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TITLE: ATF4, A Novel Mediator of the Anabolic Actions of PTH on Bone

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In this study, we have successfully demonstrated that ATF4 plays a critical role in mediating the anabolic effects of intermittent PTH on bones. ATF4 is important for intermittent PTH to stimulate bone formation in mice. ATF4 favors bone formation through, at least in part, upregulation of osteoblast proliferation and survival. Additionally, ATF4 increases osteoblast differentiation probably via osterix. At molecular level, ATF4 increases osteocalcin gene expression by cooperative interactions with TFIIAγ and Runx2. ATF4 increases the expression of cyclin D1, a key factor for cell cycle progression. We have identified and functionally characterized Erk/MAPK phosphorylation sites in Runx2, an ATF4-interacting factor. We have demonstrated that ATF4 is essential for osteoclast differentiation and bone resorption, which is increased by intermittent PTH.
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**This progress report covers research from the period 07/01/07-12/31/11**

**Introduction**

Osteoporosis, or reduced bone mass, is a metabolic bone disease that affects millions of people including many of our service women and men now in the Armed Forces and VA patients in the United States. It causes a significant amount of morbidity and mortality in patients and is often diagnosed after a fracture occurs. Reducing the risk of osteoporotic and associated fractures of these patients will greatly improve their life quality and survival. Parathyroid hormone (PTH) is the most potent anabolic treatment of osteoporosis currently available. It not only dramatically improves bone mass, but also restores bone microarchitecture and increases bone diameter. All of these mechanisms contribute to increasing bone strength and reducing the risk for fractures. However, the molecular mechanisms whereby PTH increases bone mass are not well understood and are the focus of the experiments of this proposal. This study was designed to (1) determine the mechanism whereby PTH regulates ATF4 transcriptional activity (2) establish whether the anabolic actions of PTH require ATF4 in vivo. Studies determine if ATF4 is required for the anabolic actions of PTH in vivo using ATF4-deficient mice. PTH anabolic activity is evaluated in wild type and Atf4−/− mice. PTH effects are measured using standard biochemical and histomorphometric criteria.

**Body**

*Task 1: To determine the mechanism by which PTH regulates ATF4 transcriptional activity (1-36 months).*

1) Establish that ATF4 is a novel downstream target of PTH actions in cultured osteoblasts (P1).

In order to determine the role of ATF4 in PTH actions in osteoblasts, we examined effects of PTH on ATF4 expression and activity as well as the requirement for ATF4 in the regulation of *Ocn* by PTH (see P1). PTH elevated levels of ATF4 mRNA and protein in a dose and time-dependent manner (P1-Fig. 1A-C). This PTH regulation requires transcriptional activity, but not de novo protein synthesis (P1-Fig. 2A and B). PTH also increased binding of nuclear extracts to OSE1 DNA (P1-Fig. 4A-C). PTH stimulated ATF4-dependent transcriptional activity mainly through PKA with a lesser requirement for PKC and MAPK/ERK pathways (P1-Fig. 5A-C). PTH stimulation of *Ocn* expression was lost by siRNA downregulation of ATF4 in MC-4 cells (P1-Fig. 6A and C) and in Atf4−/− bone marrow stromal cells (BMSCs) (P1-Fig. 7C). Collectively, these studies for the first time demonstrate that PTH increases ATF4 expression and activity and that ATF4 is required for PTH induction of *Ocn* expression in osteoblasts. Thus, ATF4 is a novel downstream target of PTH actions in osteoblasts.

2) Define a novel molecular mechanism through which ATF4 activates osteocalcin gene expression in osteoblasts (P2).

We defined a novel molecular mechanism mediating ATF4-Runx2 interactions (see P2). We identified general transcription factor IIAγ (TFIIAγ) as a Runx2-interacting factor in a yeast two-hybrid screen. Immunoprecipitation assays confirmed that TFIIAγ interacted with Runx2 in osteoblasts and when coexpressed in COS-7 cell or using purified GST-fusion proteins (P2-Fig. 1A-C). Chromatin immunoprecipitation (ChIP) assay of MC3T3-E1 (clone MC-4) preosteoblast cells showed that in intact cells TFIIAγ was recruited to the region of the osteocalcin promoter previously shown to bind Runx2 and ATF4 (P2-Fig. 2). A small region of Runx2 (aa 258-286) was found to be required for TFIIAγ binding (P2-Fig. 1D). While TFIIAγ interacted with Runx2, it did not activate Runx2 (P2-Fig. 3A and B). Instead, TFIIAγ bound to and activates ATF4 (P2-Fig. 3C-H). Further, TFIIAγ together with ATF4 and Runx2 stimulated osteocalcin promoter activity (P2-Fig. 5B) and endogenous mRNA expression (P2-Fig. 5A). siRNA silencing of TFIIAγ markedly reduced levels of endogenous ATF4 protein and *Ocn* mRNA in osteoblastic cells (P2-Fig. 6). Overexpression of TFIIAγ increased levels of ATF4 protein (P2-Fig. 7). TFIIAγ significantly prevented ATF4 degradation (P2-Fig. 8). Thus, TFIIAγ functions as a bridging protein linking ATF4 and Runx2. Interestingly, our preliminary studies (not shown) show that PTH increases ATF4-Runx2 interaction as measured co-immunoprecipitation assays.
1) Establish a role for ATF4 in regulating osteoblast proliferation and survival in vitro and in bones (P3 and P5).

In order to determine the mechanism where ATF4 regulates osteoblast activity, we examined the effects of ATF4 on osteoblast proliferation and survival (see P3 and P5). As shown in P3-Fig. 1B, the number of osteoblast precursor colonies (i.e., CFU-OBs) was significantly decreased in Atf4 knockout mice compared to wt mice. This decrease could be explained by a cell-autonomous defect in proliferation and/or survival or could be secondary to an impaired bone microenvironment due to ATF4 deficiency. To differentiate these possibilities, we conducted MTS and [3H] thymidine incorporation assays. Results from MTS assay showed that ATF4-deficient cells grew at significantly reduced rates compared to wt cells (P3-Fig. 1C). This result was further confirmed by the [3H] thymidine incorporation assays showing that [3H] thymidine incorporation into the DNA of Atf4 knockout bone marrow stromal cells (BMSCs) was decreased by 4-fold compared to values in wt cells (P3-Fig. 1D). We next determined if ATF4 is required for osteoblast proliferation in vivo using 6-wk-old wt and Atf4 knockout mice that were injected with bromodeoxyuridine (BrdU)/fluorodeoxyuridine (FdU) 4 h before sacrifice. 10-μm sections of calvariae were obtained, and proliferating cells from the periosteal surface were counted and normalized to total cells from the same area. In wt calvariae, periosteal osteoblasts proliferated very actively with 60% of the total cells being BrdU-positive (P3-Fig. 2A). In contrast, the percent BrdU-positive osteoblasts were significantly reduced in Atf4 knockout calvariae (P3-Fig. 2B and C). Thus, ATF4 is required for the proliferation of BMSCs or osteoblasts both in vitro and in vivo. As shown in P3-Fig. 3, ATF4-deficient cells showed a significant decrease in the cell distribution into both S and G2/M phases (32% and 45% change, respectively) when compared with values of wt control cells. In contrast, the fraction of the cells in G1 was not reduced by ATF4 deficiency. The protein level of cyclin D1 was markedly decreased in Atf4 knockout cells relative to wt control (P3-Fig. 4A). Quantitative real-time RT/PCR analysis shows that the level of cyclin D1 mRNA was reduced by 52% in Atf4 knockout cells compared to values in wt controls (P3-Fig. 4B). ATF4 is also required for the proliferation of the pre-osteoblast cell line (MC-4 cells) as demonstrated by siRNA experiments (P3-Fig. 5).

Furthermore, overexpression of ATF4 significantly increased the level of cyclin D1 protein as well as cell proliferation (P3-Fig. 6 and 7). Taken together, these results clearly establish that ATF4 increases cell proliferation probably via promoting the expression of cyclin D1 and cell cycle progression.

We also found that ATF4 is critical for osteoblast survival. The numbers of apoptotic cells including those with shrinking cytoplasm and chromatin condensation (early apoptosis) and DNA fragmentation (late apoptosis) were increased greater than 5-fold in Atf4 knockout BMSCs compared to wt cells (P3-8A-C). To confirm this finding, wt and Atf4 knockout BMSCs were stained using the ApopTag Peroxidase In Situ Apoptosis Detection Kit, a modified TUNEL staining that measures DNA fragmentation in situ. As shown in P3-Fig. 8D-F, the percent apoptotic cells in Atf4 knockout BMSCs were increased by 1.6-fold when compared to wt cells. To determine if ATF4 deficiency increases osteoblast apoptosis in vivo, 10-μm calvarial sections from wt and Atf4 knockout mice were obtained and stained using the same kit. Apoptotic cells that stained brown on the periosteal surface of calvariae were counted and normalized to total cells of the same periosteal surface. As shown in P3-Fig. 8G-I, a significant increase in apoptosis was found in Atf4 knockout mice compared to wt controls. Thus, ATF4 protects osteoblasts from apoptosis under physiological condition.

To define the mechanism whereby ATF4 regulates osteoblast function and bone formation, we further determined whether ATF4 plays a role in PTH regulation of osteoblast proliferation in the presence and absence of PTH in vivo. Sections of tibiae and calvariae from wt and Atf4 knockout mice treated with and without intermittent PTH were analyzed for in vivo cell proliferation using a Zymed BrdU immunostaining kit. As shown in Fig 4A, B, and E (P5), in wt mice, PTH increased the percentage of proliferating osteoblasts/preosteoblasts of tibial trabeculae by 2.8-fold relative to vehicle-treated control. Ablation of the Atf4 gene resulted in a 50% decline in basal proliferation. In addition, the PTH-stimulated increase in proliferation was decreased by 40 percent (Fig 4C-E, P5). Similarly, PTH-induced proliferation in calvarial periosteal osteoblasts was also significantly reduced by ATF4 deficiency (Fig 4F-J, P5). Therefore, ATF4 is critical for both basal and PTH-stimulated proliferation of osteoblasts/preosteoblasts in vivo.

Mature osteoblasts synthesize and deposit a mineralizing extracellular matrix and become osteocytes. Both osteoblasts and osteocytes can be lost through apoptosis. PTH signaling increases the survival of osteoblasts and osteocytes by reducing apoptosis. ATF4 is anti-apoptotic in osteoblasts as described above. To determine whether ATF4 plays a role in PTH-mediated anti-apoptosis, sections of tibiae were stained with TUNEL and apoptotic cells were assessed. As shown in Fig 4K and L (P5), ATF4 deficiency significantly increased the basal levels of apoptosis. As expected, PTH dramatically reduced apoptotic death of tibial trabecular osteoblasts/osteocytes by 48 percent. Importantly, the PTH-stimulated decrease in apoptotic death was completely abolished in Atf4 knockout trabeculae (Fig 4M-
O, P5). ATF4 was similarly required for PTH to inhibit apoptosis in cortical osteocytes of tibiae (Fig 4P, P5). Collectively, ATF4 is essential for PTH-mediated inhibition of apoptosis in osteoblasts/osteocytes in vivo.

4) **ATF4 is essential for PTH to increase osteoblast differentiation in vivo (P5).**

We determined the effects of ATF4 deficiency on PTH induction of osteoblast differentiation markers in vivo. Total RNA was isolated from tibiae of wt and 

\[ \text{Atf4}^-/- \] mice treated with and without PTH for 28 d and expression levels of osteoblast differentiation marker genes were measured by quantitative real-time PCR analysis. As shown in Fig 5A (P5), PTH dramatically increased the expression of genes known to be associated with osteoblast differentiation including \text{osteocalcin (Ocn)}, \text{bone sialoprotein (Bsp)}, \text{alkaline phosphatase (Alp)}, \text{α(I) collagen (Col1(I))}, \text{osteopontin (Opn)}, and \text{osterix (Osx)}. Importantly, this PTH regulation was either dramatically reduced or completely abolished in \text{Atf4}^-/- tibiae. In contrast, c-Fos and c-Jun, both early PTH-induced genes, were not induced by PTH in either wt or \text{Atf4}^-/- tibiae.

5) **ATF4 is a novel upstream transcriptional activator of osterix (Osx), a key factor for osteoblast differentiation and bone formation (P5).**

To address the molecular mechanism whereby PTH promotes osteoblast differentiation, we measured the expression of Osterix (Osx) and Runx2 proteins, two critical transcription factors that regulate osteoblast differentiation. We used immunohistochemistry (IHC) to measure Osx in the tibiae and calvariae of wt and \text{Atf4}^-/- mice with or without 28 d anabolic PTH treatment. As shown in Fig 6 (P5), in wt-vehicle-treated tibiae, Osx-positive osteoblasts were only identified in the trabeculae and cortical endosteum close to the growth plate (Fig 6A1, B1, and D1, P5) and were almost undetectable in the same regions close to the marrow (Fig 6C1, P5), indicating that cells in these areas are still in the immature (preosteoblast) state. In contrast, in the wt-PTH group, Osx-positive osteoblasts were identified on all surfaces of trabeculae and endosteum throughout the tibia. PTH increased the total number of Osx-positive cells per tibial section by 3.2-fold in wt mice (Panel 2). ATF4 deficiency reduced the numbers of Osx-positive cells by 50 percent (Panel 3). Strikingly, PTH failed to elevate the numbers of Osx-positive cells in \text{Atf4}^-/- tibiae. Similar results were obtained in calvariae (Fig 6E, P6). The IHC staining was highly specific since no signal was detected in the IgG control group (Panel 5), in which Osx antibody was omitted.

To define the mechanism whereby ATF4 regulates Osx, we examined the effect of ATF4 overexpression on the expression of Osx, a key transcription factor for osteoblast differentiation, in MC-4 preosteoblast cells. As shown in Fig 7A (P5), ATF4 dose-dependently increased levels of Osx protein (top) and mRNA (bottom). We next examined whether ATF4 up-regulates Osx by increasing gene transcription by using a -1003/+68 mouse Osx promoter (Fig 7B, P5). Using COS-7 cells, which lack detectable Runx2, ATF4 had comparable activity to Runx2 in terms of its ability to activate promoter activity (approx. 1.8-fold). Together, ATF4 and Runx2 maximally activated the Osx promoter (3.2-fold induction). To further define the region of the Osx promoter necessary for ATF4 responsiveness, several constructs containing various deletion mutants of the mouse Osx promoter were transiently transfected into COS-7 cells with and without an ATF4 expression plasmid. Results showed that luciferase activity of both control and ATF4-transfected groups decreased with progressively larger 5' deletions. However, ATF4 stimulation was abrogated when a 132-bp region between bp -215 to -83 was deleted (Fig 7C, P5). A putative ATF4-binding sequence (CTTCCTCA) at -201/-194 bp was identified in this region by using a TRANSFAC retrieval program. Introduction of a 3-bp substitution mutation to this core sequence (from CTTCCTCA to CTTgtaCA) completely abolished ATF4 activation (Fig 7D, P5). As shown in Fig 7E (P5), a DNA oligo probe from the Osx promoter that contains the TTACATCA core sequence bound to a factor(s) in nuclear extracts from COS-7 cells transfected with an ATF4 expression vector. Importantly, this binding (see arrow) was dramatically reduced by the addition of a specific antibody against ATF4 but not by normal control IgG or antibodies against cFos (an AP1 family member) or ATF2.

6) **Identification and characterization of Erk/MAPK phosphorylation sites in Runx2 (P4).**

We used a combination of in vitro and in vivo phosphorylation analysis, mass spectroscopy and functional assays to identify two sites at S301 and S319 within the proline/serine/threonine domain of Runx2 that are required for MAPK regulation. These sites are phosphorylated by activated Erk1 in vitro and in cell culture. In addition to confirming Erk-dependent phosphorylation at S319, mass spectroscopy identified two other Erk-phosphorylated sites...
at S43 and S510. Furthermore, introduction of S301,319A mutations rendered Runx2 resistant to MAPK-dependent activation and reduced its ability to stimulate osteoblast-specific gene expression and differentiation after transfection into Runx2-null calvarial cells and mesenchymal cells. In contrast, S301,319E Runx2 mutants had enhanced transcriptional activity that was minimally dependent on MAPK signaling, consistent with the addition of a negative charge mimicking serine phosphorylation. These results emphasize the important role played by Runx2 phosphorylation in the control of osteoblast gene expression and provide a mechanism to explain how physiological signals acting on bone through the ERK/MAPK pathway can stimulate osteoblast-specific gene expression. It is likely that PTH signaling impacts osteoblast function, at least in part, via ERK/MAPK-dependent phosphorylation of Runx2, an ATF4-interacting factor in osteoblasts.

7) Establish that Foxo1 mediates IGF1/insulin regulation of osteocalcin expression by antagonizing Runx2 in osteoblasts (P7).

We determined the molecular mechanisms whereby forkhead transcription factor Foxo1, a key downstream signaling molecule of insulin-like growth factor 1 (IGF1)/insulin actions, regulates Runx2 activity and expression of the mouse osteocalcin gene 2 (Bglap2) in osteoblasts in vitro. We showed that Foxo1 inhibited Runx2-dependent transcriptional activity and osteocalcin mRNA expression and Bglap2 promoter activity in MC-4 preosteoblasts. Co-immunoprecipitation assay showed that Foxo1 physically interacted with Runx2 via its C-terminal region in osteoblasts or when co-expressed in COS-7 cells. Electrophoretic mobility shift assay demonstrated that Foxo1 suppressed Runx2 binding to its cognate site within the Bglap2 promoter. IGF1 and insulin prevented Foxo1 from inhibiting Runx2 activity by promoting Foxo1 phosphorylation and nuclear exclusion. In contrast, a neutralizing anti-IGF1 antibody decreased Runx2 activity and osteocalcin expression in osteoblasts.

Chromatin immunoprecipitation assay revealed that IGF1 increased Runx2 interaction with a chromatin fragment of the proximal Bglap2 promoter in a PI3K/AKT-dependent manner. Conversely, knockdown of Foxo1 increased Runx2 interaction with the promoter. This study establishes that Foxo1 is a novel negative regulator of osteoblast-specific transcription factor Runx2 and modulates IGF1/insulin-dependent regulation of osteocalcin expression in osteoblasts. These findings are important because IGF1 was reported to mediate the anabolic actions of PTH on bone in vivo.

Task 2: To establish whether the anabolic actions of PTH require ATF4 (6-48 months).

1) The anabolic effects of PTH on bone are severely impaired in growing Atf4-deficient mice (P5).

We first evaluated our hypothesis that ATF4 mediates the anabolic actions of PTH in bone using a relatively simple “growing mouse” model system, that has been widely used for studying the anabolic actions of PTH, PTHrP, FGF2, and IGF-1 in bone (1-6). The advantages of this system are that it is less time consuming and costly versus adult ovariectomized mouse models. Furthermore, because young growing animals have relatively high osteoblast activity, they are more sensitive to PTH than adults (6-14). Mice were treated with vehicle or PTH and sacrificed 24 h after the last PTH injection. PTH-dependent anabolic activity was evaluated in these mice using standard biochemical and histomorphometric criteria. Atf4−/− mice grew more slowly than wild type (wt) animals. The growth rate was slightly but significantly increased by PTH treatment during d 6-18 in wt but not Atf4−/− animals (Fig S1A, P5). Atf4−/− femurs were also shorter than wt or Atf4+/− femurs. PTH did not alter the length of femurs (Fig S1B, P5). However, it did significantly increase the dry ash weight per femur in wt and Atf4+/− but not in Atf4−/− mice (Fig S1C, P5). Serum Pi and calcium concentration (Fig S1D and E, P5) were not markedly affected by PTH or ATF4 deficiency. Faxitron X-ray analysis of femurs revealed that wt and Atf4+/− mice responded to PTH with markedly increased radiopacity throughout the whole femur, with the most dramatic increase in the metaphyseal region (Fig S2, top and middle, P5). In contrast, PTH only slightly increased the radiopacity in the same region of Atf4−/− femurs (Fig S2, bottom, P5). As shown in Fig 1 (P5), quantitative µCT analysis of femur histomorphometric parameters showed that Atf4−/− mice had a significant reduction in bone volume/tissue volume (BV/TV), trabecular number (Tb.N), and cortical thickness (Cort.Th) and a marked increase in trabecular space (Tb.Sp) compared with the wt or Atf4+/− littermates. These data confirmed an essential role of ATF4 in bone that was previously demonstrated by the Karsenty group. As expected, in wt femurs, intermittent PTH increased BV/TV, Tb.N, and Tb.Th by 5.4-fold, 2.7-fold, and 1.5-fold, respectively, and decreased Tb.Sp by 60 percent. Similar effects were also seen in Atf4+/− mice (Fig 1B, P5). In contrast, the PTH response was greatly attenuated in Atf4−/− mice where the following PTH responses
were observed; BV/TV, 2.2-fold increase; Tb.N, 1.7-fold increase; Tb.Th, 1.1-fold increase; Tb.Sp, 36 percent decrease. In all cases, the magnitude of PTH-stimulated changes on BV/TV, Tb.N, Tb.Sp was dramatically reduced in Atf4-/- mice relative to wt or Atf4+/+ mice (P<0.05, PTH/veh-wt vs. PTH/veh-Atf4-/-). Furthermore, PTH-stimulated increases in Cort.Th and Tb.Th were completely lost in Atf4-/- femurs. Because PTH similarly affected all trabecular and cortical parameters in Atf4+/+ and Atf4-/- mice, subsequent experiments compared the PTH effects on bone only between wt and Atf4-/- mice.

We next measured effects of Atf4 gene ablation on PTH stimulation of tibiae, vertebrae, and calvariae. The anabolic effect of PTH on wt tibia was so dramatic that the majority of the bone marrow cavity was replaced by newly formed bone (Fig 2A and B, P5). In Atf4+/- tibiae, while PTH still induced a small increase in trabecular area, the magnitude of stimulation was significantly reduced (5-fold in wt vs. 2.2-fold in Atf4-/-) (P<0.05, PTH/veh-wt vs. PTH/veh-Atf4-/-) (Fig 2C-E, P5). Likewise, the PTH-stimulated increase in the trabeculae of vertebrae (L5) was markedly reduced in Atf4-/- mice (3-fold in wt vs. 2-fold in Atf4+/+) (P<0.05, PTH/veh-wt vs. PTH/veh-Atf4-/-) (Fig 2F-J). When histological sections of calvariae were compared, PTH increased the width of the calvariae by 1.8-fold in wt mice, a response that was abolished in Atf4-/- animals (Fig 2K-O, P5).

2) Ablation of the Atf4 gene impairs PTH stimulation of trabecular, but not cortical bone in 7-month-old ovariectomized (OVX) mice (P5).

The experiments described above established a critical role of ATF4 in the anabolic effects of PTH on long bones, vertebrae, and calvariae in rapidly growing mice. However, it is possible that results obtained from growing animals may be different from those in adults due to possible effects of PTH and/or ATF4 on animal growth or influences of animal growth on the anabolic response to PTH, either of which could complicate the interpretation of the results. In contrast, adult mice have a mature skeleton in which these possible complications can be avoided. Furthermore, the OVX mouse provides a model that may be more relevant to the clinical applications of PTH in the treatment of osteoporosis. For these reasons, we evaluated whether Atf4 is required for the anabolic response to PTH in 7-month-old OVX mice. OVX surgery was successful as demonstrated by significant reduction in BV/TV (65 percent), Tb.N (27 percent), and Tb.Th (18 percent) and increased Tb.Sp (37 percent) relative to sham surgery (P<0.05, wt-sham vs. wt-OVX). OVX surgery did not reduce Cort.Th, which is consistent with results from rats. As shown in Fig 3A-C (P5), similar to results from growing mice, ablation of the Atf4 gene significantly decreased BV/TV and Tb.N and increased Tb.Sp in adult OVX mice. In further agreement with results in young mice, Atf4-/- animals exhibited a clearly attenuated response to PTH. For example, while PTH increased BV/TV by 7.8-fold in wt mice, this value was only increased 4.2-fold in Atf4-/- animals. Similarly, while PTH still stimulated formation of trabecular bone in Atf4+/- trabeculae, the magnitude of this response was significantly reduced compared to wt control (P<0.05, PTH/veh-wt vs. PTH/veh-Atf4-/-). In contrast to results from growing mice, Cort.Th was not reduced by ablation of the Atf4 gene in adult OVX mice (0.21±0.01 mm in wt vs. 0.19±0.02 mm in Atf4+/+ mice, P>0.05 wt vs. Atf4+/+). Also, PTH was much less effective in stimulating Cort.Th in adult OVX mice (24%) than in growing mice (95%) (Figs 1 and 3, P5). Furthermore, no difference in stimulation of cortical thickness by PTH was observed when wt and Atf4-/- groups were compared (24% wt vs. 21% Atf4-/-) (P>0.05, PTH/veh-wt vs. PTH/veh-Atf4-/-).

3) ATF4 is critical for PTH to increase bone formation in vivo (P5).

To determine if Atf4 is required for PTH to increase bone formation in vivo, we performed calcein double labeling experiments. As shown in Fig 3A and B (P5), results from calcein double labeling of 7-month old OVX wt and Atf4-/- tibia revealed that the PTH-stimulated increase in mineral apposition rate (MAR), an indicator of osteoblast function, was significantly reduced by ATF4 deficiency (P<0.05, PTH/veh-wt vs. PTH/veh-Atf4-/-).

4) Atf4-deficient mice display a resistance to OVX-induced bone (P5).

Interestingly, as shown in Fig S3 (P5), OVX surgery significantly reduced bone mass in wt mice. This reduction was completely abolished in Atf4-/- animals (P>0.05, Atf4-/--sham vs. Atf4-/--OVX). This result suggests that ATF4 may play an important role in OVX induction of bone loss. Current study in the project laboratory is to define the molecular mechanism underlying this important finding.
5) ATF4 is critical for OCL differentiation in vitro and in vivo (P6).

Our results revealed that TRAP activity (an OCL marker) was dramatically decreased in Atf4−/− tibiae compared to wt tibiae (Figure 1C, P6). OCL surface/bone surface (Oc.S/BS) and OCL number/bone perimeter (Oc.Nb/BPm) were reduced similarly in both primary and secondary spongiosa in Atf4−/− tibiae relative to wt tibiae (Figure 1D, P6). TRAP-positive MNCs (≥3 nuclei/cell) in BMM cultures from Atf4−/− mice were dramatically reduced compared to that from wt mice (Figure 1E-F, P6). Furthermore, the number of nuclei per MNC was decreased by 75 percent in Atf4−/− BMM cultures compared to wt cultures (Figure 1G, P6), and the MNCs that formed in Atf4−/− BMM cultures were much smaller than those formed in wt cultures. Furthermore, the resorption pit area on dentin slices was dramatically reduced in Atf4−/− BMM cultures compared to wt cultures (Figure 1H-I, P6). Co-culture experiments showed that wt osteoblasts failed to rescue the defective OCL formation of Atf4−/− BMMs (Figure 2C-D, P6). Taken together, these experiments suggest that ATF4 deficiency impairs osteoclastogenesis in a cell autonomous manner. We developed transgenic mice in which the Atf4 transgene is driven by an 1846-bp mouse Trap promoter that selectively expresses ATF4 in OCLs (Figure 3A, top, P6). Transgenic ATF4 dramatically increased the protein levels of NFATc1/A but not of PU.1 or c-Fms proteins in differentiated BMM cultures. The levels of OCL differentiation marker gene mRNAs (Trap, Rank, Cat K, and Mmp9) were all dramatically elevated in differentiated BMM cultures from Trap-Atf4-tg mice compared to those from wt mice. OCL-targeted overexpression of ATF4 dramatically increased the numbers of TRAP-positive MNCs in vitro (Figure 3D, P6). We found a similar effect of transgenic ATF4 expression in vivo. The TRAP activity was markedly increased in Trap-Atf4-tg tibiae compared to wt tibiae (Figure 3E, P6). As shown in Figure 3F (P6), Oc.S/BS Oc.Nb/BPm in both primary and secondary spongiosa of tibiae were dramatically increased in Trap-Atf4-tg mice compared to wt controls. Quantitative μCT analysis of femur histomorphometric parameters showed that Trap-Atf4-tg mice had a significant reduction in bone volume/tissue volume (BV/TV) and trabecular number (Tb.N) and a marked increase in trabecular space (Tb.Sp) compared with the wt littermates (Figure 3G, P6). Because osteoclast activity was recently shown to be critical for the anabolic actions in bone, our finding that ATF4 regulates osteoclast differentiation and bone resorption has added a new layer to the molecular mechanisms underlying the PTH actions in osteoblasts and bone.

Key Research Accomplishments

- We have demonstrated that ATF4 is a novel downstream target of PTH signaling in osteoblasts (P1).
- We have demonstrated that TFIIAγ increases osteoblast-specific osteocalcin gene expression by facilitating ATF4-Runx2 interactions (P2).
- We have demonstrated that ATF4 increases the expression of cyclin D1, a key factor for cell cycle progression, and promotes osteoblast proliferation in vitro and in vivo (P3 and P5).
- We have demonstrated that ATF4 is critical for PTH to increase osteoblast differentiation in bone (P5).
- We have demonstrated that ATF4 is a novel upstream transcriptional activator of Osx, a key factor for osteoblast differentiation and bone formation (P5).
- We have demonstrated that ATF4 is important for intermittent PTH to stimulate osteoblast-mediated bone formation in vivo (P5).
- We have identified and functionally characterized Erk/MAPK phosphorylation sites in Runx2, an ATF4-interacting factor identified by the project laboratory (P4).
- We have demonstrated that ATF4 is required for the anabolic actions of PTH in bone in rapidly growing mice (P5).
- We have demonstrated that ATF4 is required for the anabolic actions of PTH in bone in adult OVX mice (P5).
- We have demonstrated that ATF4 is essential for OVX induction of bone loss in vivo (P5).
- We have demonstrated that ATF4 is essential for osteoclast differentiation and bone resorption, which is modulated by PTH (P6).
- We have demonstrated that Foxo1 mediates IGF1/insulin regulation of osteocalcin expression by antagonizing Runx2 in osteoblasts (P7).

Reportable Outcomes
Peer-reviewed papers:


Abstracts:


Conclusion

During the last four years of support, our studies establish that: i) ATF4 is essential for intermittent PTH to increase bone mass in both young and adult OVX mice. ii) ATF4 increases osteoblast proliferation in vitro and in bone probably via upregulation of cyclin D1; iii) ATF4 promotes osteoblast differentiation at least in part via upregulation of Oxs, a key osteoblast differentiation transcription factor; iv) ATF4 is critical for PTH to increase bone formation in
vivo; v) ATF4 plays a critical role in OVX induction of bone loss; and vi) ERK/MAPK, which is known to be activated by PTH signaling in osteoblasts, phosphorylates and activates Runx2; vii) ATF4 is critical for osteoclast differentiation and bone resorption; viii) ATF4 activates osteocalcin gene expression via cooperation with Runx2, a master regulator of osteoblast differentiation and bone formation; ix) Foxo1 mediates IGF1, which is critical for the anabolic actions of PTH in bone in vivo, regulation of osteocalcin expression by antagonizing Runx2 in osteoblasts. Collectively, the knowledge obtained from these studies will significantly enhance our understanding of the molecular mechanism underlying the actions of PTH in osteoblasts and bone and define new potential therapeutic targets for improved treatment of osteoporosis and other metabolic bone diseases.

References

Six peer-reviewed research papers: P1-7
Seven national meeting abstracts: A1-7

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Parathyroid Hormone Increases Activating Transcription Factor 4 Expression and Activity in Osteoblasts: Requirement for Osteocalcin Gene Expression

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PTH is an important peptide hormone regulator of calcium homeostasis and osteoblast function. However, its mechanism of action in osteoblasts is poorly understood. Our previous study demonstrated that PTH activates mouse osteocalcin (Ocn) gene 2 promoter through the osteoblast-specific element 1 site, a recently identified activating transcription factor-4 (ATF4) -binding element. In the present study, we examined effects of PTH on ATF4 expression and activity as well as the requirement for ATF4 in the regulation of Ocn by PTH. Results show that PTH elevated levels of ATF4 mRNA and protein in a dose- and time-dependent manner. This PTH regulation requires transcriptional activity but not de novo protein synthesis. PTH also increased binding of nuclear extracts to osteoblast-specific element 1 DNA. PTH stimulated ATF4-dependent transcriptional activity mainly through protein kinase A with a lesser requirement for protein kinase C and MAPK/ERK pathways. Lastly, PTH stimulation of Ocn expression was lost by small interfering RNA down-regulation of ATF4 in MC-4 cells and Atf4−/− bone marrow stromal cells. Collectively, these studies for the first time demonstrate that PTH increases ATF4 expression and activity and that ATF4 is required for PTH induction of Ocn expression in osteoblasts. (Endocrinology 149: 1960–1968, 2008)

PTH IS A MAJOR regulator of osteoblast activity and skeletal homeostasis. PTH has both catabolic and anabolic effects on osteoblasts and bone that depend on the temporal pattern of administration; continuous administration decreases bone mass, whereas intermittent administration increases bone mass (1–3). At the molecular level, PTH binds to the PTH-1 receptor (PTH1R), a G protein-coupled receptor that is expressed in osteoblasts (4–6) and activates multiple intracellular signaling pathways that involve cAMP, inositol phosphates, intracellular Ca2+, protein kinases A and C (7), and the ERK/MAPK pathway (8, 9). The ability of PTH to regulate gene expression is largely dependent on activation of specific transcription factors such as cAMP response element binding protein (CREB) (10, 11), activator protein-1 family members (12–15), pituitary-specific transcription factor-1 (16), and Runx-related transcription factor-2 (Runx2) (12, 17). A better understanding of the downstream PTH signaling events is essential to understand the mechanistic basis for the anabolic and catabolic actions of this hormone on bone.

The osteocalcin (Ocn) promoter has been the major paradigm for unraveling the mechanisms mediating osteoblast-specific gene expression and defining a number of transcription factors and cofactors (18–29). Because Ocn gene is regulated by PTH (30–32), we considered it a good model for identifying new transcriptional mediators of PTH action. Using this system, we recently showed that the osteoblast-specific element (OSE)-1 in the proximal mouse (Ocn) gene 2 (mOG2) promoter (19) is necessary and sufficient for PTH induction of this gene (33). Immediately after publication of this study, the OSE1 was identified as a binding site for activating transcription factor-4 (ATF4) (34).

AF4, also known as CREB2 (35) and tax-responsive enhancer element B67 (36), is a member of the ATF/CREB family of leucine-zipper factors that also includes CREB, cAMP response element modulator, ATF1, ATF2, ATF3, and ATF4 (37–41). These proteins bind to DNA via their basic region and dimerize via their leucine domain to form a large variety of homodimers and/or heterodimers that allow the cell to coordinate signals from multiple pathways (37–41). An in vivo role for ATF4 in bone development was established using Atf4-deficient mice (29). ATF4 is required for expression of Ocn and bone sialoprotein as demonstrated by the dramatic reduction of their mRNAs in Atf4−/− bone (29). ATF4 activates Ocn transcription through direct binding to the OSE1 site as well as interactions with Runx2 through
cooperative interactions with OSE1 and OSE2 (also known as nuclear matrix protein 2 binding sites) sites in the promoter (19, 20, 25). ATF4 activity is negatively regulated by factor inhibiting activating transcription factor-4-mediated transcription (42). Factor inhibiting activating transcription factor binds to ATF4 and represses its activity and bone formation in vivo. Although Atf4 mRNA is ubiquitously expressed, ATF4 protein preferentially accumulates in osteoblasts (34). This accumulation is explained by a selective reduction of proteasomal degradation in osteoblasts.

The purpose of this study was to determine the effects of PTH on ATF4 expression and activity and evaluate whether ATF4 mediates PTH induction of Ocn expression in osteoblasts.

Materials and Methods

Reagents

Tissue culture media and fetal bovine serum were obtained from HyClone (Logan, UT). [32P]ATP (3000 Ci/mol) and α-[32P]dCTP (3000 Ci/mol) were purchased from GE Healthcare (Piscataway, NJ). Other reagents were obtained from the following sources: H89, forskolin (FSK), GFI9202X, phorbol 12-myristate 13-acetate (PMA), cycloheximide (CHX), actinomycin D (ActD), and mouse monoclonal antibody against β-actin from Sigma (St. Louis, MO); U0124 from Promega (Madison, WI); and U0124 from Calbiochem (La Jolla, CA), PTH (1–34) from Bachem (Torrance, CA), antibodies against ATF4, Runx2, and horse-radish peroxidase-conjugated mouse or goat IgG from Santa Cruz (Santa Cruz, CA). All other chemicals were of analytical grade.

Cell cultures

Mouse MC3T3-E1 subclone 4 (MC-4) cells were described previously (43, 44) and maintained in ascorbic acid-free DMEM containing 20% FBS, 1% penicillin/streptomycin, and 1% sodium selenite (FBS), and 1% penicillin/streptomycin and were not used beyond passage 15. Rat osteoblast-like UMR106-01 cells (45) were maintained in DMEM and 10% FBS. Isolation of mouse primary bone marrow stromal cells (BMSCs) was described previously (33). Briefly, 6-wk-old male C57BL/6 mice were killed by cervical dislocation. Tibiae and femurs were isolated and the epiphyses were cut. Marrow was flushed with DMEM containing 20% FBS, 1% penicillin/streptomycin, and 10−8 M dexamethasone into a 60-mm dish, and the cell suspension was aspirated up and down with a 20-gauge needle to break clumps of marrow. The cell suspension (marrow from two mice/flask) was then aspirated up and down with a 20-gauge needle to break clumps of marrow according to the manufacturer’s instructions. Each reaction contained 1 [32P]dCTP using a random primer kit (Roche Molecular Biochemicals, Indianapolis, IN). Hybridizations were performed as previously described using a Bellco Autoblot hybridization oven (47). Same blots were reprobed with [32P]-labeled cDNA to confirm the specificity of the PCR products. Six samples were run for each primer set. The levels of mRNA were calculated by the ΔCT (the difference between the threshold cycles) method (46). Atf4, Ocn, Col1(1), Pth1r, and Omp mRNAs were normalized to Gapdh mRNA.

Western blot analysis

Twenty micrograms of total nuclear extracts were fractionated on a 10% SDS-PAGE gel and transferred onto nitrocellulose membranes (Schleicher & Schuell, Keene, NH). The membrane was blocked in 5% nonfat milk in Tris-buffered saline (TBS) containing 0.1% Tween 20. Blots were probed with antibodies against ATF4 (1:1000) followed by incubation with secondary antibodies conjugated with horseradish peroxidase (1:5000); and visualized using an enhanced chemiluminescence kit (Pierce, Rockford, IL). Finally, blots were stripped two times in buffer containing 65 mm Tris Cl (pH 6.8), 2% sodium dodecyl sulfate, and 0.7% (vol/vol) β-mercaptoethanol at 65 C for 15 min and reprobed with β-actin antibody (1:5000) for normalization.

RNA isolation and reverse transcription

Total RNA was isolated using TRIzol reagent (Invitrogen Life Technologies, Gaithersburg, MD) according to the manufacturer’s protocol. Reverse transcription was performed using 2 μg of denatured RNA and 100 pmol of random hexamers (Applied Biosystems, Foster, CA) in a total volume of 25 μl containing 12.5 U MultiScribe reverse transcriptase (Applied Biosystems) according to the manufacturer’s instructions.

Quantitative real-time PCR

Quantitative real-time PCR was performed on an iCycler (Bio-Rad, Minneapolis, MN) using a SYBR Green PCR core kit (Applied Biosystems) and cDNA equivalent to 10 ng denatured RNA and 100 pmol of random hexamers (Applied Biosystems, Foster, CA) in a total volume of 25 μl containing 12.5 U MultiScribe reverse transcriptase (Applied Biosystems) according to the manufacturer’s instructions.

Preparation of nuclear extracts and gel mobility shift assay (GMSA)

Nuclear extracts were prepared and GMSAs were conducted as previously described (43). Each reaction contained 1 μg of nuclear extracts. The DNA sequences of OSE1 oligonucleotides used for GMSA were as follows: wild-type (wt): TGC TTA CAT CAG AGA GCA; mutant (mt): TGC TTA gta CAG AGA GCA.

Northern blot

Twenty micrograms of total RNA was fractionated on a 10% agarose-formaldehyde gels and blotted onto nitrocellulose paper. The mouse Atf4 cDNA inserts were excised from plasmid DNA with the appropriate restriction enzymes and purified by agarose gel electrophoresis before labeling with α-[32P]dCTP using a random primer kit (Roche Molecular Biochemicals, Indianapolis, IN). Hybridizations were performed as previously described using a Belco Autoblot hybridization oven (47). Same blots were reprobed with [32P]-labeled cDNA to 185 rRNA for loading (48).

