Rb regulates fate choice and lineage commitment in vivo

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Abstract

Mutation of the RB-1 tumour suppressor occurs in one third of all human tumours and is particularly associated with retinoblastoma and osteosarcoma. Numerous functions have been ascribed to the product of the human RB-1 gene, pRB. The best known is pRB’s ability to promote cell cycle exit through inhibition of the E2F transcription factors and the transcriptional repression of genes encoding cell cycle regulators. In addition, pRB has been shown in vitro to regulate several transcription factors that are master differentiation inducers. Depending on the differentiation factor and cellular context, pRB can either suppress or promote their transcriptional activity. For example, pRB binds to Runx2 and potentiates its ability to promote osteogenic differentiation program in vitro. In contrast, pRB acts together with E2F to suppress PPARγ, the master activator of adipogenesis. Since osteoblasts and adipocytes can both arise from mesenchymal stem cells, these observations suggest that pRB might play a role in the choice between these two fates. However, to date, there is no evidence for this in vivo. Here we use mouse models to address this hypothesis in the context of mesenchymal tissue development and tumorigenesis. Our data show that Rb status plays a key role in establishing fate choice between bone and brown adipose tissue in vivo.
white fat, tumors fits with prior studies showing that Rb loss promotes brown fat over white fat differentiation11,12,13. Genotyping confirmed that the tumour cells had undergone Cre-mediated recombination of Rb and/or p53 (Figure 1c, data not shown). Moreover, it showed that the Prx1-Cre;Rb<sup>fl/fl</sup>;p53<sup>fl/fl</sup> tumours consistently retained the wildtype Rb allele (Figure 1c, data not shown). Thus Rb acts in a dose dependent manner to modulate the spectrum of tumours arising from p53-deficient, uncommitted mesenchymal stem cells: osteosarcomas predominant in the presence of Rb, while Rb loss strongly favours hibernoma formation.

Given that Rb loss in p53 mutant uncommitted mesenchymal cells disfavours osteosarcoma formation, we also investigated the affect of Rb loss in a bone-committed progenitor. For this, we deleted Rb and/or p53 using the Osx-Cre transgenic14 which uses Osterix promoter sequences to express Cre in the pre-oestoblast (Supplementary Figure 1d). In this model15, Osx-Cre;p53<sup>fl/fl</sup> mice specifically develop osteosarcoma (100%) while Osx-Cre;Rb<sup>fl/fl</sup>;p53<sup>fl/fl</sup> develop osteosarcoma (53%), hibernomas (46%) and sarcomas (2%). We established cell lines from multiple (≥3) independent Osx-Cre;p53<sup>fl/fl</sup> and Osx-Cre;Rb<sup>fl/fl</sup>;p53<sup>fl/fl</sup> osteosarcomas and discovered that the two genotypes have distinct differentiation properties (Figure 2, data not shown). The Rb;p53 (DKO) osteosarcoma (OS) cell lines expressed mRNAs that are characteristic of bone and fat differentiation (Figure 2a). Indeed, their expression pattern more closely resembled that of mesenchymal stem cells (MSCs) than primary osteoblasts (Supplementary Figure 3). Accordingly, culture in the appropriate differentiation media induced these DKO cells to adopt either the adipogenic or osteoblastic fate (Figure 2a). In contrast, the p53KO OS cell lines closely resembled pre-osteoblasts based on their gene expression patterns, but these cells were unable to differentiate into either bone or fat (Figure 2a and Supplementary Figure 3). Since this differentiation block occurs in the p53-null OS cell lines, but not p53-deficient primary osteoblasts16 (Figure 4a), it likely reflects their transformed state. We used these p53KOOS cells to determine whether Rb loss was sufficient to alter the differentiation potential of blocked pre-osteoblasts by introducing control (shLuc) or Rb-specific (shRb) shRNAs. pRb was readily detectable in shLuc-p53KO, but not shRb-p53KO, OS cells (Figure 2b). Strikingly, without addition of differentiation media, pRb knockdown downregulated the bone-specific mRNA Bsp and upregulated the fat regulator Pparγ (Figure 2b). Accordingly, these shRb-p53KOOS cells were now able to differentiate into either bone or fat (Figure 2c). Moreover, when transplanted into nude mice, the shRb-p53KOOS cells formed more aggressive tumors than the parental p53KOOS cells, and these were of mixed lineage (fat, bone and undifferentiated sarcomas), in stark contrast to the undifferentiated osteoblastic tumours arising from either control shLuc-p53KO or parental p53KOOS cell lines (Figure 2d, Supplementary Figure 4, and data not shown). Thus, pRB loss is sufficient to over-ride the differentiation block of these p53-deficient, tumor cell lines and also expand their fate commitment to include the adipogenic state.

