb of the *Tnfrsf11b* proximal promoter (*pOPG 3.6-luc*) was a gift of G. Karsenty, Columbia University (76). An expression plasmid for constitutively active NFATc1 (*pMSCV-caNfatc1*) was a gift of N. Clipstone, Northwestern

University, Chicago, Illinois, USA (77). 293T cells were transiently transfected using the Effectene Reagent (Qiagen) with the indicated luciferase reporter plasmids and a *Renilla* luciferase vector (*pRL-TK*) as an internal control at a ratio of 10:1, along with the indicated amounts of *pMSCV-caNfatc1*. The total amount of DNA per transfection was held constant by supplementing with empty vector DNA. Forty hours after transfection *Firefly* and *Renilla* luciferase activities were determined in cell lysates using the Dual-Luciferase Reporter Assay System (Promega). *Firefly* luciferase activity was normalized to *Renilla* luciferase to control for transfection efficiency.

**Immunofluorescence of NFATc1.** BMOcPs were cultured on chamber slides for 3 days with M-CSF and RANKL. Cells were fixed with 4% PFA, permeabilized, and stained with anti-NFATc1 and goat anti-mouse Alexa Fluor 488 (Invitrogen). Hoechst (Sigma-Aldrich) was used as a nuclear counterstain.

**Quantification of TRAP5b and OPG.** Commercially available kits were used to determine the concentrations of TRAP5b (Immunodiagnostic Systems) in mouse serum or OPG (R&D Systems) in mouse serum or tissue culture supernatants according to the manufacturer's directions. Since RANKL in the tissue culture supernatants of osteoclast cultures can interfere with the OPG ELISA, supernatants were collected from BMOcPs first incubated for 4 days with M-CSF or M-CSF and RANKL and then cultured for 2 days with M-CSF only.

**Statistics.** Unpaired, 2-tailed Student's *t* tests were used for all statistical comparisons. A *P* value of less than 0.05 was considered significant.

## Acknowledgments

The authors especially thank Stanford Peng, Geancarlo Lugo-Villarino, and Arlene Sharpe for their early participation in the NFATc1 conditional knockout project. We also acknowledge Dorothy Zhang, Kazuki Ueno, David Burr, and Keith Condon for their preparation and analysis of histologic sections; Deneen Kozoriz for cell sorting; Landy Kangaloo, Steven Swartz, Pamela Okerholm, and Jennifer Donovan for technical assistance; Zie Skobe, Roberto Fajardo, and Pat Dunning for help with radiology; Kyleigh Nevis for manuscript preparation; and Dallas Jones, Ann-Hwee Lee, Marc Wein, Jing Wu, Ellen Gravallese, Kevin McHugh, Tania Crotti, and Nicole Walsh for guidance and experimental advice. We are indebted to the staff at the Harvard Medical School Center for Animal Resources and Comparative Medicine for their excellent care of our mice. This work was supported by grants from the Charles H. Hood Foundation (to Y. Ueki), NIH AR36819 (to B.R. Olsen), NIH AI31541 (to L.H. Glimcher), and NIH AR0447129 (to M.C. Ostrowski). A.O. Aliprantis is the recipient of the Abbott Scholar Award in Rheumatology Research and the 2007 American Society for Clinical Investigation Young Investigator Award. R. Sulyanto is a recipient of the Harvard Medical School Carl W. Walter Fellowship.

Received for publication March 21, 2008, and accepted in revised form September 3, 2008.

Address correspondence to: Laurie H. Glimcher, Harvard School of Public Health, 651 Huntington Avenue, FXB 205, Boston, Massachusetts 02115, USA. Phone: (617) 432-0622; Fax: (617) 432-0084; E-mail: [lglimche@hsph.harvard.edu](mailto:lglimche@hsph.harvard.edu).

Yasuyoshi Ueki's present address is: Department of Oral Biology, School of Dentistry, University of Missouri-Kansas City, Kansas City, Missouri, USA.

Yasuyoshi Ueki and Rosalyn Sulyanto contributed equally to this work.