Small interfering RNA (siRNA)-MC-4 cells, which contain high levels of Atf4 mRNA, were seeded at a density of 25,000 cells/cm2. After 24 h, cells were transfected with lipofectAMINE 2000 (Invitrogen, Carlsbad, CA) according to the manufacturer’s instructions. Each transfection contained 0.5 μg of the indicated plasmid plus 0.05 μg of pRL-SV40, containing a CDNA for Renilla recombines luciferase to control for transfection efficiency. Cells were harvested and assayed using the dual luciferase assay kit (Promega) on a Monolight 2010 luminometer (BD Biosciences, San Diego, CA).

Preparation of nuclear extracts and gel mobility shift assay (GMSA)

Nuclear extracts were prepared and GMSAs were conducted as previously described (43). Each reaction contained 1 μg of nuclear extracts. The DNA sequences of OSE1 oligonucleotides used for GMSA were as follows: wild-type (wt): TGC TTA CAT CAG AGA GCA; mutant (mt): TGC TTA gta CAG AGA GCA.
Hughes Medical Institute and the University of Michigan School of Medicine). These mice were originally developed by Dr. Tim M. Townes (University of Alabama at Birmingham) and were used to generate \( \text{Atf4} \) wild-type (\( \text{Atf4}/\text{H11001}/\text{H11001} \)), heterozygous (\( \text{Atf4}/\text{H11001}/\text{H11002} \)), and homozygous mutant (\( \text{Atf4}/\text{H11002}/\text{H11002} \)) embryos/pups for this study. Original reports describing the phenotype of \( \text{Atf4} \) homozygote-null mutants used the identical strain of mice (50). PCR genotyping was performed on tail DNA using a cocktail of three primers (TOWNES-1: 5'-AGC AAA ACA AGA CAG CAG CCA CTA-3'; TOWNES-2: 5'-GTT TCT ACA GCT TCC TCC ACT CTT-3'; and TOWNES-3: 5'-ATA TTG CTG AAG AGC TTG GCGGC-3') obtained from the laboratories of Dr. Randal J. Kaufman. A 700-bp DNA PCR product was amplified from \( \text{Atf4}/\text{H11002}/\text{H11002} \) mouse tail DNA and a 900-bp product from wild-type mice (see Fig. 7A). The genotype of each mouse established by PCR of tail genomic DNA was confirmed by Western blotting of calvaria cell lysates and anti-ATF4 antibody. A breeding colony was established using heterozygote mice to provide littermate controls. All animal studies were approved by the Animal Care Committee of the Veterans Affairs Pittsburgh Healthcare System.

**FIG. 1.** PTH increases levels of ATF4 expression in osteoblasts. A, Effect of PTH on \( \text{Atf4} \) mRNA. MC-4 cells were seeded at a density of 50,000 cells/cm\(^2\) in 35-mm dishes and cultured in 10% FBS medium overnight. Cells were then treated with various concentration of PTH for 6 h. For each group, total RNA (20 \( \mu \)g/lane) was loaded for Northern hybridization using cDNA probes for mouse \( \text{Atf4} \) mRNA and 18S rRNAs (for normalization). B, Effect of PTH on ATF4 proteins (dose response). MC-4 cells were treated with indicated concentrations of PTH for 6 h and nuclear extracts were prepared for Western blot analysis for ATF4. C, Effect of PTH on ATF4 proteins (time course). MC-4 cells were treated with \( 10^{-7} \) M PTH for indicated time (h). Experiments were repeated three to four times, and qualitatively identical results were obtained.

**FIG. 2.** Effects of CHX/ActD treatment on PTH induction of \( \text{Atf4} \) mRNA. MC-4 cells were treated with vehicle or 10 \( \mu \)g/ml CHX (A) or ActD (B) in the absence or presence of PTH for 6 h. \( \text{Atf4} \) and Gapdh mRNAs were determined by quantitative real-time RT-PCR analysis. Experiments were repeated three times, and qualitatively identical results were obtained. *, \( P < 0.05 \) [control (ctrl) vs. PTH]; †, \( P < 0.05 \) (CHX vs. CHX/PTH); ‡, \( P < 0.05 \) (CHX vs. CHX/PTH).

**FIG. 3.** PTH increases ATF4-dependent transcriptional activity in MC-4 cells. A, Target cell specificity. Cells (MC-4, UMR106–01, and primary BMSCs) were transiently transfected with p4OSE1-luc and renilla luciferase normalization plasmid and treated with \( 10^{-7} \) M PTH for 6 h before being harvested and assayed for dual-luciferase activity. Firefly luciferase activity was normalized to renilla luciferase activity (for transfection efficiency). B, Dose dependence. MC-4 cells were transiently transfected as in Fig. 2A and treated with indicated concentration of PTH (from \( 10^{-11} \) to \( 10^{-7} \) M) for 6 h followed by dual-luciferase assay. C, Time course. MC-4 cells were transiently transfected as in Fig. 2A and treated with \( 10^{-7} \) M PTH for indicated times. Data represent mean ± SD. Experiments were repeated three to four times and qualitatively identical results were obtained. *, \( P < 0.05 \) [control (ctrl) vs. PTH].
Significant stimulatory effect first detected at a concentration with maximal stimulation at 10^(-13) to 10^(-10) M. One microgram of each nuclear extract was incubated with end-labeled double-stranded OSE1 (TGC TTA CAT CAG AGA GCA) and analyzed by electrophoresis on 4% polyacrylamide gels.

Data were analyzed with GraphPad Prism software (GraphPad, San Diego, CA). A one-way ANOVA analysis was used followed by the Dunnett's test (see Fig. 3, B and C). Student's t test was used to test for differences between two groups of data. Differences with a P < 0.05 was considered as statistically significant. Results were expressed as means ± sd.

**Results**

**PTH increases ATF4 expression in MC-4 cells**

To determine the effect of PTH on Atf4 mRNA expression, MC-4 cells were treated with increasing concentrations of PTH (from 10^(-13) to 10^(-7) M) for 6 h, and total RNA was isolated for Northern blot analysis. As shown in Fig. 1A, PTH dose-dependently increased levels of Atf4 mRNAs with a significant stimulatory effect first detected at a concentration of 10^(-10) M.

**Statistical analysis**

DNA binding to labeled wild-type OSE1 probe was analyzed in the presence of 25- to 50-fold molar excesses of cold wt (lanes 6 and 7) or mt (lanes 4 and 5) OSE1 (TGC TTA gta CAG AGA GCA) by GMSA using 1 μg of nuclear extracts from PTH-treated MC-4 cells.

**Fig. 4.** PTH increases binding of ATF4 to OSE1 DNA. A, PTH increases binding of osteoblast nuclear extracts (NE) to OSE1. Nuclear extracts were prepared from MC-4 cells with (P) (lanes 3–7) or without (C) (lane 2) PTH treatment for 6 h. One microgram of each nuclear extract was incubated with 1 μg of wild-type OSE1 probe (lanes 1–3) and mt OSE1 probe (lanes 4–6). B, Binding site specificity. Labeled wt (lanes 1–3) and mt (lanes 4–6) OSE1 probes were incubated with 1 μg nuclear extracts from MC-4 cells with and without PTH treatment. C, The nuclear complex binding OSE1 contains ATF4. Labeled wild-type OSE1 probe was incubated with 1 μg nuclear extracts from PTH-treated MC-4 cells in the presence of normal control IgG (lane 5), ATF4 antibody (lane 4), Runx2 antibody (lane 3), CREB antibody (lane 6), ATF1 antibody (lane 7), and Fra-1 antibody (lane 8). Experiments were repeated three to four times, and qualitatively identical results were obtained.

**Results**

**PTH increases ATF4-dependent transcriptional activity in osteoblasts**

The effect of PTH on Atf4-dependent transcriptional activity was evaluated in two osteoblast cell lines and primary mouse bone marrow stromal cells. Cells were transiently transfected with wt or mt p4OSE1-luc, an artificial promoter containing four copies of wt or mt OSE1 (a specific ATF4-binding element) fused to a −34 to +13 minimal mOG2 promoter, and pRL-SV40, a renilla luciferase normalization plasmid. After 42 h, cells were treated with PTH (10^(-7) M) for 6 h followed by dual-luciferase assay. Firefly luciferase activity was normalized to renilla luciferase activity as a control for transfection efficiency. As shown in Fig. 3A, PTH stimulated Atf4-dependent OSE1 activity by 17-, 2.7-, and 2.8-fold in MC-4, UMR106-01, and primary BMSCs (P < 0.05, control vs. PTH), respectively. This PTH response was completely lost with the introduction of a 3-bp point mutation in the OSE1 core sequence (from TTCACTCA to TTAGTACA).
that the earliest effect of PTH stimulation was seen within 1 h and peaked at 5–6 h (Fig. 3C).

**PTH increases ATF4 binding to OSE1 DNA**

To determine whether PTH increases ATF4 binding to OSE1 DNA, we performed GMSA using nuclear extracts from MC-4 cells with and without 10^{-7} M PTH for 6 h. Consistent with our previous observation (33), nuclear extracts from PTH-treated MC-4 cells exhibited increased binding to intact OSE1 oligonucleotides (Fig. 4A, lanes 2 and 3), and this binding was significantly reduced by the addition of 25- and 50-fold molar excesses of unlabeled wt OSE1 oligonucleotides (Fig. 4A, lanes 4 and 5) but not by unlabeled mt OSE1 oligonucleotides (Fig. 4A, lane 6 and 7). In contrast, GMSA using labeled mt OSE1 oligonucleotides as probes showed that both basal and PTH-induced binding activity was abolished by the same 3-bp point mutation (Fig. 4B, lanes 4–6). The same mutation abolished PTH activation of 647- and 116 bp mOG2 promoter fragments and 4OSE1 (33) (Fig. 3A). Importantly, PTH-induced binding to OSE1 was supershifted with an anti-ATF4 antibody (Fig. 4C, lanes 4). In contrast, normal IgG or antibodies against Runx2, CREB, ATF1, and Fra-1 did not significantly supershift the PTH-stimulated band (Fig. 4C, lanes 3–8). Taken together, these studies demonstrate that ATF4 is a component of the PTH-stimulated DNA-protein complex associating with OSE1.

[Note that PTH treatment did not alter binding of Runx2 to OSE2 DNA in the mOG2 promoter in GMSA (33).]

**Protein kinase A (PKA) is the major signaling pathway mediating the PTH response**

To identify signaling pathways mediating PTH activation of ATF4 transcriptional activity, we examined the effects of various inhibitors or activators. As shown in Fig. 5A, H89, a selective inhibitor of the PKA pathway, completely abolished PTH-stimulated ATF4 transcriptional activity (P > 0.05, control vs. PTH). GF109203X, a specific inhibitor of the protein kinase C (PKC) pathway, significantly decreased the PTH stimulation. U0126, a specific inhibitor of MAPK, partially suppressed PTH stimulation. As shown in Fig. 5B, FSK, a well-known activator of PKA, increased ATF4 activity in the absence of PTH in a dose-dependent manner. In combination with PTH, the effect of FSK was not additive, indicating that the PKA pathway was maximally stimulated. PMA, a PKC activator, did not significantly affect the PTH-induced ATF4 activity at a concentration range of 0.1–5 

![Fig. 5. PKA is the major signaling pathway mediating the PTH response.

A, Effects of inhibitors/activators on PTH-induced ATF4 transcriptional activity. MC-4 cells were transiently transfected with p4OSE1-luc and renilla luciferase normalization plasmid. After 42 h, cells were treated with 10 

B and C, Dose-response of FSK (B) and PMA (C) on PTH stimulation of ATF4 transcriptional activity. MC-4 cells were transiently transfected as in Fig. 5A and treated with indicated concentration of respective activator for 6 h in the absence and presence of 10^{-7} M PTH followed by dual-luciferase assay. Data represent mean ± SD. Experiments were repeated three times and qualitatively identical results were obtained. *, P < 0.05 [control (ctrl) vs. PTH].](https://endocrine.endojournals.org/10.1210/en.2007-1046)
**Discussion**

This study examined actions of PTH on ATF4 expression and activity in osteoblasts. Using the Ocn gene as a model system for studying PTH-dependent transcription, we found the following: 1) PTH rapidly induces Atf4 expression in MC-4 cells and mouse primary bone marrow stromal cells in a time- and dose-dependent manners; 2) PTH increases in vitro binding of ATF4 to OSE1 DNA; 3) PTH dramatically activates ATF4 transcriptional activity mainly through the PKA pathway; 4) PTH stimulation of Ocn gene expression requires ATF4 because it is abolished by ATF4 siRNA in MC-4 cells and is not seen in ATF4-deficient BMSCs. Col-

**Fig. 6.** ATF4 siRNA blocks PTH stimulation of Ocn expression. A and B, MC-4 cells were transiently transfected with Atf4 siRNA (A) or negative control (Ctrl) siRNA (B). After 48 h, total RNA was prepared for quantitative real-time RT-PCR analyses for Atf4 mRNA, which was normalized to Gapdh mRNA. C and D, MC-4 cells were transiently transfected with 40 nM Atf4 siRNA or negative control siRNAs. After 42 h, cells were treated with or without PTH (10^{-7} M) for 4 h followed by RNA preparation and quantitative real-time PCR analysis. As shown in Fig. 7B, minimal Atf4 mRNA was detected by real-time RT/PCR in the Atf4^−/−^ BMSCs. Consistent with the results of experiments with MC-4 cells, PTH significantly stimulated Atf4 mRNA in wt BMSCs (P < 0.05, control vs. PTH), but this induction was completely lost in cells from Atf4^−/−^ mice (Fig. 7B). As shown in Fig. 7C, PTH significantly increased Ocn mRNA in wt BMSCs, which was abolished in Atf4^−/−^ BMSCs (P > 0.05, control vs. PTH). The basal level of Ocn mRNA was also significantly reduced in Atf4^−/−^ BMSCs relative to wt cells (P < 0.05, wt vs. mt). In contrast, PTH did not increase Ocn mRNA in wt or mt BMSCs (P > 0.05, control vs. PTH) (Fig. 7D). However, the level of Ocn mRNA was increased in Atf4^−/−^ cells (P < 0.05, wt vs. mt), indicating that ATF4 may function as a negative regulator of Ocn expression (Fig. 7D). In addition, the levels of Pth1r mRNA were not significantly changed by either ATF4 deficiency or PTH, suggesting that PTH signaling is intact in the absence of ATF4 (Fig. 7E). Taken together, these data clearly establish that ATF4 is required for PTH induction of Ocn mRNA in primary BMSCs.

**Fig. 7.** PTH increases Ocn expression and mRNA in MC-4 cells. A, MC-4 cells were transiently transfected with siRNA and treated with or without PTH (10^{-7} M) for 4 h followed by RNA preparation and quantitative real-time PCR analysis. As shown in Fig. 7B, minimal Atf4 mRNA was detected by real-time RT/PCR in the Atf4^−/−^ BMSCs. Consistent with the results of experiments with MC-4 cells, PTH significantly stimulated Atf4 mRNA in wt BMSCs (P < 0.05, control vs. PTH), but this induction was completely lost in cells from Atf4^−/−^ mice (Fig. 7B). As shown in Fig. 7C, PTH significantly increased Ocn mRNA in wt BMSCs, which was abolished in Atf4^−/−^ BMSCs (P > 0.05, control vs. PTH). The basal level of Ocn mRNA was also significantly reduced in Atf4^−/−^ BMSCs relative to wt cells (P < 0.05, wt vs. mt). In contrast, PTH did not increase Ocn mRNA in wt or mt BMSCs (P > 0.05, control vs. PTH) (Fig. 7D). However, the level of Ocn mRNA was increased in Atf4^−/−^ cells (P < 0.05, wt vs. mt), indicating that ATF4 may function as a negative regulator of Ocn expression (Fig. 7D). In addition, the levels of Pth1r mRNA were not significantly changed by either ATF4 deficiency or PTH, suggesting that PTH signaling is intact in the absence of ATF4 (Fig. 7E). Taken together, these data clearly establish that ATF4 is required for PTH induction of Ocn mRNA in primary BMSCs.
lectively, this study establishes that ATF4 is a novel down-stream target of PTH actions in osteoblasts. It is well documented that PTH signals mainly through the PKA pathway. In the present study, we show that PKA inhibition completely blocked PTH stimulation of ATF4 activity. Furthermore, activation of the PKA pathway by FSK dramatically increased ATF4 activity in the absence of PTH. However, when combined with PTH, the effect of FSK was not additive. These results strongly suggest that PKA is the major pathway for PTH to activate ATF4 because each agent (i.e. FSK or PTH) maximally stimulates the same pathway, making additional ATF4 activation impossible. Inhibition of the PKC pathway also resulted in a significant reduction in PTH-induced ATF4 activity (data not shown), but PKC activation by PMA failed to activate both basal or PTH-induced ATF4 activity. Thus, PKC is partially required for PTH activation of ATF4. Lastly, inhibition of the MAPK/ERK pathway led to partial inhibition of the PTH stimulation. These three pathways are also required for PTH induction of both Ocn mRNA and 1.3-kb mOG2 promoter activity as previously described (33), further supporting our hypothesis that ATF4 mediates PTH induction of Ocn gene expression.

A recent study showed that ATF4 mediates β-adrenergic induction of Rankl mRNA expression via direct binding to the upstream OSE1 site in the Rankl promoter in osteoblasts (54). However, PTH stimulation of Rankl expression was not reduced in the absence of ATF4, suggesting that this catabolic action of PTH is independent of this transcription factor. Phosphorylation seems to be critical for ATF4 to elicit its function in osteoblasts and bone. A PKA phosphorylation site (serine 254) within the ATF4 molecule was recently shown to mediate β-adrenergic induction of Rankl mRNA expression in osteoblasts (54). In addition, ATF4 is phosphorylated at serine 251 by ribosomal kinase 2 (RSK2), the kinase inactivated in Coffin-Lowry syndrome, an X-linked mental retardation disorder associated with skeletal manifestations (29). Because RSK2 is an immediate downstream target of MAPK/ERK that is activated by PTH signaling (8, 9), PTH may in part activate ATF4 via the MAPK/ERK/RSK2 pathway. It remains to be determined whether the PKA and/or RSK2 phosphorylation sites are involved in the PTH activation of ATF4.

One of the major downstream factors for PTH signaling is CREB, the cAMP response element binding protein. Actions of CREB are mediated through cAMP response elements (CREs) in the regulatory regions of target genes. PTH phosphorylates CREB at serine 133. This phosphorylation event stimulates the binding of CREB to the CRE and is required for CREB to activate transcription of target genes. Through this classical pathway, PTH rapidly induces transcription of immediate-early response genes including those encoding activator protein-1 family members such as c-Fos, c-Jun, Fra-1, Fra-2, and FosB (10, 14, 15, 52, 55–57). Although CREB was shown to binding to the OSE1 site (29), overexpression of CREB was unable to activate OSE1-dependent transcription activity of the mOG2 promoter in vitro (29), suggesting

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**Fig. 7.** PTH stimulation of Ocn expression is lost in Atf4−/− BMSCs. A, PCR genotyping was performed on tail DNA using a cocktail of three primers (see Materials and Methods). A 700-bp DNA PCR product is amplified from Atf4−/− mouse tail DNA and a 900-bp product from wild-type mice. B–E, Effects of ATF4 deficiency on PTH stimulation of Atf4 (B), Ocn (C), Opn (D), and Pth1r (E) expression in BMSCs. Primary BMSCs were seeded at a density of 50,000 cells/cm² in 35-mm dishes and cultured in 10% PBS medium overnight. Cells were then treated with 10−7 M PTH for 6 h followed by RNA preparation and quantitative real-time RT/PCR for Atf4 (B), Ocn (C), Opn (D), and Pth1r (E) mRNA, which were normalized to the Gapdh mRNAs. *, P < 0.05 (ctrl vs. PTH); #, P < 0.05 (wt vs. mt). Data represent mean ± sd. Experiments were repeated three times, and qualitatively identical results were obtained.
that this site is not a major functional site for CREB. Furthermore, the OSE1 binding activity stimulated by PTH was not supershifted by an anti-CREB antibody. Instead, this complex clearly contains ATF4 protein (Fig. 4C). Thus, we were unable to obtain any evidence for the involvement of CREB in the PTH response. However, our results do not exclude the possibility that PTH/CREB activates Asf1 mRNA transcription via CREB binding to potential CRE sites in the Atf4 promoter.

PTH induction of immediate-early response genes occurs very rapidly (minutes to hours) and lasts for several hours. This PTH response is usually independent upon the presence of de novo protein synthesis but requires active cellular transcription. The time-course experiments in the present study indicate that PTH induction of Atf4 occurs within 1 h of PTH addition and peaks after 3–6 h. Furthermore, this regulation depends on active cellular transcription and does not require de novo protein synthesis. Therefore, ATF4 may be considered as an additional PTH early response gene. ATF4-deficient mice as well as humans with mutations in RSK2, an ATF4 activating kinase, exhibit striking deficits in bone formation and osteoblast activity. Because ATF4 is required for osteoblast function and bone formation in vivo, and as shown herein, ATF4 is a novel downstream target of PTH, it will be important to determine whether ATF4 is also required for the anabolic actions of PTH in bone.

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General Transcription Factor IIα-γ Increases Osteoblast-specific Osteocalcin Gene Expression via Activating Transcription Factor 4 and Runt-related Transcription Factor 2*

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ATF4 (activating transcription factor 4) is an osteoblast-enriched transcription factor that regulates terminal osteoblast differentiation and bone formation. ATF4 knock-out mice have reduced bone mass (severe osteoporosis) throughout life. Runx2 (runt-related transcription factor 2) is a runt domain-containing transcription factor that is essential for bone formation during embryogenesis and postnatal life. In this study, we identified general transcription factor IIα-γ (TFIIA-γ) as a Runx2-interacting factor in a yeast two-hybrid screen. Immunoprecipitation assays confirmed that TFIIAγ interacts with Runx2 in osteoblasts and when coexpressed in COS-7 cells or using purified TFIIA-γ-transferase fusion proteins. Chromatin immunoprecipitation assay of MC3T3-E1 (clone MC-4) preosteoblasts and when coexpressed in COS-7 cells or using purified TFIIA-γ-transferase fusion proteins. Chromatin immunoprecipitation assay of MC3T3-E1 (clone MC-4) preosteoblasts showed that in intact cells TFIIAγ is recruited to the region of the osteocalcin promoter previously shown to bind Runx2 and ATF4. A small region of Runx2 (amino acids 258–286) was found to be required for TFIIAγ binding. Although TFIIAγ interacts with Runx2, it does not activate Runx2. Instead, TFIIAγ binds to and activates ATF4. Furthermore, TFIIAγ together with ATF4 and Runx2 stimulates osteocalcin promoter activity and endogenous mRNA expression. Small interfering RNA silencing of TFIIAγ markedly reduces levels of endogenous ATF4 protein and Ocn mRNA in osteoblastic cells. Overexpression of TFIIAγ increases levels of ATF4 protein. Finally, TFIIAγ significantly prevents ATF4 degradation. This study shows that a general transcription factor, TFIIAγ, facilitates osteoblast-specific gene expression through interactions with two important bone transcription factors ATF4 and Runx2.

Skeletal integrity requires a balance between bone-forming cells (osteoblasts) and bone-resorbing cells or osteoclasts. Imbalance between bone formation and resorption results in metabolic bone diseases such as osteoporosis. Multipotent mesenchymal cells proliferate and differentiate into osteoblasts that synthesize and deposit the mineralizing extracellular matrix of bone. Osteoblast activity is regulated by a number of growth factors and hormones, including bone morphogenetic proteins, insulin-like growth factor 1, basic fibroblast growth factor 2, parathyroid hormone, tumor necrosis factor-α, and extracellular matrix signals (1–9). Runx2 is a runt domain-containing transcription factor identified as a transcriptional activator of osteoblast differentiation and the master gene for bone development in vitro and in vivo (10–14). Runx2 knock-out mice die at birth and completely lack both skeletal ossification and mature osteoblasts (10, 12). Runx2 haplo-insufficiency causes the skeletal disorder, cleidocranial dysplasia, a disease characterized by defective endochondral and intramembranous bone formation. Runx2 is expressed in mesenchymal condensations during early development at E11.5 and acts as an osteoblast differentiation factor (13).

ATF4 (activating transcription factor 4), also known as CREB2 (cAMP-response element-binding protein 2) (15) and Tax-responsive Enhancer Element B67 (TAXREB67) (16), is a member of the activating transcription factor cAMP-response element-binding protein family of leucine zipper factors that also includes cAMP-response element-binding protein 2, CREB2 (cAMP-response element-binding protein), cAMP-response element modulator (CREM) and ATF1, ATF2, ATF3, and ATF4 (17–21). These proteins bind to DNA via their basic region and dimerize via their leucine domain to form a large variety of homodimers and/or heterodimers that allow the cell to coordinate signals from multiple pathways (17–21). An in vivo role for ATF4 in bone development was established using Atf4-deficient mice (22). ATF4 is required for expression of osteocalcin (Ocn) and bone sialoprotein (Bsp) as demonstrated by a dramatic reduction of their mRNAs in Atf4−/− bone (22). ATF4 activates Ocn transcription through direct binding to the OSE1 site of the mOG2 promoter. In addition, ATF4 interacts with Runx2 in osteoblasts or when coexpressed in COS-7 cells. ATF4 and Runx2 cooperatively regulate Ocn transcription through interactions with OSE1 (osteoblast-specific element 1).

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‡ The on-line version of this article (available at http://www.jbc.org) contains supplemental Figs. S1 and S2.

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and OSE2 (osteoblast-specific element 2, also known as nuclear matrix protein 2 or NMP2-binding site) sites in the promoter (23–25).

One of the most striking characteristics of ATF4 protein is its very short half-life (30–60 min) in many cell types (26). ATF4 is rapidly degraded via a ubiquitin/proteasomal pathway. This degradation requires the presence of the serine residue 219 in the context of DSGXXXS within the ATF4 molecule and its phosphorylation by an unknown kinase. This phosphorylation was shown to be required for subsequent recognition by the SCF\(^{\text{ATF4}}\) and degradation by the 26 S proteasome (27). Although ATF4 mRNA is ubiquitously expressed, ATF4 protein preferentially accumulates in osteoblasts (28). This accumulation is explained by a selective reduction of proteasomal degradation in osteoblasts. Indeed, inhibition of the ubiquitin/proteasomal pathway by MG115, which blocks the N-terminal threonine in the active site of 26 S proteasome (29, 30), led to ATF4 accumulation and induced Ocn mRNA expression in non-osteoblastic cells (28). These observations suggest that modulation of ATF4 stability constitutes an important step to control its protein level and activity and, ultimately, osteoblast-specific gene expression and bone formation.

Transcription factor II A (TFIIA) is a general transcription factor consisting of three subunits designated TFIIA\(\alpha\), TFIIA\(\beta\), and TFIIA\(\gamma\) (31). TFIIA interacts with and stabilizes TFIID (also known as TBP, TATA box-binding protein) to DNA and activates transcription (32, 33). Although TFIIA was classified as a general transcription factor when it was first identified, more and more evidence shows that this elusive factor may play an important role in the regulation of tissue-specific gene expression via interactions with tissue- or cell type-specific transcription factors (34–36).

The Ocn promoter has been the major paradigm for unraveling the mechanisms mediating osteoblast-specific gene expression and defining a number of key transcription factors or cofactors (13, 14, 23–25, 37–41). However, very few studies have focused on how tissue-specific transcription factors interface with general transcriptional initiation factors in osteoblasts. In this study, by using a combination of a yeast two-hybrid system and pulldown assays as well as functional assays, we show that TFIIA\(\gamma\), the smallest subunit (12 kDa) of TFIIA (42), interacts with both Runx2 and ATF4. TFIIA\(\gamma\) delays ATF4 protein degradation and increases its activity. Together with ATF4 and Runx2, TFIIA\(\gamma\) enhances osteoblast-specific Ocn gene expression.

**EXPERIMENTAL PROCEDURES**

**Reagents**—Tissue culture media were purchased from Invitrogen and fetal bovine serum from HyClone (Logan, UT). Other reagents were obtained from the following sources: antibodies against TFIIA-\(\alpha\), TFIIA-\(\gamma\), ATF4, Runx2, and horseradish peroxidase-conjugated mouse or goat IgG from Santa Cruz Biotechnology (Santa Cruz, CA), mouse monoclonal antibody against \(\beta\)-actin from Sigma, and GST antibody from Amersham Biosciences. All other chemicals were of analytical grade.

**Cell Cultures**—Mouse MC3T3-E1 subclone 4 (MC-4) cells were described previously (43, 44) and maintained in ascorbic acid-free \(\alpha\)-modified Eagle’s medium, 10% fetal bovine serum (FBS), and 1% penicillin/streptomycin and were not used beyond passage 15. C2C12 myoblasts, a gift from Dr. Daniel Goldman (University of Michigan, Ann Arbor, MI), C3H10T1/2 fibroblasts (American Type Culture Collection), and 3T3-L1 mouse preadipocytes (American Type Culture Collection) were maintained in Dulbecco’s modified Eagle’s medium, 10% FBS. F9 teratocarcinoma cells (American Type Culture Collection) and rat ROS17/2.8 osteosarcoma cells (gift from Dr. Laurie McCauley, University of Michigan School of Dentistry) were grown in modified Eagle’s medium, 10% FBS.

**Yeast Two-hybrid Analysis**—A yeast pLexA two-hybrid system (Clontech) was used to identify proteins that bind to mouse Runx2. A CDNA fragment encoding the aa-263–351 region of Runx2 was subcloned into the BamHI/XhoI sites of pLexA, creating an in-frame fusion with the DNA binding domain of the LexA gene that is controlled by the strong yeast ADH1 promoter. The resultant plasmid pLexA-Runx2 (aa 263–351) was then transformed into a yeast reporter strain (YM4271), and the transformed cells (1 \(\times\) 10\(^9\)) were mated for 24 h with cells (2.5 \(\times\) 10\(^9\)) of a pretransformed two-hybrid library made from human brain cDNA. The resultant mating mixture was spread on 20 \(\times\) 10-cm plates to select for expression of the LEU2 and lacZ reporter genes. Approximately 2 \(\times\) 10\(^6\) colonies were screened. Sixty four positive colonies were isolated. The prey plasmids were extracted from the positive colonies and the cDNA inserts in the plasmids were amplified by PCR and sequenced. Of the 64 positive colonies, 5 are the full-length TFIIA-\(\gamma\) cDNAs, and the rest contained 16 different cDNAs.

**DNA Constructs and Transfection**—p657mOG2-luc, p657mOG2OSE1mt-luc, p657mOG2OSE2mt-luc, p657mOG2-OS2 (OSE1 + 2)mt-luc, p4OSE1-luc, p4OSE1mt-luc, p6OSE2-luc, p6OSE2mt-luc, pCMV/\(\beta\)-galactosidase, pCMV/ATF4, pCMV/Runx2, pCMV/FLAG-Runx2 and its deletion mutants (aa 1–300, aa 1–286, and aa 258), GST-Runx2 and GST-ATF4 fusion protein expression vectors were described previously (1, 13, 23, 25, 45). The full-length cDNA of human TFIIA-\(\gamma\) was cloned by an RT-PCR strategy using total RNA from human Saos2 osteoblastic cells as a template and specific primers (forward, 5’-ATG GCA TAT CAG TTA TAC AGA AA-3’; and reverse, 5’-TTC TGT AGT ATT GGA GCC AGT A-3’). Digested PCR products were purified and subcloned into the NotI/BamHI sites of the pFLAG-5a expression vector (Sigma). Addition of a C-terminal FLAG sequence into the TFIIA-\(\gamma\) cDNA facilitates monitoring of expression levels and immunoprecipitation using M2 antibody (Sigma). GST-TFIIA-\(\gamma\) fusion protein expression plasmid was constructed by subcloning the full-length TFIIA-\(\gamma\) cDNA into the glutathione S-transferase gene fusion vector pGEX-4T-1 (Amersham Biosciences) in correct reading frame. The accuracy of DNA sequences was verified by automatic sequencing. The size of expressed proteins was confirmed by Western blot analysis using specific antibodies. For expression and functional studies, cells were plated on 35-mm dishes at a density of 5 \(\times\) 10\(^4\) cells/cm\(^2\). After 24 h, cells were transfected with the indicated plasmid DNAs (0.01 \(\mu\)g of pRL-SV40, 0.25 \(\mu\)g of test luciferase reporter, and 1.0 \(\mu\)g of expression plasmids balanced as necessary with \(\beta\)-galactosidase expression plasmid such that the total DNA was constant).
and Lipofectamine 2000 (Invitrogen) according to manufacturer’s instructions. After 36 h, whole cell extracts were prepared and used for Western blot analysis or dual luciferase assay using the dual luciferase assay kit (Promega, Madison, WI) on a Veritas™ microplate luminometer (Turner Biosystem, Inc., Sunnyvale, CA). Firefly luciferase activity was normalized to Renilla luciferase activity for transfection efficiency.

RNA Isolation and Reverse Transcription (RT)—Total RNA was isolated using TRIzol reagent (Invitrogen) according to the manufacturer’s protocol. RT was performed using 2 μg of denatured RNA and 100 pmol of random hexamers (Applied Biosystems, Foster City, CA) in a total volume of 25 μl containing 12.5 units of MultiScribe reverse transcriptase (Applied Biosystem, Foster, CA) according to the manufacturer’s instructions.

Regular PCR—Regular PCR was performed on a 2720 Thermal Cycler (Applied Biosystem, Foster, CA), using 2.5 μl of the cDNA (equivalent to 0.2 μg of RNA) and AmpliTaq Gold DNA polymerase (Applied Biosystems, Foster City, CA) in a 25-μl reaction according to the manufacturer’s instructions. The DNA sequences of primers used for PCR were as follows: mouse/rat TFIIAγ, 5‘-ATG GCA TAT CAG TTA TCA AGA AAT ACA-3′ (forward), 5′-GGT ATT TTT ACC ATC ACA GGC T-3′ (reverse); mouse/rat Hprt, 5′-GCT CTG GAG TGG AAG ACA GAA C-3′ (reverse); mouse/rat Bsp, 5′-GTT GAG AGA TCA TCT CCA CC-3′ (forward), 5′-AGC GAT GAT GAA CCA CCT GGT TA-3′ (reverse). For all primers the amplification was performed as follows: initial denaturation at 95 °C for 30 s followed by 31 cycles of 95 °C for 15 s, 60 °C for 30 s, 72 °C for 30 s per reaction according to the manufacturer’s instructions. DNA sequences of primers used for PCR were as follows: mouse/rat TFIIAγ, 5‘-ATG GCA TAT CAG TTA TCA AGA AAT ACA-3′ (forward), 5′-GGT ATT TTT ACC ATC ACA GGC T-3′ (reverse); mouse/rat Hprt, 5′-GCT CTG GAG TGG AAG ACA GAA C-3′ (reverse); mouse/rat Bsp, 5′-GTT GAG AGA TCA TCT CCA CC-3′ (forward), 5′-AGC GAT GAT GAA CCA CCT GGT TA-3′ (reverse).

Quantitative Real Time PCR—Quantitative real time PCR was performed on an iCycler (Bio-Rad) using a SYBR® Green PCR core kit (Applied Biosystem, Foster, CA) and cDNA equivalent to 10 ng of RNA in a 50-μl reaction according to the manufacturer’s instructions. The DNA sequences of primers used for real time PCR were as follows: mouse Ocn, 5′-TAG TGA ACA GAC TCC GGC GCT A-3′ (forward), 5′-TGT AGG CGG TCT TCA AGC CAT-3′ (reverse); mouse and rat 18S rRNA, 5′-CGT CTG CCC TAT CAA CTT TCG ATG GTA G-3′ (forward), 5′-GCC TGC TGC CTT CCT TGT ATG T-3′ (reverse); mouse and rat TFIIAγ, 5′-TGG GGA ACA GTC TTT CTC AAG AGA GCC TT-3′ (forward), 5′-TTC CTG ACT CTC TCT GGC CAA AAT GCT G-3′ (reverse); rat Ocn, 5′-TGG TGA ATA GAC TCC GGC GCT ACC T-3′ (forward), 5′-CCT GGA AGA CCA TGT GGT CCG-3′ (reverse); rat Bsp, 5′-GGC TGG AGA TGA GGA GGA CCC C-3′ (forward), 5′-GGT TGG TCG TGG TGC CGT TGA CGA CCT-3′ (reverse); rat Opn, 5′-TGG TGA ATA GAC TCC GGC GCT ACC T-3′ (forward), 5′-CCT GGA AGA CCA TGT GGT CCG-3′ (reverse). For all primers the amplification was performed as follows: initial denaturation at 95 °C for 10 min followed by 40 cycles of 95 °C for 15 s and 60 °C for 60 s. Melting curve analysis was used to confirm the specificity of the PCR products. Six samples were run for each primer set. The levels of mRNA were calculated by the ∆CT method (46). Ocn, Bsp, TFIIAγ, osteopontin (Opn), and Atf4 mRNAs were normalized to 18 S rRNA mRNA.

Western Blot Analysis—Cells were washed with cold 1× phosphate-buffered saline and lysed in 1× Passive Buffer (Promega, Madison, WI) at room temperature for 20 min. Lysates were clarified by centrifugation (20 min, 13,000 × g, 4 °C). Protein concentrations were determined by the method developed by Bio-Rad. Twenty μg of total protein were fractionated on a 10% SDS-polyacrylamide gel and transferred onto nitrocellulose membranes (Schleicher & Schuell). The membrane was blocked in 5% nonfat milk in Tris-buffered saline/Tween 20 (TBST) buffer, probed with antibodies against TFIIA-γ (1:200), TFIIA-α (1:1000), ATF4 (1:1000), Runx2 (1:1000), Fra-1 (1:1000), GST (1:5000), or M2 (1:2000) followed by incubation with anti-goat-mouse or -rabbit antibodies conjugated with horseradish peroxidase (1:5000) and visualized using an enhanced chemiluminescence kit (Pierce). Finally, blots were stripped two times in buffer containing 65 mM Tris-Cl, pH 6.8, 2% SDS, and 0.7% (v/v) β-mercaptoethanol at 65 °C for 15 min and re-probed with β-actin antibody (1:5000) for normalization.

Immunoprecipitation—GST, GST-TFIIAγ, GST-ATF4, and GST-Runx2 fusion proteins were purified using the Bulk GST purification module kit (Amersham Biosciences) according to the manufacturer’s instructions. Whole cell extracts (500 μg), nuclear extracts (200 μg), or GST fusion proteins (1.0 μg) were pre-cleaned twice with 50 μl of protein A/G-agarose beads (Stratagene, La Jolla, CA) for 30 min followed by pelleting of beads. The protein A/G-agarose beads were blocked with 10 μg/ml bovine serum albumin in 1× phosphate-buffered saline for 1 h before use to reduce nonspecific binding of proteins. Five μg of respective antibody was added and incubated for 2 h at 4 °C with gentle rocking. The immune complexes were collected by addition of 30 μl of protein A/G-agarose beads and incubation for 1 h at 4 °C followed by centrifugation. Precipitates were washed five times with 1× washing buffer (20 mM HEPES, pH 7.6, 50 mM KCl, 1 mM dithiothreitol, 0.25% Nonidet P-40, 5 mM NaF, 1 mM EGTA, 5 mM MgCl2, 0.25 mM phenylmethylsulfonyl fluoride), and the immunoprecipitated complexes were suspended in SDS sample buffer and analyzed by SDS-PAGE followed by Western blot analysis using the indicated antibodies.

ChIP Assays—ChIP assays were performed as described previously (41) using a protocol kindly provided by Dr. Dwight Towler (Washington University) (47). After sonication, the amount of chromatin was quantified using the PicoGreen double-stranded DNA quantitation assay (Molecular Probes) according to the manufacturer’s instructions. The equivalent of 10 μg of DNA was used as starting material (input) in each ChIP reaction with 2 μg of the appropriate antibody (TFIIAγ or control rabbit IgG). Fractions of the purified ChIP DNA (5%) or inputs (0.02–0.05%) were used for PCR analysis. The reaction was performed with AmpliTaq Gold DNA polymerase (Applied Biosystems) for 35 cycles of 60 s at 95 °C, 90 s at 58 °C, and 120 s at 68 °C. PCR primer pairs were generated to detect DNA segments located near the Runx2-binding site at −137/−131 (primers P1 and P2), ATF4-binding site at −55/−48 (primers P3 and P4) in mouse osteocalcin gene 2 (mOG2) proximal promoter, or the Runx2-binding site located between −370 and −42 in the proximal mouse Runx2 promoter region (primers
control siRNA (low GC, catalog number 12935-200, Invitrogen) using Lipofectamine 2000 (Invitrogen) according the manufacturer's instruction. After 36 h, total RNA was harvested for quantitative real time RT-PCR analysis for TFIIAγ, Ocn, Bsp, Opn (osteopontin), and Atf4 mRNAs. A second set of mouse TFIIAγ siRNAs (sense, AUC ACA ACA CUG UGC UAU AUU; antisense, UAU AGC ACA GUG UUG UCA UUU) was designed in the project laboratory and used to confirm the results using the first set of TFIIAγ siRNA.

Statistical Analysis—Results were expressed as means ± S.D. Students' t test was used to test for differences between two groups. Differences with a p < 0.05 was considered as statistically significant.

