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Research Article

Aging

Stem cells

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FOXP1 controls mesenchymal stem cell commitment and senescence during skeletal aging

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A hallmark of aged mesenchymal stem/progenitor cells (MSCs) in bone marrow is the pivot of differentiation potency from osteoblast to adipocyte coupled with a decrease in self-renewal capacity. However, how these cellular events are orchestrated in the aging progress is not fully understood. In this study, we have used molecular and genetic approaches to investigate the role of forkhead box P1 (FOXP1) in transcriptional control of MSC senescence. In bone marrow MSCs, FOXP1 expression levels declined with age in an inverse manner with those of the senescence marker *p16^{INK4A}*. Conditional depletion of *Foxp1* in bone marrow MSCs led to premature aging characteristics, including increased bone marrow adiposity, decreased bone mass, and impaired MSC self-renewal capacity in mice. At the molecular level, FOXP1 regulated cell-fate choice of MSCs through interactions with the CEBP β / δ complex and recombination signal binding protein for immunoglobulin κ J region (RBPjk), key modulators of adipogenesis and osteogenesis, respectively. Loss of *p16^{INK4A}* in *Foxp1*-deficient MSCs partially rescued the defects in replication capacity and bone mass accrual. Promoter occupancy analyses revealed that FOXP1 directly represses transcription of *p16^{INK4A}*. These results indicate that FOXP1 attenuates MSC senescence by orchestrating their cell-fate switch while maintaining their replicative capacity in a dose- and age-dependent manner.

Introduction

Bone aging, the main risk factor for primary osteoporosis, results in a decrease in bone mass and a parallel increase of BM adiposity (1–3). At the cellular level, BM mesenchymal stem/progenitor cells (MSCs), which are the common progenitors from which osteoblasts and adipocytes arise, undergo senescence along with bone aging (4). Anatomically, MSCs, defined as PDGF- α *Sca1*CD31*CD45⁻, reside primarily within the perivascular space — a region that can be specifically targeted by *nestin-Cre* (5), *Prx1-Cre* (6), or *Lepr-Cre* (7). As compared with young cells, senescent MSCs have reduced self-renewal capacity and predominantly differentiate into adipocytes as opposed to osteoblasts (8, 9). Such osteogenic or adipogenic commitment of MSCs is controlled by several crucial transcription factors. For example, PPAR γ and CEBP α / β / δ constitute an essential cascade for the adipocyte program, whereas RUNX2 and osterix are master regulators of osteoblast differentiation (10–12).

Cellular senescence, a state of irreversible growth arrest, is mainly controlled by the *p16^{INK4A}*/Rb and *p19^{ARF}*/p53 pathways (13, 14). *p16^{INK4A}* is required for the age-associated decline in function of a number of adult stem cells (15–18), including BM MSCs (19). *p53*-Deficient mice display osteoporosis and other aging-related features (20). Alterations in the transcriptional control or activity of these pathways account for the aging-related features detected in mice deficient for *Bmi-1* (21), *Brac1* (22), *Zmpste24* (23), *Hmga2*

(24), or *BUBR1* (25). In addition, Rb determines cell-fate choice and prevents premature aging of MSCs (26, 27). However, the molecular network orchestrating adipo-osteogenic balance and cellular senescence of MSCs remains largely unknown.

Forkhead box P1 (FOXP1) is a transcriptional factor that controls multiple cell-differentiation pathways, including embryonic stem cell pluripotency (28), T and B cell development (29–31), lung epithelial cell-fate determination (32–35), cardiomyocyte proliferation (33, 36), hair follicle stem cell quiescence (37), neuronal activity (38, 39), and glucose homeostasis (40). *FOXP1* haploinsufficiency in humans leads to deformity in craniofacial structure and speech ability (41). In this study, we observed that *Foxp1* deficiency in MSCs resulted in prototypical premature bone aging. These and additional results herein suggest that age-dependent bone loss may, in part, be orchestrated by the multifaceted action of FOXP1 during differentiation and senescence of MSCs.

Results

***Foxp1* expression in BM MSCs declines with age.** Nestin⁺ MSCs in BM represent a population that constitutes an essential hematopoietic stem cell (HSC) niche (5). Immunohistochemical (IHC) examination revealed colocalization of nestin and FOXP1 in a subset of stromal cells within neonatal BM in vivo (Figure 1A). Colocalization was validated in vitro in a subpopulation of MSC in cultures (Supplemental Figure 1A; supplemental material available online with this article; <https://doi.org/10.1172/JCI89511DS1>). Of the 4 alternatively spliced FOXP1 isoforms in mouse tissues (42), we detected 3 in BM MSCs: FOXP1A, FOXP1D, and FOX-

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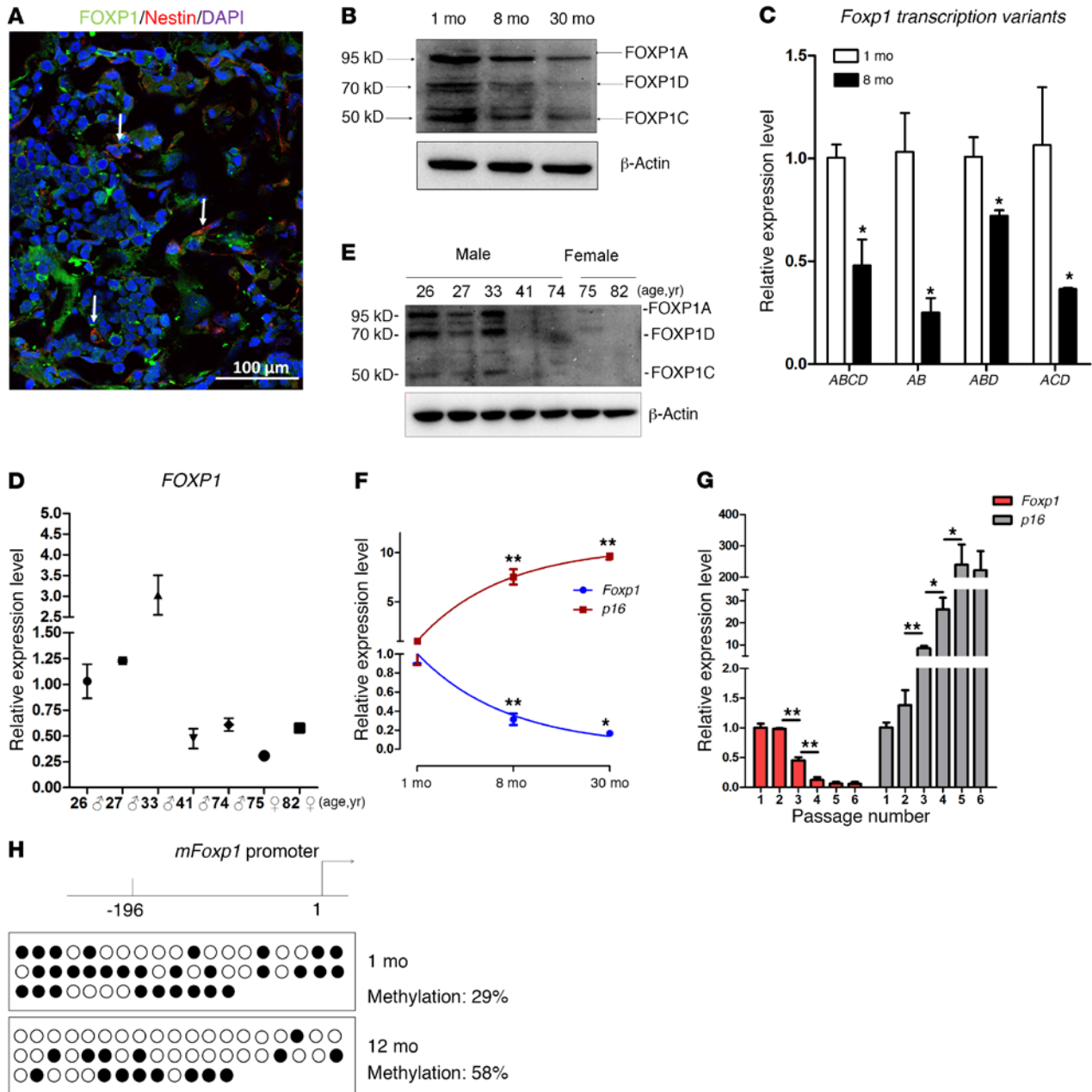


Figure 1. *Foxp1* expression declines with age in BM MSCs. (A) Representative image of IHC analysis. IHC showed overlapping of FOXP1 (green) and nestin (red) in fibroblast-like cells adjacent to endosteal (white arrows) in neonatal BM. Scale bar: 100 μ m. (B) Western blotting for the FOXP1 protein levels in BM MSCs. Western blotting for BM MSCs at 1, 8, and 30 months detected 3 major isoforms: FOXP1A (95 kD), FOXP1D (70 kD), and FOXP1C (50 kD). $n = 3$. (C) qPCR for the relative expression of *Foxp1* isoforms AB, ABD, ACD, and ABCD in MPCs from BM of 1 and 8 months old. $n = 3$. (D) Relative expression levels of *FOXP1* in young and aged hMPC as detected by qPCR. (E) Western blotting for FOXP1 protein levels in human BM MSCs from donors ages 26, 27, 33, 41, 74, 75 and 82 years. (F) Inverse expressions of *Foxp1* and *p16^{INK4A}* (*p16*) were assessed by qPCR in primitive MSCs obtained from BM of 1-, 8-, and 30-month-old mice. (G) Inverse correlation of *Foxp1* and *p16^{INK4A}* expression levels during in vitro expansion and passaging (P1–P6) of murine MSCs. $n = 3$. (H) Methylation of CpG islands within the *Foxp1* promoter variant 3 (–196 to 1) as detected by bisulfite sequencing in 1- and 12-month-old MSCs. Black circles represent methylated CpG islands and white circles unmethylated CpG islands. $n = 3$. * $P < 0.05$; ** $P < 0.01$.