We also examined the consequences of reintroducing Rb into the DKOOS cells. For this, we induced pRb in confluence-arrested DKO cells using a doxycycline-inducible expression system (DKO-Rb<sup>Dox-ON</sup>, Supplementary Figure 5). Remarkably, pRb restoration caused the DKOOS cells to adopt the differentiation state of the p53KOOS cell lines within two days: it induced down-regulation of adipogenic markers and up-regulation of osteogenic markers, and the cells were unable to differentiate into fat (Supplementary Figure 5). Thus removal or re-introduction of Rb appears sufficient to switch lineage specification between osteoblastic commitment and multipotency.

In vitro studies have shown that pRb can act with E2F to enforce transcriptional repression of Pparγ4-5, and also bind, and potentiate the transcriptional activity of, the osteogenic regulator RUNX23. We hypothesized that pRB’s role in these processes might underlie Rb’s affect on adipogenesis versus osteogenesis. Thus, we used our DKO-Rb<sup>Dox-ON</sup> cells to
determine whether the presence or absence of pRb modulated these transcriptional regulators (Figure 3). First, we used chromatin-immunoprecipitation assays to investigate promoter regulation of Pparγ and representative Runx2-responsive genes Coll1α (Figure 3a) and osteocalcin (Oc; data not shown). pRb-induction caused both pRb and E2f4, the predominant repressive E2F, to be recruited to the Pparγ promoter (Figure 3a) and this correlated with Pparγ mRNA downregulation (Figure 3a). Contemporaneously, pRb bound to Coll1α and Oc and this was accompanied by increased promoter occupancy of Runx2 and upregulation of Coll1α and Oc mRNAs (Figure 3a, data not shown). Importantly, these changes in Pparγ, Coll1α and Oc regulation were all detected within 2 days of pRb induction and without addition of differentiation-inducing media. In addition, we found that Runx2 associated with pRb in the p53KO, but not the DKO, OS cells and its transcriptional activity was 8 fold higher in the former, versus the latter, population (Figure 3b). Thus the presence or absence of pRb directly modulates the levels and activity of Pparγ and Runx2 in accordance with the preferential commitment of our OS cell lines to the osteogenic versus the adipogenic lineage.

The preceding experiments establish a clear role for pRb in fate commitment bias in vivo and in vitro. However, since this analysis was conducted in p53-deficient cells, it is unclear whether Rb alone is sufficient to determine this plasticity or whether transformation is also required. To address this, we isolated primary osteoblasts from the calvaria of Rb±/±, p53±/± or Rb0/0; p530/0 e18.5 embryos. We brought these cells to confluence, to minimize the influence of altered proliferation, infected them with adenoviruses expressing Cre or a GFP control and then assayed differentiation. As expected, the control infected osteoblasts were able to undergo osteogenesis but not adipogenesis (Figure 4a; data not shown). Similarly, p53 loss had no effect on this fate commitment16 (Figure 4a). In stark contrast, deletion of Rb, either alone or together with p53, allowed these cells to adopt either the bone or fat lineage (Figure 4a; data not shown). This switch to multipotency correlated with the significant upregulation of adipogenic markers prior to the induction of differentiation (Figure 4b). Thus, pRB-loss is sufficient to alter the fate commitment in otherwise wildtype calvarial osteoblasts.