RESULTS

TFIIAγ Interacts with Runx2 and ATF4—A yeast pLexA two-hybrid system (Clontech) was used to identify proteins that bind to mouse Runx2. cDNA fragments encoding several C-terminal regions of Runx2 were subcloned into the BamHI/XhoI sites of pLexA, creating in-frame fusions with the DNA binding domain of the LexA gene that is controlled by the strong yeast ADH1 promoter. Preliminary experiments using relatively larger regions of Runx2 (aa 232–391, aa 232–428, and aa 232–517) as baits were not successful because of their inability to autoactivate the lacZ reporter gene in yeast. In contrast, by using the aa 263–351 region of Runx2 as a bait, we identified TFIIAγ, a general transcriptional factor involved in the initiation step of eukaryotic transcription, as a Runx2-interacting factor. A diagram and a picture of a positive colony are shown in Fig. S1.

To verify the TFIIAγ-Runx2 interaction identified by yeast two-hybrid system, we conducted pulldown assays. COS-7 cells were transiently transfected with expression vectors for FLAG-TFIIAγ, Runx2, and ATF4 (a recently identified Runx2-interacting factor). After 36 h, whole cell extracts were prepared for immunoprecipitation (IP) assay using a TFIIAγ antibody followed by Western blot analysis for Runx2 and ATF4. As seen in Fig. 1A (lane 2), Runx2 protein was present in a TFIIAγ anti-
**TFIIAγ Interacts with ATF4 and Runx2**

![Diagram](image)

**FIGURE 2.** ChIP analysis of TFIIAγ interaction with Runx2/ATF4 binding sites-containing chromatin fragments of mOG2 promoter in MC-4 cells. A, schematic representation of relevant regions of the mOG2 promoter, mouse Runx2 promoter, and mOG2 gene. P1, P2, P3, P4, P5, P6, P7, and P8 indicate PCR primers used to analyze ChIP DNAs. The positions of these primers and the size of the fragments they amplify are indicated at the top or bottom of the figure. B, MC-4 cells were seeded at a density of 50,000 cells/cm² in 35-mm dishes, cultured in 10% FBS medium overnight, and cross-linked with formaldehyde for ChIP assays. IPs were conducted with TFIIAγ antibody (Ab) or normal control IgG. PCR products were run on 3% agarose gel and stained with ethidium bromide. Purified input chromatin was used to perform parallel PCRs with the respective primer sets of TFIIAγ interaction with Runx2 and ATF4 in osteoblasts or when coexpressed in COS-7 cells.

**TABLE 1**

<table>
<thead>
<tr>
<th>Oligonucleotide name</th>
<th>Sequence</th>
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<tbody>
<tr>
<td>P1</td>
<td>CGGCTTCGACGGCGACAGAC</td>
</tr>
<tr>
<td>P2</td>
<td>AGGGGATGCTGCCAGGACTAAT</td>
</tr>
<tr>
<td>P3</td>
<td>TGCATAACCTTCTTTTCAG</td>
</tr>
<tr>
<td>P4</td>
<td>AGCACTATTGTCGAAGACAGAATC</td>
</tr>
<tr>
<td>P5</td>
<td>TATGAGACACGGTCCGCACTTA</td>
</tr>
<tr>
<td>P6</td>
<td>TTATCGGCGGCTTCCAAGCAGCAT</td>
</tr>
</tbody>
</table>

As a first step to identify the TFIIAγ-binding domain, FLAG-Runx2 deletion mutant expression vectors (wild type aa 1–528, aa 1–330, aa 1–286, and aa 1–258) were transfected into COS-7 cells because of the high transfection efficiency. Nuclear extracts were prepared 36 h later, mixed with equal amounts of nuclear extracts of ROS17/2.8 (which contain large amounts of endogenous TFIIAγ), and immunoprecipitated using anti-TFIIAγ antibody followed by Western blot analysis for Runx2 (M2 antibody). As shown in Fig. 1D, deletion of Runx2 from aa 528 to aa 286 did not reduce TFIIAγ binding. However, further deletion from aa 286 to aa 258 completely abrogated TFIIAγ-Runx2 complex formation. These data clearly demonstrate the following: (i) endogenous TFIIAγ can interact with overexpressed FLAG-Runx2 proteins in vitro; and (ii) the aa 258–286 region of Runx2 is required for TFIIAγ binding. Interestingly, this same region is required for ATF4–Runx2 interactions (25).

To determine whether, in intact cells, TFIIAγ is associated with the endogenous osteocalcin gene 2 (mOG2) promoter region that has been shown to bind Runx2 and ATF4, we performed the chromatin immunoprecipitation (ChIP) assay using MC3T3-E1 (clone MC-4) preosteoblast cells. After shearing, soluble chromatin was immunoprecipitated with either an antibody against TFIIAγ or control IgG. The positions and sequences of primers used for PCR analysis of ChIP DNAs are shown in Fig. 2A and Table 1. As shown in Fig. 2B, the PCR bands amplified with primers P1/P2 and P3/P4 and corresponding to ChIP DNAs immunoprecipitated with TFIIAγ antibody revealed that TFIIAγ specifically interacts with chromatin fragments of the proximal mOG2 promoter that contain Runx2- or ATF4-binding sites. Furthermore, TFIIAγ antibody
TFIIAγ Interacts with ATF4 and Runx2

**FIGURE 3.** TFIIAγ increases ATF4 but not Runx2 transcriptional activity. A and B, 10T1/2 cells were transiently transfected with p6OSE2-luc (A) or p6OSE2mt-luc (B) and pRL-SV40 (for normalization) and expression plasmids for β-galactosidase, TFIIAγ, Runx2, or Runx2 plus TFIIAγ. After 36 h, cells were harvested for dual luciferase assay. Firefly luciferase was normalized to *Rotylenchulus reniformis* luciferase to control the transfection efficiency (*, p < 0.01; β-galactosidase versus Runx2 or Runx2+TFIIAγ). C and D, 10T1/2 cells were transiently transfected with p4OSE2-luc (C) or p4OSE1mt-luc (D) and pRL-SV40 and expression plasmids for β-galactosidase, TFIIAγ, ATF4, or ATF4 plus TFIIAγ (*, p < 0.01; β-galactosidase versus ATF4 or ATF4+TFIIAγ); #, p < 0.01 (ATF4 versus ATF4+TFIIAγ). E, 10T1/2 cells were transiently transfected with −34/−13 mOG2-luc and pRL-SV40 and expression plasmids for β-galactosidase, TFIIAγ, ATF4, or Runx2. F, dose–response experiment, 10T1/2 cells were transiently transfected with p4OSE1-luc and pRL-SV40 and ATF4 expression plasmid and increasing amounts of TFIIAγ plasmid (*, p < 0.01; β-galactosidase versus TFIIAγ). G and H, C2C12 (G) and COS-7 cells (H) were transiently transfected with p4OSE2-luc and pRL-SV40 and expression plasmids for β-galactosidase, TFIIAγ, ATF4, or ATF4 plus TFIIAγ (*, p < 0.01; β-galactosidase versus ATF4 or ATF4+TFIIAγ). Data represent mean ± S.D. Experiments were repeated three times and qualitatively identical results were obtained. Note the expanded scale for the mutant reporters (B, D, and E) because of low basal activity to enable visualization of any potential differences as a consequence of cotransfection with the expression vectors noted above.

also immunoprecipitated a Runx2-binding site-containing chromatin fragment of the proximal Runx2 promoter (primers P5/P6). In contrast, TFIIAγ antibody failed to immunoprecipitate a chromatin fragment of mOG2 gene that contains no Runx2- or ATF4-binding sites (primers P7/P8). Taken together, these data show that TFIIAγ is recruited to a chromatin fragment of the mOG2 promoter that was previously demonstrated to be bound by Runx2 and ATF4 in osteoblasts (13, 22).

**TFIIAγ Increases ATF4 but Not Runx2-dependent Transcriptional Activity**—To determine whether TFIIAγ increases Runx2- and ATF4-dependent transcriptional activity, we measured the ability of TFIIAγ to stimulate transcription of p6OSE2-luc, a reporter plasmid containing 6 copies of the Runx2-binding element OSE2 upstream of a minimal 34-bp mOG2 promoter (13, 43, 49) or p4OSE1-luc, a reporter plasmid that contains four copies of OSE1 (a specific ATF4-binding element) upstream of a minimal 34-bp mOG2 promoter (22, 25). For these studies, we used C3H10T1/2 fibroblasts because they contain undetectable levels of both endogenous Runx2 and ATF4 proteins (28, 49). As shown in Fig. 3A, as expected, Runx2 alone increased OSE2 transcriptional activity by 11-fold. This stimulation was abolished in the 6OSE2mt-luc in which the OSE2 core sequence was mutated (25) (Fig. 3B). Although we have shown above that TFIIAγ interacts with Runx2, TFIIAγ transfection did not activate basal or Runx2-dependent OSE2 transcription (Fig. 3A). As shown in Fig. 3C, ATF4 activated OSE1 activity by 2-fold (p < 0.01, β-galactosidase versus ATF4). Although TFIIAγ alone was unable to activate OSE1 activity, unexpectedly, when coexpressed with ATF4, it dramatically increased OSE1 activity 5-fold above ATF4 alone. This stimulation was abolished in 4OSE1mt-luc, in which the OSE1 core sequence was mutated from TTACATCA to TTAGTACA in the reporter plasmid (45) (Fig. 3D). Note: TFIIAγ, Runx2, or ATF4 failed to activate a minimal 34-bp mOG2 promoter that contains a TATA box (23, 50) (Fig. 3E). Fig. 3F shows that TFIIAγ activated ATF4 transcription activity in a dose-dependent manner in C3H10T1/2 cells. TFIIAγ similarly stimulated ATF4-directed OSE1 activity in C2C12 myoblasts (3-fold) and COS-7 cells (4.3-fold) (Fig. 3, G and H).

**TFIIAγ Expression in Different Cell Lines**—The levels of TFIIAγ mRNAs and proteins were determined in different cell lines by RT-PCR and Western blot analysis, respectively. As shown in Fig. 4, Western blot analysis shows that TFIIAγ protein was expressed at high levels in osteoblastic cells (MC-4
Silencing of TFIIAγ Markedly Reduces Levels of Endogenous Ocn and Bsp mRNAs and ATF4 Protein in Osteoblasts—To determine whether TFIIAγ is required for the endogenous Ocn mRNA expression in osteoblasts, we knocked down the endogenous TFIIAγ transcripts by siRNA. ROS17/2.8 osteoblast-like cells, which express high levels of TFIIAγ and Ocn and Bsp mRNAs, were transiently transfected with TFIIAγ siRNA reagent from Santa Cruz Biotechnology according to the manufacturer’s instructions. This siRNA is a pool of three specific 20–25-nucleotide siRNA targeting both mouse and rat TFIIAγ. As shown in Fig. 6A, quantitative real-time RT-PCR analysis showed that levels of TFIIAγ mRNA were efficiently reduced by TFIIAγ siRNA in a dose-dependent manner. The level of Ocn mRNA was reduced by greater than 50% by TFIIAγ siRNA (p < 0.01, control versus TFIIAγ siRNA). Interestingly, Bsp mRNA, another ATF4 downstream target gene (22), was also reduced by 50% (p < 0.01, control versus TFIIAγ siRNA). This inhibition was specific because levels of Opn and Atf4 mRNAs were not reduced by TFIIAγ siRNA. In contrast, as shown in Fig. 6B, levels of all these mRNAs were not reduced by the negative control siRNA (Invitrogen). Although Atf4 mRNA was not altered by TFIIAγ siRNA, the level of endogenous ATF4 protein was significantly reduced by silencing TFIIAγ in osteoblasts (Fig. 6C). Similar results were obtained when a different set of TFIIAγ siRNA was used (Fig. S2).

Overexpression of TFIIAγ Increases the Levels of ATF4 Protein—The above studies clearly demonstrated that TFIIAγ increased ATF4-dependent transcription activity and Ocn gene expression probably by targeting ATF4 protein. To further study the mechanism of this regulation, we determined the effect of TFIIAγ overexpression on the levels of ATF4 protein. C3H10T1/2 cells, which express undetectable level of endogenous ATF4 protein (28), were transiently transfected with indicated mRNAs and mOG2 promoter activity and Ocn mRNA expression was dependent upon the presence of Runx2 via a mechanism involving protein–protein interactions (25). To determine the effects of TFIIAγ on endogenous Ocn mRNA expression, C3H10T1/2 cells were transiently transfected with expression vectors for β-galactosidase, TFIIAγ, ATF4, Runx2, ATF4/Runx2, TFIIAγ/Runx2, TFIIAγ/ATF4, and ATF4/Runx2/TFIIAγ. After 36 h, cells were harvested for RNA preparation and quantitative real-time RT-PCR detection of Ocn mRNA. As shown in Fig. 5A, consistent with its role as a master gene of osteoblast differentiation, Runx2 alone increased endogenous Ocn expression by 3.3-fold (p < 0.01; β-galactosidase versus Runx2). TFIIAγ alone, ATF4 alone, and TFIIAγ/ATF4 were not all sufficient for activation of endogenous Ocn mRNA expression. TFIIAγ alone did not enhance Runx2-dependent Ocn expression. As demonstrated previously (25), ATF4 dramatically stimulated Runx2-dependent Ocn mRNA expression by 10-fold (p < 0.01, Runx2 versus Runx2/ATF4). Importantly, TFIIAγ further augmented Ocn mRNA expression 4.2-fold in the presence of ATF4 and Runx2 (p < 0.01, ATF4/Runx2 versus ATF4/Runx2/TFIIAγ). TFIIAγ similarly enhanced ATF4/Runx2-dependent 657-bp mOG2 promoter activity in C3H10T1/2 cells (3.6-fold) (Fig. 5B) (p < 0.01, ATF4/Runx2 versus ATF4/Runx2/TFIIAγ). This stimulation was completely ablated by point mutations in the OSE1 and/or OSE2 core sequences.

Overexpression of TFIIAγ Increases ATF4 Protein Stability—Lassot et al. (51) recently showed that acetylase p300 markedly increased the levels of ATF4 protein and ATF4-dependent transcriptional activity by inhibiting ATF4 protein degradation via a proteasomal ubiquitin pathway. As an initial step to determine whether TFIIAγ alters ATF4 protein stability, C3H10T1/2 cells were transiently transfected with expression vectors for ATF4 and Runx2. Western blot analysis was performed to determine whether TFIIAγ alters ATF4 protein stability. To further study the mechanism of this regulation, we determined the effect of TFIIAγ overexpression on the levels of ATF4 protein. C3H10T1/2 cells, which express undetectable level of endogenous ATF4 protein (28), were transiently transfected with indicated mRNAs and mOG2 promoter activity and Ocn mRNA expression was dependent upon the presence of Runx2 via a mechanism involving protein–protein interactions (25). To determine the effects of TFIIAγ on endogenous Ocn mRNA expression, C3H10T1/2 cells were transiently transfected with expression vectors for β-galactosidase, TFIIAγ, ATF4, Runx2, ATF4/Runx2, TFIIAγ/Runx2, TFIIAγ/ATF4, and ATF4/Runx2/TFIIAγ. After 36 h, cells were harvested for RNA preparation and quantitative real-time RT-PCR detection of Ocn mRNA. As shown in Fig. 5A, consistent with its role as a master gene of osteoblast differentiation, Runx2 alone increased endogenous Ocn expression by 3.3-fold (p < 0.01; β-galactosidase versus Runx2). TFIIAγ alone, ATF4 alone, and TFIIAγ/ATF4 were not all sufficient for activation of endogenous Ocn mRNA expression. TFIIAγ alone did not enhance Runx2-dependent Ocn expression. As demonstrated previously (25), ATF4 dramatically stimulated Runx2-dependent Ocn mRNA expression by 10-fold (p < 0.01, Runx2 versus Runx2/ATF4). Importantly, TFIIAγ further augmented Ocn mRNA expression 4.2-fold in the presence of ATF4 and Runx2 (p < 0.01, ATF4/Runx2 versus ATF4/Runx2/TFIIAγ). TFIIAγ similarly enhanced ATF4/Runx2-dependent 657-bp mOG2 promoter activity in C3H10T1/2 cells (3.6-fold) (Fig. 5B) (p < 0.01, ATF4/Runx2 versus ATF4/Runx2/TFIIAγ). This stimulation was completely ablated by point mutations in the OSE1 and/or OSE2 core sequences.
transiently transfected with ATF4 expression vector in the presence of β-galactosidase, TFIIAγ, or Runx2 expression vectors. After 36 h, cells were treated with 50 μg/ml of protein synthesis inhibitor cycloheximide (CHX) (i.e. to completely block de novo protein synthesis) and harvested at different time points of CHX addition (0, 0.5, 1, and 3 h) followed by Western blot analysis for ATF4 and Runx2. This technique has been widely used to study protein stability (51). As shown in Fig. 8A, in the absence of TFIIAγ overexpression, ATF4 protein was rapidly degraded and almost undetectable on Western blot by 3 h after CHX addition, which is consistent with a previous study (51). However, overexpression of TFIIAγ greatly delayed the degradation process with the levels of ATF4 protein only slightly reduced by 3 h after CHX addition. In contrast, levels of Runx2 protein were not affected by TFIIAγ (Fig. 8B).

DISCUSSION

This study identifies TFIIAγ as a bridging molecule between Runx2, ATF4, and the transcription machinery in osteoblasts. Although Runx2 and ATF4 interact in osteoblasts or when coexpressed in COS-7 cells, IPs using purified GST fusion proteins were unable to demonstrate a direct physical interaction between ATF4 and Runx2 (25). Thus, accessory factors are likely involved in bridging these two molecules. Several lines of evidence support that TFIIAγ may be a factor linking Runx2 and ATF4. (i) TFIIAγ forms complexes with both Runx2 and ATF4 in osteoblasts and when coexpressed in COS-7 cells. (ii) The same region of Runx2 (i.e. aa 258–286) is required for both TFIIAγ-Runx2 and ATF4-Runx2 interactions. (iii) Purified GST-TFIIAγ fusion protein directly binds to both purified GST-Runx2 and GST-ATF4 fusion proteins. (iv) Overexpression of TFIIAγ in 10T1/2 cells dramatically enhances endogenous Ocn gene expression and the 657-bp mOG2 promoter activity in the presence of ATF4 and Runx2. (v) siRNA knockdown of TFIIAγ mRNA markedly reduces osteoblast-specific Ocn and Bsp expression.

Accumulating evidence establishes that ubiquitin-proteasome pathways control osteoblast differentiation and bone formation. For example, the proteasome inhibitors epoxomicin and proteasome inhibitor-1, when administered systemically to mice, strongly stimulated bone volume and bone formation rates by greater than 70% after only 5 days of treatment (52). Although the mechanism of this regulation remains unclear, critical bone transcription factors seem to be targets for the ubiquitin-proteasomal pathway. Zhao and co-workers (52, 53) recently showed that Smurf1, an E3 ubiquitin-protein isopeptide ligase, accelerated Runx2 ubiquitin-proteasomal degradation and inhibited osteoblast differentiation and bone forma-
ments from this study show that overexpression of TFIIA decreases ATF4 stability in osteoblasts. Lassot et al. (51) recently found that ATF4 is similarly stabilized by cofactor p300, a histone acetyltransferase. p300 inhibits ATF4 ubiquitination and degradation through interaction with the ATF4 N terminus. Interestingly, this stabilization does not require either the acetyltransferase activity of p300 or the serine residue 219 in the context of DSGXXXS within ATF4 molecule that is known to be required for ATF4 degradation via the SCF\(^{\beta \text{TrCP}}\) and the 26 S proteasome (51).

TFIIA\(^{\gamma}\) stimulation of Ocn gene transcription is dependent on the presence of both ATF4 and Runx2. As a master regulator of osteoblast differentiation, Runx2 alone is sufficient to activate expression of many osteoblast-specific genes, including Ocn and Bsp, by direct binding to their promoters (13). In contrast, although ATF4 directly binds to the OSE1 site of the mouse Ocn gene and activates OSE1, it alone is not sufficient for activation of the endogenous Ocn gene or the 657-bp mOG2 promoter which contains sufficient information for the bone-specific expression of Ocn \textit{in vivo} (54). Instead, ATF4 stimulation of Ocn is dependent on the presence of Runx2 as demonstrated by our recent study (25). ATF4 interacts with Runx2 and activates Runx2-dependent transcriptional activity. A recent study shows that SATB2, a nuclear matrix protein that directly interacts with both ATF4 and Runx2, activates osteoblast differentiation and controls craniofacial patterning \textit{in vivo} (55). This study shows that although TFIIA\(^{\gamma}\) interacts with Runx2, it does not directly activate Runx2. Like ATF4, TFIIA\(^{\gamma}\) alone is not sufficient to activate transcription from either the Ocn gene or the 657-bp mOG2 promoter. In fact, even TFIIA\(^{\gamma}\) and ATF4 together are not sufficient for Ocn gene expression without the presence of Runx2 (Fig. 5). However, in the presence of both ATF4 and Runx2, TFIIA\(^{\gamma}\) greatly activates Ocn gene expression.

General transcription factors were originally defined as such because they were thought to be universally required for transcription. In eukaryotic cells, initiation of transcription is a complex process, which requires RNA polymerase II and many other basal transcription factors and/or cofactors, including TFIIA, TFIIH, TFID (TBP or TATA box-binding protein), TFIIIE, TFIIF, and TFIIH (56–59). Binding of TBP to the TATA box is the first step, which is regulated by TFIIA. TFIIA enhances transcription by interacting with TBP and stabilizing its binding to DNA (32, 33). More and more evidence shows that general transcription factors play unique roles in the regulation of tissue-specific gene expression under physiological and pathological conditions. For example, the androgen receptor, via its N-terminal AF1 domain, interacts with basal transcription factors TBP and TFIIA and activates tissue-specific transcription in target tissues and cells (60). Likewise, TAFII\(_{17}\) (a component of the TFID complex), via specific protein-protein interactions with the vitamin D receptor (VDR), increases osteoclast formation from osteoclast precursors in response to 1,25-dihydroxyvitamin D\(_3\) in patients with Paget disease (61). In osteoblasts, bone transcription factors such as Runx2 and ATF4 directly bind to specific DNA sequences in their target gene promoters (i.e. OSE2 or NMP2 and OSE1, respectively) and activate osteoblast-specific gene expression, osteoblast differentiation, and bone formation (1, 10–14, 24, 43). Obviously,
cooperative interactions between osteoblast-specific transcription factors and basal (general) transcriptional machinery are essential for achieving maximal transcription of osteoblast-specific genes. However, little is known about these interactions. Experiments from this study demonstrate that TFIIAγ, which is expressed at high level in osteoblasts, facilitates osteoblast-specific gene expression via two mechanisms. 1) TFIIAγ stabilizes ATF4 and increases the levels of ATF4 proteins. The increased levels of ATF4 further activate Runx2 activity and Ocn transcription (25). 2) Through its ability to directly interact with both ATF4 and Runx2, TFIIAγ could recruit these two critical bone transcription factors to the basal transcriptional machinery and greatly enhance osteoblast-specific gene expression. In support of our observation, Guo and Stein (62) showed that Yin Yang-1 (YY1) regulates vitamin D enhancement of Ocn gene transcription by interfering with interactions of the VDR with both the VDR element and TFIIB. TFIIB interacts with both VDR and YY1 (63). Likewise, Newberry et al. (64) showed that TFIIF (RAP74 and RAP30) mediates Msx2 (a homeobox transcription factor required for craniofacial development) inhibition of Ocn promoter activity. Finally, a recent study showed that TFIIB could directly bind to the transactivation domain of Osterix, another important osteoblast transcription factor (65).

TFIIAγ interacts with ATF4 and Runx2

FIGURE 7. TFIIAγ increases the levels of ATF4 protein. C3H10T1/2 (A) and COS-7 (B) cells were transfected with 1 μg of pCMV/ATF4 or pCMV/Runx2 and increasing amounts of FLAG-TFIIAγ expression vector (0, 0.5, 1, 2 μg) followed by Western blotting for ATF4, TFIIAγ, Runx2, and β-actin (top) or RNA preparation and RT-PCR for Atf4 and Hprt mRNA (bottom). ROS17/2.8 (C) and MC-4 (D) cells were transfected with increasing amounts of FLAG-TFIIAγ expression vector (0, 0.5, 1, and 2 μg). Experiments were repeated three times with similar results.

FIGURE 8. TFIIAγ increases ATF4 protein stability. C3H10T1/2 cells were transfected with 1.0 μg ATF4 (A) or Runx2 (B) expression vector with and without 1.0 μg of TFIIAγ expression vector. After 36 h, cells were treated with 50 μg/ml of protein synthesis inhibitor cycloheximide (CHX) and harvested at different time points (0, 1, and 3 h) followed by Western blot analysis for ATF4 and Runx2. Experiments were repeated three times with similar results.
and COS-7 on Western blots. The meaning of this observation remains unknown.

These findings suggest that TFIIAγ is a critical factor regulating ATF4 stability and functions as a molecular linker between ATF4 and Runx2 and the basal transcriptional machinery. TFIIAγ may play a unique role in the regulation of osteoblast-specific gene expression and ultimately osteoblast differentiation and bone formation. A working model is proposed in Fig. 9, which summarizes the role of TFIIAγ in osteoblast-specific mOG2 gene expression. Future study aimed at identifying factors that affect levels and activity of TFIIAγ will allow us to address the functional significance of TFIIAγ in osteoblast function in greater detail.

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TFIIαγ Interacts with ATF4 and Runx2
Activating Transcription Factor 4 Is Critical for Proliferation and Survival in Primary Bone Marrow Stromal Cells and Calvarial Osteoblasts

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ABSTRACT
Activating transcription factor 4 (ATF4) is essential for bone formation. However, the mechanism of its actions in bone is poorly understood. The present study examined the role for ATF4 in the regulation of proliferation and survival of primary mouse bone marrow stromal cells (BMSCs) and osteoblasts. Results showed that Atf4−/− cells display a severe proliferative defect as measured by multiple cell proliferation assays. Cell cycle progression of Atf4−/− BMSCs was largely delayed with significant G1 arrest. Expression of cyclin D1 was decreased both at the mRNA and protein level. A similar proliferation defect was observed in Atf4−/− calvarial periosteal osteoblasts when compared with wt control. Knocking down Atf4 mRNA by small interfering RNA in MC3T3-E1 subclone 4 preosteoblasts markedly reduced expression of cyclin D1 and cell proliferation. In contrast, overexpression of ATF4 increased cyclin D1 expression as well as cell proliferation in Atf4−/− BMSCs. In addition, apoptosis was significantly increased in Atf4−/− BMSCs and calvarial periosteal osteoblasts relative to wt controls. Taken together, these results for the first time demonstrate that ATF4 is a critical regulator of proliferation and survival in BMSCs and osteoblasts in vitro and in vivo. J. Cell. Biochem. 105: 885–895, 2008.

KEY WORDS: ATF4; OSTEOBLASTS; PROLIFERATION; APOPTOSIS; CELL CYCLE; CYCLIN D1

Activating transcription factor 4 (ATF4), also known as cAMP-response element-binding protein 2 (CREB2) [Karpinski et al., 1992] and Tax-responsive enhancer element B67 (TAXREB67) [Tsujiimoto et al., 1991], is a ubiquitous basic leucine-zipper transcription factor that is a member of the ATF/CREB protein family. This family includes cAMP-response element-binding protein (CREB), cAMP-response element modulator (CREM), ATF1, ATF2, ATF3, and ATF4 [Ziff, 1990; Brindle and Montminy, 1992; Meyer and Habener, 1993; Sassone-Corsi, 1994; Hai et al., 1999]. ATF4 functions as both transcriptional repressor and activator by forming homodimers and heterodimers with members of the AP-1 and C/EBP family of protein, or interacting with many other partners, such as human T-cell lymphotropic virus type 1, granulocyte colony-stimulating factor promoter element 1-binding protein, insulin-like growth factor-binding protein-1, NF-E2-related factor 2, c-maf, p300, Zhangfei, factor inhibiting ATF4-mediated transcription, special AT-rich sequence binding protein 2, transcription factor IIαγ, and runt-related transcription factor 2 [Hai and Curran, 1991; Chevray and Nathans, 1992; Nishizawa and Nagata, 1992; Vallejo et al., 1993; Vinson et al., 1993; Motohashi et al., 1997; Reddy et al., 1997; He et al., 2001; Lassot et al., 2005; Xiao et al., 2005; Yu et al., 2005, 2008b; Dobreva et al., 2006; Hogan et al., 2006]. ATF4 expression is up-regulated by several factors/stressors, including oxygen deprivation, endoplasmic reticulum stress, oxidative stress [Ameri et al., 2004; Blais et al., 2004; Roybal et al., 2005].

Abbreviations used: ATF4, activating transcription factor 4; Foxo1, forkhead box D1; BMSCs, bone marrow stromal cells; CDKs, cyclin-dependent kinases; CDKIs, cyclin-dependent kinase inhibitors.

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A role for ATF4 in bone development was established using Atf4-deficient mice [Yang et al., 2004]. Atf4−/− mice have a dramatically reduced bone formation rate and bone mineral density (severe osteoporosis) that persists throughout life. The expression of both osteocalcin (Ocn) and bone sialoprotein (Bsp), both markers for terminally differentiated osteoblasts, was markedly reduced in Atf4-deficient osteoblasts, suggesting a critical role for ATF4 in osteoblast differentiation. The Atf4−/− skeleton is much smaller relative to its wild-type (wt) control littermate, suggesting reduced numbers of bone cells. However, it is not known if ATF4 regulates osteoblast proliferation.

Cell cycle progression is a complex process that regulates cell proliferation. Cell cycle consists of several phases including G1 (from the end of the previous M phase till the beginning of DNA synthesis), S (DNA synthesis), G2 (significant protein synthesis occurs during this phase), and M phase (cell splits itself into two distinct cells) [Sherr and Roberts, 2004]. In addition, quiescent cells are in G0 phase. Cell cycle progression is a highly regulated process in that DNA replication occurs only once in each cycle. Transition from one phase to another is regulated by distinct cyclin-dependent kinases (CDKs) that are regulated by various cyclins, CDK inhibitors, and phosphorylations. Distinct cyclins and appropriate CDKs form complexes that function at different points of the cell cycle [Sherr and Roberts, 2004]. For example, formation of cyclin D1-CDK4/6 complex in early to mid G1 phase activates the kinases that phosphorylate and inactivate the tumor suppressor Rb, a critical step that is necessary for the transition from G1 to S phase [Kato et al., 1994; Sherr, 1994; Zhao et al., 2001]. The activity of cyclin E-CDK2 is periodic and maximal at the G1 to S phase transition [Roberts et al., 1994; Ohmoto et al., 1995]. Cyclin A1 activates CDK2 and is essential at the G1/S boundary and throughout S phase [Pagano et al., 1992]. In addition, cyclin-CDK complexes also have a noncatalytic role in G1 phase by sequestering proteins of the cip/kip family, including p27kip and p21cip, both CDK2 inhibitors (CDKIs) that negatively regulate cell cycle progression [Hofmann and Livingston, 1996].

Apoptosis is a form of cell death in which a programmed sequence of events leads to the destruction of cells. It occurs in both physiological and pathological conditions when the body needs to eliminate aged cells, unnecessary cells, and unhealthy cells. The caspase family, a set of cysteine proteases that cleave a variety of substrates, plays a crucial role in apoptosis. Caspases are divided into two groups: “initiator” caspses such as caspase-8 and -9 and “executioner” caspses including caspase-3, -6, and -7 [Shi, 2002]. During apoptosis, after being activated by various forms of stress such as inadequate growth factor support and different types of intracellular damage, initiator caspses proteolytically cleave executioner caspses that cause cell death events such as cytoplasm shrinkage, chromatin condensation, and DNA fragmentation. B-cell lymphoma 2 (Bcl-2) is the prototype for a family of mammalian genes encoding the proteins that control mitochondrial outer membrane permeabilization (MOMP) and can be either pro-apoptotic (Bax, BAD, Bak, and Bok) or anti-apoptotic (including Bcl-2, Bcl-XL, and Bcl-w) [Adams, 2003]. Atf4−/− mice are blind due to excessive apoptosis of cells in the lens epithelium [Kato et al., 1994; Sherr and Roberts, 2004], suggesting a critical role for ATF4 in protecting these cells from apoptosis under physiological condition.

The aim of this study was to determine the effects of ATF4 in the regulation of proliferation and apoptosis in primary BMSCs and osteoblasts.

MATERIALS AND METHODS

REAGENTS

Tissue culture media were purchased from Invitrogen (Carlsbad, CA) and fetal bovine serum from HyClone (Logan, UT). Other reagents were obtained from the following sources: Antibodies against cyclins D1, D3, p21, p27, CDK2, and horseradish peroxidase-conjugated mouse or goat IgG from Santa Cruz (Santa Cruz, CA), mouse monoclonal antibody against β-actin from Sigma (St. Louis, MO), BrdU immunostaining kit (Zymed Laboratories Inc., San Francisco, CA) from Invitrogen, ApopTag Peroxidase In Situ Apoptosis Detection Kit from Chemicon (Temecula, CA). Hoechst staining reagent was kindly provided by Dr. Rentian Feng of the University of Pittsburgh. All other chemicals were of analytical grade.

Atf4−DEFICIENT MICE

Breeding pairs of ATF4 heterozygous mice were described previously [Yu et al., 2008a] and used to generate Atf4 wild-type (wt) (Atf4+/+), heterozygous (Atf4+/−) and homozygous mutant (Atf4−−) mice for this study. All research protocols were approved by the Institutional Animal Care and Use Committee of the VA Pittsburgh Healthcare System, where this study was conducted.

HISTOLOGICAL EVALUATION

Six-week-old wt and Atf4−/− mice were euthanized and calvariae were fixed in 10% formalin at 4°C for 24 h, decalcified in 10% EDTA (pH 7.4) for 10 days, and embedded in paraffin. Calvariae were then bisected perpendicular to the sagittal suture through the central portion of the parietal bones, parallel to lamboidal and coronal sutures, and embedded in paraffin to obtain sections of a standard area according to the method described by Zhao et al. [2000].

CELL CULTURES AND ISOLATION OF PRIMARY BMSCs

Mouse MC3T3-E1 subclone 4 (MC-4) preosteoblasts were described previously [Xiao et al., 1997; Wang et al., 1999] and maintained in ascorbic acid-free alpha-modified Eagle’s medium (alpha-MEM), 10% fetal bovine serum (FBS), and 1% penicillin/streptomycin (proliferation medium). Isolation of mouse primary BMSCs was described previously [Xiao et al., 2002]. Briefly, 6-week-old male wt and Atf4−− mice were euthanized. Femurs and tibias were isolated and the epiphyses were cut. Marrow was flushed with alpha-MEM containing 20%FBS and 1% penicillin/streptomycin into a 100-mm dish and the cell suspension was aspirated up and down with a 20-gauge needle in order to break clumps of marrow. The cell suspension was then cultured in a T75 flask in the same medium. After 10 days, cells reached confluency and were ready for experiments.
ELECTROPORATION
Primary BMSCs were transfected using an Amaxa Nucleofector device (Cologne, Germany) as described by Nakashima et al. [2005]. Briefly, 2 × 10^6 BMSCs were suspended in 100 μl solution from Cell Line Nucleofector Kit V (Amaxa Biosystems, Cologne, Germany) and mixed with indicated plasmid DNAs. Cells were electroporated using the Program A-23 of the Amaxa Nucleofector device. Immediately after electroporation, cells were seeded for experiments.

MTS ASSAY
MTS assay was described previously [Singha et al., 2007]. Briefly, 1 × 10^4 cells/well were plated in a 96-well plate in 100 μl proliferation medium. Cells were incubated at 37°C for 24 h to allow attachment. The medium was changed every 48 h. At different time points, 20 μl of CellTitre96AQ solution reagent (Promega, Madison, WI) was added into each well and incubated for 2 h. Finally, the absorbance was recorded at 490 nm using a 96-well plate reader.

[H] THYMIDINE INCORPORATION
Cells were plated in 12-well plates at 5 × 10^4 cells/well in proliferation medium for 24 h and switched to 0.1% FBS alpha-MEM for another 24 h and [3H] thymidine was added to a final concentration of 5 μCi/ml and incubated at 37°C for 1 h in proliferation medium. Medium was removed by aspiration, and cells were washed twice with ice-cold serum-free alpha-MEM. Cell were extracted twice with 10% trichloroacetic acid (TCA) on ice for 5 min. TCA precipitates were solubilized by adding 10% SDS for 2 min at room temperature. Cells were harvested, and the amount of radioactivity was measured by liquid scintillation counting (Beckman Instruments, Inc., Fullerton, CA).

BROMODEOXYURIDINE (BrdU) INCORPORATION ASSAY
BrdU labeling reagent was purchased from Invitrogen. BrdU staining was performed using cells cultured in 8-well culture chambers (Nalgene Nunc, NY) or 10-μm sections of calvariae from wt and Atf4^-/- mice. Cells were cultured in 8-well chamber at a density of 10^5 cells/well in 400 μl proliferation medium. After 4 days, cells from four identically treated wells per group were labeled with BrdU (1:100 dilution) overnight in the same medium. For calvarial sections, wt and Atf4^-/- mice (6/group) were injected intraperitoneally with 100 μg bromodeoxyuridine (BrdU)/12 μg fluorodeoxyuridine (FdU) per gram of body weight 4 h before sacrifice. After sacrifice, 10-μm sections of calvariae were obtained as described previously [Demirap et al., 2002]. To identify actively proliferating cells, nuclei that have incorporated BrdU were detected using a Zymed BrdU immunostaining kit according to the manufacturer’s instructions. BrdU-positive cells on the periosteal surface of calvariae were counted and normalized to the total periosteal cell numbers in the same area [Zhao et al., 2004].

FLOW CYTOMETRIC ANALYSIS
Primary BMSCs from wt and Atf4^-/- mice were cultured in proliferation medium for 4 days and harvested. 5 × 10^5 cells were suspended in 1 ml proliferation medium and Vybrant violet dye (Invitrogen) was added to a final concentration of 5 μM and incubated at 37°C for 30 min. The distribution of BMSCs throughout the cell cycle was assessed by flow cytometry using 405 nm excitation and 440 nm emission. The fraction of the population in each phase was determined as a function of DNA content using an FACS with software.

ApopTag PEROXIDASE IN SITU APOPTOSIS DETECTION
This method was based on the classical TUNEL assay to examine apoptosis by detecting DNA fragmentation. Primary BMSCs from wt and Atf4^-/- mice were cultured in proliferation medium for 4 days and stained using the ApopTag Peroxidase In Situ Apoptosis Detection Kit according to the manufacturer’s instruction. Cells were counterstained with Hematoxylin. Sections of wt and Atf4^-/- calvariae (6/group) were prepared and stained using the same kit. Negative controls were made by omitting the terminal deoxynucleotidyl transferase (TdT). All positive (brown) and negative (blue) nuclei were counted. Apoptotic cells on the periosteal surface were counted and normalized to the total cells from the same area.

RNA ISOLATION AND REVERSE TRANSCRIPTION (RT)
Tibias (6/group) were isolated and soft tissues were removed. Tibias were frozen in liquid nitrogen and ground into powder using a mortar and pestle and total RNAs from each group were isolated using the Trizol reagent (Invitrogen) following the manufacturer’s protocol. Reverse transcription (RT) was performed using 1 μg of denatured RNA and 100 pmol of random hexamers (Applied Biosystem, Foster, CA) in a total volume of 25 μl containing 12.5 U MultiScribe reverse transcriptase (Applied Biosystem).

QUANTITATIVE REAL-TIME PCR
Quantitative real-time PCR was performed on an iCycler (BIO-RAD, Minneapolis, MN) using a SYBR® Green PCR Core Kit (Applied Biosystem) and cDNA equivalent to 10 ng RNA in a 50 microliter reaction according to the manufacturer’s instructions. The DNA sequences of mouse primers used for real-time PCR were: cyclin D1 (GenBank Accession number-NM-007631), 5’ GAG GAG GGG GAA GTG GAG GA 3’ (forward, +1,049-bp), 5’ CCT CTG TGG GGG GCC CAC TA 3’ (reverse, +1,170-bp); FazoI(GenBank Accession number-NM-019739), 5’ AGA GGC TCA CCC TGT CGC AGA 3’ (forward, +955-bp), 5’ GTG AAG GGA CAG ATT GTG GCC CAC TA 3’ (reverse, +1,080-bp); Gapdh (GenBank Accession number-NM-001001303), 5’-CAG TGC CAG CCT GTG GCC GAA 3’ (forward, +32-bp), 5’-TCG CAA ATG GCC GCA CTC GTG AC 3’ (reverse, +1,127-bp). For all primers the amplification was performed as follows: initial denaturation at 95°C for 10 min followed by 40 cycles of 95°C for 15 s and 60°C for 60 s. Melting curve analysis was used to confirm the specificity of the PCR products. Six samples were run for each primer set. The levels of mRNA were calculated by the ΔCT (the difference between the threshold cycles) method [Wang et al., 2004]. The mRNA level of each gene was normalized to Gapdh mRNA.