P1C (Figure 1B and Supplemental Figure 1B). *Foxp1* expression declined progressively in mice during their progression from adolescence (1 month) to aged (8 to 30 months; Figure 1B). To avoid the heterogeneity of MSCs in cultures, we validated decreased expression of FOXP1 in 1- and 18-month-old mouse BM sorted by flow cytometry as CD29⁺Scal⁺CD31[–]CD45[–]Ter119[–] (Supplemental Figure 1, C and D). The expression decline was observed

for all 3 FOXP1 isoforms (Figure 1C). Of note, *FOXP1* expression in MSCs from human donors in ages ranging from 1- to 82 years was consistent with *Foxp1* expression levels in mice (Figure 1, D and E). Curiously, *Foxp1* levels in MSCs of different ages inversely correlated with those of the cellular senescence marker *p16^{INK4A}* (Figure 1F). This inverse correlation was recapitulated in cultures of MSCs expanded in vitro for 6 passages (P1 to P6; Figure 1G).

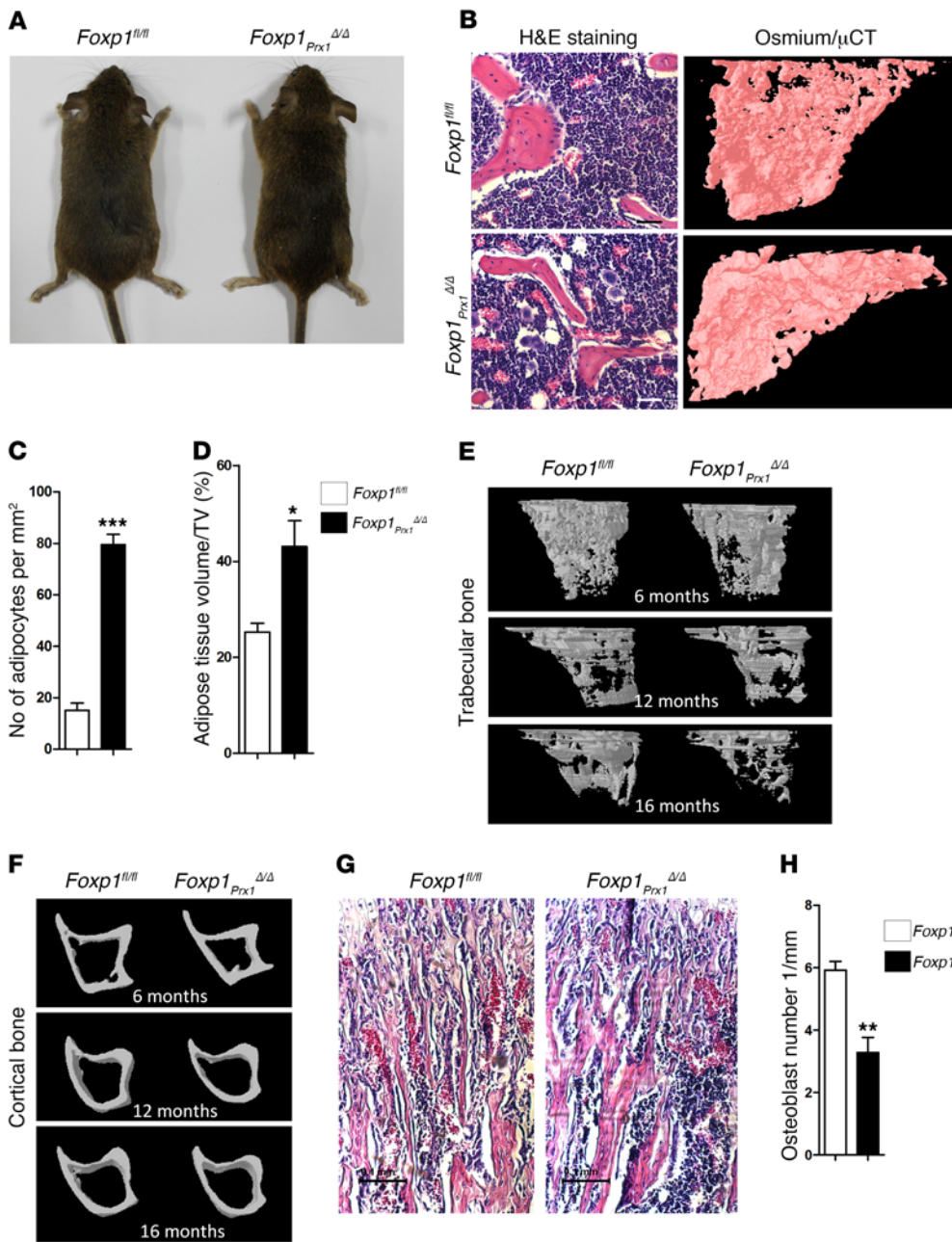


Figure 2. *Foxp1* in MSCs regulates bone mass accrual in an age-dependent manner. (A) Representative images of *Foxp1^{Prx1 Δ/Δ}* mutant and *Foxp1^{fl/fl}* mice at 3 months. (B) Left panel, representative H&E staining for adipose tissue (yellow arrowheads) in trabecular bones of tibia at 8 months of age. Right panel, representative images of osmium tetroxide staining followed by μ CT analyses in tibia BM at 6 months. Scale bar: 50 μ m. (C and D) Quantification of adipose droplet (C) and adipose tissue volume (D) in *Foxp1^{Prx1 Δ/Δ}* BM as compared with WT controls. TV, total tissue volume. $n = 5$. (E and F) Representative images of trabecular (E) and cortical bones (F) of tibia by μ CT analyses of *Foxp1^{Prx1 Δ/Δ}* mutant and *Foxp1^{fl/fl}* controls at 6, 12, and 16 months. (G) Representative H&E staining for primary ossification center of tibia bones at 6 months old. (H) Quantification of osteoblasts per trabecular bone surface in G. $n = 4$. Scale bar: 100 μ m * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

Meanwhile, the frequency of DNA methylation of CpG islands (43) within the proximal *Foxp1* promoter (nucleotides -196 to +1) nearly doubled (-29% to 58%) as MSCs aged from 1 to 12 months (Figure 1H). These observations indicated that FOXP1 expression inversely correlates with the progression of MSC aging.

Ablation of *Foxp1* in MSCs results in age-dependent bone loss. To test the role of FOXP1 in MSC differentiation and senescence, we inactivated it specifically in early BM MSCs by crossing a floxed (*fl*) allele *Foxp1^{fl/fl}* (31) with *Prx1-Cre* (44). Quantitative PCR (qPCR) and Western blotting confirmed efficient depletion of *Foxp1* expression (Supplemental Figure 2, A and B). *Prx1-Cre Foxp1^{fl/fl}* conditional KO mice (hereafter designated as *Foxp1^{Prx1 Δ/Δ}*) showed no changes in size, weight, or growth as compared with *Foxp1^{fl/fl}* controls (Figure 2A and data not shown). Notably, H&E staining revealed relative enrichment of adipose tissues within the BM of

Foxp1^{Prx1 Δ/Δ} mutants (Figure 2, B and C). In addition, the relative volume of BM adipose tissue was increased in *Foxp1^{Prx1 Δ/Δ}* mutants, as revealed by microcomputer tomography (μ CT) analyses following osmium staining (Figure 2, B and D). μ CT analyses further showed that *Foxp1^{Prx1 Δ/Δ}* mice displayed age-related bone loss as compared with *Foxp1^{fl/fl}* controls at 6, 12, and 16 months (Figure 2, E and F), as evidenced by a gradual decrease in bone mineral density (BMD) and in trabecular bone numbers (Supplemental Figure 2, C and D). This impaired bone mass accrual also was detected in 6-month-old female *Foxp1^{Prx1 Δ/Δ}* mutants (Supplemental Figure 2E). Histomorphometric analysis showed that osteoblasts per mm trabecular bone surface were decreased in mutants (Figure 2, G and H). The decline in BMD did not arise from enhanced osteoclast activity, as evidenced by tartrate-resistant acid phosphatase (TRAP) ($n = 3$) staining of osteoclasts in trabecular bones and