In vitro culture can modulate the plasticity of cells. Thus, we wished to examine Rb’s role in fate choice in vivo. For this, we employed a third transgenic strain, Meox2-Cre, which expresses Cre in the embryo proper from e6.517. Meox2-Cre;Rb±/± embryos survive to birth18. We isolated wildtype (Meox2-Cre;Rb+/+) and Rb mutant (Meox2-Cre;Rb0/0) littermates at e15.5 and e18.5 and examined both bone and brown fat development. First, there was a significant reduction in the level of calcified bone matrix in both the calvaria and long bones of Rb mutant versus wildtype embryos19 (Figure 4c). Moreover, qPCR analysis established that Runx2 mRNA was present at appropriate levels in the Rb mutant e18.5 calvarial osteoblasts, but there was a downregulation of other bone markers and a clear upregulation of fat-associated mRNAs (Figure 4c). In parallel, we found that the level of brown fat was dramatically increased in the e18.5 Rb mutant versus the wildtype controls (Figure 4d and Supplementary Figure 6). Thus, Rb loss in an, otherwise wildtype, embryo impairs bone differentiation and expands the fat compartment.

Our data establish a clear role for pRB in determining the fate choice of mesenchymal progenitors and the lineage commitment of pre-osteoblasts. This occurs both in vitro and in vivo and irrespective of whether these cells are transformed or otherwise wildtype. In vivo, Rb-loss favours adipogenesis over osteogenesis to the extent that it can reduce the levels of calcified bone and greatly increase the levels of brown fat. Moreover, Rb-loss in pre-osteoblasts is sufficient to disfavour commitment to the osteogenic state and restore multipotency. It is possible that Rb loss allows expansion of a rare multipotent progenitor population that exist within the pre-osteoblast compartment. Alternatively, Rb loss could be
actively reprogramming the pre-osteoblasts by driving either trans-differentiation to the adipogenic lineage or true de-differentiation to the multipotent progenitor stage. Between the two reprogramming models we favour de-differentiation based on both the expression of multi-lineage differentiation markers in the DKO OS cells (Supplementary Figure 3) and the broadening of the tumor spectrum from solely osteosarcomas in the Osx-Cre;p53f/f animals to include not only osteosarcomas and hibernomas but also sarcomas in the Osx-Cre;Rbfl/fl;p53f/f mice. Finally, our data offers potential insight into the cell of origin for osteosarcomas. Specifically, given the high frequency of RB-1 mutations in human osteosarcoma, we were surprised to find that Rb mutation predisposes mesenchymal cells away from the osteoblastic state. Given this finding, we speculate that RB-1 mutant osteosarcomas are likely to arise from more committed osteoblastic lineages than from uncommitted mesenchymal progenitors. In this setting, RB-1 loss could enable de-differentiation and thereby synergize with other mutations to promote tumorigenesis.

Methods Summary

Animal maintenance and histological analyses

Animal procedures followed protocols approved by MIT’s Committee on Animal Care. The Rbfl/fl (8), p53fl/fl (9), Osxl-GFP::Cre14, Prx1-Cre10, Meox2-Cre17 and Rosa26-LSL-lacZ (Jackson Labs) animals were maintained on a mixed genetic background. The transplant assays were conducted in NOD/SCID mice using 10⁴ cells. Protocols for tissue sectioning and skeletal staining are described in the additional methods section.

OS cell lines and primary osteoblast generation and analysis

Osteoblast cell lines and primary osteoblasts were generated and analyzed as described in the additional methods section or previously19. Knock-down of Rb in the p53KO-OS cells was achieved using the pMLP-miR30-based shorthairpin (Rb targeted sequence: CACGGACGTGTGAACTTATATA). Adenoviruses expressing Cre or GFP were provided by the U. of Iowa Gene Transfer Vector Core. For immunoprecipitations and immunoblotting, proteins were extracted with a Triton X-100 based buffer and quantified by the BCA assay reagent (Pierce, Inc). Antibodies were from Santa Cruz Biotechnology [pRb (H-153), E2F1 (C-20), and E2F4 (C-20)], BD Pharmingen (pRb), Ambion (GAPDH) and MBL (Runx2). Dual luciferase assays were performed as described by the manufacturer (Promega). The Runx2 reporter p6OSE2-Luc and control p4Luc were provided by Dr. Gerard Karsenty.