1. United States Department of Health and Human Services. 2004. Bone health and osteoporosis: a report of the Surgeon General. <http://www.surgeon-general.gov/library/bonehealth/content.html>.
2. Woolf, A.D., and Pfleger, B. 2003. Burden of major musculoskeletal conditions. *Bull. World Health Organ.* **81**:646–656.
3. Ueki, Y., et al. 2007. Increased myeloid cell responses to M-CSF and RANKL cause bone loss and inflammation in SH3BP2 “cherubism” mice. *Cell.* **128**:71–83.
4. Goldring, S.R., and Gravalles, E.M. 2000. Pathogenesis of bone erosions in rheumatoid arthritis. *Curr. Opin. Rheumatol.* **12**:195–199.
5. Wada, T., Nakashima, T., Hiroshi, N., and Penninger, J.M. 2006. RANKL-RANK signaling in osteoclastogenesis and bone disease. *Trends Mol. Med.* **12**:17–25.
6. Teitelbaum, S.L., and Ross, F.P. 2003. Genetic regulation of osteoclast development and function. *Nat. Rev. Genet.* **4**:638–649.
7. Simonet, W.S., et al. 1997. Osteoprotegerin: a novel secreted protein involved in the regulation of bone density. *Cell.* **89**:309–319.
8. Bucay, N., et al. 1998. Osteoprotegerin-deficient mice develop early onset osteoporosis and arterial calcification. *Genes Dev.* **12**:1260–1268.
9. Boyce, B.F., and Xing, L. 2007. Biology of RANK, RANKL, and osteoprotegerin. *Arthritis Res. Ther.* **9**(Suppl. 1):S1.
10. Hogan, P.G., Chen, L., Nardone, J., and Rao, A. 2003. Transcriptional regulation by calcium, calcineurin, and NFAT. *Genes Dev.* **17**:2205–2232.
11. Graef, I.A., Chen, F., and Crabtree, G.R. 2001. NFAT signaling in vertebrate development. *Curr. Opin. Genet. Dev.* **11**:505–512.
12. Horsley, V., Aliprantis, A.O., Polak, L., Glimcher, L.H., and Fuchs, E. 2008. NFATc1 balances quiescence and proliferation of skin stem cells. *Cell.* **132**:299–310.
13. Ishida, N., et al. 2002. Large scale gene expression analysis of osteoclastogenesis in vitro and elucidation of NFAT2 as a key regulator. *J. Biol. Chem.* **277**:41147–41156.
14. Takayanagi, H., et al. 2002. Induction and activation of the transcription factor NFATc1 (NFAT2) integrate RANKL signaling in terminal differentiation of osteoclasts. *Dev. Cell.* **3**:889–901.
15. Kim, K., Lee, S.H., Ha Kim, J., Choi, Y., and Kim, N. 2008. NFATc1 induces osteoclast fusion via up-regulation of Atp6v0d2 and the dendritic cell-specific transmembrane protein (DC-STAMP). *Mol. Endocrinol.* **22**:176–185.
16. Sharma, S.M., et al. 2007. MITF and PU.1 recruit p38 MAPK and NFATc1 to target genes during osteoclast differentiation. *J. Biol. Chem.* **282**:15921–15929.
17. Kim, Y., et al. 2005. Contribution of nuclear factor of activated T cells c1 to the transcriptional control of immunoreceptor osteoclast-associated receptor but not triggering receptor expressed by myeloid cells-2 during osteoclastogenesis. *J. Biol. Chem.* **280**:32905–32913.
18. Crotti, T.N., et al. 2008. PU.1 and NFATc1 mediate osteoclastic induction of the mouse beta3 integrin promoter. *J. Cell. Physiol.* **215**:636–644.
19. Sharma, S.M., et al. 2006. Genetics and genomics of osteoclast differentiation: integrating cell signaling pathways and gene networks. *Crit. Rev. Eukaryot. Gene Expr.* **16**:253–277.
20. Ranger, A.M., et al. 1998. The transcription factor NF-ATc is essential for cardiac valve formation. *Nature.* **392**:186–190.
21. Luis de la Pompa, J., et al. 1998. Role of the NF-ATc transcription factor in morphogenesis of cardiac valves and septum. *Nature.* **392**:182–186.
22. Asagiri, M., et al. 2005. Autoamplification of NFATc1 expression determines its essential role in bone homeostasis. *J. Exp. Med.* **202**:1261–1269.