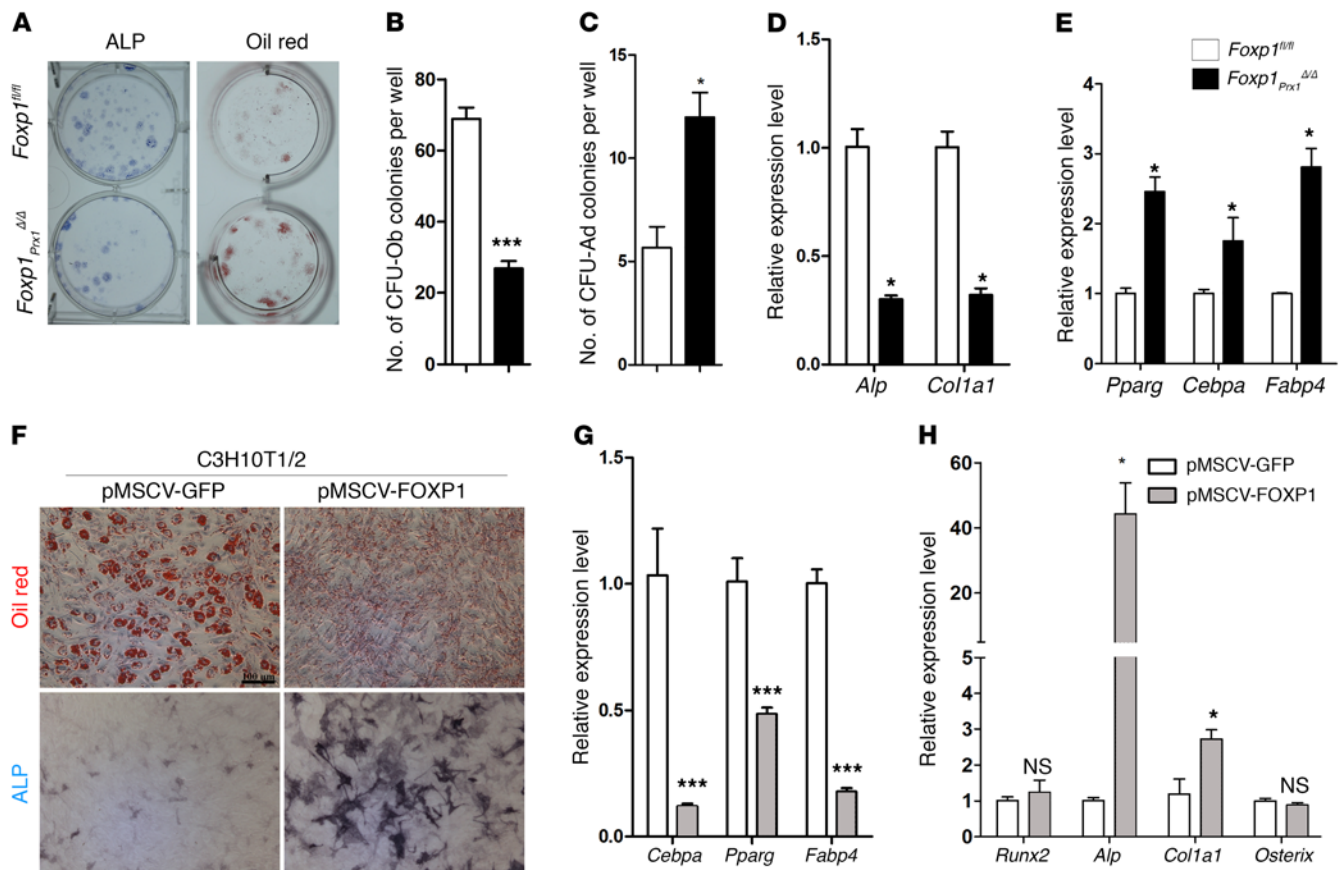


Figure 3. FOXP1 modulates cell-fate choice of MSCs between adipocytes and osteoblasts. (A) The osteogenic and adipogenic potential of MSCs from FOXP1 mutant BM was assessed 14 days after induction of differentiation by ALP and oil red O staining. $n = 5$. (B and C) Quantification of the frequencies of osteoblast clones (CFU-Ob) and adipocyte clones (CFU-Ad) following induction in A. $n = 3$. (D and E) Expression of bone (*Alp*, *Col1a1*) and fat (*Pparg*, *Cebpa*, and *Fabp4*) markers as assessed by quantitative reverse-transcriptase PCR (qRT-PCR) of uninduced MSCs in BM. $n = 3$. (F) *Foxp1* was overexpressed in C3H10T1/2 cells by retrovirus (pMSCV-FOXP1) and cultured in differentiation medium. Cell differentiation was assessed 6 days after adipogenic induction by oil red O staining or 14 days after osteogenic induction by ALP staining. $n = 3$. Scale bar: 100 μ m (G) The expression of adipogenic markers (*Cebpa*, *Pparg*, and *Fabp4*) was analyzed 6 days after adipogenic induction of FOXP1-expressing C3H10T1/2 cells. $n = 3$. (H) The expression of bone markers (*Runx2*, *Alp*, *Col1a1*, and *Osterix*) was assessed by qPCR 14 days after osteogenic culture of FOXP1-overexpressing C3H10T1/2 cells. $n = 3$. * $P < 0.05$; *** $P < 0.001$.

FACS analysis of osteoclast progenitor cells in BM at 3 months (Supplemental Figure 3). Collectively, these perturbations in bone and adipose tissue indicated that *Foxp1*-deficient bones displayed premature aging-related characteristics.

Foxp1 regulates the potential of MSCs to differentiate into adipocytes and osteoblasts. To investigate the role of *Foxp1* in cell-fate choice, MSCs were flushed from BM and induced to differentiate in vitro (detailed in Methods). Within 14 days following adipogenic or osteogenic CFU fibroblast (CFU-F) induction, the number of *Foxp1*^{*Prx1*} adipocyte CFUs (Ad-CFUs) significantly increased, whereas the number of mutant osteoblast CFUs (Ob-CFUs) decreased relative to controls (Figure 3, A–C). This fate switch in MSC differentiation potential was validated by parallel downregulation of osteoblast-specific transcripts (*Alp*, *Col1a1*) and upregulation of adipocyte-related markers (*Pparg*, *Cebpa*, and *Fabp4*) (Figure 3, D and E).

Next, we stably overexpressed *Foxp1* in C3H10T1/2 mesenchymal cells by retroviral transduction. We then assessed cell differentiation either by oil red O staining 6 days after adipogenic induction or by alkaline phosphatase (ALP) staining 14 days after

osteogenic induction (Figure 3F). As predicted by our KO results, enforced expression of *Foxp1* significantly enhanced osteogenic differentiation at the expense of adipogenic differentiation, as evidenced by increased or decreased levels, respectively, of the aforementioned osteoblast and adipocyte marker transcripts (Figure 3, G and H). Next, we transduced primary MSCs with a phosphorylated murine stem cell virus-driven (pMSCV-driven) FOXP1 retrovirus and examined differentiation by staining and expression of the same molecular markers employed for Figure 3, G and H (Supplemental Figure 4). Again, we observed that FOXP1 potentiated the osteogenic differentiation of MSCs.

To further validate the action of *Foxp1* in MSC differentiation, we generated *Nestin-Cre Foxp1*^{*fl/fl*} conditional KO mice (designated as *Foxp1*^{*Nes*}). *Nestin-Cre* targets a distinct population of MSCs within BM that constitute niches for HSC (5). *Foxp1*^{*Nes*} mutant mice displayed far more severe defects in bone growth than did *Foxp1*^{*Prx1*} mice (Supplemental Figure 5A and Supplemental Figure 6A). *Foxp1*^{*Nes*} mutants died within 6 weeks and displayed obvious growth arrest relative to controls, perhaps as a result of defects in neuronal activity (38, 39). In *Foxp1*^{*Nes*} mutants, the

osteogenic and adipogenic potential of MSCs was similarly altered, as evidenced by impaired bone formation (Supplemental Figure 5, B and C), enrichment of adipose tissue (Supplemental Figure 5, C and D), and altered numbers of Ad-CFU and Ob-CFU following MSC differentiation (Supplemental Figure 6, D–F) as well as by increased expression levels of PPAR γ and its downstream target gene, *Fabp4* (45) (Supplemental Figure 6G).

Given that *Prx1-Cre* and *nestin-Cre* target both osteoblast and osteocytes, the defects in bone mass accrual in *Foxp1_{Prx1}^{Δ/Δ}* and *Foxp1_{Nes}^{Δ/Δ}* mutants may result from loss of function of either. Thus, we crossed *Foxp1^{fl/fl}* mice with *Col2a1-Cre* (46), which targets osteoprogenitors and osteoblasts. We detected no changes in bone mass accrual as measured by μ CT analyses in *Foxp1_{Col2a1}^{Δ/Δ}* mutants as compared with WT controls at 3 months of age (Supplemental Figure 7, A and B). In addition, the loss of FOXP1 did not influence growth plate organization or chondrogenic differentiation during MSC induction (Supplemental Figure 7, C and D). These observations indicate that FOXP1 primarily exerts its influence on MSCs rather than chondrocytes or osteoblasts. Taken together, the defects in *Foxp1_{Prx1}^{Δ/Δ}* and *Foxp1_{Nes}^{Δ/Δ}* mutant mice indicate that *Foxp1* deficiency in MSCs favors adipogenesis over osteogenesis. Consequentially, FOXP1 is essential in establishing fate choice between bone and adipose tissue in vivo.