Additional Methods

Mouse genotyping

The Rb conditional band was detected using the primers 5’lox: 5’-CTCTAGATCCCTCATTCTTC-3’ and 3’lox: 5’-CCTTGACCATAGCCCAGCAC-3’. Primer Rbcrc3.2 (5’-GGTTAATGAAGGACTGGG-3’) was used in conjunction with primer 5’lox to detect the recombined band. To identify the p53 conditional allele we used primer p53A: 5’-CACAAAAACAGGTAAAACCCAG-3’ and primer p53B: 5’-AGCACATAGGAGCCAGAGAC-3’. The recombined allele was detected using primer p53A in conjunction with primer p53D: 5’-GAAGACAGAAAAGGGGAGGG-3’.

Tumor monitoring and analysis

The criteria for euthanasia (by CO2 inhalation) were a total tumor burden of 2cm³, tumor ulceration/bleeding, signs of infection, respiratory distress, impaired mobility, ≥20% reduction in body weight or general cachexia. All tissues were collected and hip bones, femurs and tibias were separated and fixed overnight in PBS with 3.7% formaldehyde. Soft
tissues were transferred into 70% ethanol and dehydrated via an ethanol series prior to embedding in paraffin for sectioning. Tissues containing bone was either decalcified in 0.46M EDTA, 2.5% Ammonium Hydroxide pH 7.2 for two weeks then processed for paraffin sectioning. All paraffin embedded sections were cut at 5µm, dewaxed and stained with H&E. Sirius red staining was performed by treating sections briefly stained with hematoxylin with 0.1% Sirius red in saturated picric acid (Electron Microscopy Sciences) for one hour, washing in 5% v/v glacial acetic acid and then dehydration in ethanol/xylene prior to mounting.

**Immunohistochemistry (IHC)**

Runx2 IHC was performed using a modified citric acid unmasking protocol. Briefly, slides were deparaffinized in xylene and rehydrated through ethanol. Slides were boiled for 30 min in citrate buffer, pH 6.0, and then cooled in running tap water. Slides were then washed in PBS for 5 min followed by inactivation of endogenous peroxidases by incubation 0.5% H$_2$O$_2$ in methanol. Slides were blocked in 10% Goat Serum for 1 h at room temperature. Primary antibody (MBL anti-Runx2 Clone 8G5) was diluted 1:200 in PBS 0.15% Triton and incubated overnight at 4°C. The following day, slides were washed three times in PBS. Secondary antibodies (Vectorstain ABC kits, Vector laboratories) were diluted 1:500 in PBS containing 0.4% Goat Serum and detected using a DAB substrate (Vector Laboratories). All samples were counterstained with hematoxylin.

**Skeletal Staining**

Embryos were sacrificed, skinned and eviscerated. The remaining tissue was fixed in 95% ethanol for 4 days, transferred to acetone for 3 days, and subsequently transferred to staining solution of 0.015% Alcian blue 8GX (Sigma), 0.005% alizarin red S (Sigma), and 5% glacial acetic acid in ethanol at 37°C for 2 days and at room temperature for a more one day. Tissue was cleared in 1% potassium hydroxide for several days and then stored in glycerol.