WESTERN BLOT ANALYSIS
Cells were washed with cold 1× PBS and lysed in 1× Passive Buffer (Promega) at RT for 20 min. Lysates were clarified by centrifugation (20 min, 13,000g, 4°C). For bone tissues, wt and Atf4^-/- tibias (6/group) were frozen in liquid nitrogen and ground into powder.
with a mortar and pestle. Samples were solubilized with 1× Passive Buffer for 20 min on ice and sonicated on ice (3 times, 5 s each) and lysates were then clarified by centrifugation (20 min, 13,000 g, 4°C). Protein concentrations were determined by the method developed by Bio-Rad Laboratories, Inc. (Hercules, CA). Twenty-five microgram of total protein were fractionated on a 10% SDS–PAGE gel and transferred onto nitrocellulose membranes (Whatman). The membrane was blocked in 5% nonfat milk in Tris-buffered saline/Tween-20 (TBST) buffer, probed with antibodies (1:1,000) against cyclins D1, D3, CDK2, p21, and p27 followed by incubation with anti-rabbit or anti-mouse antibodies conjugated with horseradish peroxidase (1:5,000) and visualized using an enhanced chemiluminescence kit (Pierce, Rockford, IL). Finally, blots were striped 2 times in buffer containing 65 mM Tris–Cl (pH 6.8), 2% SDS, and 0.7% (v/v) β-mercaptoethanol at 65°C for 15 min and re-probed with β-actin antibody (1:2,000) for normalization.

SMALL INTERFERING RNA (siRNA)

MC-4 cells, which contain high levels of Atf4 mRNA, were seeded at a density of 25,000 cells/cm². After 24 h, cells were transfected with mouse Atf4 siRNA (sense: 5’-GAG CAU UCC UUU AGU UUA GUU-3’; antisense: 5’-CUA AAC UAA AGG AAU GCU CUU-3’) or negative control siRNA (low GC, Cat #: 12935-200, Invitrogen) [Adams, 2007; Yu et al., 2008a] using LipofectAMINE 2000 (Invitrogen). After 48 h, cells from three identically treated dishes were pooled and harvested for total RNA, followed by quantitative real-time RT-PCR analysis for Atf4, cyclin D1 and Faxo1 mRNAs. A second set of mouse Atf4 siRNAs was purchased from Ambion (Cat #: AM16704, ID#: 160775 and 160776) and used to confirm the results using the first set of Atf4 siRNA.

STATISTICAL ANALYSIS

Data was analyzed with GraphPad Prism software. Students’ t-test was used to test for differences between two groups. A one-way ANOVA analysis was used followed by the Dunnett’s test for Figures 1C, 5A–D, 6A–C, and 7A,B. Differences with a P < 0.05 were considered as statistically significant. All experiments were repeated a minimum of 3 times with triplicate samples.

RESULTS

ATF4 DEFICIENCY REDUCES CELL PROLIFERATION IN PRIMARY BMSCs AND CALVARIAL PERIOSTEAL OSTEOBLASTS

To determine if ATF4 plays a role in the regulation of osteoblast proliferation, we performed several experiments. We first determined the effects of ATF4 deficiency on the numbers of total nucleated bone marrow cells, bone marrow from 6-week-old male wt and Atf4−/− mice (6/group) were isolated. After lysing the red blood cells (RBCs) using the RBC lysis buffer (Sigma), the remaining nucleated bone marrow cells were directly counted using a hemacytometer. B: Colony numbers in bone marrow. After lysing the RBCs, the remaining total nucleated bone marrow cells from each mouse (two femurs and two tibias, 6 mice/group) were diluted (1:2,000) and cultured in proliferation medium. 24 h later, the non-adhering cells were removed by washing the cells 3 times with 1× PBS. Cells were then cultured in proliferation medium for 10 days. The numbers of colonies were then counted using a hemacytometer. C: MTS assay. Wt and Atf4−/− BMSCs were seeded at a density of 10⁴ cells/well in 96-well plate and cultured in proliferation medium for 0, 2, 4, 6, and 8 days followed by incubation with 20 µl of CellTiter96AQU solution reagent for 2 h. The absorbance was recorded at 490 nm using a 96-well plate reader. D: [3H] thymidine incorporation. wt and Atf4−/− BMSCs were plated in 12-well plates at 5× 10⁴ cells/well in proliferation medium for 24 h and switched to 0.1% FBS alpha-MEM for 24 h and [3H] thymidine was then added to the culture (proliferation medium) to a final concentration of 5 µCi/ml and incubated at 37°C for 1 h. *P < 0.05 (wt vs. Atf4−/−). Data represent mean ± SD. Experiments were repeated 3 times and qualitatively identical results were obtained.

Fig. 1. ATF4 is required for primary BMSC proliferation. A: Total nucleated bone marrow cells. Long bones (two femurs and two tibias) from 6-week-old male wt and Atf4−/− mice (6 mice per group) were isolated. Bone marrow was flushed out and red blood cells (RBC) were lysed using the RBC lysis buffer. Total nucleated bone marrow cells were counted using a hemacytometer. B: Colony numbers in bone marrow. After lysing the RBCs, the remaining total nucleated bone marrow cells from each mouse (two femurs and two tibias, 6 mice/group) were diluted (1:2,000) and cultured in proliferation medium. 24 h later, the non-adhering cells were removed by washing the cells 3 times with 1× PBS. Cells were then cultured in proliferation medium for 10 days. The numbers of colonies were then counted using a hemacytometer. C: MTS assay. Wt and Atf4−/− BMSCs were seeded at a density of 10⁴ cells/well in 96-well plate and cultured in proliferation medium for 0, 2, 4, 6, and 8 days followed by incubation with 20 µl of CellTiter96AQU solution reagent for 2 h. The absorbance was recorded at 490 nm using a 96-well plate reader. D: [3H] thymidine incorporation. wt and Atf4−/− BMSCs were plated in 12-well plates at 5× 10⁴ cells/well in proliferation medium for 24 h and switched to 0.1% FBS alpha-MEM for 24 h and [3H] thymidine was then added to the culture (proliferation medium) to a final concentration of 5 µCi/ml and incubated at 37°C for 1 h. *P < 0.05 (wt vs. Atf4−/−). Data represent mean ± SD. Experiments were repeated 3 times and qualitatively identical results were obtained.
positive in both wt and Atf4−/− mice) that permeate the bone matrix were stained BrdU− (Fig. 2B,C). It should be noted that very few osteocytes (mature and ATF4 is required for osteoblast proliferation in vivo, 6-week-old wt proliferation measured is primarily that of BMSCs. To determine if proliferative under the culture conditions used, consequently, the however, macrophages and their precursors would be less significant reduction in the number of total bone marrow cells between wt and Atf4−/− mice, surprisingly, we found no significant difference in the numbers of total nucleated bone marrow cells from each mouse (two femurs and two tibias, 6 mice/group) were diluted (1:2,000) and cultured in proliferation medium. Twenty-four hours later, the non-adhering cells were removed by washing the cells 3 times with 1 × PBS. Cells were then cultured in proliferation medium for 10 days. The numbers of colonies from each dish were then counted using a microscope. As shown in Figure 1B, the number of BMSC colonies per mouse (from two femurs and two tibias) was significantly reduced in Atf4−/− group compared to values in the wt group (P < 0.05, wt vs. Atf4−/−). This decrease could be explained by a cell-autonomous defect in proliferation and/or survival or could be secondary to an impaired bone microenvironment due to ATF4 deficiency. To differentiate these possibilities, we conducted MTS and [3H] thymidine incorporation assays. For the MTS assay, 104 cells/well of wt and Atf4−/− BMSCs were seeded in 96-well plate and were cultured in proliferation medium for 0, 2, 4, 6, and 8 days. Cell numbers were then measured as described previously [Singha et al., 2007]. As shown in Figure 1C, ATF4-deficient cells grew at significantly reduced rates compared to wt cells. For the [3H] thymidine incorporation assays, wt and Atf4−/− BMSCs were cultured in proliferation medium for 4 days and labeled with [3H] thymidine for 1 h before harvesting. As shown in Figure 1D, [3H] thymidine incorporation into the DNA of Atf4−/− BMSCs was decreased by fourfold compared to values in wt cells (P < 0.05, wt vs. Atf4−/−). It should be noted that the adherent murine BMSC population is contaminated with macrophages, however, macrophages and their precursors would be less proliferative under the culture conditions used, consequently, the proliferation measured is primarily that of BMSCs. To determine if ATF4 is required for osteoblast proliferation in vivo, 6-week-old wt and Atf4−/− mice were injected with bromodeoxyuridine (BrdU)/fluorodeoxyuridine (FdlU) 4 h before sacrifice, 10-μm sections of calvariae were obtained, and proliferating cells from the perisetal surface were counted and normalized to total cells from the same area. As shown in Figure 2A, in wt calvariae, periosteal osteoblasts proliferated very actively with 60% of the total cells being BrdU-positive. In contrast, the percent BrdU-positive osteoblasts were significantly reduced in Atf4−/− calvariae (P < 0.05, wt vs. Atf4−/−) (Fig. 2B,C). It should be noted that very few osteocytes (mature osteoblasts) that permeate the bone matrix were stained BrdU-positive in both wt and Atf4−/− calvariae. Thus, ATF4 is required for the proliferation of BMSCs or osteoblasts both in vitro and in vivo.

**ATF4 IS REQUIRED FOR CELL CYCLE PROGRESSION AND CYCLIN D1 EXPRESSION**

To determine whether ATF4 is required for cell cycle progression, we performed flow cytometric analysis to compare the cell distribution across different phases of the cell cycle between wt and Atf4−/− BMSCs. As shown in Figure 3, ATF4-deficient cells showed a significant decrease in the cell distribution into both S and G2/M phases (32% and 45% change, respectively) when compared with values of wt control cells (P < 0.05, wt vs. Atf4−/−). In contrast, the fraction of the cells in G1 was not reduced by ATF4 deficiency (P > 0.05, wt vs. Atf4−/−). Cell-cycle progression from one phase to another is controlled by cyclin-dependent kinases (CDKs) whose activity is mainly regulated by distinct cyclins and cyclin-dependent kinase inhibitors (CDKIs). We examined the effects of ATF4 deficiency on the expression levels of critical cyclin D1, a major regulator of cell cycle progression and cell proliferation, and CDKIs. Wt and Atf4−/− BMSCs were cultured in proliferation medium for
4 days followed by Western blot analysis for cyclin D1, CDK2, p21, and p27. As shown in Figure 4A, the protein level of cyclin D1 was markedly decreased in Atf4−/− cells relative to wt control. In contrast, the level of p21 protein was slightly increased in Atf4−/− cells. Levels of both p27 and CDK2 proteins were low in BMSCs and their relative abundance did not display any significant difference between wt and Atf4−/− cells. Quantitative real-time RT/PCR analysis shows that the level of cyclin D1 mRNA was reduced by 52% in Atf4−/− cells compared to values in wt controls (P < 0.05, wt vs. Atf4−/−) (Fig. 4B). Note: as expected, minimal Atf4 mRNA was detected by real-time RT/PCR in Atf4−/− cells (Fig. 4B, top).

To determine whether ATF4 is required for proliferation of the pre-osteoblast cell line, MC-4, which expresses high levels of Atf4 mRNA and protein, the cells were transiently transfected with the indicated concentrations of Atf4 siRNA or negative control siRNA (Invitrogen). As shown in Figure 5A, levels of Atf4 mRNA were efficiently reduced by Atf4 siRNA in a dose-dependent manner. In contrast, the negative control (40 nM) did not reduce Atf4 mRNA. Importantly, the level of cyclin D1 mRNA was significantly reduced by Atf4 siRNA in a dose-dependent manner (Fig. 5B). Conversely, the level of Foxo1 mRNA, a factor of the forkhead transcription factor family, was not reduced by Atf4 siRNA (Fig. 5C). Furthermore, knocking-down Atf4 mRNA markedly reduced MC-4 proliferation as measured by both MTS (Fig. 5D) and BrdU incorporation assays (Fig. 5E–G) (P < 0.05, ctrl siRNA vs. Atf4 siRNA). Similar results were obtained when a different set of Atf4 siRNAs was used in MC-4 cells (data not shown).

OVEREXPRESSING ATF4 RESCUES THE DEFECT IN CELL PROLIFERATION IN Atf4−/− BMSCs

We next determined whether overexpression of ATF4 could increase cell proliferation in BMSCs. To this end, we used a Cell Line
Nucleofector Kit V and a Nucleofector Device from Amaxa Biosystem (Cologne, Germany) because primary mouse BMSCs were transfected with higher than 40% efficiency by using this method (data not shown). To determine the effects of ATF4 overexpression on cyclins expression, Atf4−/− BMSCs were electroporated with increasing amounts of FLAG-ATF4-YFP expression vector (0, 0.5, 1, and 2 μg) [Xiao et al., 2005]. Thirty hours later, cells were harvested for the preparation of total RNA or protein. In addition, as shown in Figure 6A, Atf4 mRNA was efficiently expressed in primary BMSCs in a dose-dependent manner. As shown in Figure 6B, ATF4 dose-dependently stimulated cyclin D1 expression both at the mRNA and protein level. In contrast, overexpressing ATF4 failed to elevate the level of cyclin D3 mRNA and protein (Fig. 6C,D). As shown in Figure 7, ATF4 overexpression significantly increased the rate of proliferation in Atf4−/− BMSC as measured by direct cell count, MTS assay, and BrdU staining (P < 0.05, ctrl vs. ATF4). Overexpressing ATF4 similarly increased cyclin D1 expression and cell proliferation in wt BMSCs (data not shown).

Taken together, these results clearly establish that ATF4 increases cell proliferation probably via promoting expression of cyclin D1 and cell cycle progression.

**ATF4 DEFICIENCY INCREASES CELL APOPTOSIS**

ATF4 is known to prevent lens epithelium from apoptosis [Tanaka et al., 1998; Hettmann et al., 2000]. As an initial step to determine if ATF4 regulates apoptosis in BMSCs, wt and Atf4−/− cells were cultured in proliferation media for 4 days and stained by the Hoechst method [Yamanaka et al., 2003]. As shown in Figure 8A–C, the numbers of apoptotic cells including those with shrinking cytoplasm and chromatin condensation (early apoptosis) and DNA fragmentation (late apoptosis) were increased greater than fivefold in Atf4−/− BMSCs compared to wt cells (P < 0.05, wt vs. Atf4−/−). To confirm this finding, wt and Atf4−/− BMSCs were stained using the ApopTag Peroxidase In Situ Apoptosis Detection Kit, a modified TUNEL staining that measures DNA fragmentation in situ. As shown in Figure 8D–F, the percent apoptotic cells in Atf4−/− BMSCs were increased by 1.6-fold when compared to wt cells (P < 0.05, wt vs. Atf4−/−). To determine if ATF4 deficiency increases osteoblast apoptosis in vivo, 10-μm calvarial sections from wt and Atf4−/− mice were obtained and stained using the same kit. Apoptotic cells that stained brown on the periosteal surface of calvariae were counted and normalized to total cells of the same periosteal surface. As shown in Figure 8G–I, a significant increase in apoptosis was found in Atf4−/− mice compared to wt controls (P < 0.05, wt vs.
Thus, ATF4 protects osteoblasts from apoptosis under physiological condition.

DISCUSSION

In this study, we used two complementary approaches to establish the requirement for ATF4 in the regulation of proliferation in primary BMSCs and a preosteoblast cell line (MC-4 cells): (i) loss-of-function studies using Atf4−/− or Atf4 siRNA-treated cells demonstrate that cell proliferation is significantly reduced in the absence of ATF4; and (ii) gain-of-function experiments via overexpression of ATF4 show that ATF4 enhances osteoblast proliferation in both wt and Atf4−/− BMSCs. In addition, this study reveals that ATF4 protects osteoblasts against apoptosis. Thus, the reduced bone mass

Fig. 6. Overexpression of ATF4 increases cyclin D1 expression. A–C: Quantitative real-time-RT/PCR. Atf4−/− BMSCs (2 × 10^5 cells/group) were electroporated with increasing amounts of FLAG-ATF4-YFP expression vector (0, 0.5, 1, and 2 μg). The amount of plasmid DNAs was balanced as necessary with beta-galactosidase expression plasmid such that the total DNA was constant in each group. Thirty hours later, cells were harvested for the preparation of total RNA and quantitative real-time RT/PCR analysis. D: Western blot analysis. Cells were treated as in (A) and harvested for whole cell extracts preparation and Western blot analysis. *P < 0.05 (beta-gal vs. ATF4). Data represent mean ± SD. Experiments were repeated at least 3 times and qualitatively identical results were obtained.

Fig. 7. Overexpression of ATF4 increases BMSC proliferation. A: Direct cell count. Atf4−/− BMSCs were electroporated as in Figure 6A. 5 × 10^3 cells/well were seeded in 96-well plates in proliferation medium for 4 days followed by direct cell count using a hemacytometer. B: MTS assay. Cells were electroporated as in Figure 6A. After electroporation, cells were used for the MTS assay as in Figure 1C. C–E: BrdU staining. Cells were electroporated as in Figure 6A. After electroporation, cells were used for the BrdU staining. *P < 0.05 (beta-gal vs. ATF4). Data represent mean ± SD. Experiments were repeated at least 3 times and qualitatively identical results were obtained. Arrows indicate BrdU-positive (proliferating) cells and Δ indicates BrdU-negative (nonproliferating) cells. [Color figure can be viewed in the online issue, which is available at www.interscience.wiley.com.]
compared to that in Atf4 proliferation by MC-4 cells (Figs. 1C and 5D), the magnitude of inhibition of cell proliferation by the Atf4 inhibitory effect of the siRNA may not completely knock down the Atf4 mRNA in MC-4 cells. This discrepancy is likely due to our observation that the control siRNA treatment of MC-4 cells non-specifically reduces cell proliferation, which could mask the inhibitory effect of the Atf4 siRNA on cell proliferation. In addition, the Atf4 siRNA may not completely knock down the Atf4 mRNA in MC-4 cells.

Previous studies established that ATF4 is critical for osteoblast differentiation as demonstrated by dramatically reduced expression of osteocalcin and bone sialoprotein mRNA in ATF4-deficient mice [Yang et al., 2004]. The present study demonstrates an essential role for ATF4 in the regulation of BMSC and osteoblast proliferation. Atf4−/− BMSCs and osteoblasts proliferate at a dramatically reduced rate compared to wt cells both in vitro and in vivo. Overexpressing ATF4 in Atf4−/− BMSCs rescues the proliferative defect. Expression of cyclin D1 is highly dependent upon the presence of ATF4 as demonstrated by the dramatic reduction in its mRNA and protein in Atf4−/− BMSCs. Reduced expression of cyclin D1 and possibly other cyclins in Atf4−/− cells leads to a lack of progression from G1 into the S and G2/M phases of the cell cycle. Since the percentage of the population in G1 is not affected by ATF4 deficiency, there must be concomitantly an increase in the percentage of G0 cells (quiescent cells) and/or cells undergoing cell death, thereby reducing overall cell proliferation.

Cyclin D1 is a key sensor and integrator of extracellular signals of cells and plays a critical role in cell cycle progression and proliferation [Stacey, 2003]. The expression level of cyclin D1 has been shown to be rate-limiting in cell proliferation induced by a variety of stimuli [Zhao et al., 2001]. Our results show that ATF4 is a key regulator of cyclin D1 expression in BMSCs or osteoblasts. Levels of cyclin D1 mRNA and protein are significantly reduced in the absence of ATF4. Forced expression of ATF4 efficiently augmented the level of cyclin D1 mRNA as well as cell proliferation in both wt and Atf4−/− cells. Mechanisms whereby ATF4 increases cyclin D1 mRNA remain to be determined. An ATF/CRE site located in the proximal region of cyclin D1 promoter was reported to bind to CREB and AP1 proteins and mediate active transcription of the gene [Sabbah et al., 1999; Nagata et al., 2001; Datta et al., 2007]. It is likely that ATF4 directly binds to this ATF/CRE site and activates transcription. It is also probable that ATF4 activates the cyclin D1 promoter through interactions with CREB and AP1 proteins. Finally, ATF4 can stabilize cyclin D1 mRNA through post-transcriptional mechanisms. Future study will differentiate these possibilities.

The periosteal surface of calvariae is mainly occupied by four cell types: osteoblasts or osteoprogenitors, bone lining cells, and osteoclasts. In rapidly growing animals, osteoblasts or osteoprogenitors are the major cells that cover the surface. In contrast, bone lining cells occupy the majority of the surface in adult or aged bones that have reached peak bone mass. Calvaria, which does not involve endochondral bone formation, has a relatively simple structure relative to other bones such as long bones and vertebrae. In addition, it is easy to histologically localize the osteoprogenitors/osteoblasts on the periosteal surface of calvaria. Therefore, it provides a unique model system for studying the functions of osteoblasts in vivo.
Using this model system, we found that osteoblasts from wt 6-week-old mice proliferate very actively on the periosteal surface (Fig. 2A). Conversely, proliferation of Atf4−/− osteoblasts is significantly reduced (Fig. 2B,C). Thus, an in vivo role for ATF4 in osteoblast proliferation is established. In support of our findings, Masuoka and Townes (2002) showed that Atf4−/− mice have severe fetal anemia due to impaired fetal-liver definitive hematopoiesis associated with a proliferative defect in fetal-liver cells. Furthermore, primary murine embryonic fibroblasts (MEFs) from Atf4−/− mice also display a defect in proliferation [Masuoka and Townes, 2002]. Lastly, transgenic overexpression of ATF4 in the developing lens results in hyperproliferation of lens fiber cells [Hettmann et al., 2000]. These results suggest that ATF4 is critical for proliferation of rapidly growing cells (i.e., BMSCs, osteoblasts, fetal-liver cells, MEFs, and lens fiber cells). Consistent with this notion, ATF4 is usually expressed at high level in rapidly growing tissues or cells [Tanaka et al., 1998; Hettmann et al., 2000; Masuoka and Townes, 2002; Yang et al., 2004].

The numbers of osteoblasts are eventually determined by the relative rate of cell proliferation and death by apoptosis. Experiments from this study establish that ATF4 is anti-apoptotic in BMSCs and osteoblasts. Nevertheless, ATF4 is not a global anti-apoptotic factor since no increase in apoptosis has been observed in Atf4−/− fetal liver cells although the ability of these cells to proliferate is impaired. ATF4 may elicit its anti-apoptotic function in combination with other factors in specific tissue and cells types. Thus, ATF4 deficiency increases apoptosis in lens fiber cells in a p53-dependent manner. The embryonic lens in double homozygous p53/Atf4−/− mice does not undergo apoptosis [Tanaka et al., 1998; Hettmann et al., 2000]. Interestingly, ATF4 expression is usually induced by oxygen deprivation, endoplasmic reticulum stress, and the oxidative stressor arsenite, all of which are known to induce cell apoptosis [Ameri et al., 2004; Blais et al., 2004; Roybal et al., 2005]. Therefore, it is reasonable to speculate that expression of ATF4 induced by apoptosis-inducing factors in fact provides a protection mechanism for cells to antagonize apoptosis. The molecular mechanism whereby ATF4 regulates apoptosis in osteoblasts remains to be determined in future study.

In summary, this study for the first time establishes that ATF4 is essential for cell proliferation and anti-apoptosis in BMSCs and osteoblasts.

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Identification and Functional Characterization of ERK/MAPK Phosphorylation Sites in the Runx2 Transcription Factor*†‡

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The Runx2 transcription factor is required for commitment of mesenchymal cells to bone lineages and is a major regulator of osteoblast-specific gene expression. Runx2 is subject to a number of post-transcriptional controls including selective proteolysis and phosphorylation. We previously reported that Runx2 is phosphorylated and activated by the ERK/MAPK pathway (Xiao, G., Jiang, D., Thomas, P., Benson, M. D., Guan, K., Karsenty, G., and Franceschi, R. T. (2000) J. Biol. Chem. 275, 4453–4459). In this study, we used a combination of in vitro and in vivo phosphorylation analysis, mass spectroscopy, and functional assays to identify two sites at Ser301 and Ser319 within the proline-serine/threonine domain of Runx2 that are required for this regulation. These sites are phosphorylated by activated ERK1 in vitro and in cell culture. In addition to confirming ERK-dependent phosphorylation at Ser319, mass spectroscopy identified two other ERK-phosphorylated sites at Ser43 and Ser510. Furthermore, introduction of S301A,S319A mutations rendered Runx2 resistant to MAPK-dependent activation and reduced its ability to stimulate osteoblast-specific gene expression and differentiation after transfection into Runx2-null calvarial cells and mesenchymal cells. In contrast, S301E,S319E Runx2 mutants had enhanced transcriptional activity that was minimally dependent on MAPK signaling, consistent with the addition of a negative charge micromgging serine phosphorylation. These results emphasize the important role played by Runx2 phosphorylation in the control of osteoblast gene expression and provide a mechanism to explain how physiological signals acting on bone through the ERK/MAPK pathway can stimulate osteoblast-specific gene expression.

The bone cell lineage is controlled by a hierarchy of transcription factors that are expressed in a defined temporal sequence. Runx2, an essential factor for both hypertrophic cartilage and bone formation, is expressed very early in skeletal development, first appearing coincident with the formation of mesenchymal condensations (1). Subsequent development of the osteoblast lineage requires at least two additional factors; Osterix, which is essential for subsequent progression of the osteoblast lineage, and ATF4, which regulates osteoblast activity, particularly in postnatal animals (2, 3). Runx2 expression continues during the later stages of bone development and persists in regions of active bone remodeling throughout life. Skeletal development in Runx2-deficient mice fails to progress beyond the cartilage anlage stage, whereas dominant-negative suppression of Runx2 even in postnatal animals inhibits osteoblast activity and bone formation (4). Thus, Runx2 is required for both the initial formation of osteoblasts and hypertrophic chondrocytes during development and for sustained osteoblast differentiation during bone remodeling.

Consistent with its multiple roles in bone formation, Runx2 is highly regulated. In addition to transcriptional control by factors such as bone morphogenetic proteins (5), Runx2 activity is controlled both by its interaction with a number of accessory nuclear factors and by post-translational modifications, including phosphorylation. We have been particularly interested in this latter regulation and proposed that Runx2 is phosphorylated and activated by a ERK/MAPK-dependent pathway initiated by the interaction of osteoprogenitors with a type I collagen-containing extracellular matrix (ECM) via α2β1 integrins (6, 7). This collagen-integrin interaction is necessary for subsequent osteoblast-specific gene expression and differentiation (7–9). Consistent with this model, steady-state Runx2 phosphorylation and DNA binding activity increase with osteoblast differentiation, whereas pharmacological inhibition of the ERK/MAPK pathway rapidly inhibits ECM and BMP-induced gene expression (10–12). In related studies, FGF2 treatment of osteoblasts, which is known to stimulate both ERK/MAPK and protein kinase C pathways, increases Runx2 phosphorylation and Ocn expression in a MAPK-dependent manner (13). Furthermore, manipulation of the MAPK pathway by overexpression of constitutively active or dominant-negative mutants of MEK1, respectively, increases or decreases osteocalcin gene expression and Runx2 phosphorylation (6). ERK/MAPK signaling is also important for in vivo bone development. Transgenic overexpression of constitutively active or dominant-negative MEK1 in mouse osteoblasts, respectively, stimulates or inhibits Runx2 phosphorylation and skeletal maturation. Furthermore,

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† The on-line version of this article (available at http://www.jbc.org) contains supplemental Fig. S1.
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3 The abbreviations used are: ERK, extracellular signal-regulated kinase; MAPK, mitogen-activated protein kinase; ECM, extracellular matrix; MEK, mitogen-activated protein kinase/extracellular signal-regulated kinase; OSE2, osteoblast-specific element 2; Mek(sp), constitutively active MEK1 mutant; FGF, fibroblast growth factor; HA, hemagglutinin; aa, amino acid; FBS, fetal bovine serum; MS, mass spectrometry.
the cleidocranial dysplasia phenotype of Runx2 heterozygous null mice can be partially rescued by crossing these animals with mice expressing constitutively active MEK1, consistent with the \textit{in vivo} actions of the ERK/MAPK pathway being at least in part mediated by Runx2 (14).

In addition to the work from our laboratory cited above (6–14), a number of studies from other groups support the concept that ECM-integrin binding, MAPK activation, and Runx2 phosphorylation are important for osteoblast differentiation. The requirement for α1β1 and α2β1 collagen-binding integrins in osteoblast differentiation and BMP responsiveness was demonstrated by both \textit{in vitro} and \textit{in vivo} analysis (15–18). Also, ERK/MAPK signaling was shown to be necessary for differentiation of human osteoblasts and marrow stromal cells (19, 20). A number of groups also confirmed that Runx2 can be phosphorylated and activated by MAPK inducers. During the osteoblastic differentiation of human marrow stromal cells, Runx2 levels remain relatively unchanged, but DNA binding increases as does Runx2 phosphorylation (21). Also, mechanobiological osteoblasts, mediated in part through α2β1 integrins, induces MAPK activity (22, 23). Similarly, loading of periodontal ligament cells (osteoprogenitor-like cells) increases Runx2 phosphorylation and binding to OSE2 DNA via an ERK/MAPK-dependent process (24). In osteosarcoma prostate cancer cells, differentiation is accompanied by ERK1/2 activation, increased Runx2-DNA binding, and Ocn expression, responses that were all blocked by MAPK inhibition (25). Last, insulin-like growth factor-1, which activates phosphatidylinositol 3-kinase and, subsequently, ERK/MAPK pathways, stimulates Runx2-OSE2 binding and phosphorylation in vascular endothelial cells (26) as well as differentiation of marrow stromal cells (27, 28). Thus, ERK/MAPK-dependent phosphorylation of Runx2 likely plays an important role in the response of osteoblasts to a variety of signals initiated by cell-ECM binding, hormone/growth factor signaling, and mechanical loading.

To further understand how the ERK/MAPK pathway regulates Runx2 transcriptional activity, in the present study we identify amino acid residues in Runx2 that are phosphorylated in a ERK/MAPK-dependent manner and show that these sites are necessary for osteoblast-specific gene expression and differentiation.

**EXPERIMENTAL PROCEDURES**

**Reagents**—The reagents used in this study were obtained from the following sources: tissue culture medium and fetal bovine serum from Invitrogen; U0126 from Calbiochem; mouse anti-Runx2 antibody from MBL; phosphoserine antibody from ABCam; and M2 and M2 horseradish peroxidase-conjugated antibody from Sigma.

**DNA Constructs and Viral Expression Vectors**—The 0.6-kb mouse osteocalcin gene 2-Luc reporter plasmid and a constitutively active MEK1 expression vector were previously described (29–31). A series of Runx2 expression plasmids encoding HA- or FLAG-tagged full-length type II Runx2 (N-terminal sequence: MASN) or several N- and C-terminal deletions were generated by PCR and subcloning into the pCMV5 expression vector. Serine mutants of full-length and aa 1–330 Runx2 were generated using the QuikChange site-directed mutagenesis kit (Stratagene). A cDNA encoding Runx2 with a biotinylation tag was generated by adding the sequence, MASSLQRILDSQK-MEWRSNAGGS, to the N terminus of mouse Runx2. This sequence is specifically recognized by bacterial BirA biotin ligase (32). Plasmids containing cDNAs encoding the biotinylation tag and BirA were a generous gift from Dr. John Strouboulis (Alexander Fleming Biomedical Sciences Research Center, Athens, Greece). Adenoviruses encoding wild-type and mutant Runx2, and biotinylation tagged Runx2, BirA, and Mek(sp) (constitutively active MEK1) were constructed by first subcloning the respective cDNA into pAdlox and then generating viruses using Cre-Lox recombination as previously described (33).

**Cell Culture**—C3H10T1/2, COS7, and HEK293 cells were obtained from the American Type Culture Collection and maintained in Dulbecco’s modified Eagle’s medium containing 10% FBS and 1% antibiotics. An mTERT-immortalized calvarial cell line from Runx2−/− mice (34) was a generous gift from Drs. Jane Lian and Gary Stein (University of Massachusetts Medical Center, Worcester, MA) and maintained in minimal essential medium-α, 10% FBS. MC3T3-E1 clone 4 (MC-4) cells, previously developed in this laboratory (35), were also maintained in minimal essential medium-α, 10% FBS. To induce differentiation, C3H10T1/2, mTERT cells, and MC-4 cells were grown in minimal essential medium-α, 10% FBS containing 50 μg/ml ascorbic acid as previously described (12, 34, 36).

**Transfections**—COS7 cells were plated at a density of 5 × 10⁴ cells/cm² on 35-mm dishes and transfected using Lipofectamine (Invitrogen). Each transfection contained 0.5 μg of the indicated plasmid and 0.05 μg of pRL-SV40 containing a cDNA for \textit{Renilla reniformis} luciferase to control transfection efficiency. Cells were harvested and assayed using a dual luciferase assay kit (Promega) with a Monolight 2010 luminometer (Pharmingen). For mTERT cells, transfection was accomplished using FuGENE 6 reagent (Roche). For studies where effects of more sustained Runx2 expression were analyzed, C3H10T1/2 cells were transduced with adenovirus expression vectors as previously described (36).

**Western Blot Analysis**—Whole cell extracts were prepared by dissolving cell layers in SDS sample buffer. Samples were fractionated by SDS-PAGE on 4–12% precast minigels (Invitrogen) and electrophoretically transferred to nitrocellulose membranes (Schleicher & Schuell). Primary antibodies were used at the following dilutions (Runx2, 1:500; M2, 1:2000; phosphoserine, 1:500). Secondary antibody was used at a dilution of 1:10,000. Immunoreactivity was detected using ECL chemiluminescence reagents (Amersham Biosciences).

**RNA Analysis**—RNA was isolated using TRIzol reagent (Invitrogen) and further purified by DNase I treatment and an RNeasy kit (Qiagen). Reverse transcriptase reactions were conducted with 2 μg of total RNA, TaqMan reverse transcriptase reagents, and an oligo(dT) primer (Applied Biosystems). PCR was performed using an ABI Prism 7700 sequence detection system. Glyceraldehyde-3-phosphate dehydrogenase mRNA was used as an endogenous control.

**Metabolic Labeling and Immunoprecipitation of Runx2**—COS7 cells were transfected with Runx2 expression plasmids, cultured for 30 h, and preincubated in phosphate-free Dulbec-
co’s modified Eagle’s medium, 0.1% FBS for 12 h. Labeling was conducted for 4 h in phosphate-free Dulbecco’s modified Eagle’s medium containing 200 μCi/ml [γ-32P]orthophosphate (phosphorus-32 or [35S]methionine/cysteine Tran35S-label, Amersham Biosciences). Nuclear extracts were prepared as previously described (12) and precleared twice with 50 μl of protein A/G-agarose beads. Appropriate antibodies were added and incubated for 2 h at 4 °C with gentle rocking. Immune complexes were then collected by the addition 30 μl of protein A/G-agarose beads and incubation for 1 h at 4 °C followed by centrifugation. Precipitates were washed five times with 1× washing buffer (20 mM Tris-HCl, pH 7.4, 50 mM KCl, 1 mM dithiothreitol, 0.25% Nonidet P-40, 5 mM sodium fluoride, 1 mM EGTA, 5 mM MgCl2). The immunoprecipitated complexes were suspended in SDS sample buffer and analyzed by SDS-PAGE and autoradiography or Western blot analysis using the indicated antibodies. 32P incorporation was measured using a Packard A2024 InstantImager.

In Vitro Phosphorylation—Synthetic peptides were prepared by the University of Michigan Protein Structure Facility. Each peptide (1 μg) was incubated with 10 μCi of [γ-32P]ATP and 1 unit of activated MAPK (Calbiochem) in a buffer containing 25 mM Tris-HCl, pH 7.4, 10 mM MgCl2, 1 mM dithiothreitol, 40 μM ATP, and 0.5 mM EGTA in a final volume of 25 μl. Samples were incubated at 25 °C for 30 min. Reactions were terminated by the addition of 30 μl of protein A/G-agarose beads and incubation for 1 h at 4 °C with gentle rocking. Immune complexes were then collected by the addition 30 μl of protein A/G-agarose beads and incubation for 1 h at 4 °C followed by centrifugation. Precipitates were washed five times with 1× washing buffer (20 mM HEPES, pH 7.6, 50 mM KCl, 1 mM dithiothreitol, 0.25% Nonidet P-40, 5 mM sodium fluoride, 1 mM EGTA, 5 mM MgCl2). The immunoprecipitated complexes were suspended in SDS sample buffer and analyzed by SDS-PAGE and autoradiography or Western blot analysis using the indicated antibodies. 32P incorporation was measured using a Packard A2024 InstantImager.

Runx2 Purification and Identification of Phosphorylation Sites by Mass Spectroscopy—An adenovirus expression system for biotinylation tagging of Runx2 was developed (32). COS7 cells were transduced with adenovirus encoding Runx2 cDNA with an N-terminal biotinylation sequence and AdBirA (adenovirus expressing bacterial biotin protein ligase) with or without AdMek(sp). After 48 h, cell lysates were adsorbed to streptavidin magnetic beads and purified Runx2 was resolved by SDS-PAGE using a 4–12% gradient gel. The Runx2 gel band was alkylated with iodoacetamide and peptide fragments were generated by in-gel digestion with trypsin. Samples were analyzed by LC/MS/MS using a ThermoFisher LTQ Orbitrap XL (NextGenSciences, Ann Arbor, MI). The Orbitrap MS scan was performed at 60,000 full-width at half-maximum resolution and searched using a local copy of Mascot. Phosphorylated peptides containing P-serine were then identified. In certain cases, product ion data were used to confirm identification of the phosphorylation site.

Statistical Analysis—All statistical analyses were performed using SPSS 16.0 Software. Unless indicated otherwise, each reported value is the mean ± S.D. of triplicate independent samples. Statistical significance was assessed using a one-way analysis of variance.

RESULTS

ERK/MAPK-dependent Phosphorylation and Activation of Runx2 Require a Specific Region of the C-terminal Pro/Ser/Thr

FIGURE 1. Identification of a region in Runx2 necessary for ERK/MAPK-dependent transcriptional activation and phosphorylation. A, schematic of the domain structure of Runx2 with relevant serine residues indicated. AD1–3, transcriptional activation domains; QA, glutamine/alanine-rich domain; RUNT, runt/DNA-binding domain; NLS, nuclear localization sequence; P/S/T domain, proline/serine/threonine-rich domain; RD, repressor domain (from Ref. 30). B, MAPK-dependent transcriptional activity. COS7 cells were transfected with wild type (WT) Runx2 or the indicated C-terminal deletions in the presence of control (LacZ) or Mek(sp) expression vectors and a 6OSE2-luc reporter as described under “Experimental Procedures.” Firefly luciferase activity was normalized for transfection efficiency using a R. reinformis luciferase plasmid. Asterisk, significantly different from corresponding control, p < 0.01; brackets indicate comparisons made; error bars, ± S.D. C and D, Runx2 phosphorylation. COS7 cell cultures treated as in B were metabolically labeled with [32P]orthophosphate or Tran35S-label as described under “Experimental Procedures.” Runx2 was immunoprecipitated (C) and 32P incorporation was normalized to total 35S-labeled protein in each group and expressed as fold-increase with Mek(sp) stimulation (D).

Domain—Fig. 1A shows a schematic of the domain structure of Runx2 with positions of potential phosphorylation sites to be discussed in this study. Activation of the ERK/MAPK pathway by overexpression of a constitutively active form of MEK1 or by treatment with FGF2 was previously shown to stimulate Ocn mRNA expression and promoter activity via a mechanism requiring Runx2 (6, 13). An initial deletion analysis of the Runx2 coding sequence showed that removal of the entire C-terminal proline/serine/threonine-rich (Pro/Ser/Thr) domain (contains AD3 and RD regions encompassing amino acid residues 258–528 in the mouse sequence) rendered Runx2 completely resistant to MAPK regulation and phosphorylation. In contrast, deletion of the N-terminal AD1, AD2, and QA-rich regions (amino acids 1–108) lowered basal transcriptional
activity without affecting MAPK-dependent activation (6).
Similarly, deletion of the Pro/Ser/Thr domain also rendered Runx2 unresponsive to activation by FGF2 (13).

To more precisely define regions of Runx2 necessary for MAPK responsiveness, we carried out a more detailed deletion analysis of the Pro/Ser/Thr domain (Fig. 1, B–D). Wild type Runx2 or several C-terminal deletions (to residues 410, 330, and 286) were transfected into COS7 cells in the presence or absence of constitutively active MEK1 (Mek(sp)) and a p6OSE2-luc Runx2 reporter gene. Samples were either assayed for luciferase activity (panel B) or were metabolically labeled with [32P]orthophosphate and assayed for Runx2 phosphorylation by immunoprecipitation and autoradiography (panels C and D). C-terminal Runx2 deletions gradually reduced MAPK activation of Runx2 transcriptional activity from ~9-fold with wild type Runx2 to ~3-fold with the amino acid 330 deletion. Mek(sp) stimulation was completely lost with deletion to residue 286. Similarly, Mek(sp) stimulated total 32P incorporation into wild type Runx2 by ~2-fold (panel C). This stimulation gradually decreased in the 410 and 330 deletions and was completely lost after deletion to residue 286. These results indicate that the minimal region for MAPK phosphorylation and activation of Runx2 is between amino acids 286 and 330. Subsequent analysis was restricted to this region although it is possible that more C-terminal sites may also participate in this regulation.