Foxp1 controls adipocyte and osteoblast differentiation by interacting with CEBP β / δ and RBP κ . Adipogenic differentiation of MSCs is initiated by a program of sequential activation of the transcriptional factors CEBP β / δ , CEBP α , and PPAR γ (10). We observed via coimmunoprecipitation (co-IP) assays that FOXP1 interacts with CEBP β and CEBP δ in 3T3-L1 cells (Figure 4, A and B) and in BM MSCs (Figure 4C). FOXP1 and CEBP β / δ colocalize within nuclei following transfection of C3H10T1/2 cells with vectors encoding either FOXP1-His/CEBP β / δ or FOXP1/CEBP β / δ -Flag (Figure 4D and Supplemental Figure 8B).

CEBP α and CEBP β / δ act upstream to induce *Pparg* transcription during terminal adipocyte differentiation (47). Luciferase reporter assays in C3H10T1/2 cells revealed that transactivation of *Pparg-Luc* by either CEBP α or CEBP β / δ was repressed by FOXP1 (Figure 4E and Supplemental Figure 8A). In contrast, expression of PPAR γ and FABP4 was relatively elevated in *Foxp1*-deficient MSCs compared with controls (Figure 4, F and G). We identified a consensus FOXP1-binding site –1112 bp upstream of the *Pparg* transcriptional start site (data not shown). ChIP-PCR indicated strong FOXP1 binding to that site (Figure 4, H and I), indicative of direct FOXP1 repression of *Pparg* transcription. Consistent with this interpretation, overexpression of FOXP1 repressed *Pparg* and *Fabp4* transcription following adipogenic differentiation of C3H10T1/2 cells (Figure 3G). Collectively, our observations suggest that a FOXP1-CEBP β / δ complex attenuates *Pparg* transcription to restrain adipogenic differentiation of MSCs.

Notch/recombination signal binding protein for immunoglobulin κ J region (RBP κ) signaling maintains MSC identity by suppressing osteoblast differentiation (48, 49). Via co-IP, we detected interaction of FOXP1 with RBP κ in both C3H10T1/2 mesenchymal cells and BM MSCs (Figure 4, J and K). Colocalization of FOXP1 and RBP κ protein within C3H10T1/2 nuclei was also detected (Figure 4D and Supplemental Figure 8B). Next, we observed that, through this interaction, FOXP1 repressed the acti-

vation of *RBP κ -Luc* via the intracellular domain of Notch (NICD) (Figure 4L). Additionally, *Foxp1_{Prx1}^{Δ/Δ}* BM MSCs exhibited elevated expression of *Hey1* and *Heyl* — 2 quintessential downstream targets of Notch signaling (Figure 4M). Accordingly, enforced expression of FOXP1 in C3H10T1/2 cells repressed expression of *Hey1* and *Heyl* (Figure 4N). These findings suggest that FOXP1 promotes osteogenic differentiation of MSCs through repression of Notch signaling during postnatal skeletal aging. However, further studies are needed to more robustly confirm this hypothesis.

Foxp1 attenuates MSC senescence through repression of p16^{INK4A} transcription. In terms of stem cell replication capacity, explants of *Foxp1_{Prx1}^{Δ/Δ}* MSCs displayed marked arrest in population doubling measured in vitro at 1 and 6 months (Figure 5A). Following 48-hour BrdU pulse-chase, FACS analyses revealed a smaller proportion of BrdU⁺ MSCs (CD31⁺CD45⁺CD29⁺Sca1⁺) in the BM of *Foxp1_{Prx1}^{Δ/Δ}* mice (Figure 5, B and C). We further observed that expression levels of cell cycle inhibitors *p16^{INK4A}*, *p21*, and *p27* were relatively increased in MSCs following isolation from *Foxp1*-deficient BM (Figure 5D). However, the number of CFU-F colonies began to progressively decrease at 6 months (Figure 5, E and F), consistent with a reduction in self-renewal ability upon loss of *Foxp1*. Expression levels of *p16^{INK4A}*, a marker for cellular senescence (18), were also significantly increased (Figure 5G). Yet expression of another 2 aging-associated markers, H3K9me3 and LAP2 β (50), was decreased in *Foxp1*-deficient MSCs (Figure 5G). Whereas second-passage MSC cultures of *Foxp1_{Prx1}^{Δ/Δ}* BM had relatively fewer Ki67⁺ cells than *Foxp1^{fl/fl}* controls (Figure 5, H and I), a greater number of γ H2AX⁺ and senescence-associated β -galactosidase activity (SA- β -gal⁺) cells as well as increased ROS levels were detected in mutant MSC cultures (Figure 5, H–K, and Supplemental Figure 8D). This decreased distribution of nuclear LAP2 β in *Foxp1*-deficient MSCs was confirmed by high-resolution IHC (Figure 5H) and was consistent with the Western blot data (Figure 5G). These in vivo and in vitro experiments indicated that *Foxp1*-deficient MSCs undergo a decline in self-renewal capacity. This, in turn, leads to swift accumulation of DNA damage and premature senescence during expansion.

Tight transcriptional control of the *p16^{INK4A}* locus is important in regulating senescence of a number of adult stem cell lineages during aging (13, 14, 51, 52). Promoter occupancy analysis identified FOXP1-binding sites within the *p16^{INK4A}* promoter (–1701 to –1695, Figure 6A). Luciferase reporter assays employing *p16-Luc* (driven by its essential 2.8 kb promoter) revealed that FOXP1 repressed activity of the *p16^{INK4A}* WT, but not the FOXP1-binding site–mutated *p16^{INK4A}* reporter (Figure 6, B and C). These findings suggested that FOXP1 restrains MSC senescence through repression of *p16^{INK4A}* transcription during skeletal aging.

To further test this hypothesis, we investigated the phenotypes of doubly deleted *Foxp1* and *p16^{INK4A}* conditional KO mice (*Foxp1_{Prx1}^{Δ/Δ}* *p16^{INK4A}*^{–/–}). Even though singular loss of *p16^{INK4A}* had shown no effect on MSC expansion or bone growth at 3 months of age as described previously (52), we observed a partial rescue of the replicative capacity in *Foxp1_{Prx1}^{Δ/Δ}* *p16^{INK4A}*^{–/–} double-mutant mice (Figure 6D). μ CT analyses of bone parameters showed that, as compared with single *Foxp1_{Prx1}^{Δ/Δ}* mutants, double-mutant mice displayed significantly increased volume, number, and BMD of trabecular bones (Figure 6, E and F). We also observed decreased anatomical separation of trabecular,