**Generation of osteosarcoma cell lines**

Osteosarcomas were dissected, minced, filtered through a 70µm filter, and plated in normal growth medium (10% FBS in DME, 1% P/S, L-glutamine) to generate the OS cell lines. Cells were passaged as they reached confluence. For RNA purification, cells were rinsed 2x with PBS, and RNA extraction was performed using RNAeasy kit (QIAGEN). First-strand cDNA was transcribed from 1 µg of RNA using Superscript III reverse transcriptase (Invitrogen) following the manufacturer’s instructions. Quantitative RT-PCR with 20 to 100 ng cDNA was done using SYBR Green (Applied Biosystems). Reactions were run on the ABI Prism 7000 Sequence Detection System and analyzed using the 7000 SDS software. Primers used for qPCR were: Alkaline Phosphatase (F) TCTTCAGACCTGCAACCTC and (R) CATCTGAGCAGACCTGGT; Col1a1 (F) CGAGTCACACCCGAACTTGG and (R) GCAGGCGGGAATGTGCTCT; Osteocalcin (F) CTCTGTCTCTGACCTCACAG and (R) CAGGTCTAAAATAGTAGAAGCAC; Osteopontin (F) TGGCTTTTGCCTTTTGGCAT and (R) TCTGTGGCCAGCAAGGGAGATT; Runx2 (F) TGAAGATTTTGCTGGGCGGA and (R) TCTGTGCTTCTTGTCCTCCC; Ap2 (F) ATCCCTTGTGGAACCTCGGA and (R) ACGTGATGATCATGGTGTGCT; C/ebp α (F) CAAGAACAAGCAAGAGCTTGG and (R) GTCACTGTGCAACTCCAGAC; Pparγ (F) GAGCTGACCCCAATCGTGCT and (R) GCTTCAATCGGAGTGTCTTC; Srebp-1c (F) GGAGCCATGGATTGAGATT (R) GCTTCCAGAGGGAGGCAG; Ucp-1 (F) AGCCGGCTTAATGACTGGAG and (R) TCTGTAGCTGGCCAATGAAC; Pgc-1 (F) GTCCACACAGAGACCTGGA and (R) TGGTCTGTGCACTGGAC; Nbrf-1 (F) CCGCACTAGCGCCCGG and (R) CGGCACTAGCGCCCGG; MyoD (F) CGCCACTCCGGGACATAG and (R) GAAGTCGTCTGCTGCTCTAAGG; Prdm16 (F) GACCACGTTGAACGGCATT and (R) GCGTGCATCAGCTGTGCT; Taz (F) Calo et al. Page 5

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GTCACCAACAGTGCTCAGATC and (R) AGTGATTACAGCCAGGTTAGAAG; and Gapdh (F) CAAGGTGGCAGAGGCCTT and (R) TCCAGCTGCTCAATGGACGCACTTT.

**Calvarial Osteoblasts Preparation and Culture**

Calvaria from embryonic day 18.5 embryos were removed and carefully cleaned in sterile PBS from contaminating tissue. Then treated with several rounds of collagenase/trypsin digestion at 37°C, and plated onto six-well plates for 2 days in αMEM with 10% fetal bovine serum and penicillin/streptomycin. For differentiation, 3.5×10⁵ cells were plated onto a well of a 6-well tissue culture plates. Upon reaching confluence, calvarial osteoblasts were treated with medium supplemented with 50 µg/mL of ascorbic acid and 10 mmol/L of β-glycerol-phosphate. Adenovirus (University of Iowa Gene Transfer Vector Core) was added to the medium at 100 plaque-forming units per cell and washed away 24 h later. To assay for calcium deposits, plates were stained with 1% alizarin red S solution (pH 5.0).