Identification of Runx2 Phosphorylation Sites—Inspection of the Runx2 peptide sequence in the 286–330 region (Fig. 2A) identified two putative proline-directed serine phosphorylation sites at residues 301 and 319 and an adjacent site at residue 282. A similar proline-directed threonine site was also seen (Thr326). Incubation of peptides spanning the 264 to 337 region with active ERK1 and [γ-32P]ATP revealed that Ser282, Ser301, and Ser319 were all phosphorylated in vitro, whereas Ser/Ala substitution at each site prevented phosphorylation. In contrast, Thr326 was not phosphorylated under these conditions (i.e. Introduction of an S319A mutation in the 316–337 peptide blocked phos-
phorylation). Peptides containing residues 264–280 and 515–528 were also not phosphorylated.

More extensive analysis of Runx2 phosphorylation was conducted using mass spectroscopy. COS7 cells were transduced with adenovirus vectors expressing a biotinylation tagged Runx2, BirA (bacterial biotin protein ligase), and Mek(sp). Runx2 was then purified from cell lysates using streptavidin magnetic beads and LC/MS/MS analysis was carried out on pepsin-digested samples. As shown in Fig. 2B, five peptides were identified containing phosphoserine at residues 28, 43, 282, 319, and 510. The probability of a correct identification for each peptide was 95% with the exception of the peptide containing Ser28, which had a correlation of 74%. In contrast, only phosphoserine 28 was identified in Runx2 purified from cells pretreated with the MAPK inhibitor, U0126. This indicates that other sites at Ser43, Ser282, Ser319, and Ser510 are directly or indirectly dependent on ERK/MAPK activity for phosphorylation.

To identify direct MAPK targets, nonadenatured Runx2 purified from U0126-treated cells was phosphorylated directly on streptavidin beads with activated P-ERK in vitro and MS was repeated. In this case, phosphate was detected on Ser24, Ser43, Ser319, and Ser347, and Ser510. From this analysis, we conclude that the in vitro phosphorylation sites identified by MS, Ser24, Ser319, and Ser510 are probably direct ERK substrates, whereas Ser282 is likely phosphorylated by a second kinase activated by ERK. Because we also obtained functional evidence for the importance of Ser319 in Runx2 regulation (below), more extensive confirmation that this site is phosphorylated was obtained by analyzing product ion data for the LSQMTSPEHSTTPLSSTRGTL peptide (residues 313–337). Both [M+H]+ charge states were analyzed ([M+H]+ is shown in the left panel of Fig. 2C). Analysis of product ion data (Fig. 2C, right) confirmed that this peptide was phosphorylated at Ser319 (Ser319) within a mass accuracy of 2 ppm.

Surprisingly, although MS analysis consistently identified peptides containing 56–69% of the entire Runx2 sequence (62–76% if the N-terminal Gln/Ala region is excluded), peptides were never identified spanning the Ser301 region. Identified peptides contained 11 of the 12 proline-directed serine/threonine sites in Runx2, making Ser301 the only site not included in our analysis. Similar results were obtained using alternative protease digestions (trypsin/AspN versus pepsin). This suggests that the Ser301 region contains some abnormality in secondary structure, possibly due to post-translational modification, which prevents normal fragmentation and identification.

To further explore the possibility that the Ser301 site is phosphorylated in vivo, we used an indirect approach that takes advantage of the observation that activation of MAPK signaling stimulates a shift in the electrophoretic mobility of a truncated Runx2-(1–330) (Fig. 3A). For this experiment, wild type truncated Runx2-(1–330) or Runx2 containing an S301A mutation, an S319A mutation, or combined S301A,S319A mutations was transfected into COS7 cells with or without constitutively active Mek(sp) and ERK1 expression vectors and analyzed by SDS-PAGE. Activation of MAPK signaling clearly reduced the electrophoretic mobility of wild type Runx2-(1–330) and this mobility change was eliminated by treatment of samples with alkaline phosphatase. In contrast, no detectable MAPK-dependent change in mobility was seen with S301A, S319A mutations or the S301A,S319A double mutation. This is the result that would be expected if a detectable mobility shift requires phosphorylation on both Ser301 and Ser319.

Although the larger size of full-length Runx2 (528 amino acid residues) precluded conducting mobility shift analysis, we obtained additional evidence for Ser301 and Ser319 phosphorylation using immunoprecipitation/immunoblotting with Runx2 and Ser(P) antibodies (Fig. 3B). COS7 cells were transfected with FLAG-tagged wild type, S301A, S319A, or S301A,S319A full-length Runx2 mutants in the presence or absence of Mek(sp) expression vector or in the presence of the MAPK inhibitor, U0126. Samples were then either immunoprecipitated with a specific anti-Runx2 (M2) antibody followed by Western blotting with antiphosphoserine antibody or, alternatively, immunoprecipitated with antiphosphoserine followed by probing with anti-Runx2 antibody. Cells transfected with WT Runx2 displayed a strong Runx2-associated P-serine signal that was further increased by Mek(sp). S301A or S319A mutations each reduced the P-serine signal to a similar extent, whereas the combined S301A,S319A double mutant displayed even weaker P-serine immunoreactivity. MAPK inhibition (+U0126) greatly reduced the P-serine signal in all groups and eliminated differences between WT and mutant Runx2 as would be expected if Ser301 and Ser319 were both phosphorylated in a MAPK-dependent manner. However, the fact that the double mutant still displayed a Mek(sp)-dependent increase in P-serine indicates that Runx2 contains additional direct or indirect MAPK phosphorylation sites, in agreement with our MS data.

In summary, multiple phosphorylation sites were identified in Runx2 including two sites in the aa 286–330 region. Definitive identification of Ser319 as a direct ERK substrate was established using a combination of in vitro peptide phosphorylation, MS/MS, and electrophoresis mobility shift analysis. Strong evidence was also obtained that Ser301 is phosphorylated by ERK (in vitro peptide phosphorylation, electrophoresis mobility shift analysis, and Ser(P)/Runx2 co-precipitation). However, it
**MAPK Regulation of Runx2 Phosphorylation**

![Graph and Table]

**FIGURE 3.** Additional evidence for Runx2 phosphorylation at Ser<sup>301</sup>. A, analysis of electrophoretic mobility of wild type (WT) truncated Runx2-(1–330) and Ser<sup>301</sup> and Ser<sup>319</sup> mutants. The indicated truncated Runx2 mutants were expressed in COS7 cells in the presence or absence of Mek(sp) and ERK1 expression vectors. Cell lysates were analyzed by SDS-PAGE with or without prior treatment with calf intestinal alkaline phosphatase (CIAP). Runx2 was detected by Western blotting. B, anti-P-serine antibody reactivity with full-length Runx2. FLAG-tagged WT Runx2 or the indicated mutants were expressed in COS7 cells in the presence (+) or absence (−) of Mek(sp) or in the presence of the MAPK inhibitor (+ U0126). Nuclear extracts were then either immunoprecipitated with M2 antibody and probed with an anti-P-serine monoclonal antibody or immunoprecipitated (IP) with anti-P-serine and probed with M2.

Inspection of the runt domain region of Runx2 revealed that it contains a consensus ERK docking “D” site (GKSFTLTITVFTNPP) at aa 201–215 (37). To confirm its role in ERK complex formation, this region of Runx2 was deleted, resulting in complete loss of ERK binding (Fig. 4D). The D site deletion also almost completely blocked the ability of Mek(sp) to stimulate Runx2-dependent transcription of a 6OSE2-luc reporter (supplemental Fig. S1).

### Functional Analysis of Phosphorylation Sites

We initially assessed the functionality of the above phosphorylation sites in the context of the Runx2-(1–330) fragment because this contained the minimal sequence for MAPK responsiveness. Ser/Ala mutants described above or Ser/Glu mutations were generated at each site individually or in combination, and wild type or mutated expression plasmids were transfected into COS7 cells with 6OSE2-luc reporter (+/− Mek(sp)). As shown in Fig. 5A, the ability of Mek(sp) to stimulate transcriptional activity of Runx2-(1–330) was totally blocked with the MEK1/2 inhibitor, U0126. However, individual Ser/Ala mutations at residues 301 or 319 only slightly inhibited MAPK stimulation, whereas the S282A mutation was without effect. On the other hand, introduction of a S301E, S319E double mutation completely eliminated MAPK responsiveness. In contrast, the S301E, S319E mutant exhibited high basal transcriptional activity in the absence of MAPK stimulation that was not affected by Mek(sp). This is consistent with previous studies showing that addition of the charged amino acid is able to mimic a phosphorylated serine residue (38).

To evaluate the role of phosphorylation sites in the context of the native Runx2 protein, S301A, S319A or S301E, S319E mutations were also introduced into full-length Runx2 and evaluated for Mek(sp) (Fig. 5B) or FGF2 responsiveness (Fig. 5C) using the same 6OSE2-luc reporter used above. As was the case with Runx2-(1–330), the Mek(sp)-dependent induction of luciferase activity was blocked with U0126, whereas the S301E, S319E mutation resulted in constitutive activation of reporter activity that was not further stimulated by Mek(sp). In contrast to the result obtained with Runx2-(1–330), S301A, S319A mutations only partially blocked the MAPK response (approximately 40% inhibition).

As noted above, FGF2 induction of Ocn expression also requires ERK/MAPK activity and is associated with Runx2 phosphorylation (13). A preliminary deletion analysis (not shown) also indicated that FGF2 responsiveness was lost after deletion of the aa 286–330 Runx2 region. To evaluate whether Ser<sup>301</sup> and Ser<sup>319</sup> are required for the FGF2 response, wild type...
and artificial reporter constructs often do not mimic those obtained with endogenous genes, we considered it important to show that these mutations also reduce the ability of Runx2 to stimulate endogenous osteoblast gene expression and differentiation. Two approaches were taken to address this issue. In the first, mTERT-immortalized calvarial cells from Runx2−/− mice (34) were transfected with wild type Runx2, S301A,S319A, or S301E,S319E mutants (Fig. 6). After 5 days of growth in ascorbic acid–containing medium to stimulate ECM-dependent differentiation, total RNA was isolated and levels of osteocalcin (A) and bone sialoprotein mRNAs (B) were measured by quantitative reverse transcription-PCR. As expected, wild type Runx2 transfection strongly induced both mRNAs and this induction was suppressed by the MAPK inhibitor, U0126. In contrast, the S301A,S319A mutant only weakly stimulated mRNA expression and its activity was resistant to further inhibition by U0126. The S301E,S319E mutant, on the other hand, induced Ocn and Bsp mRNAs to higher levels than those obtained with wild type Runx2 via a mechanism that was largely resistant to MAPK inhibition. As shown in panel C, these results cannot be explained by differences in expression levels of wild type and mutant Runx2 proteins.

To examine the requirement for Runx2 phosphorylation sites in osteoblast differentiation over a more prolonged time period, we used an adenovirus expression system in C3H10T1/2 cells. This mesenchymal cell line contains no detectable endogenous Runx2, but will undergo osteoblast differentiation after transduction with an adeneno-Runx2 expression vector. Previous studies showed that AdRunx2 vectors continue to produce active transcription factors for at least 7–10 days in this system (36). Wild type and S301A,S319A Runx2 adenoviruses were constructed and titered in C3H10T1/2 cells to produce equivalent amounts of Runx2 protein as measured on Western blots. As shown in Fig. 7, wild type Runx2 clearly induced osteoblast differentiation in this system. Sustained induction of alkaline phosphatase activity (panel A) as well as Ocn and Bsp mRNAs (panels B and C) was observed over a 12-day period. In contrast, cells expressing the Ser/Ala mutant had less than 25% the alkaline phosphatase activity of wild type at all times examined. Induction of Ocn and Bsp mRNAs was similarly attenuated. Western blot results revealed little or no differences in Runx2 protein levels in the two groups at all times examined, making it unlikely that differences in Runx2 expression or stability could explain these results.

**DISCUSSION**

In this study, we identified two phosphorylation sites in Runx2 at Ser301 and Ser319 that are required for MAPK-dependent activation of Runx2 transcriptional activity and osteoblast differentiation. These sites are phosphorylated in a MAPK-dependent manner in intact cells. As shown by MS/MS analysis, at least one, Ser319, is a direct ERK1 substrate. Furthermore, Runx2 can bind ERK using a D site between amino acids 201 and 215, and artificial reporter constructs often do not mimic those obtained with endogenous genes, we considered it important to show that these mutations also reduce the ability of Runx2 to stimulate endogenous osteoblast gene expression and differentiation. Two approaches were taken to address this issue. In the first, mTERT-immortalized calvarial cells from Runx2−/− mice (34) were transfected with wild type Runx2, S301A,S319A, or S301E,S319E mutants (Fig. 6). After 5 days of growth in ascorbic acid–containing medium to stimulate ECM-dependent differentiation, total RNA was isolated and levels of osteocalcin (A) and bone sialoprotein mRNAs (B) were measured by quantitative reverse transcription-PCR. As expected, wild type Runx2 transfection strongly induced both mRNAs and this induction was suppressed by the MAPK inhibitor, U0126. In contrast, the S301A,S319A mutant only weakly stimulated mRNA expression and its activity was resistant to further inhibition by U0126. The S301E,S319E mutant, on the other hand, induced Ocn and Bsp mRNAs to higher levels than those obtained with wild type Runx2 via a mechanism that was largely resistant to MAPK inhibition. As shown in panel C, these results cannot be explained by differences in expression levels of wild type and mutant Runx2 proteins.

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In this study, we identified two phosphorylation sites in Runx2 at Ser301 and Ser319 that are required for MAPK-dependent activation of Runx2 transcriptional activity and osteoblast differentiation. These sites are phosphorylated in a MAPK-dependent manner in intact cells. As shown by MS/MS analysis, at least one, Ser319, is a direct ERK1 substrate. Furthermore, Runx2 can bind ERK using a D site between amino acids 201 and 215 in the runt domain region. Consistent with Ser301 and Ser319 being important for osteoblast function, inactivating Ser to Ala mutations greatly reduced the ability of Runx2 to stimulate expression of Ocn and Bsp mRNAs in Runx2−/− calvarial cells and blocked Runx2-dependent induction of osteoblast differentiation.
MAPK Regulation of Runx2Phosphorylation

blast gene expression and differentiation in a mesenchymal cell line. In contrast, Ser to Glu mutations, which mimic the charge density of phosphorylated amino acids, activated Runx2-dependent transcription. Taken together, these studies emphasize the importance of MAPK-dependent phosphorylation as a means of controlling Runx2 transcriptional activity in bone.

As one of the major signal transduction pathways in bone, the ERK/MAPK pathway is able to integrate stimuli from growth/differentiation factor binding to receptor tyrosine kinases (39), ECM-integrin binding and focal adhesion kinase activation (40), certain non-genomic actions of estrogens (41), and mechanical stimulation mediated by FAK activation (42) and connexin 43 up-regulation (43). It also has important functions in the differentiation of post-mitotic mesenchymal and neuronal cells (44, 45) and regulates the activity of several tissue-specific transcription factors including MyoD (muscle (46)), Sox9 (cartilage (47)) and peroxisome proliferator-activated receptor γ (adipose (48)). Furthermore, as previously shown by this laboratory, in vivo transgenic stimulation of ERK/MAPK signaling in osteoblasts accelerates bone development and is able to partially rescue the cleidocranial dysplasia pheno-

FIGURE 5. Functional analysis of Runx2 phosphorylation sites. A, identification of phosphorylation sites necessary for MAPK responsiveness using Runx2-(1–330). Specific mutations were created in Runx2-(1–330) to generate S282A, S301A, or S319A mutants or the indicated combinations as well as an S301E,S319E mutant. Runx2 expression plasmids were transfected into COS7 cells in the presence or absence of Mek(sp) vector and luciferase reporters as described in the legend to Fig. 1. B, evaluation of requirement for Ser131 and Ser919 sites in the context of full-length Runx2 protein. S301A,S319A or S301E,S319E mutations were generated in full-length Runx2 and evaluated for Mek(sp) responsiveness as in panel A. C, FGF2 responsiveness of wild type (WT) and mutant Runx2. COS7 cells were transfected with wild type full-length Runx2 or the S301A,S319A mutant. After 24 h, cells were treated for an additional 24 h with FGF2 (50 ng/ml) before luciferase activity was measured. Statistically significant differences are indicated: a, p < 0.05; b, p < 0.01. Error bars, ± S.D.

FIGURE 6. Induction of osteoblast differentiation markers by wild type (WT) Runx2 and phosphorylation site mutants. An mTERT-immortalized cell line derived from Runx2−/− calvaria was transfected with wild type and mutant Runx2 expression vectors and grown for 5 days in ascorbic acid-containing medium before measurement of Ocn (A) and Bsp (B) mRNAs by real-time reverse transcription-PCR. Indicated samples were also treated with U0126 12 h before harvest. The mRNA levels were normalized to glyceraldehyde-3-phosphate dehydrogenase (Gapdh) mRNA in each sample. Runx2 protein in each group was measured on Western blots (WB) (C). S/A mut, S301A,S319A mutant; S/E mut, S301E,S319E mutant. Statistically significant differences are indicated: a, p < 0.05; b, p < 0.01. β-gal, β-galactosidase. Error bars, ± S.D.
MAPK Regulation of Runx2 Phosphorylation

Possible that phosphorylation sites in the region between residues 330 and 528 may also participate in the MAPK response, perhaps functioning as secondary sites after priming phosphorylations occur at Ser\(^{301}\) and Ser\(^{319}\). Consistent with this idea, our MS/MS analysis, which identified peptides containing all proline-directed serines in Runx2 except Ser\(^{301}\), detected two additional direct MAPK phosphorylation sites at Ser\(^{313}\) and Ser\(^{510}\) as well as an indirect MAPK site at Ser\(^{282}\). Also, the S301A,S319A mutant Runx2 still retained a MAPK-dependent increase in total P-serine on Western blots as would be expected if additional sites were present (Fig. 3). However, the observation that the high constitutive activity of the S301E,S319E Runx2 mutant was refractory to MAPK inhibition (Fig. 6) argues against these other sites being major contributors to the MAPK response because inhibiting their phosphorylation was not able to prevent the transcriptional activation seen when 301 and 319 sites were in an activated state.

MAPK-dependent phosphorylation of Runx2 is also clearly required for FGF2-dependent induction of Ocn expression (13). However, S301A,S319A mutations only partially blocked FGF2 activation of the GOSE2-luc reporter (Fig. 5C). This partial inhibition may be explained by the involvement of additional MAPK sites as well as other kinase sites. In this regard, a recent study by Kim and co-workers (49) reported that FGF2 also activates Runx2 via phosphorylation by PKC\(\gamma\) at Ser\(^{247}\). Thus, it is possible that FGF2 activates Runx2 transcriptional activity by phosphorylating both ERK/MAPK and PKC sites.

Runx family members exhibit a high degree of amino acid sequence conservation, particularly in the DNA binding or Runt domain. Although the C-terminal Pro/Ser/Thr domain of Runx proteins is not highly conserved, the two phosphorylation sites we identified in Runx2 are as well present in Runx1 (but not in Runx3). Interestingly, EGF and phorbol ester activation of the ERK/MAPK pathway can stimulate Runx1 transcriptional activity via phosphorylation on these sites (50, 51). Runx1 is essential for hematopoietic cell differentiation (52) and is also a frequent site for chromosomal translocations in acute myelogenous leukemia. Consistent with this oncogenic activity, transfection of Runx1 into fibroblasts stimulates anchorage-independent growth and transformation. Interestingly, Ser/Ala mutations at Ser\(^{249}\) and Ser\(^{266}\) in Runx1 (equivalent to Ser\(^{301}\) and Ser\(^{319}\) in Runx2) inhibit growth of NIH 3T3 fibroblasts in soft agar, a common assay for cell transformation (50). Runx2 can also function as an oncogene under certain conditions and has been associated with cell proliferation and migration of breast cancer cells (53, 54). It is, therefore, possible that ERK/MAPK-dependent phosphorylation of Runx2 at Ser\(^{301}\) and Ser\(^{319}\) could also be associated with metastasis-related behavior.

It is not presently understood how phosphorylation of Runx2 stimulates transcription. We and others previously observed that the apparent affinity of Runx2 for OSE2-containing DNA increases with differentiation and this increase can be blocked with MAPK inhibition (12, 26, 55). However, it is not known if this is a direct consequence of Runx2 phosphorylation. Interestingly, using chromatin immunoprecipitation assays, we find Runx2 associated with Ocn and Bsp chromatin in both differentiated and undifferentiated MCT3-T3-E1 cells even in the pres-
enence of MAPK inhibition (56). Thus, Runx2 does not dissociate from its binding sites on chromatin even though its in vitro affinity for DNA may be lower in the unphosphorylated state. Runx2 is also known to serve as a docking site for many nuclear factors that can form active or inactive transcription complexes on chromatin (57). In this regard, we recently showed that the physical association of Runx2 with ERK reported in the present study can also be detected on the chromatin of Runx2 target genes in vivo (56). In this case, P-ERK binding to Ocn and Bsp chromatin required Runx2 and intact Runx2 binding sites in the DNA. Furthermore, this binding was dependent on the elevated MAPK activity associated with osteoblast differentiation. Runx2 can therefore be viewed as providing a docking site for P-ERK on the chromatin of target genes. In addition to phosphorylating Runx2, chromatin-bound P-ERK may also initiate subsequent events such as phosphorylation of other chromatin substrates or the recruitment of additional factors including histone acetyltransferases like p300/cAMP-response element-binding protein to modify chromatin structure, thereby allowing the initiation of transcription. Interestingly, the ERK/MAPK-dependent phosphorylation of Runx1 discussed above is associated with the dissociation of the histone deacetylase co-factor, mSin3a, from Runx1, thereby allowing subsequent increases in histone acetylation (58). Because Runx1 phosphorylation sites are conserved in Runx2, this observation provides a plausible mechanism for how ERK/MAPK phosphorylation could alter Runx2-dependent transcription. This possibility is currently being pursued by this laboratory. Last, Afzal and coworkers (59) showed that MAPK-mediated phosphorylation of Runx2 is also necessary for complex formation with Smads.

In addition to the ERK/MAPK-dependent regulation of Runx2 described herein, several other types of post-translational modifications have been described for this molecule. Phenylthiohydantoin/protein kinase A-mediated phosphorylation of a C-terminal Runx2 site was correlated with induction of MMP13 (60). More recently, Cdk4-mediated phosphorylation at Ser272 was shown to target Runx2 for ubiquitination and proteosomal degradation during the cell cycle (61), whereas cdc2 phosphorylation at Ser451 was shown to be necessary for cell cycle progression of endothelial cells (62). Also, glycogen synthase kinase 3β-dependent phosphorylation of Runx2 at Ser369–Ser373, Ser377 was shown to reduce transcriptional activity (63). Last, Runx2 can be acetylated on critical lysine residues by p300 acetyltransferase. This modification, which is stimulated by BMP2, increases transcription and stabilizes Runx2 against proteosomal degradation (64). Thus, post-translational modification appears to be a common mechanism for regulating Runx2 activity and stability.

In summary, phosphorylation of Runx2 at Ser301 and Ser319 clearly has an important regulatory role in Runx2-dependent transcription because mutation of these sites in the context of the intact Runx2 molecule severely attenuated the ability of Runx2 to stimulate osteoblast-specific gene expression during differentiation. Ongoing in vivo studies will be necessary to assess the full impact of Runx2 phosphorylation to the overall activity of this molecule during skeletal development and remodeling.

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IDENTIFICATION AND FUNCTIONAL CHARACTERIZATION OF EXTRACELLULAR-REGULATED KINASE/MAPK PHOSPHORYLATION SITES IN THE RUNX2 TRANSCRIPTION FACTOR

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Supplementary Figure S1

Supplementary Figure Legend

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A supplementary figure is used to describe the effect of a D site deletion on MAPK-stimulated Runx2 transcriptional activity. Wild type full-length Runx2 (WT-FL) or Runx2 containing a 15 amino acid residue deletion (Δ201-215) spanning the Erk-binding D site (D site-FL) was transfected into COS7 cells with a 6OSE2-luc reporter and renilla normalization vector in the presence or absence of constitutively-active Mek1 (Meksp). After 48 h, cells were harvested for measurement of luciferase activity (A) or total transfected Runx2 protein as measured by immunoblotting (B).
Critical Role of Activating Transcription Factor 4 in the Anabolic Actions of Parathyroid Hormone in Bone

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Abstract

Parathyroid hormone (PTH) is a potent anabolic agent for the treatment of osteoporosis. However, its mechanism of action in osteoblast and bone is not well understood. In this study, we show that the anabolic actions of PTH in bone are severely impaired in both growing and adult ovariectomized mice lacking bone-related activating transcription factor 4 (ATF4). Our study demonstrates that ATF4 deficiency suppresses PTH-stimulated osteoblast proliferation and survival and abolishes PTH-induced osteoblast differentiation, which, together, compromise the anabolic response. We further demonstrate that the PTH-dependent increase in osteoblast differentiation is correlated with ATF4-dependent up-regulation of Osterix. This regulation involves interactions of ATF4 with a specific enhancer sequence in the Osterix promoter. Furthermore, actions of PTH on Osterix require this same element and are associated with increased binding of ATF4 to chromatin. Taken together, these experiments establish a fundamental role for ATF4 in the anabolic actions of PTH on the skeleton.

Introduction

Parathyroid hormone (PTH) is a major regulator of calcium homeostasis and has both catabolic and anabolic effects on osteoblasts and bone that depend on the temporal pattern of administration. Continuous administration of PTH decreases bone mass whereas intermittent administration increases bone mass [1–6]. The mechanism(s) responsible for these differing effects are poorly understood. The anabolic activity of PTH has been attributed to both direct actions of this hormone on osteoprogenitor cells as well as indirect effects mediated by the production of growth factors such as insulin-like growth factor-1 (IGF-1) and basic fibroblast growth factor-2 (FGF-2) [7,8]. Most cellular actions of PTH are mediated by the PTH-1 receptor, a G protein-coupled receptor that is expressed in osteoblasts [9,10]. Binding of PTH to its receptor activates multiple intracellular signaling pathways that involve cAMP, inositol phosphates, intracellular Ca2+ and protein kinases A and C [11], and the extracellular signal-related (ERK)/mitogen-activated protein kinase (MAPK) pathway [12,13]. Activation of these signal transduction pathways ultimately affects cellular behavior. In this regard, the anabolic actions of PTH on bone have been attributed to increased proliferation of osteoprogenitors/osteoblasts [2,14,15] and/or decreased osteoblast apoptosis [6,16,17].

Although a number of transcription factors including cAMP response element binding protein (CREB) [18,19], API family members [20–22], and Runx2 [20,23] have been implicated in the molecular actions of PTH in osteoblasts, genetic studies have not strongly linked any of these factors in the anabolic actions of this hormone. To better understand the anabolic actions of PTH, it is essential that the downstream signals induced by this hormone be identified and evaluated for possible roles in bone formation. The osteocalcin (Ocn) promoter has been an important tool for unraveling the mechanisms mediating osteoblast-specific gene expression and was used to identify a number of important transcription factors and cofactors involved in Ocn gene expression [24–28]. Because the Ocn gene is regulated by PTH [29,30], we have used it as a model system for identifying new transcriptional mediators of PTH action. We previously demonstrated that the OSE1 (osteoblast-specific element 1) in the proximal mOg2 promoter [24] is necessary and sufficient for PTH induction of this gene [31]. The OSE1 core sequence (TTACATCA) was subsequently identified as a DNA binding site for the ATF4 transcription factor. The critical role of ATF4 in osteoblast differentiation and bone development was established using Aft4-deficient mice [26]. At the cellular level, ATF4 is critical for proliferation and differentiation as well as survival in osteoblasts [32,33]. We recently showed that ATF4 is also required for PTH induction of Ocn expression in osteoblasts [34]. Specifically, PTH elevated levels of ATF4 mRNA and protein in a dose and time-dependent manner and increased binding of ATF4 to OSE1 DNA. Furthermore, PTH stimulation of Ocn expression was lost by siRNA downregulation of ATF4 in


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MC-4 cells and in primary bone marrow stromal cells from *Atf4<sup>−/−</sup>* mice. Collectively, these studies demonstrate that ATF4 is a novel downstream mediator of PTH signaling.

Osterix (Osx, or Sp7), a zinc-finger-containing transcription factor of the sp family, is essential for osteoblast differentiation and bone formation [35]. Since Osx is not detected in mice lacking Runx2 [35], a master regulator of osteoblast differentiation [25,36–30], it functions downstream of Runx2. However, the molecular mechanisms whereby the Osx gene is transcriptionally regulated are not well understood.

In the present study, we used ATF4-deficient mice to determine whether ATF4 is more generally required for the in vivo anabolic actions of PTH in bone as well as explore the mechanism used by PTH to regulate ATF4 activity. As will be shown, loss of ATF4 greatly attenuated the anabolic effects of PTH. Furthermore, ATF4 may participate in the PTH response by regulating the expression of the Osterix transcription factor.

**Results**

The anabolic effects of PTH on bone are severely impaired in growing Atf4-deficient mice

We first evaluated our hypothesis that ATF4 mediates the anabolic actions of PTH in bone using a relatively simple “growing mouse” model system, that has been widely used for studying the anabolic actions of PTH, PTH/PTHrP, FGF2, and IGF-1 in bone [2–4,8,39,40]. The advantages of this system are that it is less time consuming and costly versus adult ovariectomized mouse models. Furthermore, because young growing animals have relatively high osteoblast activity, they are more sensitive to PTH than adults [2,11,14,16–18,41–43]. Mice were treated with vehicle or PTH and sacrificed 24 h after the last PTH injection. PTH-dependent anabolic activity was evaluated in these mice using standard biochemical and histomorphometric criteria. *Atf4<sup>−/−</sup>* mice grew more slowly than wild type (wt) animals. The growth rate was slightly but significantly increased by PTH treatment during d 6–18 in wt but not *Atf4<sup>−/−</sup>* animals (Fig. SIA). *Atf4<sup>−/−</sup>* femurs were also shorter than wt or *Atf4<sup>+/+</sup>* femurs. Consistent with results from a previous study [3], PTH did not alter the length of femurs (Fig. S1B). However, it did significantly increase the dry ash weight per femur in wt and *Atf4<sup>−/−</sup>* but not in *Atf4<sup>+/+</sup>* mice (Fig. S1C). Serum Pi and calcium concentration (Fig. S1D and E) were not markedly affected by PTH or ATF4 deficiency. Fasxtron X-ray analysis of femurs revealed that wt and *Atf4<sup>−/−</sup>* mice responded to PTH with markedly increased trabeculae throughout the whole femur, with the most dramatic increase in the metaphyseal region (Fig. S2, top and middle). In contrast, PTH only slightly increased the radiopacity in the same region of *Atf4<sup>−/−</sup>* femurs (Fig. S2, bottom). As shown in Fig. 1, quantitative μCT analysis of femur histomorphometric parameters showed that *Atf4<sup>−/−</sup>* mice had a significant reduction in bone volume/tissue volume (BV/TV), trabecular number (Tb.N), and cortical thickness (Cort.Th) and a marked increase in trabecular space (Tb.Sp) compared with the wt or *Atf4<sup>+/+</sup>* littermates. These data confirmed an essential role of ATF4 in bone that was previously demonstrated by the Karsenty group [26]. As expected, in wt femurs, intermittent PTH increased BV/TV, Tb.N, and Tb.Sp by 5.4-fold, 2.7-fold, and 1.5-fold, respectively, and decreased Tb.Sp by 60 percent. Similar effects were also seen in *Atf4<sup>−/−</sup>* mice (Fig. 1B). In contrast, the PTH response was greatly attenuated in *Atf4<sup>−/−</sup>* mice where the following PTH responses were observed; BV/TV, 2.2-fold increase; Tb.N, 1.7-fold increase; Tb.Th, 1.1-fold increase; Tb.Sp, 36 percent decrease. In all cases, the magnitude of PTH-stimulated changes on BV/TV, Tb.N, Tb.Sp was dramatically reduced in *Atf4<sup>−/−</sup>* mice relative to wt or *Atf4<sup>+/+</sup>* mice (*P*<0.05, PTH/veh-wt vs. PTH/veh-*Atf4<sup>−/−</sup>*/). Furthermore, PTH-stimulated increases in Corr.Th and Tb.Th were completely lost in *Atf4<sup>−/−</sup>* femurs. Because PTH similarly affected all trabecular and cortical parameters in *Atf4<sup>−/−</sup>* and *Atf4<sup>+/+</sup>* mice, subsequent experiments compared the PTH effects on bone only between wt and *Atf4<sup>−/−</sup>* mice.

We next measured effects of *Atf4* gene ablation on PTH stimulation of tibiae, vertebrae, and calvariae. The anabolic effect of PTH on wt tibiae was so dramatic that the majority of the bone marrow cavity was replaced by newly formed bone (Fig. 2A and B). In *Atf4<sup>−/−</sup>* tibiae, while PTH still induced a small increase in trabecular area, the magnitude of stimulation was significantly reduced (5-fold in wt vs. 2.2-fold in *Atf4<sup>−/−</sup>*; <0.05, PTH/veh-wt vs. PTH/veh-*Atf4<sup>−/−</sup>*/). Similarly, the PTH-stimulated increase in the trabeculae of vertebrae (L5) was markedly reduced in *Atf4<sup>−/−</sup>* mice (3-fold in wt vs. 2-fold in *Atf4<sup>−/−</sup>*; <0.05, PTH/veh-wt vs. PTH/veh-*Atf4<sup>−/−</sup>*/). When histological sections of calvariae were compared, PTH increased the width of the calvariae by 1.8-fold in wt mice, a response that was abolished in *Atf4<sup>−/−</sup>* animals (Fig. 2K–O).

Ablation of the Atf4 gene impairs PTH stimulation of trabecular, but not cortical bone in 7-month-old ovariectomized (OVX) mice

The experiments described above clearly establish a critical role of ATF4 in the anabolic effects of PTH on long bones, vertebrae, and calvariae in rapidly growing mice. However, it is possible that results obtained from growing animals may be different from those in adults due to possible effects of PTH and/or ATF4 on animal growth or influences of animal growth on the anabolic response to PTH, either of which could complicate the interpretation of the results. In contrast, adult mice have a mature skeleton in which these possible complications can be avoided. Furthermore, the OVX mouse provides a model that may be more relevant to the clinical applications of PTH in the treatment of osteoporosis. For these reasons, we next evaluated whether ATF4 is required for the anabolic response to PTH in 7-month-old OVX mice. OVX surgery was successful as demonstrated by significant reduction in BV/TV (65 percent), Tb.N (27 percent), and Tb.Th (18 percent) and increased Tb.Sp (37 percent) relative to sham surgery (*P*<0.05, wt-sham vs. wt-OVX) (Fig. S3). OVX surgery did not reduce Corr.Th, which is consistent with results from rats [44]. Interestingly, OVX surgery did not significantly reduce bone parameters in *Atf4<sup>−/−</sup>* animals (*P>*0.05, *Atf4<sup>−/−</sup>*-sham vs. *Atf4<sup>−/−</sup>*-OVX). As shown in Fig. 3A–C, similar to results from growing mice, ablation of the *Atf4* gene significantly decreased BV/TV and Tb.N and increased Tb.Sp in adult OVX mice. In further agreement with results in young mice, *Atf4<sup>−/−</sup>* animals exhibited a clearly attenuated response to PTH. For example, while PTH increased BV/TV by 7.8-fold in wt mice, this value was only increased 4.2-fold in *Atf4<sup>−/−</sup>* animals. Similarly, while PTH still stimulated formation of trabecular bone in *Atf4<sup>−/−</sup>*-trabeculae, the magnitude of this response was significantly reduced compared to wt control (*P>*0.05, PTH/veh-wt vs. PTH/veh-*Atf4<sup>−/−</sup>*/). In contrast to result from growing mice, Corr.Th was not reduced by ablation of the *Atf4* gene in adult OVX mice (0.21±0.01 mm in wt vs. 0.19±0.02 mm in *Atf4<sup>−/−</sup>*; >0.05 wt vs. *Atf4<sup>−/−</sup>*). Also, PTH was much less effective in stimulating Corr.Th in adult OVX mice (24%) than in growing mice (95%) (Figs. 1 and 3). Furthermore, no difference in stimulation of cortical thickness by PTH was observed when wt and *Atf4<sup>−/−</sup>* groups were compared (24% wt vs. 21% *Atf4<sup>−/−</sup>*; >0.05, PTH/veh-wt vs. PTH/veh-*Atf4<sup>−/−</sup>*/). As shown in Fig. 3D and E, results from calcein double labeling of 7-month old OVX wt and *Atf4<sup>−/−</sup>* tibia revealed that the PTH-stimulated increase in mineral
apposition rate (MAR), an indicator of osteoblast function, was significantly reduced by ATF4 deficiency ($P<0.05$, PTH/veh-wt vs. PTH/veh-Atf4<sup>2/2</sup>).

ATF4 deficiency significantly reduces basal and PTH-stimulated proliferation in osteoblasts/preosteoblasts

PTH and PTHrP are both known to increase the proliferation and numbers of osteoblasts [14,15,17,45]. ATF4 is also a positive regulator of osteoblast proliferation and can be up-regulated by PTH in these cells [33,34]. To determine whether ATF4 plays a role in PTH regulation of osteoblast proliferation, sections of tibiae and calvariae from wt and Atf4<sup>2/2</sup> mice treated with and without intermittent PTH were analyzed for in vivo cell proliferation using a Zymed BrdU immunostaining kit. As shown in Fig. 4A, B, and E, in wt mice, PTH increased the percentage of proliferating osteoblasts/preosteoblasts of tibial trabeculae by 2.8-fold relative to vehicle-treated control. Ablation of the Atf4 gene resulted in a 50% decline in basal proliferation. In addition, the PTH-stimulated increase in proliferation was decreased by 40 percent (Fig. 4C–E). Similarly, PTH-induced proliferation in calvarial periosteal osteoblasts was also significantly reduced by ATF4 deficiency (Fig. 4F–J). As expected, very few osteocytes were BrdU-positive in both tibiae and calvariae. Note: basal proliferation rate of calvarial periosteal osteoblasts was significantly higher than that of tibial trabecular osteoblasts (28% vs. 4%). Therefore, ATF4 is critical for basal and PTH-stimulated proliferation of osteoblasts/preosteoblasts in vivo.

**Figure 1.** PTH-stimulated bone was significantly reduced or lost in Atf4<sup>2/2</sup> femurs. A, two-dimensional (2D) reconstruction from μCT scan of femurs from growing wt, Atf4<sup>++</sup> and Atf4<sup>2/2</sup> mice treated with and without intermittent PTH for 28 d. B, quantitative analysis of bone volume/tissue volume (BV/TV), trabecular number (Tb. N), trabecular thickness (Tb.Th), trabecular space (Tb.Sp), and cortical thickness (Cort. Th). *$P<0.05$ (veh vs. PTH), †$P<0.05$ (wt-veh vs. Atf4<sup>2/2</sup>-veh), ‡$P<0.05$ (PTH/veh-wt vs. PTH/veh-Atf4<sup>2/2</sup>).

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ATF4 and PTH Anabolism

**Table 1.**

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N = 6-12


**Figure 1.** PTH-stimulated bone was significantly reduced or lost in Atf4<sup>2/2</sup> femurs. A, two-dimensional (2D) reconstruction from μCT scan of femurs from growing wt, Atf4<sup>++</sup> and Atf4<sup>2/2</sup> mice treated with and without intermittent PTH for 28 d. B, quantitative analysis of bone volume/tissue volume (BV/TV), trabecular number (Tb. N), trabecular thickness (Tb.Th), trabecular space (Tb.Sp), and cortical thickness (Cort. Th). *$P<0.05$ (veh vs. PTH), †$P<0.05$ (wt-veh vs. Atf4<sup>2/2</sup>-veh), ‡$P<0.05$ (PTH/veh-wt vs. PTH/veh-Atf4<sup>2/2</sup>).

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the survival of osteoblasts and osteocytes by reducing apoptosis [6,16,17,45]. Our recent study shows that ATF4 is anti-apoptotic in osteoblasts [33]. To determine whether ATF4 plays a role in PTH-mediated anti-apoptosis, sections of tibiae were stained with TUNEL and apoptotic cells were assessed. As shown in Fig. 4K and L, ATF4 deficiency significantly increased the basal levels of apoptosis. As expected, PTH dramatically reduced apoptotic death of tibial trabecular osteoblasts/osteocytes by 48 percent, which is consistent with results from previous studies [6,16,17,45]. Importantly, the PTH-stimulated decrease in apoptotic death was completely abolished in Atf4−/− trabeculae (Fig. 4M–O). ATF4 deficiency also reduced basal Ocn and Osx mRNA levels. Consistent with our previous report [34], PTH increased Atf4 mRNA 2.2-fold in wt tibiae, while Atf4 was undetectable in Atf4−/− animals. In contrast, c-Fos and c-Jun, both early PTH-induced genes, were not induced by PTH in either wt or Atf4−/− tibiae. As shown in Fig. 5B, the levels of IGF-1 and FGF-2 which have both been implicated in the anabolic actions of PTH in bone [7,8] were markedly reduced in plasma from Atf4−/− mice compared to wt mice (P<0.05, wt vs. Atf4−/−). However, their levels were not significantly elevated by the treatment of intermittent PTH in both wt or Atf4−/− animals (P>0.05, veh vs. PTH).