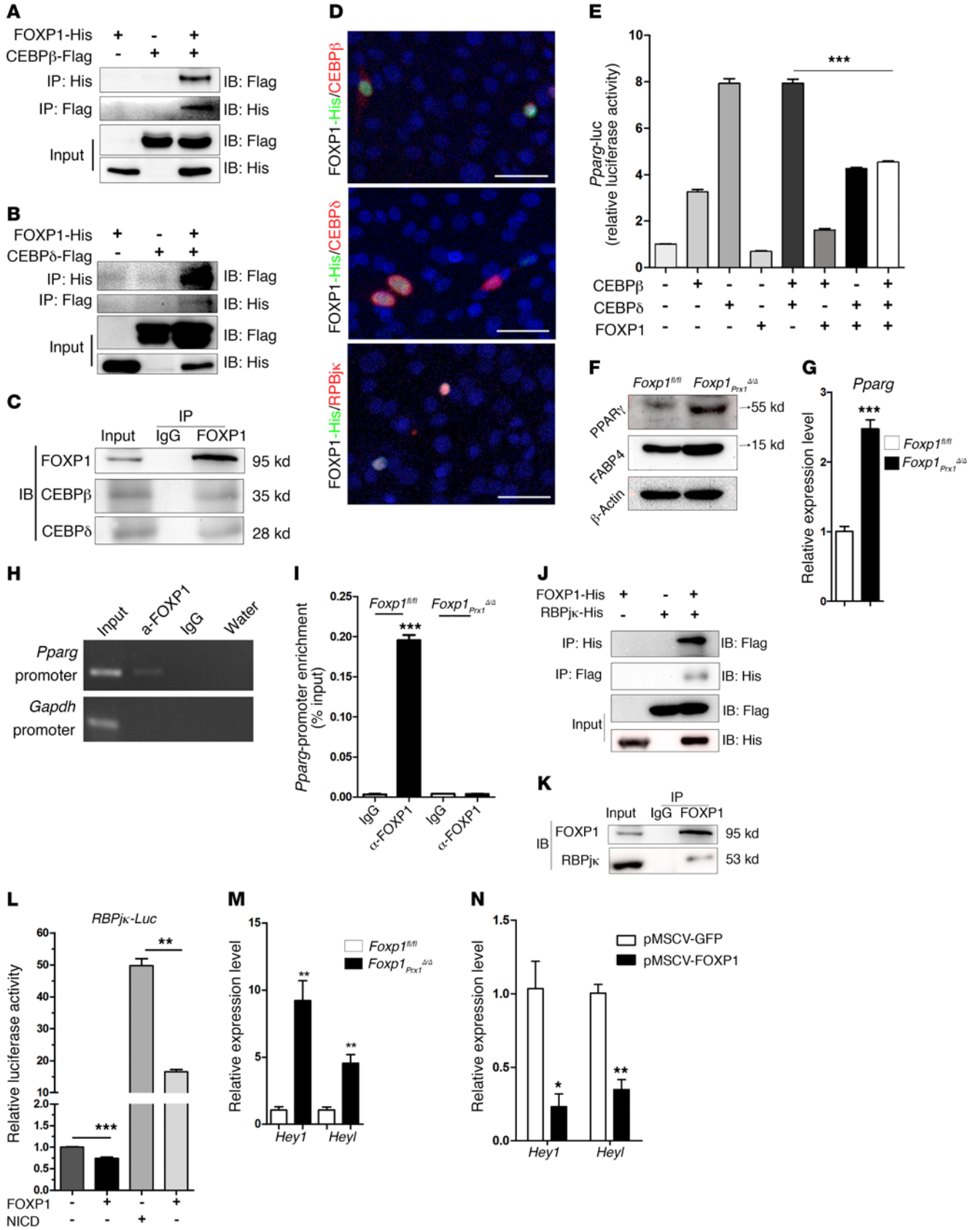


Figure 4. FOXP1 controls adipocyte and osteoblast differentiation by interacting with CEBP β / δ and RBPjk. (A and B) In vitro co-IP of FOXP1-His with CEBP β -Flag (A) or CEBP δ -Flag (B) following transfection into 3T3-L1 cells. (C) Western blotting detection of CEBP β and CEBP δ in FOXP1 immunoprecipitates from primary BM MSCs. (D) Colocalization of FOXP1 and CEBP β , CEBP δ or RBPjk in the nuclei of C3H10T1/2 cells transfected with FOXP1-His and CEBP β / δ - or RBPjk-expressing vectors. Green, anti-His antibody; red, antibodies for CEBP β , CEBP δ , or RBPjk; blue, DAPI staining for nucleus. Scale bar: 50 μ m. (E) FOXP1 represses the transactivation ability of CEBP β and CEBP δ in inducing *Pparg-Luc* luciferase activity in C3H10T1/2 cells. $n = 3$. (F) Western blot detection of PPAR γ and FABP4 levels in *Foxp1*^{Prx1^{-/-}/Δ/Δ} mutant BM at 8 months. (G) qRT-PCR of *Pparg* mRNA levels in MSCs. $n = 3$. (H and I) Promoter occupancy of *Pparg* gene as assessed by anti-FOXP1 ChIP-PCR in MSCs. (J and K) Co-IP of FOXP1 with RBPjk in C3H10T1/2 cells (J) and BM MSCs (K). (L) FOXP1 repression of luciferase activity as judged by *Rbpjk-Luc* reporter activity following the induction by NICD in C3H10T1/2 cells. $n = 3$. (M and N) qPCR confirms relatively higher expression levels of *Hey1* and *Heyl* in MSCs from *Foxp1*^{Prx1^{-/-}/Δ/Δ} mutant BM (M) or C3H10T1/2 cells transfected with pMSCV-FOXP1 (N). $n = 3$. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

but not of cortical, bones (data not shown). These results suggest that deletion of *p16^{INK4A}* partially compensates for defects in the replicative function of *Foxp1*-deficient MSCs.

Overexpression of FOXP1 in human mesenchymal progenitors augments their replication capacity. Given that *Foxp1* expression correlated with the aging progress of murine MSCs, we tested whether overexpression of FOXP1 could compromise senescence in human mesenchymal progenitor cells (hMPCs). Overexpression of lentiviral FOXP1 (Supplemental Figure 8E) within hMPCs collected from donors of both sexes at ages ranging from 27 to 82 years significantly augmented their replicative capacity, as assessed by measurement of population doubling (Figure 7, A-D). Of note, overexpression of FOXP1 hMPCs from a 74-year-old donor increased their expansion capacity to levels comparable to that of hMPCs from a 27-year-old donor (Figure 7D). This was coupled with a dramatic repression of *p16^{INK4A}* transcription within the fifth passage of FOXP1-overexpressing hMPCs (Figure 7E). We also observed that hMPCs transduced with *Foxp1* had greater osteogenic potential and reduced adipogenic potential, as evidenced by ALP and oil red O staining, by qPCR for osteogenic markers (*ALP*, *COL1A1*, *HEY1* and *HEYL*), and by Western blotting for adipogenic markers (PPAR γ and FABP4) (Figure 7, F-H). Collectively, our results indicate that FOXP1 is capable of augmenting the replication capacity of hMPCs irrespective of their age.

Discussion

Studies of mesenchymal progenitor-associated transcription factors in humans first reported that FOXP1 expression was higher in hMPCs than in their differentiated progenies (53). Expanding upon that observation, we identified FOXP1 as a dose-dependent orchestrator of MSC senescence and differentiation potency during skeletal aging. Our data suggest a model in which high levels of FOXP1 expression in young MSCs prevent their premature senescence, leading to an outcome that favors bone formation over adipogenesis (Figure 7I). In aged MSCs, a decline in FOXP1 levels alleviates negative regulation of *p16^{INK4A}* transcription and adipogenesis to promote the aging of MSCs (Figure 7I).

In a previous study, neither FOXP1 nor *p16^{INK4A}* expression was found to significantly decrease in MSCs from middle-aged to elderly donors (54). Perhaps FOXP1 expression in MSCs from middle-aged donors had already begun to decrease when their measurements were made. In support of this interpretation, *Foxp1* expression in 8-month-old murine MSCs decreased approximately 60% as compared with that in 1-month-old mice (Figure 1, B and F).

Epigenetic modifications, such as promotor-associated DNA methylation, appear to be a hallmark of MSC senescence

(55). DNA methylation within the *Foxp1* promoter region is increased with age, thereby potentially contributing to the progressive downregulation of *Foxp1* expression. That FOXP1 directly represses *Pparg* transcription in MSCs may, at least in part, explain the chronic increase of PPAR γ expression in senescent MSCs (56). PPAR γ acts as a master regulator of adipocyte differentiation by suppressing osteoblast differentiation (57–59). Thus, we contend that a decrease in FOXP1 levels, coupled with augmentation of PPAR γ expression, contributes to the increased adiposity in aged bones. We also found that FOXP1 may restrain the osteogenic potential of MSCs by suppressing Notch signaling (Figure 4, J–N). Collectively, our data indicate that FOXP1 acts as a critical effector of aging by controlling adipo-osteogenic balance during MSC differentiation.

A previous study claimed that siRNA-mediated knockdown (KD) of FOXP1 impaired adipogenesis, but not osteogenic differentiation, when tested in vitro (53). However, the efficacy and specificity of FOXP1 KD — a critical parameter for drawing this conclusion — were not presented in this study. Given the highly conserved sequence and redundant functions of FOXP1/2/4 isoforms (60), one cannot exclude the possibility that inefficient and/or mistargeted KD led to this conclusion.

Our results, derived both from in vivo and in vitro analyses, demonstrated a critical function for FOXP1 in fate choice of MSC differentiation. In addition, elevation of FOXP1 expression in aged hMPCs reversed their decline in expansion capacity and osteogenic potential. The current anabolic targets for osteoporosis, such as parathyroid hormone (PTH) or Wnt agonists, mainly promote osteoblastogenesis in bone formation (61). The multiple actions of FOXP1 in regulating MSC plasticity and senescence engender it as a potential anabolic target for osteoporosis therapy.

A central finding of our study is that FOXP1 attenuates MSC aging by directly regulating *p16^{INK4A}* transcription. Most relevant to this, FOXP1 overexpression has been associated with a variety of cancers (62–71). Given that the state of senescence can protect cells against the development of cancer (14), we submit that our findings may be instructive in understanding FOXP1-associated mechanisms of multiple tumorigenesis. Further studies are needed to address how the FOXP1/*p16* cascade protects multiple cell lineages against premature senescence or unlimited overgrowth.

Methods

Mice. *Foxp1^{fl/fl}* (29), *p16^{-/-}* (17), *Prx1-Cre* (72), and *nestin-Cre* (5) mice were generated as described elsewhere. The genetic backgrounds of all KO mice were uniform mixtures of 129S1/SvIMJ and C57BL/6. Mice were bred in pathogen-free conditions.