**Chromatin Immunoprecipitation assay**

Protein complexes were cross-linked to DNA by adding formaldehyde (Sigma, Inc.) to live cells to a final concentration of 1%. After incubation for 10 min at 37°C, glycine was added to give a final concentration of 0.125 M for 5 min. The cells were washed twice with PBS containing 1mM PMSF, scraped and pelleted. Nuclei were extracted with a 20mM Tris pH 8, 3 mM MgCl₂, 20 mM KCl buffer containing protease inhibitors, pelleted by microcentrifugation and lysed with SDS lysis buffer (1% SDS, 10 mM EDTA, 50 mM Trischloride pH 8.1), containing protease inhibitors. The resulting chromatin solution was sonicated to generate 500–1000 bp DNA fragments. After microcentrifugation, the supernatant was diluted 1:10 with dilution buffer (0.01% SDS, 1.1% Triton X-100, 1.2 mM EDTA, 16.7 mM Trischloride pH 8.1, 167 mM NaCl, containing protease inhibitors), precleared with blocked protein A-positive Staph cells (Santa Cruz, Inc), and split into aliquots. These were incubated at 4°C for 12 to 16 hours with 5 µg antibody with rotation. Antibody-protein-DNA complexes were isolated by immunoprecipitation with blocked protein A-positive Staph A cells. Following extensive washing, bound DNA fragments were eluted and analyzed by Quantitative RT-PCR using the following primers: Ppar γ proximal E2F site (F) ACGCGGAAGAAGAGACCT and (R) TCCAGCTGCTCAATGGACGCACTTT; Ppar γ distal E2F site (F) TCGCAGTCAGACGCCGAG and (R) AGGTCTCTTCCGCGTCCCT; Coll1a1 Runx2 site (F) TGCTTCCACGTTTACAGCTTAAAG and (R) GTCAGGAAAGGGTGTAGTATATCG.

**Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

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Figure 1. Rb cooperates with p53 and modulates mesenchymal tumor fate in a dosage-dependent manner

a. Mesenchymal tumor distribution (percentage of animals analyzed up to 24 months of age) for Prx1-Cre;Rb and/or p53 compound mutant animals. b. H+E staining of representative sarcomas (20× magnification). c. PCR genotyping to detect Rb wildtype (wt) and recombined conditional mutant (loxp) alleles in control Rb<sup>fl/+</sup>;p53<sup>fl/+</sup> tissues (lane 1) or cell lines derived from Prx1-Cre;Rb<sup>fl/fl</sup>;p53<sup>fl/fl</sup> (DKO) or Prx1-Cre;Rb<sup>fl/+</sup>;p53<sup>fl/fl</sup> osteosarcomas. Cell lines were cultured for ≥20 passages prior to genotyping to eliminate stromal cell contribution.
Figure 2. Rb regulates osteosarcoma-cell lineage plasticity in vitro and in vivo

The differentiation potential of 3 different Osx-Cre;Rb<sup>fl/fl</sup>;p53<sup>fl/fl</sup> (DKO) and Osx-Cre;p53<sup>fl/fl</sup> (p53KO) OS cell lines was assessed 0, 7, 14, or 21 days after addition of differentiation media. a, Representative staining for: (left lane) alkaline phosphatase prior to differentiation induction; (middle lane) Alizarin Red to detect bone mineralization 14 days after culture in osteogenic-induction media and (right lane) Oil-Red-O to detect lipid droplets 14 days after culture in adipogenic-induction media. Expression of bone (Runx2, Alp, Coll1a and Bsp) and fat (Ap2, Pparγ, C/ebpα and Pgc1α) markers was assessed by qPCR of un-induced DKO (orange) and p53KO (black) OS cells. Bars represent the mean of
three independent experiments (+/− SD). NS = not significantly expressed. b, *Rb* or control (Luc) shRNAs were expressed in the *p53KO* cell lines. *Rb* knockdown was confirmed by immunoprecipitation and qPCR showed that this caused downregulation of bone markers *Bsp* (and also *Coll1a* and *Alp*, data not shown), and upregulation of fat markers *Pparγ* (and also *Ap2* and *Cebpα*, data not shown) without culture in differentiating media. Bars represent the mean of three independent experiments (+/− SD). c, The osteogenic and adipogenic potential of *shLuc*- and *shRb-p53KO* cell lines was assessed 0, 7, 14 and 21 days after differentiation induction by Alizarin Red and Oil-Red-O staining. A representative timepoint (14 days) is shown. d, H+E staining of representative tumors derived from *shLuc*- and *shRb-p53KO* cell lines injected subcutaneously into immunocompromised mice. *shRb-p53KO* OS cells consistently (10/10 injections) yielded tumors that arose faster, and were more aggressive, than those arising from the parental *p53KO* OS controls (10 injections). Moreover, the *shRb-p53KO* OS derived tumors were frequently (6/10 injections) mixed lineage (top inset shows fat neoplasm; bottom inset bone/undifferentiated sarcoma), while the control *shLuc-p53KO* tumors were uniformly (10/10 injections) osteosarcomas. Additional analysis of these tumors (H+E, sirius red staining and Runx2 IHC) is shown in Supplementary Figure 4.
Figure 3. pRb modulates the activity and the expression of the master lineage regulators Runx2 and Pparγ