**Figure 2.** PTH-stimulated bone is severely impaired in Atf4−/− tibiae, vertebrae, and calvariae. Representative H&E stained sections of tibiae (A–E), vertebrae (L–J), and calvariae (K–O) are shown. Trabecular bone area versus total area of tibiae (E) and vertebrae (J) was measured using an Image Pro Plus 6.2 software. The calvarial width was obtained from 20 random measurements throughout the whole calvaria using a SPOT Advanced imaging software (O). *P<0.05 (veh vs. PTH), † P<0.05 (wt-veh vs. Atf4−/−-veh), ‡P<0.05 (PTH/veh-wt vs. PTH/veh-Atf4−/−).

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**Intermittent PTH increases in vivo Osx expression in osteoblasts through a pathway requiring ATF4**

Our above results demonstrate that ATF4 is essential for the major anabolic actions of PTH on bone and is also required for PTH-dependent induction of osteoblast differentiation. To begin to address the mechanism underlying this response, we measured the expression of Osterix (Osx) and Runx2 proteins, two critical transcription factors that regulate osteoblast differentiation. Initially, we used immunohistochemistry (IHC) to measure Osx in the tibiae and calvariae of wt and Atf4−/− mice with or without 28 d anabolic PTH treatment. As shown in Fig. 6, in wt-vehicle-treated tibiae, Osx-positive osteoblasts were only identified in the trabecular and cortical endosteum close to the growth plate (Fig. 6A1, B1, and D1) and were almost undetectable in the same
regions close to the marrow (Fig. 6C1), indicating that cells in these areas are still in the immature (preosteoblast) state. In contrast, in the wt-PTH group, Osx-positive osteoblasts were identified on all surfaces of trabeculae and endosteum throughout the tibia. PTH increased the total number of Osx-positive cells per tibial section by 3.2-fold in wt mice (Panel 2). ATF4 deficiency reduced the numbers of Osx-positive cells by 50 percent (Panel 3). Strikingly, although PTH slightly increased bone volume in 

\(\text{Atf4}^{2/2}\) bone (Panels 2 and 3), it failed to elevate the numbers of Osx-positive cells in \(\text{Atf4}^{-/-}\) tibiae. Similar results were obtained in calvariae (Fig. 6E). The IHC staining was highly specific since no signal was detected in the non-immune IgG control group (Panel 5).

Consistent with IHC results, as shown in Fig. 6I, Western blot analysis using protein extracts showed that PTH dramatically elevated the level of Osx protein in wt tibiae. In contrast, Osx was not detected by Western blot in extracts from \(\text{Atf4}^{2/2}\) animals (Fig. 6I). The level of Runx2 protein was slightly up-regulated by PTH in the wt group, but not in \(\text{Atf4}^{-/-}\) group. Unlike Osx, the basal level of Runx2 was not reduced by ATF4 deficiency.

\(\text{PTH1R}^{\text{protein}}\), the major receptor for PTH and PTHrP signaling in osteoblasts, was expressed in endosteal osteoblasts of tibiae (Fig. 6F) and periosteal osteoblasts of calvariae and hypertrophic chondrocytes in the growth plate area (unpublished data). The signal for PTH1R protein was weak in trabecular osteoblasts that actively form new bone (unpublished data). In contrast to results from a previous study showing that PTH1R is down-regulated by PTH in cultured osteoblasts \([46]\), PTH1R was slightly increased by intermittent PTH treatment in vivo as measured by IHC and Western blot analysis. Importantly, ATF4 deficiency did not reduce the level of PTH1R (Fig. 6F and I). In addition, primary calvarial osteoblasts from wt and \(\text{Atf4}^{2/2}\) mice displayed an identical cAMP accumulation curve in response to treatment of increasing concentrations of PTH in vitro. Taken together, these results indicate that the impaired anabolic response of skeleton to PTH observed in \(\text{Atf4}^{2/2}\) animals cannot be explained by a reduction in the level of PTH1R and/or cAMP production.

These results clearly demonstrate that: i) intermittent PTH stimulates the expression of Osx and, to a lesser extent,
Runx2, ii) PTH fails to stimulate Osx/Runx2 expression in the absence of ATF4, and iii) ATF4 is also required for basal level Osx expression.

Identification of a 132-bp ATF4-response element in proximal Osx promoter

Osx is not detected in Runx2 2/2 mice [35], indicating that Runx2 functions upstream of this factor and is essential for Osx expression. However, the results described above showed that ATF4 deficiency dramatically reduced the level of Osx protein without decreasing Runx2, suggesting that Runx2 is not sufficient for the maximal expression of Osx and that ATF4 has an important role in Osx expression. To define the mechanism whereby ATF4 regulates Osx, we examined the effect of ATF4 overexpression on Osx expression in MG-6 preosteoblast cells. As shown in Fig. 7A, ATF4 dose-dependently increased levels of Osx protein (top) and mRNA (bottom). We next examined whether ATF4 up-regulates Osx by increasing gene transcription by using a −1003/+68 mouse Osx promoter (Fig. 7B). Using COS-7 cells, which lack detectable Runx2, ATF4 had comparable activity to Runx2 in terms of its ability to activate promoter activity (approx. 1.8-fold). Together, ATF4 and Runx2 maximally activated the Osx promoter (3.2-fold induction). To further define the region of the Osx promoter necessary for ATF4 responsiveness, several constructs containing various deletion mutants of the mouse Osx promoter were transiently transfected into COS-7 cells with and without an ATF4 expression plasmid. Results showed that luciferase activity of both control and ATF4-transfected groups decreased with progressively larger 5’ deletions. However, ATF4 stimulation was abrogated when a 132-bp region between bp −215 to −83 was deleted (Fig. 7C). A putative ATF4-binding sequence (CTTCCTCA) at −201/−194 bp was identified in this
region by using a TRANSFAC retrieval program. Introduction of a 3-bp substitution mutation to this core sequence (from CTTCCTCA to CTTGtCA) completely abolished ATF4 activation (Fig. 7D). As shown in Fig. 7E, a DNA oligo probe from the Osx promoter that contains the TTACATCA core sequence bound to a factor(s) in nuclear extracts from COS-7 cells transfected with an ATF4 expression vector. Importantly, this binding (see arrow) was dramatically reduced by the addition of a specific antibody against ATF4 but not by normal control IgG or antibodies against cFos (an AP1 family member) or ATF2.

**P**H stimulation of Osx gene transcription requires an ATF4 response element

To study the mechanism whereby PTH regulates Osx, we next evaluated the effect of PTH on mouse -1003/+68 Osx promoter activity in MC-4 preosteoblast cells. As illustrated in Fig. 7F, PTH stimulated promoter activity in a dose-dependent manner with a detectable response seen at a PTH concentration of 10^{-8} M (significance at P<0.01). Measurable activation of the Osx promoter was observed 0.5 h after PTH addition with maximal induction occurring between 4–6 h (Fig. 7G). PTH-stimulated Osx protein expression was entirely blocked by PKA inhibition (Fig. 7H). The 132-bp ATF4-responsive element identified above was also required for PTH induction of promoter (Fig. 7I). Furthermore, the same 3-bp substitution mutation that abrogates ATF4 activation dramatically reduced PTH-dependent activation (Fig. 7J), indicating that this element is critical for the actions of PTH on this promoter.

**ATF4 is recruited to the endogenous Osx promoter in a PTH-dependent manner**

To determine whether ATF4 is associated with the endogenous Osx promoter in vivo, we performed chromatin immunoprecipitation (ChIP) assays using MC-4 cells with and without PTH treatment. As shown in Fig. 7K and L, ATF4 specifically interacted with a chromatin fragment of the proximal Osx promoter that contains the ATF4-binding site identified above. This interaction was not detected in primary calvarial osteoblasts from Atf4^-/- mice (data not shown). Furthermore, this interaction was dramatically stimulated by PTH treatment. Supporting our previous demonstration that the OSE1 site in the mOG2 promoter mediates PTH induction of the gene [31,34], ATF4 also bound to an ATF4-binding site (OSE1)-containing chromatin fragment of the proximal mOG2 promoter in a PTH-dependent manner (primers P3/P4). In contrast, ATF4 antibody failed to immunoprecipitate a 3' chromatin fragment in the transcribed region of the mOG2 gene that contains no ATF4-binding sites (primers P5/P6).

**Discussion**

Our goals in this study were: 1) to determine whether the bone-related transcription factor ATF4 plays a role in the anabolic effects of PTH in bone, and 2) if so, to define the relevant mechanisms. Our results clearly show that PTH-stimulated increases in osteoblast proliferation, volume of long bones, vertebrae, and calvariae as well as decreases in apoptosis are all dramatically reduced or completely abolished in Atf4-deficient mice. Equally importantly, PTH-induced bone in Atf4^-/- mice cannot mature due to a severe defect in osteoblast differentiation as manifested by a defect in Osx expression. Therefore, this study establishes a critical role for ATF4 in the anabolic actions of PTH in bone.

In agreement with results from previous studies [14,47], we find that intermittent PTH dramatically increases trabecular and cortical bone volume. This PTH response is achieved at least in part by stimulating the proliferation of osteoblasts and preosteoblasts and/or by inhibiting apoptotic death of osteoblasts and osteocytes in vivo [2,6,45,47]. Effects of PTH on osteoblast proliferation and survival can be reproduced in cultured osteoblasts [15,16,48]. Importantly, this study demonstrates that ATF4 plays a pivotal role in PTH stimulation of cell proliferation in osteoblasts and preosteoblasts and attenuation of apoptosis in osteoblasts and osteocytes in vivo. PTH can increase osteoblast proliferation at least in part through ATF4-mediated expression of cyclin D1 [33] because both factors are up-regulated by PTH in osteoblasts [15,34]. The mechanism whereby ATF4 blocks apoptosis in osteoblasts remains unknown. ATF4 deficiency also increases apoptosis in lens fiber cells in a p53-dependent manner. The embryonic lens in double homozygous p53/Atf4^-/- mice does not undergo apoptosis, which suggests possible involvement of p53 in this process [49,50].

Accumulating evidence supports the concept that, in addition to increasing osteoblast cell number, intermittent PTH also stimulates osteoblast differentiation [4,51–53]. In agreement with these results, the present study clearly demonstrates that: i) PTH dramatically increases the in vivo expression of osteoblast differentiation marker genes, including Ocn, Bsp, Alp, Opg, and Col1(I); and ii) PTH strikingly elevates the numbers of Osx-positive osteoblasts (i.e., mature or differentiating osteoblasts) and level of Osx protein as demonstrated by both IHC and Western blot analysis. Because PTH is able to stimulate the expression of many osteoblast differentiation marker genes in cultured osteoblast-like cells [12,30,31,54–57], it is likely that intermittent PTH also activates these genes in vivo via a similar molecular mechanism. Although part of the increased osteoblast activity in PTH-treated animals is likely explained by a PTH-dependent increase in...
Figure 6. PTH fails to promote osteoblast maturation/differentiation in the absence of ATF4. A–E, IHC analysis of Osx expression, sections of tibiae (A–D) and calvariae (E) were immunohistochemically stained using a specific antibody against Osx protein. The nuclei of Osx-positive cells (i.e., osteoblasts) were stained brown. The nuclei of preosteoblasts and other cells are stained blue. The total numbers of Osx-positive osteoblasts per tibial (G) or calvarial (H) section were counted under microscope. F, sections of tibiae were stained using an antibody against PTH1R protein. I, Western blot analysis, protein extracts were isolated from tibiae and analyzed for Osx, Runx2, and PTH1R proteins. *P < 0.05 (veh vs. PTH), †P < 0.05 (wt-veh vs. Atf4−/−-veh), ‡P < 0.05 (PTH/veh-wt vs. PTH/veh-Atf4−/−). J, cAMP assay, primary calvarial osteoblasts from 3-d-old wt or Atf4−/− mice were isolated, seeded at density of 5×10^4 on 96-well plate, and treated with vehicle or increasing concentrations of human recombinant PTH(1-34) for 5 min followed by measurement of cAMP.

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Figure 7. PTH activates Osx gene transcription via an ATF4-responsive element in the proximal Osx promoter. A, MC-4 cells were electroporated with indicated amount of ATF4 expression plasmid followed by Western blot. B, COS-7 cells were transfected with p1060mOsx-luc, pRL-SV40, and indicated expression vectors followed by dual luciferase assays. C, COS-7 cells were transfected with various deletion constructs and pRL-SV40 with and without ATF4 expression plasmid. D, COS-7 cells transfected with p215mOsx-luc or the same plasmid containing a 3-bp substitution mutation in the putative ATF4-binding site and pRL-SV40 with and without ATF4 expression plasmid. E, EMSA, labeled wild-type DNA probe was incubated with 2 μg nuclear extracts from COS-7 cells transfected with pCMV/ATF4 plasmid in the presence of normal control IgG (lane 3), ATF4 antibody (lane 4), cFos antibody (lane 5), and ATF2 antibody (lane 6). Experiments were repeated 3–4 times and qualitatively identical results were obtained. F and G, MC-4 cells transfected with p215mOsx-luc and pRL-SV40 were treated with indicated concentration of PTH for 6 h (F) or with 10^{-7} M PTH for indicated times (G). H, MC-4 cells were treated with and without 10^{-7} M PTH in the presence and absence of 10 μM of H89 for 6 h. I, MC-4 cells transfected as in Fig. 7C were treated with and without 10^{-7} M PTH for 6 h. J, MC-4 cells transfected as in Fig. D were treated with and without 10^{-7} M PTH for 6 h. K, a schematic illustration of putative ATF4 binding sites in the 5' flanking regions of the Osx and osteocalcin gene promoters and osteocalcin gene. L, ChiP assay of the Osx promoter in MC-4 cells treated with and without 10^{-7} M PTH for 6 h. *P<0.05 (β-gal vs. ATF4, Runx2, and ATF4 plus Runx2, or veh vs. PTH), **P<0.01 (ATF4 plus Runx2 vs. β-gal, ATF4, or Runx2).

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osteoblast proliferation and/or reduction of apoptosis, our studies suggest that PTH also increases osteoblast differentiation by rapidly up-regulating Osx.

Importantly, experiments from the current study show that effects of PTH on osteoblast differentiation are mediated by ATF4. Although PTH increased osteoblast proliferation in Atf4−/− animals as evidenced by significant increases in bone volume and nuclear BrdU labeling (Figs. 1–3), expression of osteoblast marker genes and numbers of Ocn-positive osteoblasts were either dramatically reduced or completely abolished in Atf4−/− animals. Furthermore, since the basal levels of osteoblast differentiation markers are either slightly reduced or not changed at all in Atf4−/− animals, the impaired differentiation response to PTH cannot be explained by a nonspecific blockage in osteoblast differentiation associated with ATF4 deficiency. Instead, ATF4 appears to have a unique role in the PTH-dependent component of osteoblast differentiation.

Further support for the concept that PTH actions are mediated by ATF4 comes from mechanistic studies. Specifically, we showed that PTH is a potent inducer of Osx expression in vivo and this response is completely abolished by ATF4 deficiency. Furthermore, PTH directly activated Osx gene transcription in cultured osteoblast-like cells, a response that required an ATF4 response element located between −201 and −194 bp in the proximal mouse Osx promoter. Introduction of a 3-bp substitution mutation into this ATF4-binding site essentially eliminated the PTH response. ChIP assays demonstrated that ATF4 binds to an endogenous chromatin fragment near the putative ATF4-binding site in the proximal Osx promoter in MC-3T3 cells. Of particular significance, ATF4 binding to the Osx promoter is dramatically enhanced by PTH. Collectively, these studies establish a unique role for ATF4 in PTH-mediated induction of Osx and osteoblast differentiation.

Runx2 is absolutely required for Osx expression, osteoblast differentiation, and bone formation [35,58]. Nevertheless, ATF4 deficiency dramatically reduced the level of Osx protein without altering Runx2, suggesting that Runx2 is not sufficient for maximal Osx expression. ATF4 stimulated Osx gene transcription in COS-7 cells that lack Runx2 protein to an extent similar to that seen with Runx2. In addition, an ATF4 response element was identified in the proximal Osx promoter of MC-4 cells. Of particular significance, ATF4 binding to the Osx promoter is dramatically enhanced by PTH. Collectively, these studies establish a unique role for ATF4 in PTH-mediated induction of Osx and osteoblast differentiation.

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It should be noted that ATF4 deficiency did not completely block the anabolic actions of PTH since this hormone increased bone volume in both growing and adult OVX bones from Atf4−/− mice (Figs. 1–3). This suggests that other factors must be involved for the PTH response. c-Fos [3], cAMP response element binding protein (CREB) [16], a major downstream target for PTH/cAMP and calcium signals, cAMP-response element modulator (CREM) [60], and Runx2 [16,23], a master regulator of osteoblast differentiation and bone formation, have all been shown to mediate components of the PTH anabolic response. Interestingly, these factors are either structurally related to ATF4 (c-Fos and CREM) or can interact with this factor (Runx2 and c-Fos) [59,61–65]. Most recently, low-density lipoprotein-related protein 6 (LRP6) [66], a major component of the Wnt signaling pathway, has been implicated in the anabolic actions of PTH in bone. It would be interesting to determine whether ATF4 mediates the PTH anabolic response via interactions with these factors or signaling pathways.

PTH signaling may regulate ATF4 via several mechanisms. First, PTH up-regulates Atf4 gene expression in cultured osteoblasts as demonstrated by our recent study [34] as well as in vivo (Fig. 5). Second, PTH post-translationally activates ATF4 via PKA [34], a major route for PTH signaling in osteoblasts. PKA phosphorylation of ATF4 at its Ser254 residue mediates β-adrenergic induction of Rankl mRNA expression in osteoblasts [67]. ATF4 can also be directly phosphorylated and activated by RSK2 [26], a growth factor-regulated serine-threonine protein kinase activated by the Ras-Mitogen-Activated Protein Kinase (MAPK) pathway. This phosphorylation is critical for ATF4 activity as well as bone formation [26]. Because PTH signaling activates Erk/MAPK [48], an immediate upstream activator of RSK2, ATF4 can be activated via the PTH-MAPK-RSK2 signaling pathway. Lastly, PTH promotes ATF4-Runx2 interactions which are critical for osteoblast function and bone formation [59,61,63]. This notion is supported by the fact that PTH up-regulates both factors in osteoblasts [23,34].

Based on findings from this and other studies, we proposed a working model for ATF4 to mediate PTH stimulation of osteoblast function and bone formation (Fig. 8). Binding of PTH to its receptor, PTH1R, activates PKA and probably other intracellular signaling pathways, leading to up-regulation/activation of ATF4. ATF4 subsequently increases proliferation of osteoblasts and/or preosteoblasts via modulation of cyclin D1 protein, and attenuates apoptotic death in osteoblasts and osteocytes, resulting in a significant increase in the numbers of osteoblasts and/or osteocytes. At the same time, ATF4 together with Runx2 maximally activates Osx expression and increases osteoblast differentiation. The resulting increases in osteoblast number and

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**Figure 8. Proposed model for ATF4 mediation of PTH stimulation of bone formation.** Binding of PTH to PTH1R activates PKA and leads to up-regulation of ATF4. ATF4 subsequently increases proliferation and survival of osteoblasts. At the same time, ATF4 together with Runx2 maximally activates Osx expression and increases osteoblast differentiation. These increases in osteoblast number and differentiation lead to massive bone formation. Osx also negatively regulates osteoblast proliferation, thus preventing excess bone formation.

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differentiation lead to bone formation. Osterix also negatively regulates osteoblast proliferation, thus preventing excess bone formation [68].

Since ATF4 is known to regulate the expression of RANKL in osteoblasts and thereby osteoclast differentiation [67,69], it will be interesting to determine if ATF4 is also required for the catabolic actions of PTH in bone.

In summary, this study establishes a critical role for ATF4 in the anabolic actions of PTH in bone. ATF4 is necessary for PTH to increase both osteoblast numbers and differentiation. Therefore, ATF4 may provide a potential new therapeutic target for improving bone mass and for treating metabolic bone diseases such as osteoporosis.

Materials and Methods

Reagents

Tissue culture media and fetal bovine serum were obtained from HyClone (Logan, UT). H89, DMSO, PTH1R antibody, mouse monoclonal antibody against β-actin were purchased from Sigma (St Louis, MO). Other reagents were obtained from the following sources: antibodies against ATF4 (for Western blot), Runx2, normal control IgGs and horseradish peroxidase-conjugated mouse or goat IgG from Santa Cruz (Santa Cruz, CA), and Osterix antibody from Abcam Inc. (Cambridge, MA). ATF4 antibody used for EMSA was raised against epitope QETN-KEPPQTVNPIGHLPESLIK (St Louis, MO). All other chemicals were of analytical grade.

Atf4-deficient mice

Breeding pairs of Atf4 heterozygous mice were described previously [34] and used to generate Atf4 wild-type (wt) (Atf4+/+), heterozygous (Atf4+/−) and homozygous mutant (Atf4−/−) mice for this study. All research protocols were approved by the Institutional Animal Care and Use Committee of the VA Pittsburgh Healthcare System, where this study was conducted.

In vivo PTH administration

For the “growing mouse model”, five-day-old mice were given daily subcutaneous injections of vehicle (saline) or hPTH(1–34) (60 ng/g body weight, Bachem, Torrance, CA) for 28 d. For the “adult OVX mouse model”, four-month-old female mice were anesthetized and ovariotomized as follows: A 1-cm midline incision was made through the skin. The “white line” will be visualized on the peritoneum and a second incision was made along the white line through the peritoneum. Using long, straight forceps, the left ovary was isolated and the connective tissue between the ovary and kidney was dissected away. Straight forceps with flat ends was used to pinch the uterine horn while another set of straight forceps was used to tear the ovary away from the uterine horn. The same was done on the right ovary. 6-0 PDS (Polydioxanone Sutures, 6/0) was used to tie around the uterine horn. The same was done on the right ovary. 6-0 PDS was used to tie around the uterine horn to provide hemostasis if necessary. 6-0 PDS was used to close the innermost layer. Sterile surgical staples were used to close the incision. Two months later mice were given daily subcutaneous injections of vehicle or hPTH (100 ng/g body weight) for 28 d. Mice were euthanized 24 h after last PTH injection. The effects of these PTH dosing regimens on bone were determined by both biochemical and histomorphometric criteria.

Gross evaluation and serum biochemistry

Body weight was recorded every other day. The length of the femurs was measured using an electronic digital caliper. Faxitron X-ray analysis of femurs was conducted at 27 kV and 7.5 seconds (Faxitron X-Ray Corp., Wheeling, IL). Femurs were ashed at 800°C for 4 h and weighed. Serum calcium and Pi concentrations were determined using kits from Pointe Scientific, Inc (Canton, MI) following the manufacture’s instructions (Sigma Diagnostics).

Bone morphometric analyses by micro-computerized tomography (μCT)

Upon termination of PTH or vehicle treatment, mice were sacrificed and femurs were isolated. Fixed non-decalcified femurs were used for μCT analysis at the Center for Bone Biology using a VIVACT40 (SCANCO Medical AG) following the standards of techniques and terminology recommended by American Society for Bone and Mineral Research [70]. For trabecular bone parameters, transverse CT slices were obtained in the region of interest in the axial direction from the trabecular bone 0.1 mm below the growth plate (bottom of the primary spongiosa) to the mid-femur. Contours were defined and drawn close to the cortical bone. The trabecular bone was then removed and analyzed separately. 3D analysis was then performed on trabecular bones slices. A 3-mm section was used to obtain mid-femoral cortical bone thickness. The analysis of the specimens involves the following bone measurements: bone volume fraction (BV/TV, %), trabecular number (Tb. N), trabecular thickness (Tb. Th), trabecular spacing (Tb. Sp), and cortical thickness (Cort. Th).

Histological evaluation

Tibiae, lumbar vertebrae (L5), and calvariae were fixed in PBS buffered 10% formalin at 4°C for 24 h, decalcified in 10% EDTA (pH 7.4) for 10–14 d, and embedded in paraffin. Longitudinal sections of tibiae and vertebrae were cut at 4 µm and stained with hematoxylin and eosin (H&E). Trabecular area of tibial sections was measured in the proximal metaphysis beginning immediately below the chondro-osseous junction to the mid-tibia. Calvariae were bisected perpendicular to the sagittal suture through the central portion of the parietal bones, parallel to lamboidal and coronal sutures, and embedded in paraffin to obtain sections of a standard area according to the method described by Zhao et al. [71]. Trabecular area versus total bone area was measured using an Image Pro Plus 6.2 software (Media Cybernetics, Inc, Bethesda, MD). The calvarial width was the average value from 20 random measurements of each calvaria (at least 6 samples per group) using a SPOT Advanced imaging software (provided with the purchase of the Olympus BX41 microscope).

Measurement of mineral apposition rate (MAR)

Mice were injected with calcine subcutaneously (20 mg/kg) at 6 and 2 d before sacrifice. Undecalcified tibia were fixed in 70% ethanol, embedded in methylmethacrylate and sectioned at 10 µm. Calcine labeling was visualized using a Nikon E800 fluorescence microscope. The metaphyseal trabecular bone projected into the marrow space was evaluated and the distance between the all double-labeled areas was measured at a magnification of 200x. MAR was calculated as mean distance between the double labels divided by the number of the days between the calcine injections. Histomorphometric analysis was performed using BioQuanti image analysis software (R&M Bio Metrics, Nashville, TN, USA).

In vivo proliferation assay

Mice were injected intraperitoneally with 100 µg bromodeoxyuridine (BrdU)/12 µg fluorodeoxyuridine (FdU) per gram of body weight 12 h before sacrifice. After sacrifice, sections of tibiae and calvariae were obtained. To identify actively proliferating cells,
nuclei that have incorporated BrdU were detected using a Zymed BrdU immunostaining kit according to the manufacturer’s instruction (Invitrogen, Carlsbad, CA). BrdU-positive cells (brown) on the calvarial periosteal surface or in the osteoid of tibiae were counted and normalized to the total numbers in the same area [33]. BrdU-positive hematopoietic cells in marrow were not counted.

In situ apoptosis detection
This assay is based on the classical TUNEL assay to examine apoptosis by detecting DNA fragmentation. 4-μm sections of tibiae were prepared and stained using the ApopTag Peroxidase In Situ Apoptosis Detection Kit according to the manufacturer’s instructions (Millipore, Billerica, MA). Apoptotic osteoblasts and osteocytes in tibiae were counted and normalized to the total cells from the same area.

Immunohistochemistry (IHC)
Tibiae and calvariae were fixed, decalcified, and embedded in paraffin. Sections of tibiae and calvariae were stained with antibodies against Osx (Abcam Inc, Cambridge, MA) and PTH1R (Sigma, St. Louis, MO) using the EnVision System-HRP (DAB) kit (Dako North America, Inc, Carpinteria, CA) according to the manufacturer’s instructions. Briefly, slides were baked at 55°C for 45 min, deparaffinized in three washes of xylene, and rehydrated in a decreasing ethanol gradient. Antigen retrieval was performed using 0.1% trypsin for 10 min at 37°C in a humidified chamber. Endogenous peroxidases were deactivated with 3% H2O2 in 1x PBS for 10 min, and sections were blocked in blocking solution for 30 min at room temperature. Sections were incubated with primary antibody (1:200 dilution for both Osx and PTH1R) in blocking solution for 2 hours at 4°C. Sections were washed in PBS three times and incubated with a donkey-anti-rabbit IgG-HRP secondary antibody solution for 30 min at room temperature. After washing with PBS three times, HRP activity was detected using a DAB substrate solution for 5 min at room temperature. Sections were counter-stained with a Mayer’s hematoxylin solution.

Measurement of plasma levels of IGF-1 and FGF-2 by ELISA
Blood plasma samples were prepared from whole-blood samples from mice of each group and plasma levels of IGF-1 and FGF-2 were measured by using ELISA kits (human FGF basic Quantikine Kit, cat #: DFB50, and mouse IGF-I Quantikine ELISA Kit, cat #: MG100, both from R&D Systems Inc, Minneapolis, MN) according to the manufacturer’s instructions.

cAMP assay
Primary osteoblasts from calvariae of 3-d-old wt or Afp−/− mice were isolated as described previously [72]. Cells were seeded at a density of 5 × 10^4/well on 96-well plate and treated with vehicle or increasing concentrations of human recombinant PTH(1-34) or increasing concentrations of human recombinant PTH(1-34) for 5 min. Cells were then lysed with lysis reagent 1B and cell lysates used for cAMP assay using a cAMP Biotrak Enzymeimmunoassay (EIA) kit (cat #: RPN225, GE Healthcare Biosciences Corp, Piscataway, NJ) according to the manufacturer’s instructions and the protein concentrations were measured using a BCA protein assay kit (Pierce). cAMP was normalized to total protein.

Quantitative real-time RT/PCR and Western blot analysis
RNA isolation, reverse transcription (RT), regular PCR, and quantitative real-time PCR analysis were performed as previously described [34]. The DNA sequences of mouse primers used for real-time PCR were summarized in Table 1. Western blot analysis was performed as previously described [33]. RNAs or protein extracts from at least six specimens in each group were used.

**DNA constructs and site-directed mutagenesis, transfection**

pCMV/β-gal, pCMV/ATF4, and pCMV/Ranx2 were previously described [62]. mOx-luc containing different mouse Osx promoter elements (−1003/+68, −567/+68, −215/+68, and −83/+68) driving a firefly luciferase reporter gene were constructed in the project laboratory by PCR subcloning promoter fragments using mouse tail DNA as a template into pGL3-luc vector (Promega, Madison, WI). Mutant p215mOsx-luc which contains a 3-bp substitution mutation in a putative ATF4-binding site at positions −198, −197 and −196 (from CTTCCTCA to CTTgtaGA) was generated from the wild-type p215mOsx-luc by PCR amplification using a QuickChange™ XL Site-Directed Mutagenesis Kit (Stratagene, La Jolla, CA) using the following primers: 5′-GGT ACC CCT CCG TCT CTC GGC TTT TaC ATT GGA TCC GGA GTC TTC TCC TCC GC-3′ (forward); 5′-GGG GAG AAG ACT CCG GAT CCA ATG tac AAG GCG AGA GAG GGA GGC GTT ACC-3′ (reverse). Sequence accuracy was confirmed by automatic DNA sequencing. For all transfection experiments, the amount of plasmid DNAs (reporter plasmid, 0.25 μg; normalization plasmid pRL-SV4, 10 ng; and expression plasmid, 1.0 μg) was balanced as necessary with β-galactosidase expression plasmid such that the total DNA was constant in each group. Experiments were performed in triplicates and repeated 3–4 times.

**Nuclear extracts preparation and electrophoretic mobility shift assay (EMSA)**
Nuclear extracts were prepared from COS-7 cells transfected with pCMV/ATF4 plasmids as previously described [73]. The DNA sequences of the oligonucleotides used for EMSA were as follows: GAT CCC TGC CCT CCT CCT CCT GTC CCT TCA TTG GAT CCG GAG TCT TC GAG TCG. DNA oligonucleotide was labeled using a Biotin 3′ end DNA Labeling Kit (cat #: 89818, Pierce Biotechnology Inc., Rockford, IL). Two μg of nuclear extracts and 20 fmol biotin-labeled DNA probe were incubated in 1x binding buffer for 30 min at room temperature.

**Table 1. real-time PCR primers.**

<table>
<thead>
<tr>
<th>Gene name</th>
<th>5′ primer</th>
<th>3′ primer</th>
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<td>Alp</td>
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<tr>
<td>Atf4</td>
<td>GAGCTTCCCGAAGGGCAGGAGTG</td>
<td>TGCCCACTCCCAAGATATGCATCATC</td>
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<td>Bip</td>
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<td>cFos</td>
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<td>cJun</td>
<td>GCCAAACATGGTGTCCAGAGCAATG</td>
<td>GCCACCTACGGCTAGCAGT</td>
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<tr>
<td>Col I(1)</td>
<td>AGATGGAGAAACATCCGCAACGGC</td>
<td>TCCAGATCTTGCTCCTTGCA</td>
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<tr>
<td>Gapdh</td>
<td>CAGTCGCCAGCTTCGGTGGTA</td>
<td>CTGCAAATGGGCCCCGTCCTT</td>
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<td>Ocn</td>
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<td>Opn</td>
<td>CCAATGGAACGTCAATCCACCA</td>
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doi:10.1371/journal.pone.0007583.t001
For supershift assay, 1 μg of IgG or indicated antibodies were first incubated with nuclear extracts prior to addition of DNA probe. Protein–DNA complexes were separated on 4% polyacrylamide gels in 1x TBE buffer, and transferred onto Biodyne B Nylon Membrane (cat #: 77016, Pierce). The membrane was blocked in 1x blocking buffer, washed five times with 1x wash buffer, and visualized by a Chemiluminescent Nucleic Acid detection Module (cat #: 89880, Pierce, Rockford, IL).

Chromatin immunoprecipitation (ChIP)

ChIP assays were performed using ATF4 antibody or control IgG as described previously [62]. PCR primer pairs (Table 2) were generated to detect DNA segments located near a putative ATF4-binding site (CTTCCTCAT) at −201/−193 (primers P1 and P2) determined by the TRANSFAC retrieval program in the 5' flanking region of the Otx promoter, a previously identified ATF4-binding site (OSE1) in osteocalcin gene 2 (mOG2) promoter (primers 3 and 4), and a mOG2 gene region (+177/+311) that contains no ATF4-binding sites (primers P5 and P6) [62]. PCR products were run on 3% agarose gel and stained with ethidium bromide. Purified input chromatin was used to perform parallel PCRs with the respective primer pairs.

Statistical analysis

Data was analyzed with a GraphPad Prism software (4.0). A one-way ANOVA analysis was used followed by the Tukey test. Student’s t-test was used to test for differences between two groups of data as needed. Data of Figs. 1, 2, 4, and 5 and S1-2 were from students’ one-way ANOVA analysis was used followed by the Tukey test. Data of Figs. 1, 2, 4, and 5 and S1-2 were from students’ one-way ANOVA analysis was used followed by the Tukey test. Data of Figs. 1, 2, 4, and 5 and S1-2 were from students’ one-way ANOVA analysis was used followed by the Tukey test. Data of Figs. 1, 2, 4, and 5 and S1-2 were from students’ one-way ANOVA analysis was used followed by the Tukey test.

Table 2. PCR primers used in ChIP assay.

<table>
<thead>
<tr>
<th>Oligo name</th>
<th>Sequence</th>
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<tr>
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<td>P2</td>
<td>GGCTCTCTCCGTCCTAGGG</td>
</tr>
<tr>
<td>P3</td>
<td>CACAGCATCTTTGAGTTAGC</td>
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<td>P4</td>
<td>TATGCTCCTCCGTCCTCTGA</td>
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<td>P5</td>
<td>TAGTGGACAGACTCCCGCGTA</td>
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<td>P6</td>
<td>TGAGCCCGCTTCTCAAGCAT</td>
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doi:10.1371/journal.pone.0007583.t002

References


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<tr>
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<th>Sham(+/+)</th>
<th>OVX(+/+)</th>
<th>Sham(-/-)</th>
<th>OVX(-/-)</th>
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</thead>
<tbody>
<tr>
<td>BV/TV(%)</td>
<td>12±1.2</td>
<td>3.4±0.5*</td>
<td>1.8±0.08†</td>
<td>2.0±0.12</td>
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<tr>
<td>Tb.N(/mm)</td>
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<td>1.5±0.06*</td>
<td>1.4±0.07†</td>
<td>1.3±0.01</td>
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<tr>
<td>Tb.Th(µm)</td>
<td>77.2±6.0</td>
<td>62.4±1.2*</td>
<td>59.3±11†</td>
<td>57.4±0.8</td>
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<tr>
<td>Tb.Sp(mm)</td>
<td>0.46±0.04</td>
<td>0.62±0.24*</td>
<td>0.71±0.03†</td>
<td>0.83±0.23</td>
</tr>
<tr>
<td>Cort.Th(mm)</td>
<td>0.2±0.05</td>
<td>0.21±0.01</td>
<td>0.17±0.01†</td>
<td>0.18±0.01</td>
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</table>

N = 3
Activating transcription factor 4 regulates osteoclast differentiation in mice

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Kenneth Patrene,1 G. David Roodman,1 and Guozhi Xiao1,7

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Activating transcription factor 4 (ATF4) is a critical transcription factor for osteoblast (OBL) function and bone formation; however, a direct role in osteoclasts (OCLs) has not been established. Here, we targeted expression of ATF4 to the OCL lineage using the Trap promoter or through deletion of Atf4 in mice. OCL differentiation was drastically decreased in Atf4−/− bone marrow monocyte (BMM) cultures and bones. Coculture of Atf4−/− BMMs with WT OBLs or a high concentration of RANKL failed to restore the OCL differentiation defect. Conversely, Trap-Atf4-tg mice displayed severe osteopetrosis with dramatically increased osteoclastogenesis and bone resorption. We further showed that ATF4 was an upstream activator of the critical transcription factor Nfatc1 and was critical for RANKL activation of multiple MAPK pathways in OCL progenitors. Furthermore, ATF4 was crucial for M-CSF induction of RANK expression on BMMs, and lack of ATF4 caused a shift in OCL precursors to macrophages. Finally, ATF4 was largely modulated by M-CSF signaling and the PI3K/AKT pathways in BMMs. These results demonstrate that ATF4 plays a direct role in regulating OCL differentiation and suggest that it may be a therapeutic target for treating bone diseases associated with increased OCL activity.

Introduction
Skeletal integrity requires a delicate balance between bone-forming osteoblasts (OBLs) and bone-resorbing osteoclasts (OCLs). Abnormal osteoclastogenesis results in bone destruction, such as osteoporosis, metastatic ostteolytic lesions, Paget disease of bone, and rheumatoid arthritis. In contrast, reduced osteoclastogenesis causes osteopetrosis, a disorder characterized by significantly increased skeletal mass and lack of a marrow space. Osteopetrosis is usually observed in animals or humans in which genes encoding cytokines, receptors, and signal transduction and transcription factors critical for OCL differentiation—such as RANKL; its receptor, RANK; the M-CSF receptor CSF1R (also known as c-Fms); TNF receptor-associated factor 6 (TRAF6); Src; PU.1, encoded by Spi1; p50/p52 NF-kB subunits; c-Fos; or v-ATPase V0 subunit—are deleted or mutated (1–10). Defining the molecular mechanisms underlying osteoclastogenesis is essential to advance the understanding of the molecular basis for the pathogenesis of bone diseases with altered OCL activity. This knowledge will be important for the prevention and treatment of these diseases.

OCLs originate from cells in the monocyte/macrophage lineage (11). OCL formation and maturation are tightly regulated by OBL/stromal cell/hypertrophic chondrocyte-derived factors such as M-CSF, RANKL, and osteoprotegerin (OPG), a soluble decoy receptor that blocks RANKL binding to RANK and thereby inhibits OCL differentiation (12–15). M-CSF binds to its receptor, CSF1R, on early macrophage progenitor cells and activates the Rank gene to generate OCL progenitors (16). The PI3K/AKT signaling pathways, which are strongly activated by M-CSF, play a critical role in activating OCL differentiation and bone resorption in normal and diseased states (17–24). However, little is known about the downstream molecular events that result from M-CSF–PI3K/AKT signaling and their relationship to osteoclastogenesis.

RANKL, a member of the TNF superfamily, binds to RANK on OCL precursors and recruits TRAF6, resulting in the activation of multiple signaling pathways including IKK complexes (IKKα, IKKβ, IKKγ, and NIK-IKKα) and MAPKs (Erk1/2, p38, and JNK) (2, 3, 25), which leads to activation of critical transcription factors such as NF-kB and c-Fos. RANKL activation of NF-kB and c-Fos results in induction of initial expression of the key transcription factor NFATc1 (also designated as NFAT2 or NFATc), which is activated by the Ca2+/calmodulin-regulated phosphatase calcineurin (3, 26–29). Eventually, calcium signaling occurs and activates the existing NFATc1, which triggers NFATc1 autoamplification (3, 30) required for further OCL differentiation. Inactivation of the Nfatc1 gene in vivo is embryonic lethal due to a heart valve defect (31, 32). However, the lack of rescue of osteopetrosis in OCL-deficient c-Fos−/− mice by adoptive transfer of Nfatc1−/− (as opposed to Nfatc1−/−) hematopoietic stem cells provided evidence that NFATc1 is essential for osteoclastogenesis in vivo (33). Accordingly, transgenic overexpression of a constitutively active form of NFATc1 (caNFATc1) results in a skeletal phenotype of osteopetrosia associated with increased osteoclastogenesis and bone resorption (34). Importantly, overexpression of NFATc1 activates osteoclastogenesis in the absence of RANKL (29, 35, 36). In addition, cTnFt can restore osteoclastogenesis in both the p50/p52 double knockout (37) and the c-Fos knockout mouse (38). These results suggest that the major OCL differentiation signals converge on NFATc1. However, the molecular mechanisms that control the expression of this key factor are not completely understood.