23. Winslow, M.M., et al. 2006. Calcineurin/NFAT signaling in osteoblasts regulates bone mass. *Dev. Cell.* **10**:771–782.
24. Liu, B., Yu, S.F., and Li, T.J. 2003. Multinucleated giant cells in various forms of giant cell containing lesions of the jaws express features of osteoclasts. *J. Oral Pathol. Med.* **32**:367–375.
25. Ueki, Y., et al. 2001. Mutations in the gene encoding c-Abl-binding protein SH3BP2 cause cherubism. *Nat. Genet.* **28**:125–126.
26. Tiziani, V., et al. 1999. The gene for cherubism maps to chromosome 4p16. *Am. J. Hum. Genet.* **65**:158–166.
27. Lietman, S.A., Kalinchinko, N., Deng, X., Kohanski, R., and Levine, M.A. 2006. Identification of a novel mutation of SH3BP2 in cherubism and demonstration that SH3BP2 mutations lead to increased NFAT activation. *Hum. Mutat.* **27**:717–718.
28. Weitzmann, M.N., and Pacifici, R. 2005. The role of T lymphocytes in bone metabolism. *Immunol. Rev.* **208**:154–168.
29. Peng, S.L., Gerth, A.J., Ranger, A.M., and Glimcher, L.H. 2001. NFATc1 and NFATc2 together control both T and B cell activation and differentiation. *Immunity.* **14**:13–20.
30. Kaminuma, O., et al. 2008. Differential contribution of NFATc2 and NFATc1 to TNF-alpha gene expression in T cells. *J. Immunol.* **180**:319–326.
31. Kuhn, R., and Torres, R.M. 2002. Cre/loxP recombination system and gene targeting. *Methods Mol. Biol.* **180**:175–204.
32. Kuhn, R., Schwenk, F., Aguet, M., and Rajewsky, K. 1995. Inducible gene targeting in mice. *Science.* **269**:1427–1429.
33. Rodda, S.J., and McMahon, A.P. 2006. Distinct roles for Hedgehog and canonical Wnt signaling in specification, differentiation and maintenance of osteoblast progenitors. *Development.* **133**:3231–3244.
34. Kawata, T., et al. 1999. Recruitment of osteoclasts in the mandibular condyle of growing osteopetrotic (op/op) mice after a single injection of macrophage colony-stimulating factor. *Arch. Oral Biol.* **44**:81–88.
35. Tolar, J., Teitelbaum, S.L., and Orchard, P.J. 2004. Osteopetrosis. *N. Engl. J. Med.* **351**:2839–2849.
36. Jacquin, C., Gran, D.E., Lee, S.K., Lorenzo, J.A., and Aguila, H.L. 2006. Identification of multiple osteoclast precursor populations in murine bone marrow. *J. Bone Miner. Res.* **21**:67–77.
37. Leenen, P.J., de Bruijn, M.F., Voerman, J.S., Campbell, P.A., and van Ewijk, W. 1994. Markers of mouse macrophage development detected by monoclonal antibodies. *J. Immunol. Methods.* **174**:5–19.
38. Clausen, B.E., Burkhardt, C., Reith, W., Renkawitz, R., and Forster, I. 1999. Conditional gene targeting in macrophages and granulocytes using LysMcre mice. *Transgenic Res.* **8**:265–277.
39. Kenner, L., et al. 2004. Mice lacking JunB are osteopenic due to cell-autonomous osteoblast and osteoclast defects. *J. Cell Biol.* **164**:613–623.
40. Kim, H.J., et al. 2006. Glucocorticoids suppress bone formation via the osteoclast. *J. Clin. Invest.* **116**:2152–2160.
41. Kurihara, N., Tatsumi, J., Arai, F., Iwama, A., and Suda, T. 1998. Macrophage-stimulating protein (MSP) and its receptor, RON, stimulate human osteoclast activity but not proliferation: effect of MSP distinct from that of hepatocyte growth factor. *Exp. Hematol.* **26**:1080–1085.
42. Sato, M., and Grasser, W. 1990. Myosin II antibodies inhibit the resorption activity of isolated rat osteoclasts. *Cell. Motil. Cytoskeleton.* **17**:250–263.
43. Wu, H., et al. 2007. Down syndrome critical region-1 is a transcriptional target of nuclear factor of activated T cells-c1 within the endocardium during heart development. *J. Biol. Chem.* **282**:30673–30679.