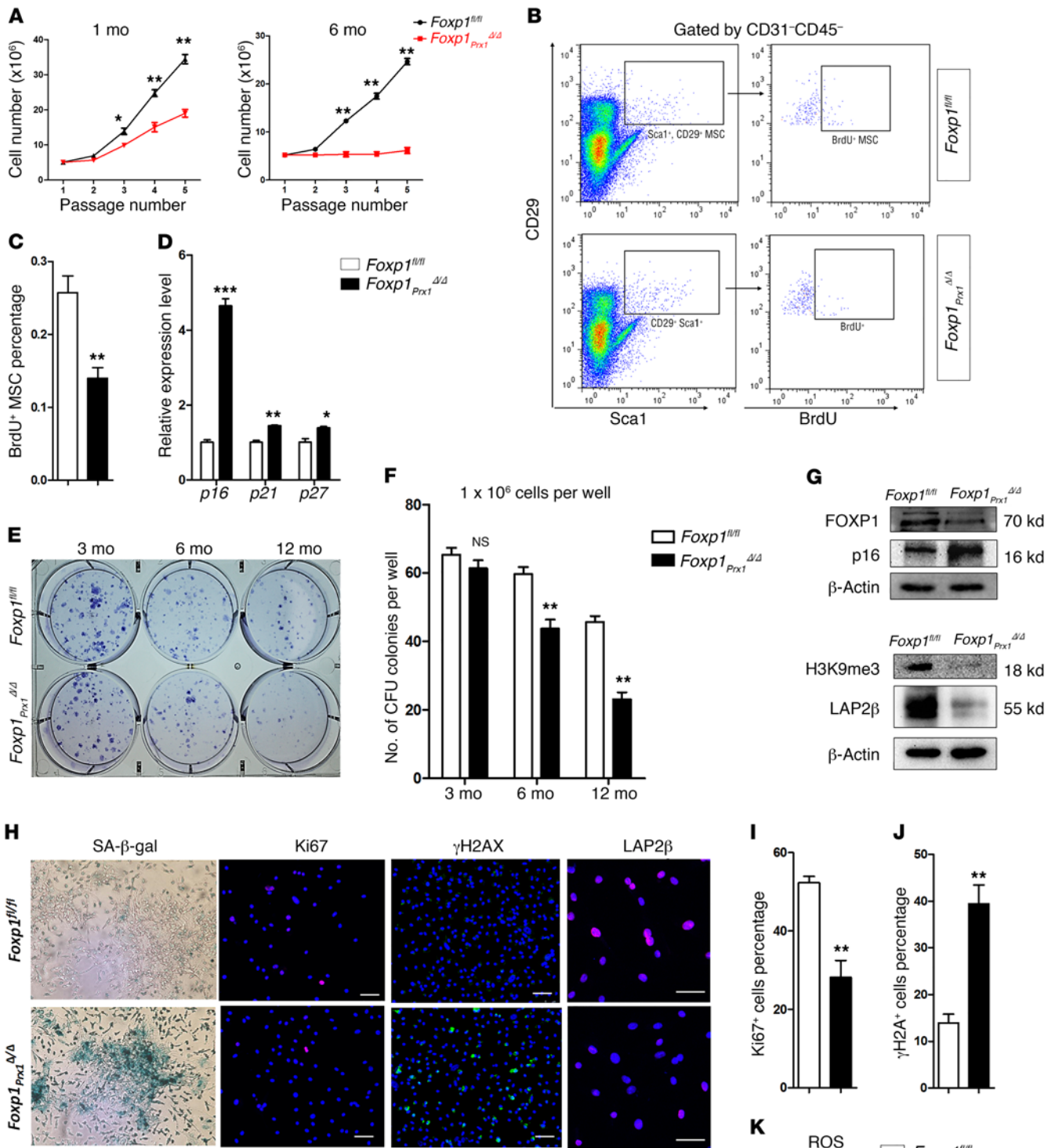


Figure 5. Ablation of *Foxp1* results in premature senescence of MSCs. (A) In vitro population doublings of MSCs from $Foxp1_{Prx1}^{\Delta/\Delta}$ BM at 1 and 6 months. $n = 4$. (B) FACS analysis dot plot of $BrdU^+$ labeled MSCs (defined as $CD31^+CD45^-CD29^+Sca1^+$) following 48-hour pulse-chase in BM of 3-month-old mice. $n = 3$. (C) Quantification of the percentages of $BrdU^+$ MSCs in 3-month-old mice. $n = 4$. (D) qPCR assessment of cell-cycle inhibitor ($p16^{INK4A}$, $p21$, and $p27$) expression in BM MSCs. $n = 3$. (E) Giemsa staining for the CFU-F colonies of BM MSCs from $Foxp1_{Prx1}^{\Delta/\Delta}$ mutant mice at 3, 6, and 12 months. $n = 3$. (F) Quantification of the number of CFU-F colonies in E. $n = 3$. (G) Western blot detection of FOXP1, $p16^{INK4A}$, H3K9me3, and LAP2 β protein levels in BM MSCs. (H) SA- β -gal staining, Ki67, γ H2AX, and LAP2 β immunostaining of the second passage of MSCs. Scale bars: 50 μ m. (I and J) Quantification of Ki67⁺ and γ H2A⁺ frequencies in MSCs from H. $n = 3$. (K) Quantification of the DCFDA fluorescence intensities reveals increased ROS levels in MSCs ($CD31^+CD45^-CD29^+Sca1^+$) from $Foxp1_{Prx1}^{\Delta/\Delta}$ BM at 3 months old. $n = 4$. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

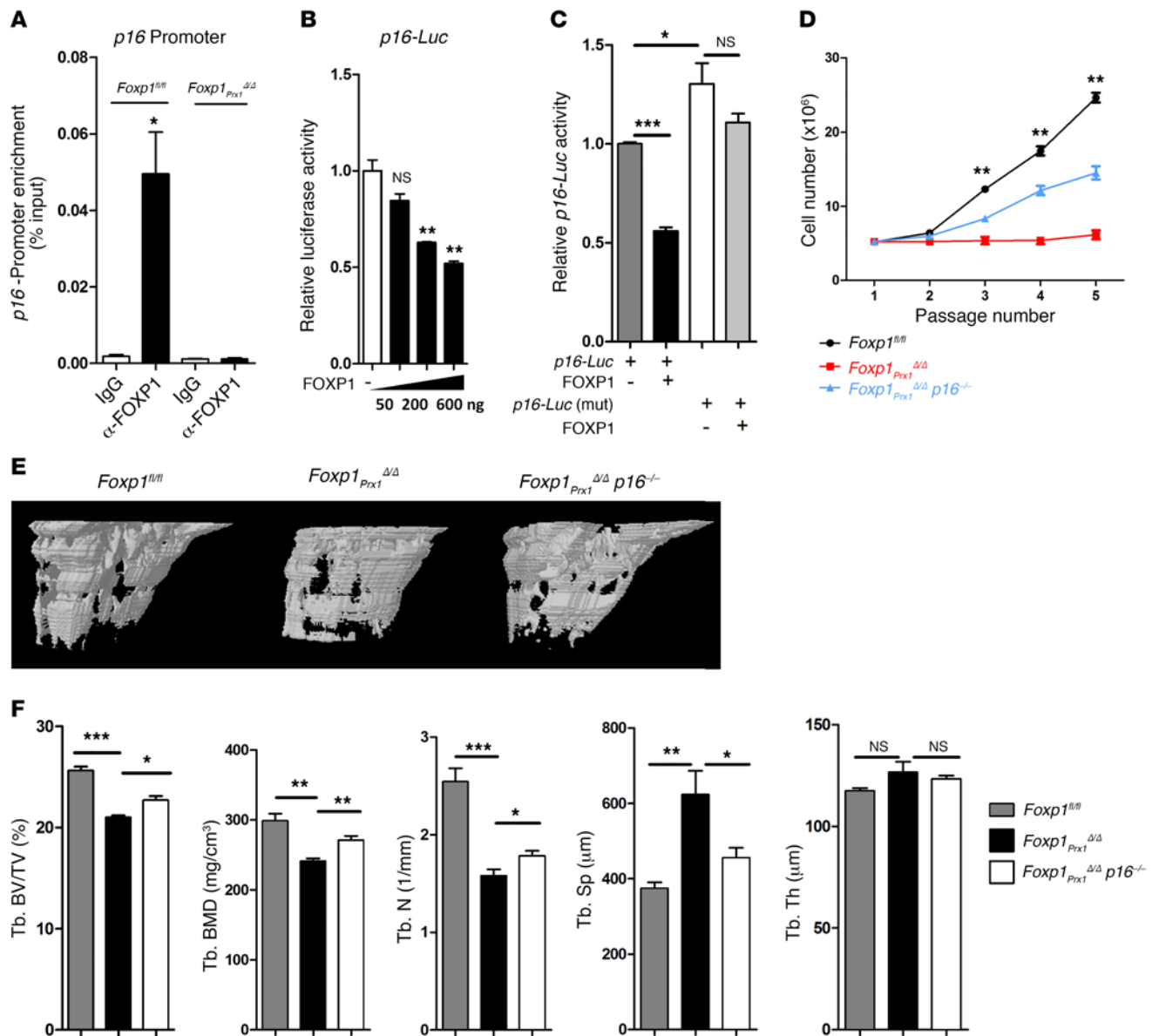


Figure 6. *Foxp1* attenuates MSC senescence through repression of *p16^{INK4A}* transcription. (A) Promoter occupancy of *p16^{INK4A}* as assessed by anti-FOXP1 ChIP-PCR in MSCs. Sequence analysis identified a consensus FOXP1-binding site within the *p16^{INK4A}* promoter (1701-1695). (B) FOXP1 repression of *p16^{INK4A}* transcription as revealed by *p16-Luc* reporter assays in C3H10T1/2 cells transfected with the indicated levels of cotransfected FOXP1. (C) FOXP1 repressed transcription of *p16-Luc* reporter, but not the mutant reporter *p16-Luc* (mut). (D) Doubling times of BM MSCs expanded in vitro. The replication capacity of *Foxp1^{Prx1}^{Δ/Δ} p16^{-/-}* double mutants is significantly higher than that of *Foxp1^{Prx1}^{Δ/Δ}* single mutants, but lower than that of control mice. $n = 4$. (E) Representative images of μ CT analyses of trabecular bones of tibia in 3-month single (*Foxp1^{Prx1}^{Δ/Δ}*) and double (*Foxp1^{Prx1}^{Δ/Δ} p16^{-/-}*) mutants. (F) Bone volume, BMD, and number of trabecular bones are significantly rescued in *Foxp1^{Prx1}^{Δ/Δ} p16^{-/-}* double-mutant compared with *Foxp1^{Prx1}^{Δ/Δ}* single-mutant mice. $n = 4$. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$. Tb. BV/TV, trabecular bone volume/total volume; Tb. N, trabecular number; Tb. Sp, trabecular spacing; Tb. Th, trabecular thickness.

qPCR. Total RNA was extracted with TRIzol reagent (Invitrogen), and cDNA was generated using the GoScript Reverse Transcription System (Promega). Real-time qPCR was performed with the ABI 7500 System (Applied Biosciences) using SYBR Green (Roche). The primers used for qPCR are listed in Supplemental Table 1.