Critical roles for ATF4 in OBLs and bone formation are well established. ATF4 favors bone formation by promoting OBL-specific gene expression, amino acid import, and the synthesis of type I...
collagen, and proliferation and survival of OBLs (39, 40). Furthermore, our most recent work showed that ATF4 is critical for osteoblastic responses to parathyroid hormone (PTH) to increase bone formation (41). Elefteriou and coworkers showed that ATF4 mediates β-adrenergic induction of Rankl mRNA expression via direct binding to the upstream OSE1 site in the Rankl promoter in OBLs (42). Work from the same group further showed that OBL-targeted expression of ATF4 increased osteoblastic Rankl expression and thereby OCL differentiation (43). Although these studies clearly demonstrate that increased OBL expression of ATF4 enhances OCL differentiation via RANKL production in OBLs, to our knowledge, the possibility of an OCL-intrinsic direct role for ATF4 in regulating OCL differentiation has not previously been addressed.

Using biochemical, cellular, and genetic approaches, the present study demonstrates that ATF4 is an osteoclastic transcription factor, which we believe to be novel, that is essential for OCL differentiation. We showed that both in vitro and in vivo OCL differentiation was severely impaired by lack of ATF4 in a cell-autonomous manner and increased by OCL-targeted transgenic ATF4 expression. Our results revealed that, mechanistically, ATF4 functioned as a direct upstream activator of the gene encoding the critical transcription factor NFATc1. Further, ATF4 modulated RANKL activation of MAPK pathways, a key molecular event in OCL differentiation. Additionally, we demonstrated that ATF4 was critical for M-CSF induction of RANK expression, a key step to generate OCL progenitors, and the level of ATF4 protein was largely modulated by M-CSF and the PI3K/AKT pathways in BMMs. Therefore, these results indicate that ATF4 has important OCL-intrinsic functions both upstream and downstream of RANKL signaling during OCL differentiation.

Results
Inactivation of the Atf4 gene severely impairs OCL differentiation in vitro and in vivo in a cell-autonomous manner. We first tested whether ATF4, a previously known OBL-enriched transcription factor (39), is expressed in OCL-like cells. As shown in Supplemental Figure 1 (supplemental material available online with this article; doi:10.1172/JCI42106DS1), the levels of ATF4 protein in primary mouse BMMs and RAW264.7 cells (a mouse monocyte/macrophage cell line) were comparable to those in OBL-like cells (mouse
MC-4 preosteoblastic cells, primary mouse bone marrow stromal cells [BMSCs], and rat UMR106-01 osteoblastic cells). As shown in Figure 1A, ATF4 protein was phosphorylated in primary mouse BMMs (note loss of upper bands with phosphatase treatment). Furthermore, a strong ATF4 signal was detected in the cytoplasm of the large multinuclear OCLs by immunohistochemical (IHC) staining using a specific ATF4-antibody (Figure 1B).

To determine whether ATF4 is required for osteoclastogenesis in vivo, the tibiae of 4-week-old WT and Atf4−/− mice were decalcified, and histological sections were stained for the OCL enzyme tartrate-resistant acid phosphatase (TRAP). We found that TRAP activity throughout the tibiae, including both the metaphyseal and the epiphyseal regions, was dramatically reduced in Atf4−/− compared with WT mice (Figure 1C). We next measured the effect of ATF4 deficiency on OCL differentiation in both primary and secondary spongiosa. OCL surface/bone surface (Oc.S/BS) and OCL number/bone perimeter (Oc.Nb/BPm) were reduced similarly in both primary and secondary spongiosa of tibiae in Atf4–/– mice (Figure 1D and Table 1). Given the dramatic decreases in Oc.S/BS and Oc.Nb/BPm in the Atf4−/− bone, we next determined whether ATF4 is intrinsically required in BMMs for OCL differentiation by assessing whether OCL differentiation was normal upon addition of exogenous RANKL to Atf4−/− compared with WT BMM cultures in vitro by measuring the number of TRAP+ multinucleated cells (MNCs; defined as having 3 or more nuclei per cell) generated by each. We found that TRAP+ MNCs in BMM cultures from Atf4−/− mice were dramatically reduced compared with those from WT mice (Figure 1, E and F). Furthermore, the number of nuclei per MNC was decreased by 75% in Atf4−/− versus WT BMM cultures (Figure 1G), and the MNCs that formed in Atf4−/− BMM cultures were much smaller than those formed in WT cultures. Similar results were obtained in purified CD11b+ BMM cultures (Supplemental Figure 2, A and B). The ability to form TRAP+ MNCs was almost completely lost in BMM cultures from 15-month-old Atf4−/− mice (Supplemental Figure 2, C and D), which suggests that ATF4 is even more important for OCL differentiation in old animals. Furthermore, the resorption pit area on dentin slices was dramatically reduced in Atf4−/− versus WT BMM cultures (Figure 1, H and I). Although MNC number and size were both dramatically reduced in Atf4−/− BMM cultures relative to WT cultures, the ratio of total resorption pit area to total TRAP+ MNC area per slice was not significantly different in WT and Atf4−/− BMM cultures (WT, 0.46 ± 0.11; Atf4−/−, 0.56 ± 0.05), which suggests that the bone-resorbing activity of Atf4−/− TRAP+ MNCs is not impaired. Time-course experiments showed that the percentage of TRAP+ mononuclear OCLs was dramatically reduced in Atf4−/− versus WT BMM cultures in the first 4 days in OCL differentiation media. At day 5, this difference disappeared (Figure 1J), suggestive of a substantial delay rather than absolute decrease in early OCL differentiation. However, we noted that although the number of Atf4−/− and WT TRAP+ mononuclear cells was equivalent at day 5, the Atf4−/− cells did not go on to efficiently form MNCs at day 9 as described above (Figure 1, E–G). To further investigate the delayed early OCL differentiation, we assessed the in vitro formation of CFU-GMs, which are

<table>
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<tr>
<th>Table 1</th>
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<td><strong>Bone histomorphometry in WT and Atf4−/− tibiae</strong></td>
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<tr>
<td><strong>WT</strong></td>
</tr>
<tr>
<td><strong>Primary spongiosa</strong></td>
</tr>
<tr>
<td>Oc.S/BS</td>
</tr>
<tr>
<td>Oc.Nb/BPm</td>
</tr>
<tr>
<td><strong>Secondary spongiosa</strong></td>
</tr>
<tr>
<td>Oc.S/BS</td>
</tr>
<tr>
<td>Oc.Nb/BPm</td>
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Oc.S/BS and Oc.Nb/BPm in primary and secondary spongiosa of tibia in Figure 1C were measured as described in Methods. *P < 0.01 versus WT.
known to contain the earliest OCL precursors (44), and found that it was severely compromised in Atf4−/− BMMs (Figure 2, A and B). We next examined whether coculture with WT OBLs rescues the OCL formation of Atf4−/− BMMs (Figure 2, C and D). As expected, cocultures of WT OBLs with WT BMMs significantly induced the formation of TRAP+ MNCs. Although WT OBLs did induce some TRAP+ mononuclear OCLs in coculture with Atf4−/− BMMs, there were almost no TRAP+ MNCs observed. Furthermore, primary calvarial OBLs from 3-day-old Atf4−/− mice failed to induce TRAP+ MNC formation when cocultured with either WT or Atf4−/− BMMs (H. Cao and G. Xiao, unpublished observation), in support of the notion that osteoblastic ATF4 plays a role in OCL differentiation, probably via upregulation of RANKL expression (42). In addition, high concentrations of RANKL (up to 200 ng/ml) did not restore the OCL differentiation defect in BMMs or purified CD11b+ BMMs of Atf4−/− mice (Figure 2, E and F). Collectively, these findings suggest that ATF4 deficiency impairs osteoclastogenesis in a cell-autonomous manner.

OCL-targeted overexpression of ATF4 dramatically increases OCL differentiation and bone resorption and results in a severe osteopenic phenotype. To further examine the OCL-intrinsic role of ATF4 in regulating osteoclastogenesis in vivo, we developed transgenic mice in which the Atf4 transgene is driven by an 1,846-bp mouse Trap promoter that selectively expresses ATF4 in OCLs (referred to herein as Trap-Atf4-tg mice; Figure 3A). This promoter has been successfully used to target OCL expression of many transgenes (45–47). Trap-Atf4-tg mice were stained for TRAP activity (Figure 3B). TRAP+ OCLs (arrows) on trabecular surfaces of WT and Trap-Atf4-tg tibiae. Oc.S/BS and Oc.Nb/BPm values for primary and secondary spongiosa are shown in Table 2. (G) μCT analysis. Fixed nondemineralized femurs from 3-month-old male WT and Trap-Atf4-tg mice were used for μCT analysis as previously described (41). BV/TV, Tb.N, and Tb.Sp values are shown in Table 3. n = 3–7. *P < 0.01 versus WT. Original magnification, ×100 (D and E); ×200 (F).
transgenic mRNA was highly expressed in RANKL-differentiated BMMs, but was minimal in undifferentiated BMMs, calvarial OBLs, or BMSCs (Figure 3A). As shown in Figure 3B, the level of ATF4 protein was dramatically increased in RANKL-differentiated BMM cultures from Trap-Atf4-tg mice compared with BMM from the control littermates. Transgenic ATF4 dramatically increased the protein levels of NFATc1/A, but not of PU.1 or CSF1R1, in differentiated BMM cultures. The levels of OCL differentiation marker gene mRNAs (Trap, Rank, Cat K, and Mmp9) were all dramatically elevated in differentiated BMM cultures from Trap-Atf4-tg mice compared with those from WT mice. In contrast, like the proteins, the levels of Spi1 and Csfr1 mRNAs were not increased by transgenic ATF4 (Figure 3C). Using BMMs from 3 different transgenic lines, we found that OCL-targeted overexpression of ATF4 dramatically increased the number of TRAP⁺ MNCs in vitro (WT, 56 ± 11 TRAP⁺ MNCs/well; Trap-Atf4-tg, 270 ± 17 TRAP⁺ MNCs/well; P < 0.01; Figure 3D). We found a similar effect of transgenic ATF4 expression in vivo. TRAP activity was markedly increased in Trap-Atf4-tg compared with WT tibiae (Figure 3E). Oc.S/BS and Oc.Nb/BPm in both primary and secondary spongiosa of tibiae were dramatically increased in Trap-Atf4-tg versus WT mice (P < 0.01; Figure 3F and Table 2). The serum level of C-telopeptide (CTX), an indicator of in vivo bone turnover, was increased in Trap-Atf4-tg mice compared with WT (Figure 4A and Supplemental Figure 3). As shown in Figure 4G, introduction of a 4-bp substitution mutation to the known API binding site located at –644/–637 (from TGACTTC A to TGCGAAC A) decreased ATF4 activation by 50% to 25 ± 1.8 (WT) and 12 ± 2.4 (Trap-Atf4-tg) (Figure 4H). Consistent with results from previous studies (33, 38), both c-Fos and NFATc1 itself were also recruited to the same region of the Nf1 promoter in a RANKL-dependent manner. These data demonstrate that ATF4 is a critical upstream activator of the Nf1 gene and indicate that ATF4 not only regulates the number of OCL progenitors, but also has a direct role in activating genes downstream of RANK signaling.

We next examined whether NFATc1 can rescue the defective OCL differentiation of Atf4−/− BMMs. WT and Atf4−/− BMMs were infected with increasing amounts of retrovirus expressing a constitutively active form of NFATc1 (50) and differentiated for 7 days, followed by TRAP staining. Although cNFATC1-dose-dependently increased the number of TRAP⁺ MNCs in Atf4−/− BMM cultures (Figure 4I), surprisingly, at even the highest dose, it only slightly increased the number of nuclei per MNC (WT plus empty virus, 24 ± 3.8; WT plus NFATc1 virus, 32.6 ± 5.5; P < 0.01; KO plus empty virus, 3.3 ± 0.42; KO plus NFATc1 virus, 4.8 ± 0.71; P < 0.01).

**ATF4 modulates RANKL activation of MAPKs, but not IκBα pathways, in OCL progenitors.** Because activation of the MAPK and IκBα/NF-κB pathways by RANKL is crucial for NFATc1 expression and OCL differentiation, we next examined whether ATF4 deficiency affects NFATc1 expression.

**Table 2**

<table>
<thead>
<tr>
<th>Primarily spongiosa</th>
<th>WT</th>
<th>Trap-Atf4-tg</th>
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</thead>
<tbody>
<tr>
<td>Oc.S/BS</td>
<td>9.1 ± 0.36</td>
<td>16 ± 0.71a</td>
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<tr>
<td>Oc.Nb/BPm</td>
<td>9.2 ± 1.2</td>
<td>16 ± 0.94a</td>
</tr>
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</table>

**Secondary spongiosa**

| Oc.S/BS             | 9.2 ± 0.51 | 16.5 ± 0.72a |
| Oc.Nb/BPm           | 9.3 ± 1.03 | 16.6 ± 1.33a |

Oc.S/BS and Oc.Nb/BPm in both primary and secondary spongiosa of tibiae in Figure 3E were measured as described in Methods. *AP < 0.01 versus WT.*

**Table 3**

<table>
<thead>
<tr>
<th>Quantitative μCT analysis of WT and Trap-ATF4-tg femur histomorphometric parameters</th>
</tr>
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<tbody>
<tr>
<td><strong>WT</strong></td>
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<tr>
<td>BV/TV (%)</td>
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<tr>
<td>Tb.N (per mm)</td>
</tr>
<tr>
<td>Tb.Sp (mm)</td>
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</table>

Fixed nondemineralized femurs from 3-month-old male WT and Trap-ATF4-tg animals were used for μCT analysis as previously described (41).
RANKL activation of these important pathways in OCL progenitors. As shown in Figure 5, RANKL rapidly induced the phosphorylation of Erk1/2 in a time-dependent manner, which was delayed and reduced in Atf4−/− cells. ATF4 deficiency also slightly reduced the basal phosphorylation level of Erk1/2. Lack of ATF4 similarly compromised RANKL activation of p38 and JNK without markedly affecting their basal levels. In contrast, no difference was seen in RANKL-induced phosphorylation of IκBα in cells of the 2 genotypes. Collectively, lack of ATF4 reduced the ability of RANKL to activate the Erk1/2, p38, and JNK MAPK pathways in OCL progenitors.

We next determined whether ATF4 deficiency affects M-CSF-dependent signaling in BMMs. As shown in Supplemental Figure 5, M-CSF rapidly stimulated the phosphorylation of AKT and JNK in WT cells, which was slightly reduced in Atf4−/− cells. The levels of both phosphorylated and total Src were similar in WT and Atf4−/− cells with or without M-CSF. In contrast to RANKL, M-CSF similarly activated Erk1/2 and p38 in WT and Atf4−/− cells. Thus, ATF4 deficiency did not dramatically impact M-CSF signaling in BMMs.

Levels of ATF4 protein in BMMs are modulated by M-CSF and PI3K/AKT, and ATF4 is required for M-CSF induction of RANK expression. To determine whether ATF4 is regulated by M-CSF in early OCL differentiation, BMMs were cultured in the presence and absence of 30 ng/ml M-CSF for 0, 6, 12, 24, and 48 hours, followed by Western blot for ATF4. The results showed that the level of ATF4 protein was dramatically reduced in the absence of M-CSF in a time-dependent manner. However, this reduction was completely prevented by M-CSF (Figure 6A). M-CSF did not alter the level of Atf4 mRNA (Figure 6B), which suggests that a posttranscriptional mechanism is involved in this regulation.

To define the signaling pathways through which M-CSF regulates ATF4, BMMs were treated with and without inhibitors or activators for various pathways in the presence of M-CSF for 24 hours. As shown in Figure 6C, LY294002, a specific inhibitor of the PI3K/AKT pathway, dramatically reduced total and phosphorylated ATF4. In contrast, the p38 inhibitor SB203580, the Erk1/2 inhibitor U0126, the PKA inhibitor H89, the PKC inhibitor GF109203X, and the PKA activator FSK did not markedly decrease or increase ATF4 or
alter its phosphorylation. Importantly, exposure to LY294002 for only 24 hours prior to the addition of differentiation media inhibited in vitro OCL differentiation in a dose-dependent manner (Figure 6D). The concentrations of the inhibitors or activators used in this study are in the ranges previously reported to selectively affect M-CSF–induced RANK expression is essential for generating OCL progenitors. We next evaluated whether ATF4 is required for this regulation. To this end, highly purified CD11b+ BMMs were cultivated in M-CSF–containing medium for 72 hours and stained with an anti-RANK antibody or control IgG. As shown in Figure 6, G and H, the RANK signal was dramatically decreased in Atf4−/− relative to WT cells, as measured by both IHC and Western blot using specific antibodies. M-CSF time-dependently induced Rank mRNA expression in WT BMM cultures (Figure 6I), consistent with results from a previous study (16). However, this induction was dramatically reduced in Atf4−/− cells. Taken together, these results suggest that ATF4 regulates early OCL differentiation at least in part by facilitating M-CSF induction of the Rank gene.

ATF4 deficiency results in increased CD11b+ cells in bone marrow and spleen and reduced CD3− CD45R− CD11b−/lo c-kit+CD115hi cells in bone marrow. Because OCLs and macrophages share the same precursor, we next determined whether lack of ATF4 affects monocyte/macrophage precursors by measuring the CD11b+ cell population in splenocytes and bone marrow from WT and Atf4−/− mice. In 5 independent experiments, the percentage of CD11b+ cells was dramatically increased in Atf4−/− splenocytes compared with WT cells, as measured by flow cytometry (WT, 3.04% ± 0.64%; KO, 9.6% ± 2.1%; P < 0.01; Figure 7A). This increase in CD11b+ cells was specific, because the percentages of both T lymphocytes (CD3+; Figure 7A) and dendritic cells (CD11c+; Supplemental Figure 6, A and B) were not increased by ATF4 deficiency (WT, 27.63% ± 1.9%; CD3+; 1.59% ± 0.06%; KO, 21.04% ± 4.1%; CD3+; 1.52% ± 0.05%; P > 0.05). Likewise, in bone marrow, the number of CD11b+ cells was similarly increased in Atf4−/− mice (WT, 41.5% ± 0.08%; KO, 63.6% ± 3.4%; P < 0.01), but the number of CD3+ cells (WT, 1.04% ± 0.2%; KO, 0.8% ± 0.06%; P > 0.05) and CD11c+ cells (Supplemental Figure 6C) was not. At the same time, the percentage of CD11b+ cells, the osteoclastogenic population in the bone marrow, was dramatically reduced in Atf4−/− marrow and spleen. Since the CD3− CD45R− CD11b−/lo c-kit+CD115hi population (approximately 2% of fresh murine bone marrow preparations) contains the highest in vitro osteoclastogenic activity (55), we next determined whether ATF4 inactivation affects this OCL precursor population in bone marrow cells and splenocytes. In 5 independent experiments, the percentage of this cell population was slightly but significantly reduced in bone marrow cells by ATF4 deficiency (WT, 2.3% ± 0.11%; KO, 2.0% ± 0.02%; P < 0.01; Figure 7A). However, the percentage of this cell population in splenocytes was much lower than in bone marrow (approximately 0.25%) and was not significantly changed by ATF4 deficiency. It should be noted that although Atf4−/− bones are smaller and thinner than WT bones, total nucleated bone marrow cells per bone were not reduced in Atf4−/− compared with WT mice (Supplemental Figure 6D). ATF4 deficiency did not alter the proliferation and survival of CD11b+ BMMs (Figure 7, B–E). Therefore, the lack of...
ATF4 causes a lineage shift between OCLs and macrophages, resulting in an increase in macrophages. Interestingly, a similar increase in macrophages was observed in mice lacking c-Fos, whose deficiency also reduces NFATc1 and OCL differentiation (7–9).

Discussion
The results of our present study establish, for the first time to our knowledge, that ATF4, a transcription factor previously shown to be important in OBLs, also plays a direct and critical role in regulating OCL differentiation both in vitro and in bones. Drastically reduced OCL differentiation in BMM cultures from Atf4–/– animals was not rescued by coculture with WT OBLs or high concentrations of RANKL, which suggests that ATF4 plays an intrinsic role in OCLs that is indispensable for RANKL-induced OCL differentiation. Dramatic reduction in the formation of CFU-GMs in Atf4–/– BMM cultures suggests that ATF4 deficiency impairs the formation of OCL precursors. To examine the in vivo actions of ATF4 in OCLs, this study used the mouse Trap promoter to drive expression of ATF4 in OCLs. Using this approach, ATF4 was shown to stimulate expression of NFATc1 and other OCL-specific genes and OCL differentiation in BMM cultures. Of particular significance, Trap-Atf4-tg animals displayed a striking in vivo effect on OCL differentiation and bone resorption, resulting in a severe osteopenic phenotype. Because the Trap promoter is not active in OBLs, we were able to discriminate between the effects of ATF4 in OCLs and those in OBLs or BMSCs (i.e., via ATF4-dependent production of RANKL). Furthermore, as shown in Figure 1C and Figure 3E, TRAP activity in hypertrophic chondrocytes close to primary spongiosa was very weak, which suggests that the Trap promoter is not active in these cells. Therefore, Atf4 transgene expression driven by this promoter should be low in the hypertrophic chondrocytes; consequently, the potential contribution of the Atf4 transgene expression in these cells to the observed bone phenotype in Trap-Atf4-tg mice (i.e., osteopenia, increased OCL differentiation, and bone resorption) should be minimal.
One striking result in the present study, which we believe to be novel, is the finding that ATF4 is a direct upstream activator of the \textit{Nfatc1} gene, a master regulator of OCL differentiation. Levels of NFATc1 were drastically reduced in \textit{Atf4}\textsuperscript{−/−} OCLs and bones. Conversely, OCL-specific expression of ATF4 in transgenic mice greatly increased the expression of NFATc1 and its downstream target genes as well as OCL differentiation. ATF4 activated \textit{Nfatc1} gene transcription via interaction with the P1 promoter. Interestingly, a previously known API binding site located at −644/−637 was critical for ATF4 activation of the \textit{Nfatc1} P1 promoter, which suggests that ATF4 directly binds to this site and/or functions via interactions with API factors. Future study will differentiate among these possibilities. Of particular significance, ATF4 interaction with the P1 promoter was stimulated by RANKL. In addition to its direct regulation, ATF4 may also indirectly increase NFATc1 expression by promoting RANKL-RANK signaling. This notion is supported by our findings that (a) M-CSF induction of RANK, the receptor for RANKL in OCL precursors, was severely impaired in \textit{Atf4}\textsuperscript{−/−} BMMs; and (b) RANKL activation of the Erk1/2, JNK, and p38 MAPK pathways in OCL progenitors, which is crucial for the subsequent expression of NFATc1, was greatly compromised by the lack of ATF4. Although ATF4 is essential for its expression,
retroviral transduction of NFATc1 cDNA into BMM only partially rescued the OCL differentiation defect in Atf4−/− BMM cultures, as large OCLs were still not formed. Possible explanations include: (a) the magnitude and duration of NFATc1 expression was inappropriate for full rescue; and (b) other OCL differentiation–related genes and/or factors are also regulated by ATF4.

Our results established that ATF4 is a critical downstream target of M-CSF–PI3K/akt signaling in early OCL differentiation. The level of ATF4 protein was largely dependent upon the presence of M-CSF and the PI3K/akt pathway in BMMs. M-CSF upregulated ATF4 at least in part by stabilizing its protein, which involves the PI3K/akt pathway. Strikingly, short-term pharmacologic inhibition of the PI3K/akt pathway dramatically reduced the level of ATF4 protein in undifferentiated BMMs and subsequent OCL differentiation. These results strongly suggest that the M-CSF–PI3K/akt–ATF4 axis identified in the present study plays a crucial role in regulating early OCL differentiation. PI3K/akt signaling has been implicated in OCL activation and bone resorption in neurofibromatosis type I (NF1), a congenital disorder resulting from loss of function of the tumor suppressor gene NFI, which encodes neurofibromin, a GTPase-activating protein for Ras. NFI patients have a significantly higher incidence of osteoporosis and osteopenia (21–23). Recent studies showed that increased OCL activity and osteoporosis is caused by increased AKT signaling in OCLs in murine and human NFI haploinsufficiency (18, 19). It would be interesting to test whether ATF4 plays a role in the AKT-induced OCL differentiation and bone resorption in NF1 patients. Notably, mice selectively lacking NFI in OBLs displayed increased OCL activity, probably via upregulation of ATF4 in OBLs, and thereby exhibited increased RANKL expression (43).

Results from the present study demonstrated that ATF4 is not required for early OCL lineage commitment and development. First, CSFR1 and PU.1 expression, which is required for the generation of the common progenitors for both macrophages and OCLs, was not altered in Atf4−/− BMMs or bones. Second, M-CSF activation of AKT and MAPKs was not markedly different in WT and Atf4−/− BMMs. Third, both proliferation and survival of highly purified CD11b+ BMMs of the 2 genotypes were similar. Finally, the number of macrophages was increased in Atf4−/− animals. Therefore, ATF4 deficiency impairs OCL, but not macrophage, differentiation, and the OCL differentiation defect in Atf4−/− mice occurs later than in Spi1- or Csf1r-deficient mice.

Although ATF4 is crucial for OCL differentiation, Atf4−/− mice did not display an osteopetrosic phenotype, which is often observed in mice and humans in which genes encoding key OCL-regulating factors—such as CSFR1, RANK, RANKL, TRAF6, Src, PU.1, NF-κB (p50/p52 double knockout), and c-Fos—are inactivated or mutated (1–9). Lack of osteopetrosis in Atf4−/− mice could be explained, at least in part, by the fact that ATF4 is also important for OBL function and bone formation (39, 40, 56). Therefore, it is expected that Atf4−/− mice have a low–bone turnover osteoporosis.

ATF4 can be specifically regulated by different signal transduction factors in OBLs and OCLs. For examples, ATF4 is directly phosphorylated and activated by RSK2 in OBLs, which is critical for ATF4 activity as well as bone formation (56). Furthermore, PTH—via its receptor, PTHR1, which is expressed in OBLs—upregulates Atf4 gene expression and activity, which is required for this hormone to induce osteocalcin gene expression as well as bone formation (41, 53). PKA phosphorylation of ATF4 at its Ser254 residue mediates β-adrenergic induction of RANKl mRNA expression in OBLs (42). Additionally, ATF4 increases OBL function and bone formation through interactions with the OBL-specific transcription factor Runx2 (57–60). In contrast, in OCLs, ATF4 is regulated by M-CSF and the PI3K/AKT pathway, mediates M-CSF induction of RANK, and facilitates RANKL induction of MAPKs and expression of NFATc1. In contrast to the effects of M-CSF and PI3K/AKT on ATF4 protein levels in BMMs and OCLs, both RSK2 and PKA do not affect the ATF4 protein level in OBLs (42, 56). Furthermore, ATF4Ser219 and ATF4Ser224 phosphorylation is involved in binding to βTrCP ubiquitin ligase, thus increasing proteolytic degradation in HeLa cells (61, 62), whereas the activation of AKT in BMMs enhances ATF4 stability. This suggests that ATF4 is modulated by differential phosphorylations in OBLs and OCLs and/or that M-CSF–PI3K/akt signaling regulates a protein that modulates ATF4 protein stability in BMMs/OCLs. Therefore, it should be possible to design therapeutic agents that selectively inhibit ATF4 in OCLs.

The results of the present study establish a direct and important role for ATF4 in regulating multiple steps in OCL differentiation and suggest that manipulation of ATF4 in OCLs may be a strategy for increasing bone mass in diseased states.

Methods

Reagents. Tissue culture media and FBS were obtained from Thermo Scientific HyClone. LY294002, SB209580, U0126, H89, FSK, GF109203X, and DMSO were purchased from Sigma-Aldrich. Calf intestinal phosphatase and calf liver alkaline phosphatase (CIP) were purchased from Promega. All other chemicals were of analytical grade.

Act4-deficient and Trap-Atf4-tg mice. Breeding pairs of Act4-heterozygous mice (Swiss black), as described previously (40, 53), were used to generate WT Atf4−/−, heterozygous Atf4+/−, and homozygous mutant Atf4−/− mice for this study. 4- to 8-week-old mice were sacrificed for BMMs. Mice selectively expressing ATF4 in OCLs were developed at the Transgenic & Chimeric Mouse Facility of the University of Pittsburgh using an 1,846-bp mouse Atf4 cDNA sequence (5′-ATGACCCACCTGGAGTTAGTTGACA-3′) and a 3′ primer in the bGH-Poya sequence (5′-GCAGTGGGGGAGGTCACA-3′). Expression of Atf4 transgene was measured by Western blot analysis using ATF4 antibody. 5 founders (animal no. 2330, 2333, 2335, 2359, and 2360) were found to express Atf4 transgene at different levels. These transgenic animals were in C57BL/6 background. All research protocols were approved by the Institutional Animal Care and Use Committee of the VA Pittsburgh Healthcare System, where this study was conducted.

In vitro OCL assays and serum CTX assay. Nonadherent BMMs were isolated from total bone marrow cells cultured on tissue culture dishes for 48 hours. CD11b+ or CD11c+ cells were isolated from BMMs using the CD11b or CD11c microbeads and the LS columns according to the manufacturer’s instructions (Miltenyi Biotec Inc). For differentiation, cells were first cultured in proliferation medium (α-MEM containing 10% FBS and 10 ng/ml human recombinant M-CSF) for 3 days, then switched to differentiation medium (proliferation medium plus 50 ng/ml human recombinant RANKL) for 4–9 days. The TRAP+ MNCs were scored using an inverted microscope. For Pit assay, BMMs (5 × 10^5 cells/well) were seeded on dentin slices in 24-well plates in proliferation medium for 3 days and switched to differentiation medium for 9 days. Bone resorption pits were stained with hematoxylin solution. Pit area versus total bone area and pit area versus TRAP+ MNC area of each dentin slice were measured using Image Pro Plus 6.2 software (Media Cybernetics Inc). Coculture experiments were performed as previously described (64). Briefly, primary calvarial OBLs (3.2 × 10^5 cells/well) were seeded in 24-well plates and cultured in α-MEM containing 10% FBS and 50 ng/ml ascorbic acid. BMMs (4 × 10^5 cells/well) were seeded on top of the OBLs. The medium was changed daily.
supplemented with 10 ng/ml M-CSF and 10^8 M 1,25 dihydroxyvitamin D₃. OCLs were identified by TRAP staining and counted. Serum levels of CTX, degradation products from type I collagen during osteoclastic bone resorption, were measured using the RatLaps ELISA Kit according to the manufacturer’s instruction (Immunodiagnostic Systems Limited).

Histological
evaluation, bone histomorphometry, and IHC. WT and Atf4⁺/⁻ mice were euthanized, and tibiae were fixed in 10% formalin at 4°C for 24 hours, dealcified in 10% EDTA (pH 7.4) for 10–14 days, and embedded in paraffin. Sections of tibiae from WT and Atf4⁺/⁻ mice were used for TRAP staining as described previously (65). Bone histomorphometry, such as Oc.S/BS and Oc.Nb/Bpm, in both primary and secondary spongiosa of tibiae was measured using Image Pro Plus 6.2 software (Media Cybernetics Inc.) as previously described (43, 66). Cells cultured in 8-well culture chambers (Nalgene Nunc), or 5-μm sections of tibiae, were subjected to IHC staining with antibodies against ATF4, NFATc1, or RANK (Santa Cruz Biotechnology) using the EnVision System–HRP (DAB) kit (Dako North America Inc.) as described previously (41). Fixed nondemineralized femurs were used for PCR subcloning promoter fragment using mouse tail DNA as a template into pGL3-luc vector (Promega). Mutant plasmid pRL-SV4, and 0.5–1.5 μg expression plasmid such that the total DNA was constant in each group. Experiments were performed in triplicate and repeated 3–4 times. Adenoviruses expressing ATF4 and β-gal were described previously (57). Retrovirus expressing cnNFATc1 and its control empty retrovirus were described previously (38). The amount of adenovirus or retrovirus was balanced as necessary with a control adenovirus expressing β-gal or an empty retrovirus such that the total amount was constant in each group.

ChIP. ChIP assays were performed as described previously (41, 58). The equivalent of 10 μg DNA was used as starting material (input) in each ChIP reaction with 2 μg of the appropriate antibody (ATF4, c-Fos, or NFATc1). Fractions of the purified ChIP DNA (5%) or inputs (0.02%–0.05%) were used for PCR analysis. The reaction was performed with AmpliTaq Gold DNA Polymerase (Applied Biosystems) for 35 cycles of 60 seconds at 95°C, 90 seconds at 58°C, and 120 seconds at 68°C. A PCR primer pair (P1, CCGGGAGC-GCCATGCAATGTCGTTAATT; P2, CCGGGTGCCCGAGGAAGC-TACTCCTCCCT) was generated to detect DNA segments located near the AP-1–binding site at –64/–637 and 2 NFATc1-binding sites at –689/–684 (TTTCCC) and –680/–676 (TTTCCC), respectively, in mouse Nfatc1 proximal promoter (33). The PCR products were separated on 3% agarose gels and visualized with ultraviolet light. All ChIP assays were repeated at least 3 times.

Flow cytometry. Cell surface fluorescence was determined using FITC-conjugated CD3, bio-strept–PB- or APC-conjugated CD11b, Pecy5-conjugated c-kit (i.e., CD117), PE-conjugated CD45R, APC-conjugated CD115, and FITC-conjugated CD11c with appropriate isotype controls (eBioscience Inc.). For all transfection experiments, the equivalent of 10 μg DNA was used as starting material (input) in each ChIP reaction with 2 μg of the appropriate antibody (ATF4, c-Fos, or NFATc1). Fractions of the purified ChIP DNA (5%) or inputs (0.02%–0.05%) were used for PCR analysis. The reaction was performed with AmpliTaq Gold DNA Polymerase (Applied Biosystems) for 35 cycles of 60 seconds at 95°C, 90 seconds at 58°C, and 120 seconds at 68°C. A PCR primer pair (P1, CCGGGAGC-GCCATGCAATGTCGTTAATT; P2, CCGGGTGCCCGAGGAAGC-TACTCCTCCCT) was generated to detect DNA segments located near the AP-1–binding site at –64/–637 and 2 NFATc1-binding sites at –689/–684 (TTTCCC) and –680/–676 (TTTCCC), respectively, in mouse Nfatc1 proximal promoter (33).

Acknowledgments

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Figure S2
Figure S3
Figure S4
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| β-actin   | ![Image](image21.png) | ![Image](image22.png) |

Figure S5
Figure S6

Spleen

A

B

Bone Marrow

C

D
**Table S1**: real-time PCR primers  

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Legends for Supplementary Figures:

**Fig. S1. ATF4 is expressed in OCL-like cells.** Mouse MC-4 preosteoblastic cells, rat UMR106-01 osteoblastic cells, primary mouse bone marrow stromal cells (BMSCs), RAW264.7 cells (a mouse monocyte/macrophage cell line), and primary mouse bone marrow-derived monocytes (BMMs) were cultured in 10%FBS media for 24 h. Whole cell extracts were used for Western blot analysis for ATF4.

**Fig. S2. TRAP staining in CD11b+ BMMs and BMMs from aged mice.** (A) TRAP staining. Purified CD11b+ BMMs from 4-week-old wt and Atf4−/− mice were differentiated for 5d followed by TRAP staining and TRAP-positive multinucleated cells (MNCs) were scored. (B) Statistical analysis of TRAP staining in (A). Magnification: 100X. (C) TRAP staining. Primary BMMs from 15-month-old wt and Atf4−/− mice were differentiated for 5d followed by TRAP staining. (D) Statistical analysis of TRAP staining in (C). Magnification: 40X.

**Fig. S3. RANKL induction of Nfatc1 mRNA expression in wt and Atf4−/− BMMs (dose-response).** Wt and Atf4−/− BMMs were treated with indicated concentrations of RANKL (25-200 ng/ml) for 7d followed by quantitative real-time RT/PCR for Nfatc1 mRNA. Gapdh mRNA was used for loading. *P<0.01 wt vs. Atf4−/−.
Fig. S4. ATF4 but not Runx2 activates the Nfatc1 P1 promoter. COS-7 cells were transfected with 0.125 µg 0.8-kb Nfatc1-luc and 1.0 ng pRL-SV40 with 1.0 µg expression plasmid for ATF4 or Runx2. *P<0.01 (β-gal vs. ATF4).

Figure S5. Effects of M-CSF on activation of the AKT, Src, and MAPKs pathways in wt and Atf4−/− BMMs. Wt and Atf4−/− BMMs were cultured in proliferation medium for 3 d and switched to 2% FBS α-MEM without M-CSF overnight. Cells were then exposed to 30 ng/ml M-CSF for the indicated times. Cells were then lysed, fractionated by SDS–PAGE, and analyzed by Western blot analysis using antibodies recognizing phosphorylated and total ERK1/2, p38, JNK, AKT, and Src. β-actin served as the loading control. Similar results were obtained from three independent experiments.

Fig. S6. Atf4−/− mice display an increase in macrophage. (A-B) Flow cytometry. Splenocytes from wt and Atf4−/− mice were stained with CD11b-APC and CD11c-FITC antibodies and analyzed with flow cytometry as described in Experimental Procedures. (C) CD11c+ cells were purified and counted from wt and Atf4−/− BMMs as described in Experimental Procedures. (D) Total nucleated bone marrow cells were counted from long bones (two femurs and two tibias) from 6-week-old wt and Atf4−/− mice (6 mice per group).
Foxo1 Mediates Insulin-like Growth Factor 1 (IGF1)/Insulin Regulation of Osteocalcin Expression by Antagonizing Runx2 in Osteoblasts

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In this study, we determined the molecular mechanisms whereby forkhead transcription factor Foxo1, a key downstream signaling molecule of insulin-like growth factor 1 (IGF1)/insulin actions, regulates Runx2 activity and expression of the mouse osteocalcin gene 2 (Bglap2) in osteoblasts in vitro. We showed that Foxo1 inhibited Runx2-dependent transcriptional activity and osteocalcin mRNA expression and Bglap2 promoter activity in MC-4 preosteoblasts. Co-immunoprecipitation assay showed that Foxo1 physically interacted with Runx2 via its C-terminal region in osteoblasts or when co-expressed in COS-7 cells. Electrophoretic mobility shift assay demonstrated that Foxo1 suppressed Runx2 binding to its cognate site within the Bglap2 promoter. IGF1 and insulin prevented Foxo1 from inhibiting Runx2 activity by promoting Foxo1 phosphorylation and nuclear exclusion. In contrast, a neutralizing anti-IGF1 antibody decreased Runx2 activity and osteocalcin expression in osteoblasts. Chromatin immunoprecipitation assay revealed that IGF1 increased Runx2 interaction with a chromatin fragment of the proximal Bglap2 promoter in a PI3K/AKT-dependent manner. Conversely, knockdown of Foxo1 increased Runx2 interaction with the promoter. This study establishes that Foxo1 is a novel negative regulator of osteoblast-specific transcription factor Runx2 and modulates IGF1/insulin-dependent regulation of osteocalcin expression in osteoblasts.

Foxo1 is a forkhead transcription factor that is defined by its amino forkhead DNA binding domain and carboxyl trans-activation domain. Foxo1 plays a pivotal role in mediating the effect of insulin or insulin-like growth factor 1 (IGF1)2 on the expression of genes involved in cell growth, differentiation, metabolism, and longevity (1–5). Insulin or IGF1 exerts its inhibitory effect on gene expression via a highly conserved sequence (TG/ATTTT/G), termed the insulin response element (IRE), in the promoter of genes that are negatively regulated by insulin/IGF1. In the absence of insulin/IGF1, Foxo1 resides in the nucleus and binds as a trans-activator to the IRE, enhancing promoter activity. In response to insulin, Foxo1 is phosphorylated at three highly conserved phosphorylation sites (Thr-24, Ser-256, and Ser-319) through the PI3K-dependent pathway, resulting in its nuclear exclusion and inhibition of target gene expression (1, 2, 6). Except for Foxo6 (7), all members of the Foxo superfamily undergo insulin/IGF1-dependent phosphorylation and nuclear exclusion. Failure to phosphorylate Foxo1 results in its permanent nuclear localization and constitutive trans-activation of gene expression. This phosphorylation-dependent Foxo1 translocation has been viewed as an acute mechanism for insulin or growth factors to inhibit gene expression, as insulin-induced Foxo1 phosphorylation is kinetically coupled to its subsequent translocation to the cytoplasm (1, 8).