44. Sundaram, K., et al. 2007. RANK ligand signaling modulates the matrix metalloproteinase-9 gene expression during osteoclast differentiation. *Exp. Cell Res.* **313**:168–178.
45. Sato, T., et al. 1997. Identification of the membrane-type matrix metalloproteinase MT1-MMP in osteoclasts. *J. Cell Sci.* **110**:589–596.
46. Wu, H., Peisley, A., Graef, I.A., and Crabtree, G.R. 2007. NFAT signaling and the invention of vertebrates. *Trends Cell Biol.* **17**:251–260.
47. Kieslinger, M., et al. 2005. EBF2 regulates osteoblast-dependent differentiation of osteoclasts. *Dev. Cell.* **9**:757–767.
48. Atkins, G.J., et al. 2001. Osteoprotegerin inhibits osteoclast formation and bone resorbing activity in giant cell tumors of bone. *Bone.* **28**:370–377.
49. Kim, N., et al. 2005. Osteoclast differentiation independent of the TRANCE-RANKL-TRAF6 axis. *J. Exp. Med.* **202**:589–595.
50. Ruocco, M.G., et al. 2005. I{kappa}B kinase (IKK){beta}, but not IKK{alpha}, is a critical mediator of osteoclast survival and is required for inflammation-induced bone loss. *J. Exp. Med.* **201**:1677–1687.
51. Begg, S.K., et al. 1993. Delayed hematopoietic development in osteopetrotic (op/op) mice. *J. Exp. Med.* **177**:237–242.
52. Kim, K., et al. 2005. Nuclear factor of activated T cells c1 induces osteoclast-associated receptor gene expression during tumor necrosis factor-related activation-induced cytokine-mediated osteoclastogenesis. *J. Biol. Chem.* **280**:35209–35216.
53. Asagiri, M., and Takayanagi, H. 2007. The molecular understanding of osteoclast differentiation. *Bone.* **40**:251–264.
54. Mizuno, A., et al. 1998. Severe osteoporosis in mice lacking osteoclastogenesis inhibitory factor/osteoprotegerin. *Biochem. Biophys. Res Commun.* **247**:610–615.
55. Li, J., et al. 2000. RANK is the intrinsic hematopoietic cell surface receptor that controls osteoclastogenesis and regulation of bone mass and calcium metabolism. *Proc. Natl. Acad. Sci. U. S. A.* **97**:1566–1571.
56. Yun, T.J., et al. 1998. OPG/FDCR-1, a TNF receptor family member, is expressed in lymphoid cells and is up-regulated by ligating CD40. *J. Immunol.* **161**:6113–6121.
57. Schoppert, M., et al. 2007. Osteoprotegerin expression in dendritic cells increases with maturation and is NF-kappaB-dependent. *J. Cell. Biochem.* **100**:1430–1439.
58. Naito, A., et al. 1999. Severe osteopetrosis, defective interleukin-1 signalling and lymph node organogenesis in TRAF6-deficient mice. *Genes Cells.* **4**:353–362.
59. Gohda, J., et al. 2005. RANK-mediated amplification of TRAF6 signaling leads to NFATc1 induction during osteoclastogenesis. *EMBO J.* **24**:790–799.
60. Baksh, S., et al. 2002. NFATc2-mediated repression of cyclin-dependent kinase 4 expression. *Mol. Cell.* **10**:1071–1081.
61. Hodge, M.R., et al. 1996. Hyperproliferation and dysregulation of IL-4 expression in NF-ATp-deficient mice. *Immunity.* **4**:397–405.
62. Tsytysykova, A.V., and Goldfeld, A.E. 2000. Nuclear factor of activated T cells transcription factor NFATp controls superantigen-induced lethal shock. *J. Exp. Med.* **192**:581–586.
63. Pettit, A.R., et al. 2001. TRANCE/RANKL knockout mice are protected from bone erosion in a serum transfer model of arthritis. *Am. J. Pathol.* **159**:1689–1699.
64. Redlich, K., et al. 2002. Osteoclasts are essential for TNF-alpha-mediated joint destruction. *J. Clin. Invest.* **110**:1419–1427.
65. Xu, X., et al. 2001. Direct removal in the mouse of a floxed neo gene from a three-loxP conditional knockout allele by two novel approaches. *Genesis.*