Osmium tetroxide staining and μ CT analysis. Femurs or tibias were dissected from mice and fixed in 70% ethanol at 4°C. To visualize and quantify BM fat, bones were decalcified and stained by osmium tetroxide as described previously (73). μ CT scanning of bone and fat was performed on SkyScan 1176 (Bruker). A 3D model was reconstructed, and structural indices were calculated using

CTAn software. The region of interest (ROI) selected was 5 mm below the growth plate of bones.

Cell cultures. Mouse MSCs were enumerated and expanded using the mouse MesenCult Proliferation Kit (STEMCELL Technologies) according to the manufacturer's protocols. Briefly, mouse BM cells from tibia and femur were flushed out with 2 ml IMDM with 2% FBS and filtered through a 70- μ m cell strainer (BD Falcon). Nucleated cells were counted using acetic acid with methylene blue. Cell numbers and volumes were adjusted using MesenCult medium (STEMCELL Technologies) according to assay requirements. Cell lines C3H10T1/2 and 3T3-L1 were obtained from ATCC.

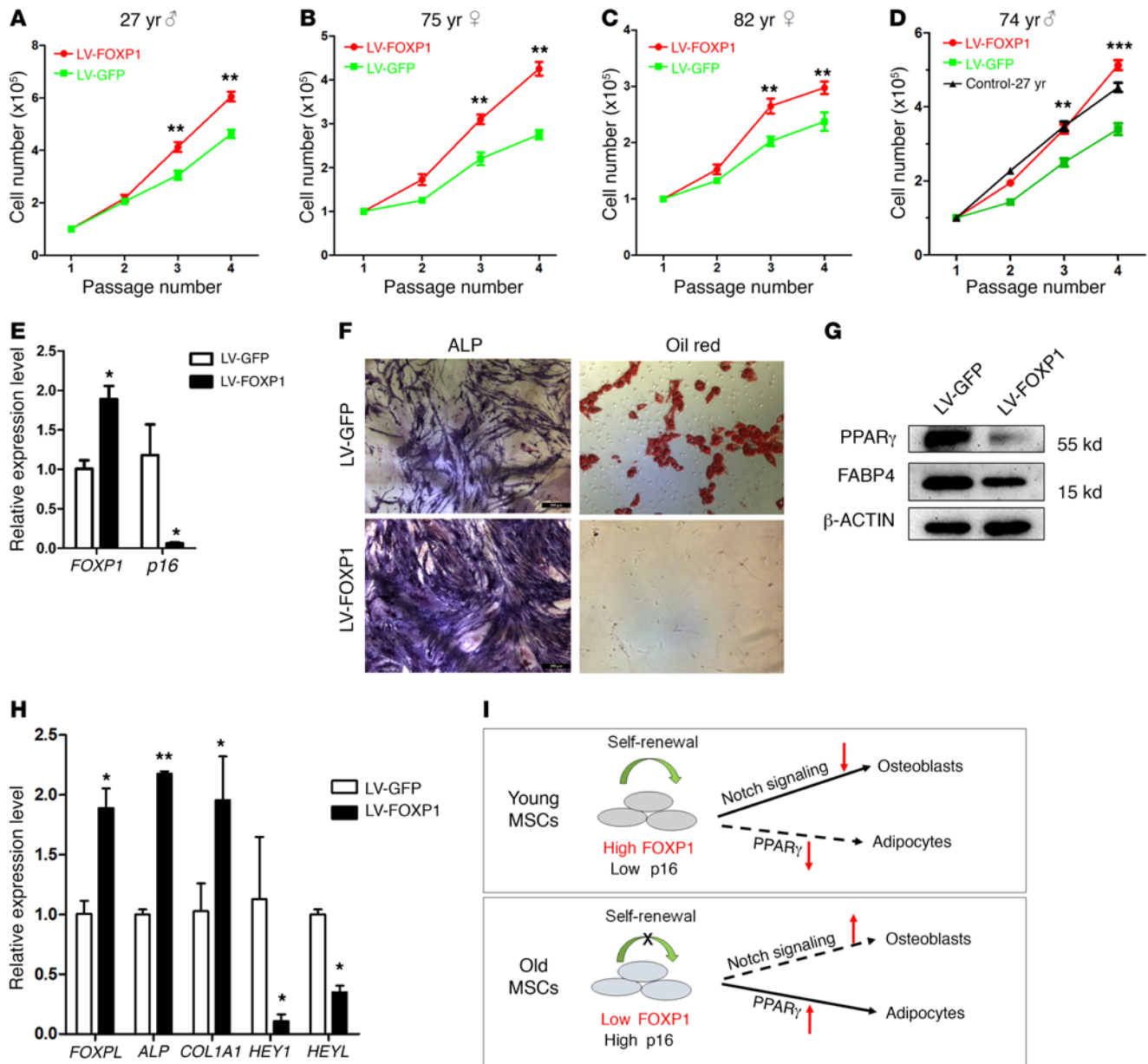


Figure 7. Overexpression of FOXP1 in hMPCs augments their expansion capacity. (A–D) In vitro expansion of hMPCs isolated from BM donors ages 27 (male, A), 75 (female, B), 82 (female, C), and 74 (male, D) years. MPCs were transfected with lentiviral LV-FOXP1 or LV-GFP. Control 27 yr, hMPCs were from a 27-year-old male donor. (E) Expression of *FOXP1* and *p16^{INK4A}* in the fifth passage of hMPCs with *FOXP1* overexpression from a 74-year-old donor. (F) Representative images of ALP and oil red O staining 14 days after osteogenic differentiation of the 74-year-old donor’s hMPCs transfected with LV-FOXP1. Scale bar: 100 μ m. (G) Western blot for PPAR γ and FABP4 expression in adipogenic cultures of hMPCs in F. (H) qPCR of *FOXP1*, *ALP*, *COL1A1*, *HEY1*, and *HEYL* expression in osteogenic cultures of FOXP1-expressing hMPCs in F. *n* = 3. (I) Model by which FOXP1 regulates BM MSC aging. In young MSCs, high expression of *Foxp1* restrains the activation of *p16^{INK4A}* to maintain the replication capacity of MSCs while sustaining osteogenic rather than adipogenic differentiation. In old MSCs, reduction of *Foxp1* expression releases *p16^{INK4A}* repression, leading to impaired MSC replication capacity while promoting adipogenic differentiation at the expense of osteogenic differentiation. **P* < 0.05; ***P* < 0.01; ****P* < 0.001.

For mouse CFU-F assays, 2 million BM cells were plated per well in 6-well plates, each group was plated in duplicate or triplicate, and cells were cultured for 14 days at 37°C in 5% CO₂. CFU-F-derived colonies were stained by Giemsa staining solution and enumerated. For CFU-Ob assays, 1 million BM cells were plated on 35-mm dishes in 2 ml complete MesenCult medium. After approximately 7 to 9 days, osteogenic medium (10 nM dexamethasone [Sigma], 10 mM β -glycerophosphate, and 50 μ M ascorbic acid [Sigma] in α -MEM

[Hyclone] containing 10% FBS [Hyclone]) was added. At 14 days, cells were fixed and stained for ALP using TRACP & ALP Double-Stain Kit (Clonotect) or Alizarin red S. For CFU-Ad assays, 2 million BM cells were plated per well in 24-well plates with 0.5 ml Complete MesenCult Medium (STEMCELL Technologies). Culture conditions were changed to adipogenic medium (DMEM supplemented with 10% FBS, 1 μ M dexamethasone, 10 μ g/ml insulin, 0.5 mM 3-isobutyl-1-methylxanthine, and 200 μ M indomethacin) when cells reached 80% to 90%

confluence. Cells were cultured for 1 week before oil red O staining. Oil red O staining was performed according to standard protocols. For population doubling assays, the first passaged mouse MSCs were seeded in 10-cm plates with 5×10^6 cells and hMPCs were seeded in 35-mm dishes with 1×10^5 cells. Medium changes were performed every 3 days and cells passaged at weekly intervals. Cells were counted at each passage to calculate population doublings.

hMPCs were isolated from vertebrae or femur bones of trauma patients between 20 and 80 years old. hMPCs were then separated using the Human Marrow Lymphocyte Medium Kit (Sangon Biotech) and cultured using the MesenCult Proliferation Kit (human) (STEM-CELL Technologies). For osteogenic differentiation of human mesenchymal stem cells, 0.1 million hMPCs were seeded per well of 24-well plates with the same medium for CFU-Ob assays above. After 9 days of induction, cells were fixed and stained for ALP.