Osteoblasts, the bone-forming cells, originate from multipotential mesenchymal cells. Osteoblast activity and function are regulated by a number of growth factors and hormones including IGF1, insulin, bone morphogenetic proteins (BMPs), basic fibroblast growth factor 2 (FGF-2), parathyroid hormone (PTH), tumor necrosis factor-α (TNF-α), and extracellular matrix signals (9–21). At the molecular level, osteoblast function is controlled by several key transcription factors including Runx2, osterix, and ATF4 (22–29). Runx2 is a runt domain-containing transcription factor that is characterized as a transcriptional activator of osteoblast differentiation and master gene for bone development (22–26). Runx2 expression and activity are controlled by a number of factors including IGF1, BMPs, FGF-2, PTH, TNF-α, and extracellular matrix signals (9, 12–14, 30, 31) as well as by nuclear factors via protein-protein interactions (29, 32–46).

Recent studies showed that osteocalcin, an osteoblast-specific product encoded by the Bglap2 (bone γ-carboxyglutamate protein) gene plays a critical role in regulating glucose metab-
olism (10, 11, 47). Using mice lacking Foxo1 selectively in osteoblasts, Rached et al. (48) recently showed that Foxo1 expressed in osteoblasts regulates glucose homeostasis through an osteocalcin-dependent mechanism. Specifically, osteoblast-conditional inactivation of Foxo1 increased β-cell proliferation and insulin secretion and sensitivity. Importantly, osteoblastic osteocalcin protein, which is active in the absence of β-carboxylation, was found to be responsible for the metabolic actions of Foxo1 in regulating glucose homeostasis. Foxo1 decreases osteocalcin mRNA expression and increases osteocalcin carboxylation. Foxo1 achieves the latter by increasing the expression of Esp, a gene that encodes a protein that decreases osteocalcin function (i.e. increases its carboxylation). Therefore, Foxo1 negatively regulates both osteocalcin production/expression and function. However, the molecular mechanism whereby Foxo1 suppresses the Bglap2 gene is undefined. In this study, we hypothesized that Foxo1 inhibits Bglap2 gene expression and promoter activity, at least in part, via suppression of Runx2, a major transcriptional activator of the Bglap2 gene (25, 49). To address this hypothesis, we analyzed the molecular interplay between Runx2 and Foxo1 in Bglap2 gene expression in osteoblasts. We demonstrate that Foxo1 physically binds to and functionally antagonizes Runx2 from driving Bglap2 expression. Furthermore, we demonstrate that IGF1/insulin prevents Foxo1 from inhibiting Runx2, probably by promoting Foxo1 phosphorylation and nuclear exclusion in osteoblasts. This effect results in inhibition of Foxo1 action, which contributes to increased osteocalcin expression in osteoblasts and favors glucose homeostasis.

**EXPERIMENTAL PROCEDURES**

Reagents and Cell Lines—Tissue culture media and fetal bovine serum were obtained from HyClone (Logan, UT). Mouse MC3T3-E1 subclone 4 (MC-4) cells were described previously (31, 50) and maintained in AA (ascorbic acid)-free α-modified Eagle’s medium (α-MEM), 10% fetal bovine serum (FBS), and 1% penicillin/streptomycin and were not used beyond passage 15. COS-7 cells and rat UMR106–01 osteosarcoma cells were described (46, 51, 52). IGF1 was purchased from R&D Systems, Inc (Minneapolis, MN). Insulin and LY294002 were purchased from Sigma Aldrich.

DNA Constructs—Wild-type (wt) or mutated (mt) 6XRUNX2-LUC (also known as p6OSE2-luc (49)) or BGLAP2-LUC (also known as p657mOG2-luc (49)), which contains 2-bp substitution mutations in the OSE2 sites that abolishes Runx2 binding, 4XATF4-LUC (p4OSE1-luc (53, 54)), pCMV5/β-gal, pCMV/Runx2 expression plasmids (wt, amino acids 1–410, amino acids 1–330, amino acids 1–286, amino acids 1–258) containing cDNAs encoding either wt Runx2 or C-terminal deletions under CMV promoter control, and full-length GST-Runx2 and GST-Foxo1 fusion protein expression vectors were previously described (29, 46, 49, 53, 55, 56). HA-tagged pCMV/Runx2 expression plasmids containing cDNAs encoding wt or N-terminal deletions (ΔN97, ΔN242, and ΔN326) under CMV promoter control were previously described (57). To generate pCMV/Foxo1 expression plasmids expressing truncated forms of Foxo1 (amino acids 1–558, amino acids 1–456, amino acids 1–360, amino acids 1–258), a stop code (TAA, TAG, or TGA) that results in premature stop of Foxo1 protein at indicated amino acid residues was introduced into Foxo1 cDNA by PCR using pCMV/Foxo1 as a template. The pCMV/Foxo1(3A) expression plasmid expressing a mutant Foxo1 protein in which three insulin/akt-dependent phosphorylation sites (Thr-24, Ser-256, and Ser-319) were mutated from Thr or Ser to Ala, was described previously (58). Adenovirus expressing Foxo1 under control of a CMV promoter (AdCMV/Foxo1) was constructed by subcloning full-length Foxo1 cDNA into pAdlox plasmid followed by CRE-mediated recombination as previously described (55, 56). Expression of Foxo1 protein was confirmed by Western blot analysis (data not shown). All sequences were verified by automatic DNA sequencing.

Transfection and Dual Luciferase Assay—Cells were plated on 35-mm dishes at a density of 5 × 10^4 cells/cm². After 24 h, cells were transfected with Lipofectamine 2000 (Invitrogen) according to manufacturer’s instructions. Each transfection contained 0.25 μg of the indicated reporter plasmids plus 0.01 μg of pRL-SV40, containing a cDNA for Renilla Reformis luciferase to control for transfection efficiency. Cells were harvested and assayed using the Dual Luciferase Assay kit (Promega, Madison, WI) on a Module™ Microplate Multimode Reader (Turner Biosystem, Sunnyvale, CA). For all transfection experiments, the amount of plasmid DNAs was balanced as necessary with β-galactosidase (β-gal) expression plasmid such that the total DNA was constant in each group. Experiments were performed in triplicates and repeated 3–4 times.

Adenoviral Infection—Adenoviral vectors for β-galactosidase (Ad-β-gal), Foxo1 (Ad-Foxo1), Foxo1 RNAi (Ad-Foxo1i), or control RNAi (USi) were described previously (29, 55, 56, 59). MC-4 cells were infected with adenovirus as described previously (29, 55, 56). Briefly, virus was added to cells in 1% FBS and incubated for 1 h at 37°C. Dishes were rotated every 5 min for the first 15 min to ensure that all of the cells were exposed to virus. After 1 h, media were aspirated, and cultures were rinsed twice with serum-free medium, and then fresh media supplemented with 10% FBS were added to the dishes. The amount of adenovirus was balanced as necessary with a control adenovirus expressing β-galactosidase such that the total amount was constant in each group.

RNA Isolation, Reverse Transcription (RT), Quantitative RT-PCR, and Western Blot Analyses—RNA isolation, RT, and quantitative real-time RT-PCR were performed to measure the relative mRNA levels using SYBR Green kit (Bio-Rad) as previously described (46, 54, 60). Samples were normalized to Gapdh expression. The DNA sequences of mouse primers used for real-time PCR were as follows: Bglap2: 5’-TAG TGA ACA GAC TCC GGC GCT A-3’ (forward), 5’-TG TGG AGG CGG TCT TCA AGC CAT-3’ (reverse); Runx2: 5’-TAA AGT GAC AGT GGA CGG TCC C-3’ (forward), 5’-TG GCC CTA AAT CAC TGA GG-3’ (reverse); Adenovirus: 5’-CAG TGC CAG CCT CGT CCC GTA GA-3’ (forward), 5’-CTG CAA ATG GCA GCC CTG GTG AC-3’ (reverse). Western blot analysis was performed as previously described (46, 54). Antibodies used in this study were from the following sources: antibodies against Foxo1, Runx2, HA, anti-IGF1 neutralizing antibody, normal control IgG, and anti-rabbit or anti-mouse antibodies conjugated with horseradish peroxidase from Santa Cruz Biotech-
nology, Inc., and mouse monoclonal antibody against β-actin and M2 antibody from Sigma Aldrich.

**Nuclear Extracts (NE) Preparation and Electrophoretic Mobility Shift Assay (EMSA)—** Nuclear extracts were prepared from rat UMR106 – 01 osteosarcoma cells, which express high level of Runx2 protein, and EMSA was performed as previously described (21, 31). GST-Runx2, GST-Foxo1, and GST proteins were purified using the Bulk GST Purification Module kit (GE Healthcare/Amersham Biosciences) as previously described (46). The DNA sequences of the oligonucleotides used for EMSA were as follows: OSE2 (Runx2 binding site) (49): 5-GAT CCG CTG CAA TCA TCA CCA ACC ACA GCA-3. IRE (Insulin Response Element that contains a Foxo1 binding site) (55): 5-TGT AGT TTG TTT TGT TTT GTT GGC ATG-3. DNA oligonucleotide was labeled using a biotin 3’-end DNA labeling kit (cat: 89818, Pierce Biotechnology Inc.). 2 μg of nuclear extracts and 20 fmol biotin-labeled DNA probe were incubated in the presence and absence of indicated amounts of purified GST-Foxo1 or GST protein in 1x binding buffer for 30 min at room temperature. For supershift assay, 1 μg of IgG or anti-Runx2 antibody was first incubated with nuclear extracts prior to addition of DNA probe. Protein-DNA complexes were separated on 4% polyacrylamide gels in 0.5x TBE buffer, and transferred onto Biodyne B Nylon Membrane (cat: 77016, Pierce). The membrane was blocked in 1x blocking buffer, washed five times with 1x wash buffer, and visualized by a Chemiluminescent Nucleic Acid Detection Module (cat: 89880, Pierce).

**Immunoprecipitation (IP) and Chromatin Immunoprecipitation (ChIP) Assays—** Whole cell extracts and purified GST fusion proteins were used for immunoprecipitation using specific antibodies as previously described (46). ChIP assays were performed as described previously (21, 46). Briefly, the equivalent of 10 μg of DNA was used as starting material (input) in each ChIP reaction with 2 μg of the appropriate antibody (Runx2 or control rabbit IgG). Fractions of the purified ChIP DNA (5%) or inputs (0.02–0.05%) were used for PCR analysis. The reaction was performed with AmpliTaq Gold DNA Polymerase (Applied Biosystems) for 30 cycles of 15 s at 95 °C, 30 s at 60 °C, and 15 s at 72 °C. PCR primer pairs were generated to detect DNA segments located near the Runx2-binding site at –137/-131 (primers P1 and P2) and the Bglap2 cDNA region (primers P3 and P4) (46). The PCR products were separated on 3% agarose gels and visualized with ultraviolet light. All ChIP assays were repeated at least three times.

**Statistical Analysis—** Results were expressed as means ± standard deviation (S.D.). Students’ t test was used to test for differences between two groups. Differences with a p < 0.05 was considered as statistically significant. All experiments were repeated a minimum of three times with triplicate samples.

**RESULTS**

**Foxo1 Decreases Bglap2 Expression and Promoter Activity and Runx2-dependent Transcriptional Activity—** To determine the effect of Foxo1 on Bglap2 expression in osteoblasts, MC3T3-E1 subclone 4 (MC-4) preosteoblasts were infected with equal amount of adenovirus expression vectors for Foxo1 (Ad-Foxo1) or β-galactosidase (Ad-β-gal), followed by RNA preparation and quantitative real-time RT/PCR for Bglap2 and Runx2 mRNAs, which were normalized to Gapdh mRNA. Consistent with the result from a recent study (48), adenoviral overexpression of Foxo1 dramatically reduced the level of Bglap2 mRNA in the MC-4 cells (Fig. 1A). Surprisingly, Foxo1 did not alter the mRNA level of Runx2, a major upstream transcriptional activator of the Bglap2 (25, 26, 49, 61). A similar result was observed in rat UMR106 – 01 osteoblast-like cells (data not shown). Therefore, we next evaluated whether Foxo1 modulated Runx2 activity. To this end, COS-7 cells were co-infected with 6XRUNX2-LUC and pRL-SV40, and FLAG-Runx2 plasmid with and without increasing amounts of Foxo1 expression plasmid. *, p < 0.05, versus Runx2; E, COS-7 cells were transfected with BGLAP2-LUC or BGLAP2mt-LUC in which two previously defined Runx2-binding sites were mutated (29), pRL-SV40, and FLAG-Runx2 plasmid with and without increasing amounts of Foxo1 expression plasmid. *, p < 0.05, versus 0 μg Foxo1; #, p = 0.05, versus wt.
Foxo1 Inhibits Runx2

Foxo1 Physically Interacts with Runx2—To define the molecular mechanism through which Foxo1 inhibits Runx2, we next determined whether Foxo1 interacted with Runx2 by performing immunoprecipitation (IP) assays using nuclear extracts from the rat UMR106–01 osteoblastic cells. As shown in Fig. 2A, Runx2 protein was present in an anti-Foxo1 antibody immunoprecipitate (lanes 1 and 2). Reciprocal IP assay showed that an anti-Runx2 antibody, but not control IgG, immunoprecipitated the Foxo1 protein (lanes 3 and 4). As expected, ATF4 protein, a known Runx2-interacting factor (29), was present in the anti-Runx2 antibody immunoprecipitate (lanes 5 and 6). In contrast, an Nfatc1 antibody failed to immunoprecipitate Runx2 (lanes 7 and 8). As shown in Fig. 2B, Foxo1 and Runx2 can also be co-immunoprecipitated from COS-7 cells exogenously expressing both factors. To determine if Foxo1 directly interacted with Runx2 in the absence of other nuclear proteins, we conducted IP assays using purified GST-Foxo1 and GST-Runx2 and GST proteins. As shown in Fig. 2C, GST-Runx2 protein was immunoprecipitated by an anti-Foxo1 antibody (lanes 1 and 2) and, vice versa, GST-Foxo1 was immunoprecipitated by an anti-Runx2 antibody (lanes 3 and 4). In contrast, GST protein was not immunoprecipitated by antibodies against Runx2 or Foxo1 or normal IgG (lanes 5–8), thus demonstrating a direct interaction between Foxo1 and Runx2.

Deletion Analysis of the Runx2 cDNA—To identify the Foxo1-binding domain within the Runx2 molecule, COS-7 cells were co-transfected with the Foxo1 expression plasmid and wt FLAG-Runx2 or various FLAG-Runx2 C-terminal deletion mutant expression vectors. Forty hours later, nuclear extracts were prepared for IP using an anti-Foxo1 antibody, followed by Western blot analysis using a M2 antibody. As shown in Fig. 3B, deletion of Runx2 from amino acid 528 (wt) to amino acid 258 did not abolish the Foxo1 binding. This result was confirmed by the reciprocal IP using a M2 antibody, followed by Western blot for Foxo1 (Fig. 3C). As shown in Fig. 3E, the deletion of the N-terminal 242 amino acid of Runx2 did not reduce its ability to bind to Foxo1. However, further deletion from amino acid 242 to amino acid 326 completely abrogated the Foxo1-Runx2 interaction. Collectively, these results suggest that the amino acid 242–258 region of Runx2 is critical for interaction with Foxo1.

Deletion Analysis of the Foxo1 cDNA—Several C-terminal Foxo1 deletion mutant expression vectors were generated and tested for their ability to suppress Runx2 activity in COS-7 cells. As shown in Fig. 4A (top), the deletion of Foxo1 cDNA from amino acids 653 (wt) to 456 did not alter its inhibition of Runx2 activity. However, further deletion from amino acids 456 to 360 completely abolished its Runx2 inhibitory activity. As shown in Fig. 4A (bottom), Western blot analysis showed that both wt and mutant Foxo1 proteins were expressed at comparable levels. Likewise, co-expression of wt or mutant Foxo1 expression vectors did not markedly alter the level of Runx2 protein. Consistent with the results from the functional study, IP assays revealed that the deletion from amino acids 456 to 360 of Foxo1 also abolished the Foxo1-Runx2 interaction (Fig. 4, B and C). These results suggest that the C-terminal region (amino acids 360–456) is essential for Foxo1 to bind to and inhibit Runx2 activity.

Foxo1 Inhibits Runx2 Binding to Its Cognate Site (OSE2) within the Bglap2 Promoter—To study the mechanism whereby Foxo1 inhibits Runx2 as demonstrated above, we next determined whether Foxo1 affected Runx2 DNA binding activity by performing EMSA using the wild-type (wt) OSE2 (Osteoblast-Specific Element 2) oligo, a well-established Runx2 binding element from the Bglap2 promoter (49), as probe and 2 μg of nuclear extracts (NE) from UMR106-01 cells in the presence and absence of increasing amounts of purified GST-Foxo1 protein. As shown in Fig. 5A, while GST-Foxo1 protein itself did not bind to the OSE2 oligo (lane 6), it dose-dependently inhibited the binding of Runx2 to the OSE2 oligo (lanes 3–5). This inhibition was specific because the GST protein neither altered the Runx2 DNA binding activity (lanes 7 and 8) nor bound to
FIGURE 3. Deletion analysis of the Runx2 cDNA. A, schematic showing the domain structure of Runx2 and the C-terminal deletion mutants. Runx2 contains three transcriptional activation domains (AD1, 2, 3), a transcriptional repression domain (RD) containing a VWRPY motif at C-terminal, a Runt domain responsible for DNA binding, a nuclear localization sequence (NLS), and a large C-terminal PST domain which is rich in serine/threonine/tyrosine residues. B and C, whole cell extracts from COS-7 cells cotransfected with expression vectors for Foxo1 and wt FLAG-Runx2 or various FLAG-Runx2 C-terminal deletion mutants (amino acid 1–410, amino acid 1–330, amino acid 1–286, and amino acid 1–258) were immunoprecipitated with Foxo1 antibody, followed by Western blot using M2 antibody (B). In reciprocal IP, the same extracts were immunoprecipitated with control IgG (lanes 1) or M2 antibody (lanes 2–6), followed by Western blot using Foxo1 antibody. D, schematic showing the domain structure of Runx2 and its N-terminal deletion mutants. E, whole cell extracts from COS-7 cells cotransfected with expression vectors for Foxo1 and wt HA-Runx2 or various HA-Runx2 N-terminal deletion mutants (∆N97, ∆N242, and ∆N326) were immunoprecipitated with control IgG (lane 1) or Foxo1 antibody (lanes 2–5), followed by Western blot using HA antibody.
Foxo1 Inhibits Runx2

FIGURE 4. Deletion analysis of the Foxo1 cDNA. A, COS-7 cells were transiently transfected with 6XRUNX2-LUC and pRL-SV40 with and without Runx2 plasmids as well as with and without wt or various deletion mutant Foxo1 plasmids (amino acids 1–558, amino acids 1–456, amino acids 1–360, and amino acids 1–258), followed by dual luciferase assay (top) and Western blot (bottom). *, p < 0.05, versus 0 μg Foxo1. B and C, whole cell extracts from COS-7 cells overexpressing FLAG-Runx2 and wt or two C-terminal deletion Foxo1 mutants (amino acids 1–558 and amino acids 1–360) were immunoprecipitated with control IgG (lane 1) or Foxo1 antibody (lanes 2–4), followed by Western blot using an anti-Foxo1 antibody (B). In reciprocal IP, the same extracts were immunoprecipitated with control IgG (lane 1) or Foxo1 antibody (lanes 2–4), followed by Western blot using a M2 antibody (C).

To determine whether this also occurs in osteoblasts, MC-4 cells were electroporated with expression plasmids for Fox1-GFP or GFP proteins as previously described (63). Thirty hours later, cells were treated with and without 10 ng/ml mouse recombinant IGF1 or 100 nM insulin for the indicated times. As shown in Fig. 6A, top and middle, GFP-Foxo1 protein was rapidly translocated from the nucleus to the cytoplasm 0.5–1 h after IGF1 or insulin addition to the MC-4 cells, which contain a high level of endogenous Runx2 (12, 20). Conversely, IGF1 treatment did not alter subcellular distribution of the GFP protein in the same cells (Fig. 6A, bottom). A similar result was observed in insulin-treated MC-4 cells (data not shown). Based on these observations, we reasoned that IGF1 or insulin should abolish or reduce Foxo1 inhibition of Runx2 by phosphorylation and subsequent translocation of Foxo1 to the cytoplasm from the nucleus where Runx2 is located and activates Bglap2 transcription. To test this, MC-4 cells were co-transfected with 6XRUNX2-LUC (Fig. 6B) or BGLAP2-LUC (Fig. 6C) and pRL-SV40 with and without expression vector for Foxo1. Twenty hours later, cells were treated with and without 10 ng/ml IGF1 as well as with and without 0.5 μg/ml of IGF1 neutralizing antibody or control IgG for 6 h, followed by dual luciferase assay. Importantly, Foxo1-mediated inhibition of Runx2 activity was completely reversed by the addition of IGF1 (Fig. 6B) or insulin (Fig. 7E). Furthermore, the effect of IGF1 was completely blocked by the addition of a specific IGF1 neutralizing antibody, but not by the control IgG. A similar result was obtained when the 657-bp BGLAP2-LUC was used in MC-4 cells (Fig. 6C). The IGF1 neutralizing antibody significantly diminished the level of Bglap2 but not Runx2 mRNAs in the MC-4 cells (Fig. 6D and E), which suggests that endogenous IGF1 signaling, probably via an autocrine mechanism, plays a critical role for maintaining the Bglap2 mRNA expression in osteoblasts. Taken together, these results suggest that IGF1/insulin favors Bglap2 expression probably by preventing Foxo1 from inhibiting Runx2 in osteoblasts.

IGF1 Increases Runx2 Interaction with the Bglap2 Promoter in a PI3K/AKT-dependent Manner in the MC-4 Cells—To determine whether Runx2 associates with the endogenous Bglap2 promoter in vivo, we performed ChIP assays using MC-4 cells with and without IGF1 treatment for the indicated times. Consistent with our previous observation (46), Runx2 specifically interacted with a chromatin fragment of the proximal Bglap2 promoter that contains the Runx2-binding site (primers P1/P2) (Fig. 7A). Importantly, this interaction was markedly stimulated by IGF1 treatment. In contrast, Runx2 antibody failed to immunoprecipitate a 3′ chromatin fragment from the transcribed region of the Bglap2 that contains no Runx2-binding sites (primers P3/P4) (46) (Fig. 7A, bottom). Further, the IGF1-induced increase in Runx2 binding to the Bglap2 promoter was abolished by treatment with LY294002, a specific inhibitor of the PI3K/AKT pathway (Fig. 7B). We next determined the effect of Foxo1 knockdown on Runx2 interaction with the Bglap2 promoter in MC-4 cells. Adenoviral overexpression of a mouse Foxo1 RNAi (Foxo1i) dramatically reduced the level of Foxo1 protein in MC-4 cells (supplemental Fig. S3). As shown in Fig. 7C, knockdown of Foxo1 markedly increased Runx2 binding to the Bglap2 promoter. As shown in Fig. 7D, the PI3K/AKT inhibition abrogated IGF1-induced reversal of
Foxo1 inhibition of Runx2 activity. Foxo1 is phosphorylated at three highly conserved phosphorylation sites (Thr-24, Ser-256, and Ser-319) through the PI3K/AKT-dependent pathway, resulting in its nuclear exclusion and inhibition of target gene expression (1, 2, 6). To further study the role of the PI3K/AKT pathway in Foxo1 modulation of Runx2 activity, we compared the effects of wt Foxo1 and a mutant Foxo1(3A), in which the PI3K/AKT-dependent phosphorylation sites were mutated from either Thr or Ser to Ala, on Runx2 activity in the presence and absence of IGF1 or insulin. Significantly, in contrast to result from the wt Foxo1 group, neither IGF1 nor insulin prevented the mutant Foxo1(3A) from inhibiting Runx2 in the MC-4 cells (Fig. 7E).

DISCUSSION

This study identifies Foxo1 as a negative regulator of the bone transcription factor Runx2 in osteoblasts. Foxo1 physically interacts with Runx2 and inhibits Runx2 activity and decreases expression of the Bglap2 gene encoding osteocalcin. Most importantly, we demonstrated that both IGF1 and insulin, which phosphorylate and export Foxo1 from the nucleus to the cytoplasm, prevent Foxo1 from inhibiting Runx2 and increase Runx2 activity, thereby favoring osteocalcin expression, an osteoblast-secreted hormone that plays a critical role in regulating glucose metabolism via its actions in β-cells in the pancreas.

Results from the present study established that Foxo1 indirectly down-regulates osteocalcin expression, at least in part, by inhibiting Runx2, a major upstream transcriptional activator of the Bglap2 gene in osteoblasts. Foxo1 interaction with Runx2 in osteoblasts, or when coexpressed in COS-7 cells, requires the presence of the Foxo1 C-terminal 360–456 amino acid region.
Insulin signaling in osteoblasts was recently shown to be critical for postnatal bone acquisition and remodeling (10, 11). Fulzele et al. (10) recently showed that the insulin receptor (IR) is expressed in osteoblasts and that osteoblast-specific deletion of the IR severely impairs osteoblast differentiation. Mice lacking the IR in osteoblasts have reduced bone mass with increased adiposity and insulin resistance. Importantly, this study further revealed that the metabolic dysregulation in these mice was caused by reduced osteocalcin expression/function. Our results from the present study suggest that insulin favors osteocalcin expression by preventing Foxo1 from inhibiting Runx2, a key activator of the Bglap2 gene. This notion is strongly supported by: (i) the well-established role of IGF1/insulin signaling via the PI3K/AKT pathway that phosphorylates and exports Foxo1 from the nucleus to the cytoplasm, where it binds to 14-3-3 proteins; (ii) Foxo1 inhibition of Runx2 activity and Bglap2 expression was completely prevented by IGF1 treatment in osteoblasts, which expresses both the IR and IGF1 receptors; (iii) IGF1 neutralizing antibody reduced Runx2 activity and Bglap2 expression in osteoblasts; and (iv) IGF1 increased Runx2 interaction with the Bglap2 promoter in osteoblasts, which was abolished by PI3K/AKT inhibition. Interestingly, IR deficiency in osteoblasts caused increased expression of Twist2, a known Runx2 inhibitor (45). However, the molecular mechanism whereby insulin signaling modulates the expression of Twist2 is still unknown. Additionally, IGF1 and probably insulin were shown to activate Runx2 activity via the Erk1/2 MAPK pathway (30), a major signaling route in osteoblasts (14, 19). Erk1/2 phosphorylation sites in Runx2 were recently identified.
and characterized (57). This phosphorylation is critical for Runx2 activity and osteoblast differentiation as well as bone formation (57, 64). Collectively, these studies suggest that IGFl/insulin signaling favors Runx2 activity and Bglap2 expression via at least three distinct molecular mechanisms. Therefore, our findings from this study add a new layer to the molecular mechanisms through which IGFl/insulin signaling in osteoblasts modulates glucose homeostasis as well as bone metabolism.

Based on the findings from this and other studies, we propose the following molecular model for Foxo1 modulation of osteocalcin expression in osteoblasts and the effect of IGFl/insulin signaling on this function of Foxo1 (Fig. 8). Binding of IGFl/insulin to their receptors, which are expressed in osteoblasts (10, 11), activates the PI3K/AKT pathway, which phosphorylates and translocates Foxo1 from the nucleus to the cytoplasm, where it binds to 14-3-3 chaperone proteins (65). This nuclear exclusion prevents Foxo1 from binding to and inhibiting Runx2, which results in enhanced Bglap2 expression. In the meantime, insulin and probably IGFl down-regulate the expression of Twist2 (10), a known Runx2 suppressor, and thereby increase Runx2 activity (not shown), which further up-regulates osteocalcin expression. Finally, the uncarboxylated form of osteocalcin, via the circulation, regulates glucose homeostasis (i.e. insulin secretion and production and sensitivity) (47). It should be noted that Foxo1 can also bind independently to its target genes (not shown). In summary, we, for the first time to our knowledge, demonstrate that Foxo1 is a novel negative regulator of Runx2 and mediates IGFl/insulin actions in regulating osteocalcin expression in osteoblasts.

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FIGURE 8. Molecular model of Foxo1 and IGFl/insulin regulation of osteocalcin expression. In osteoblasts, in the absence of IGFl/insulin signaling, Foxo1 binds to Runx2 and inhibits Runx2 DNA binding to the OSE2 sites of the Bglap2 promoter, resulting in a reduction of osteocalcin (Ocn) expression. In the presence of IGFl/insulin signaling, Foxo1 is phosphorylated via the PI3K/AKT pathway and translocated from the nucleus to the cytoplasm where it binds to the 14-3-3 protein, thus eliminating the inhibition of Runx2. Runx2 serves as a transcriptional activator that binds to the Bglap2 promoter and increases osteocalcin expression, which favors glucose metabolism.

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ATF4 Is Required for the Anabolic Actions of PTH on Bone in vivo

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Parathyroid hormone (PTH) is a potent stimulator of bone formation and a proven anabolic agent for the treatment of osteoporosis. However, the mechanism whereby PTH increases bone formation remains poorly understood. Activating transcription factor 4 (ATF4) is a critical factor for bone formation during development and throughout postnatal life. This study examined if ATF4 is required for the anabolic actions of PTH on bone using an Atf4⁻/⁻ mouse model. Five-day-old wt and Atf4⁻/⁻ mice were given daily subcutaneous injections of vehicle (saline) or hPTH(1-34) (0.04 µg/g body weight) for 28 days. In wt mice, µCT analyses of femurs show that this PTH regimen significantly increased bone volume/tissue volume (BV/TV, 4.3-fold), trabecular thickness (Tb.Th, 50%), trabecular numbers (Tb.N, 1.5-fold), cortical thickness (Cort.Th, 77%), and cross-sectional area (CSA, 24%) and decreased trabecular spacing (Tb.Sp, 1.7-fold). These PTH effects were dramatically reduced or completely abolished in the absence of ATF4. Histological analyses show that PTH displayed potent anabolic effects on tibiae, vertebrae, and calvariae, which were significantly reduced in Atf4⁻/⁻ mice. At the molecular level, PTH markedly increased levels of osteocalcin (Ocn) and bone sialoprotein (Bsp) mRNA of long bones as measured by quantitative real-time RT/PCR. This increase was completely abolished in the absence of ATF4. This study demonstrates that ATF4 is required for the anabolic actions of PTH on bone in vivo and also suggested that modulation of the levels and activity of ATF4 may have therapeutic significance for the treatment of metabolic bone diseases such as osteoporosis.
TFIIA, ATF4, and Runx2 Synergistically Activate Osteoblast-specific Osteocalcin Gene Expression

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Runx2, a member of the runt homology domain family of transcription factors, is a master regulator of osteoblast function and bone formation. Mice lacking Runx2 have no mineralized skeleton due to a complete lack of mature osteoblasts. The expression level of Runx2 protein is regulated by a number of factors including BMPs, FGF-2, IGF-1, TNF-α, TGF-β, and PTH, all of which play important roles in osteoblasts and bone both in vitro and in vivo. In addition, the activity of Runx2 protein is positively or negatively modulated through protein-protein interactions. Activating transcription factor 4 (ATF4) is an osteoblast-enriched factor which regulates the terminal differentiation and function of osteoblasts. ATF4 knock-out mice have reduced bone mass and bone mineral density (severe osteoporosis) throughout their life. To identify proteins interacting with Runx2, we used a yeast two-hybrid system and identified TFIIA, a general transcriptional factor, as a Runx2-interacting factor. While pull-down assays confirmed that TFIIA physically interacted with Runx2 when both factors were coexpressed in COS-7 cells, surprisingly, it did not activate or inhibit Runx2-dependent transcriptional activity. In contrast, TFIIA unexpectedly activated ATF4, which we recently identified as a Runx2-interacting protein, in a dose-dependent manner. Deletion analysis found that this activation required the presence of the C-terminal 15 amino acid residues of ATF4 molecule. Finally, TFIIA, ATF4, and Runx2 synergistically stimulated the 0.657-kb mOG2 (mouse osteocalcin gene 2) promoter activity and endogenous osteocalcin mRNA expression. In summary, this study demonstrates a novel mechanism through which bone-specific transcription factors and general transcription factors cooperate in regulating osteoblast-specific gene expression.
Role of ATF4 in Osteoblast Proliferation and Survival

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ATF4 (activating transcription factor 4) is essential for bone formation. However, the mechanism of its actions in bone is poorly understood. The present study examined the role for ATF4 in the regulation of proliferation and survival of primary mouse bone marrow stromal cells (BMSCs) and osteoblasts. Results showed that Atf4−/− cells display a severe proliferative defect as measured by multiple cell proliferation assays. Cell cycle progression of Atf4−/− BMSCs was largely delayed with a significant G1 arrest. Expression of cyclins A1, D1, and D3 was dramatically decreased both at the mRNA and protein level. A similar proliferation defect was observed in Atf4−/− calvarial periosteal osteoblasts in vivo. Knocking down Atf4 mRNA by small interfering RNA in MC3T3-E1 subclone 4 preosteoblasts markedly reduced expression of the cyclins and cell proliferation. In contrast, overexpression of ATF4 increased cyclin D1 expression as well as cell proliferation. In addition, apoptosis was significantly increased in cultured Atf4−/− BMSCs and calvarial periosteal osteoblasts in vivo relative to wt controls. Taken together, this study demonstrates that ATF4 is a critical regulator of proliferation and survival in primary BMSCs and osteoblasts.
Activating Transcription Factor 4 Mediates the Anabolic Actions of Parathyroid Hormone in Bone. S. Yu¹, R.T. Franceschi⁴,⁵, M. Luo¹, J. Fan², D. Jiang⁴, H. Cao¹, Y. Lai³, J. Zhang¹, K. Patrene¹, K. Hankenson⁶, G. D. Roodman¹, G. Xiao¹. Department of Medicine¹, Surgery², and Pharmacology and Chemical Biology³, University of Pittsburgh, Pittsburgh, PA 15240; ⁴Department of Periodontics and Oral Medicine, School of Dentistry; ⁵Department of Biological Chemistry, School of Medicine, University of Michigan, Ann Arbor, MI 48109; ⁶Department of Animal Biology, School of Veterinary Medicine, University of Pennsylvania, Philadelphia, PA 19104-6010.

Parathyroid hormone (PTH) is a potent anabolic agent for the treatment of osteoporosis. However, its mechanism of action in osteoblast and bone is not completely understood. In this study, we show that the anabolic actions of PTH in bone are severely impaired in both growing and adult ovariectomized mice lacking bone-related activating transcription factor 4 (ATF4). Our study demonstrates that inactivation of the Atf4 gene in mice i) suppresses PTH-stimulated osteoblast proliferation and survival; and ii) abolishes PTH-induced osteoblast differentiation, which, together, compromise the anabolic response. We further demonstrate that intermittent PTH increases osteoblast differentiation in vivo at least in part through an ATF4-dependent up-regulation of Osterix (Osx). ATF4 stimulates Osx expression by activating Osx gene transcription. PTH activates Osx transcription through an ATF4 responsive element in the proximal promoter. ATF4 binds to an endogenous Osx promoter in a PTH-dependent manner. Taken together these experiments establish a novel role for ATF4 in the regulation of the anabolic response of osteoblast and bone to PTH.
CRITICAL ROLE OF ACTIVATING TRANSCRIPTION FACTOR 4 IN PARATHYROID HORMONE-MEDIATED BONE FORMATION

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University of Pittsburgh Medical Center

**Background and Objectives:** Parathyroid hormone (PTH) is a potent stimulator of bone formation and a proven anabolic agent for the treatment of osteoporosis. However, the mechanism whereby PTH increases bone formation remains poorly understood. The objective of this study was to test our hypothesis that activating transcription factor 4 (ATF4) mediates the anabolic response of skeleton to intermittent PTH.

**Materials and Methods:** Five-day-old wt and Atf4-/- mice were given daily subcutaneous injections of vehicle (saline) or hPTH(1–34) (60 ng/g body weight) for 28 days. Mice were euthanized 24 hours after the last injection. The effects of this PTH-dosing regimen on long bones (femurs and tibiae), vertebrae, and calvariae were determined by both biochemical and histomorphometric criteria.

**Results:** Quantitative uCT analysis of femurs showed that Atf4-/- mice had a significant reduction in bone volume/tissue volume (BV/TV), trabecular number (Tb.N), and cortical thickness (Cort.Th) and a marked increase in trabecular space (Tb.Sp) compared with the wt or Atf4+//- littermates. In wt femurs, intermittent PTH increased BV/TV, Tb.N, and trabecular thickness (Tb.Th) by 4.4-fold, 1.7-fold, and 50%, respectively, and decreased Tb.Sp by 1.5-fold. PTH-stimulated bone formation were dramatically reduced (BV/TV, Tb.N, Tb.Sp) or completely abolished (Cort.Th and Tb.Th) in Atf4-/- mice relative to wt mice. Histological analyses show that PTH displayed potent anabolic effects on tibiae, vertebrae, and calvariae, which were markedly reduced in Atf4-/- mice. ATF4 deficiency resulted in a drastic reduction in osteoprogenitors in bone marrow, which impairs the anabolic response of skeleton to PTH. This is probably related to the ability of ATF4 to promote proliferation of osteoblasts/osteoprogenitors as well as survival of osteoblasts and osteocytes, two critical steps for intermittent PTH to increase bone formation. In addition, ATF4 is required for PTH stimulation of osteoblast differentiation.

**Conclusion and Impact:** This study demonstrates that ATF4 is required for the anabolic actions of PTH in bone and suggests that modulation of the levels and activity of ATF4 may have therapeutic significance for the treatment of metabolic bone diseases such as osteoporosis.

This work was supported by the U.S. Army Medical Research and Materiel Command under W81XWH-07-1-0160 and by a National Institutes of Health grant DK072230.
Critical Role of ATF4 in Regulating Osteoclast Differentiation and Bone Resorption

Abstract Body

Activating Transcription Factor 4 (ATF4) has a critical role in regulating osteoblast differentiation and bone formation. However, its direct role in osteoclasts differentiation has not been addressed yet. In order to define the role of ATF4 in osteoclasts, we deleted ATF4 expression in mice or targeted expression of ATF4 to the osteoclast lineage using the tartrate-resistant acid phosphatase (TRAP) promoter. Osteoclast differentiation was severely impaired in primary bone marrow monocyte (BMM) cultures and bones from Atf4-/- mice. Coculture with wt osteoblasts or a high concentration of receptor activator of NF-kappaB ligand (RANKL) failed to restore the osteoclast differentiation defect in Atf4-/- BMM cultures. Conversely, Trap-ATF4-tg mice display a severe osteopenia with dramatically increased osteoclastogenesis and bone resorption. We further demonstrated that ATF4 is a novel upstream activator of the critical transcription factor nuclear factor of activated T cells c1 (NFATc1) gene and is critical for RANKL activation of multiple MAPK pathways in osteoclast precursors. Furthermore, ATF4 is crucial for M-CSF induction of receptor activator of NF-kappaB (RANK) expression on BMMs, and lack of ATF4 caused a shift in osteoclast precursors to macrophages, thereby increasing macrophages. Finally, ATF4 is largely modulated by M-CSF signaling and the PI3K/AKT pathways in BMMs. These results for the first time demonstrate that ATF4 plays a critical direct osteoclast-intrinsic role in regulating osteoclast differentiation and suggest that it may be a therapeutic target for treating bone diseases associated with increased osteoclast activity.

Keywords

ATF4, NFATc1, RANK

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Foxo1 Mediates IGF1/Insulin Regulation of Osteocalcin Expression by Antagonizing Runx2 in osteoblasts

Abstract Body
In this study, we determined the molecular mechanisms whereby forkhead transcription factor Foxo1, a key downstream signaling molecule of insulin-like growth factor 1 (IGF1)/insulin actions, regulates Runx2 activity and expression of the mouse osteocalcin gene 2 (Bglap2) in osteoblasts in vitro. We showed that Foxo1 inhibited Runx2-dependent transcriptional activity and osteocalcin mRNA expression and Bglap2 promoter activity in MC-4 preosteoblasts. Co-immunoprecipitation assay showed that Foxo1 physically interacted with Runx2 via its C-terminal region in osteoblasts or when coexpressed in COS-7 cells. Electrophoretic mobility shift assay demonstrated that Foxo1 suppressed Runx2 binding to its cognate site within the Bglap2 promoter. IGF1 and insulin prevented Foxo1 from inhibiting Runx2 activity by promoting Foxo1 phosphorylation and nuclear exclusion. In contrast, a neutralizing anti-IGF1 antibody decreased Runx2 activity and osteocalcin expression in osteoblasts. Chromatin immunoprecipitation assay revealed that IGF1 increased Runx2 interaction with a chromatin fragment of the proximal Bglap2 promoter in a PI3K/AKT-dependent manner. Conversely, knockdown of Foxo1 decreased Runx2 interaction with the promoter. This study establishes that Foxo1 is a novel negative regulator of osteoblastspecific transcription factor Runx2 and modulates IGF1/insulin-dependent regulation of osteocalcin expression in osteoblasts.

Keywords
Runx2, Foxo1, Osteoblast, IGF1/Insulin

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