- 30:1–6.
66. Xanthoudakis, S., et al. 1996. An enhanced immune response in mice lacking the transcription factor NFAT1. *Science*. **272**:892–895.
67. Lugo-Villarino, G., Maldonado-Lopez, R., Possemato, R., Penaranda, C., and Glimcher, L.H. 2003. T-bet is required for optimal production of IFN-gamma and antigen-specific T cell activation by dendritic cells. *Proc. Natl. Acad. Sci. U. S. A.* **100**:7749–7754.
68. Erlebacher, A., and Derynck, R. 1996. Increased expression of TGF-beta 2 in osteoblasts results in an osteoporosis-like phenotype. *J. Cell Biol.* **132**:195–210.
69. Takahashi, N., Udagawa, N., Tanaka, S., and Suda, T. 2003. Generating murine osteoclasts from bone marrow. *Methods Mol. Med.* **80**:129–144.
70. Fajardo, R.J., Muller, R., Ketcham, R.A., and Colbert, M. 2007. Nonhuman anthropoid primate femoral neck trabecular architecture and its relationship to locomotor mode. *Anat. Rec. (Hoboken)*. **290**:422–436.
71. McHugh, K.P., et al. 2000. Mice lacking beta3 integrins are osteosclerotic because of dysfunctional osteoclasts. *J. Clin. Invest.* **105**:433–440.
72. Jones, D.C., et al. 2006. Regulation of adult bone mass by the zinc finger adapter protein Schnurri-3. *Science*. **312**:1223–1227.
73. Jankila, A.J., Takahashi, K., Sun, S.Z., and Yam, L.T. 2001. Naphthol-ASBI phosphate as a preferred substrate for tartrate-resistant acid phosphatase isoform 5b. *J. Bone Miner. Res.* **16**:788–793.
74. Smith, C.W., and Goldman, A.S. 1972. Effects of concanavalin A and pokeweed mitogen in vivo on mouse peritoneal macrophages. *Exp. Cell Res.* **73**:394–398.
75. Loots, G., and Ovcharenko, I. 2007. ECRbase: database of evolutionary conserved regions, promoters, and transcription factor binding sites in vertebrate genomes. *Bioinformatics*. **23**:122–124.
76. Glass, D.A., 2nd, et al. 2005. Canonical Wnt signaling in differentiated osteoblasts controls osteoclast differentiation. *Dev. Cell*. **8**:751–764.
77. Hirotsani, H., Tuohy, N.A., Woo, J.T., Stern, P.H., and Clipstone, N.A. 2004. The calcineurin/nuclear factor of activated T cells signaling pathway regulates osteoclastogenesis in RAW264.7 cells. *J. Biol. Chem.* **279**:13984–13992.
78. Wang, X., and Seed, B. 2003. A PCR primer bank for quantitative gene expression analysis. *Nucleic Acids Res.* **31**:e154.