Viral infection. To obtain transformants stably expressing FOXP1, pMSCV-FOXP1-puro and pMSCV-GFP-puro control retroviruses were transfected into Plat-E packaging cells using FuGENE HD (Promega). After 48 hours, culture supernatants were collected and used for infection. Retroviruses were introduced into C3H10T1/2 cells, and stable transformants were selected with $5 \mu\text{g/ml}$ puromycin. For osteoblast and adipocyte differentiation, cells were cultured in osteogenic medium and adipogenic medium described above, respectively. For lentivirus infections, hMPCs were plated at a density of 1×10^5 cells in 3.5-mm dishes. LV-FOXP1 or LV-GFP virus was added at a multiplicity of infection of 20, and cells were exposed to virus for 24 hours and selected with $1 \mu\text{g/ml}$ puromycin for 3 days. Lentiviruses were obtained from Hanbio.

ChIP assay. Briefly, after MSCs were isolated from mouse BM and cultured for 2 weeks, chromatin was crosslinked with 1% formaldehyde and sheared to 300- to 500-bp fragments by sonication (Sonics VCX130, 30% amplitude, 5 seconds on and 10 seconds off for 16 cycles). FOXP1 antibody (Millipore, ABE68, 1:200) and normal rabbit IgG (Santa Cruz Biotechnology Inc., sc-2027, 1:80) were used to immunoprecipitate the relevant protein-DNA complex. The cross-linked DNA and protein were reversed by addition of 5 M NaCl; DNA was extracted by phenol/chloroform and precipitated with ethanol and glycogen. Purified DNA was quantified using real-time PCR. The primer sequences are listed in Supplemental Table 1.

Luciferase reporter assay. Luciferase assays were performed in HEK293T, C3H10T1/2, or 3T3-L1 cells. The reporter plasmid *Pparg-Luc*, containing a 2.2-kb fragment of the 5' flanking region of the *Pparg* gene, was obtained from Hiroshi Takayanagi (Tokyo Medical and Dental University, Tokyo, Japan). The RBP-JK luciferase reporter plasmid and expression plasmid with NICD2 were provided by Lianping Xing (University of Rochester Medical Center, Rochester, New York, USA). The *p16^{INK4A}-luc* reporter plasmid, which contains a 2.8-kb fragment of the 5' flanking region of the *p16^{INK4A}* gene, was constructed in our lab. The mutant *p16^{INK4A}-luc* reporter plasmid was generated by replacing the putative FOXP1-binding site (TATTTAT) at position -1701 with the random sequence GCGGGCG. The expression plasmids of FOXP1, CEBP α , CEBP β , and CEBP δ were constructed into the pcDNA3.0 vector. Cells were transfected using Lipofectamine 2000 (Invitrogen) in 24-well plates. The transfection amount of each plasmid was 200 μg , and the total amount of transfected DNA was kept consistent across each transfection condition of pcDNA3.0 empty plasmids where necessary. After 32 hours, dual luciferase assay was performed according to the manufacturer's protocols (Promega).

co-IP. For in vitro co-IP, His- or FLAG-tagged proteins were produced in HEK293T cells transfected by Lipofectamine 2000 with corresponding plasmids. For in vivo co-IP, BM MSCs were isolated and cultured from WT mice. Total cell lysates were incubated overnight at 4°C with antibodies (described below) or normal IgG (Santa Cruz Biotechnology Inc., sc-2027) as control. Antibody-antigen complexes were precleared with Protein A/G PLUS-Agarose (Santa Cruz Biotechnology Inc., sc-2003). After several washes, samples were boiled and analyzed by immunoblot.

Western blot. Cells were lysed by NP40 lysis buffer (Beyotime Biotechnology) supplemented with protease inhibitor and phosphatase inhibitor cocktails (Selleck). Protein samples were separated by SDS-PAGE, transferred to a nitrocellulose filter membrane (Bio-Rad), blocked with 5% nonfat milk (in TBST), and incubated with primary antibodies against FOXP1 (Millipore, ABE68, 1:1000), C/EBP β (Santa Cruz Biotechnology Inc., sc-150, 1:500), C/EBP δ (Santa Cruz Biotechnology Inc., sc-151, 1:500), FABP4 (Abcam, ab13979, 1:1000), RBPj κ (Santa Cruz Biotechnology Inc., sc-28713, 1:500), γ H2A antibody (GeneTex, GTX11174, 1:1000), Bmi (Abcam, ab14389, 1:1000), PPAR γ (Santa Cruz Biotechnology Inc., sc-22020, 1:500), p16^{INK4a} (Millipore, 04-239, 1:1000), H3K9me3 (Abcam, ab8898, 1:1000), LAP2 β (BD, 611000, 1:1000), His-Tag (MBL, M136-3, 1:2000), FLAG (Agilent, 200471, 1:2000), or β -actin (Selleck, A1016, 1:2000) at 4°C overnight. Proteins were visualized using horseradish peroxidase-conjugated (HRP-conjugated) secondary antibody and chemiluminescent HRP substrate (Millipore).

Immunohistochemistry and confocal microscope. Cultured MPCs were fixed in 4% PFA for 10 minutes at room temperature. After blocking with 10% normal goat serum, cells were then incubated with γ H2A antibody (GeneTex, GTX11174, 1:500), Ki67 antibody (Abcam, ab15580, 1:500), and LAP2 β (BD, 611000, 1:500), followed by incubation with Alexa Fluor 488- or Alexa Fluor 594-conjugated secondary antibodies. For SA- β -gal staining, cells were fixed with 0.5% glutaraldehyde in PBS (pH 7.2) for 5 minutes at room temperature, then washed with PBS containing MgCl₂ (pH 7.2, 1 mM MgCl₂) and stained with X-gal solution (1 mg/ml X-gal, 0.12 mM K₃Fe[CN]₆, 1 mM MgCl₂ in PBS, pH 6.0) overnight at 37°C.

Bones were fixed in 4% PFA for 24 hours at 4°C, decalcified in 10% EDTA (pH 7.4) for 21 days, embedded in paraffin or tissue-freezing medium (Leica), and sectioned to 8 μm . For immunofluorescence, heat-induced antigen retrieval with sodium citrate buffer (10 mM sodium citrate, 0.05% Tween 20, pH 6.0) was performed before bone sections were blocked with 10% normal serum containing 1% BSA in TBST (pH 7.6) for 2 hours at room temperature, then incubated overnight at 4°C with primary antibodies to mouse FOXP1 (Millipore, ABE68, 1:100) and Nestin (Abcam, ab11306, 1:100) diluted in TBST with 1% BSA. Subsequently, sections were incubated with secondary fluorescent-conjugated antibodies at room temperature for 2 hours in the dark. Samples were imaged by the Leica TCS SP5 confocal microscope, the Leica DM2500, or the 3000B microscope. H&E staining was conducted according to standard protocols.

Flow cytometry analysis. For BrdU labeling of MSCs, mice were intraperitoneally injected with BrdU (100 $\mu\text{g/g}$ of body mass, Sigma-Aldrich) and sacrificed 48 hours later. BM cells were flushed out and collected in DMEM with 2% FBS, then diluted to 8 million cells in 400 μl of medium and incubated with anti-mouse CD45-APC (17-0451, eBioscience), CD11b-APC (17-0112, eBioscience), Ter119-APC (17-5921, eBioscience), CD31-APC (17-0311, eBioscience), Sca1-PerCP (45-5981,

eBioscience), and CD29-PE antibodies (12-0291, eBioscience) in the dark for 30 minutes at 4°C. After surface staining, cells were fixed with 1% PFA and permeabilized with 0.5% Tween-20. DNA was fragmented by DNase I; then cells were stained with anti-BrdU-FITC (11-5071, eBioscience) antibody or DCFDA for 45 minutes at room temperature. Cells were acquired on a BD FACSCalibur, and analysis was performed with FlowJo software, version 7.6. For MSC sorting, BM cells were stained with CD45-APC, Ter119-APC, CD31-APC, Sca1-PerCP, and CD29-PE antibodies and sorted on a BD FACSAria II.

Genomic DNA isolation and bisulfite sequencing PCR. MSC genomic DNA was extracted with the TIANamp genomic DNA kit (TIANGEN), and bisulfite sequencing PCR was performed with Oebiotec reagents. The sequencing primers were 5'-ATAGTAATTAAGAG-GAGTTGTTGGGG-3' and 5'-CCTAACACTCTCCATATAACCRC-3'.

Statistics. All data are presented as mean ± SEM. Error bars indicate SEM. Nonlinear regression analysis with exponential 2-phase decay was used for *Foxp1* and *p16* expression. Two-tailed Student's *t* tests were used for comparisons between 2 groups, and 1-way ANOVA with Bonferroni's post-hoc analysis was used for multiple comparisons (3 or more groups). *P* < 0.05 was considered significant.

Study approval. All murine studies were conducted under the approved Shanghai Jiao Tong University IACUC protocol SYXK 2011-0112. For all human studies, written informed consent was received

from participants prior to inclusion in the study where required. All samples were obtained in accordance with standard protocols of the Bioethics Committee of Bio-X Institutes at Shanghai Jiao Tong University.

Author contributions

HL, PL, SX, ZY, and XG designed and performed experiments. YL, ZZ, YH, GY, TT, YF, and YR provided human biospecimens. BL, JDD, and HOT provided mouse lines. XG wrote the manuscript.

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