a. The stable DKO-Rb<sup>Dox-ON</sup> OS cells were generated by drug selection of pools of DKO cells transfected with the doxycycline inducible construct pCW22-Rb. These were cultured for two days in the absence (Rb Off) or presence (Rb On) of doxycycline and then analyzed. Results are representative of three independent experiments. Promoter occupancy was assessed by chromatin immunoprecipitation. Sequence analysis identified two potential E2f binding sites (−278 and −160) within the Pparγ promoter. pRb induction caused a dramatic upregulation of both pRb and E2F4 binding to the proximal site. (No binding was observed at the distal element.) Similarly, pRb induction allowed pRb to bind to the known Runx2
response element of Col1α 20 and also increased the binding of Runx2. These changes correlated with the downregulation of Pparγ mRNA and upregulation of Col1α mRNA as judged by qPCR. Bars represent the mean of three independent experiments (+/- SD). b, Western blotting detected Runx2 in pRb-immunoprecipitates from p53KO-OS cell lines (left, top panel). Western blotting of whole cell extracts confirmed that Runx2 was expressed in both DKO and p53KO-OS cell lines (left, bottom panel). MSCs and osteoblasts were used as a positive control. Right panel: Runx2 transcriptional activity was shown to be higher in the p53KO- versus the DKO OS cell lines as judged by activation of the artificial Runx2-responsive reporter p6OSE2-Luc. Results are the average of six independent samples. Error bars indicate S.E.
Figure 4. Rb maintains the osteoblastic fate commitment in normal osteoblasts and regulates fate choice during normal development in vivo

a. Calvarial osteoblasts were prepared from e18.5 Rb<sup>fl/fl</sup> or p53<sup>fl/fl</sup> embryos and infected with Ad-GFP or Ad-Cre at P1. Five days later, the cells were induced with differentiation media and then assayed for osteogenesis and adipogenesis at 0, 14 and 25 days by staining with Alizarin Red and Oil-Red-O. A representative timepoint (25 days) is shown.

b. qPCR was also used to assess osteogenic and adipogenic markers in the un-induced Rb<sup>fl/fl</sup> (wt) versus Rb<sup>fl/fl</sup>+Ad-Cre (Rb<sup>−/−</sup>) osteoblasts. Bars represent the mean of three independent experiments (+/- SD).

c. Alizarin Red (bone mineralization) and Alcian Blue (cartilage) staining of e15.5 skeletons (top panel), e18.5 calvaria (middle panel) and e18.5 limbs.
(botten panel) from Meox2-Cre;Rb<sup>+/+</sup> and Meox2-Cre;Rb<sup>fl/fl</sup> littermate embryos. Arrows mark visible skeletal defects. qPCR was used to assess osteogenic (Runx2, Alp, and Bsp) and adipogenic (Ap2 and C/ebpα) markers in mRNA extracted from the calvarial bones of e18.5 Meox2-Cre;Rb<sup>+/+</sup> and Meox2-Cre;Rb<sup>fl/fl</sup> embryos. Bars show the mean of three embryos arising in two independent crosses (+/- SD). Brown adipose tissue (BAT) was dissected from the backs of Meox2-Cre;Rb<sup>fl/fl</sup> embryos (n=10) and their Meox2-Cre;Rb<sup>+/+</sup> littermate controls. All 10 showed a dramatic expansion of the brown fat compartment. A representative example is shown (upper two panels). Introduction of the LSL-LacZ reporter into this model, and LacZ staining confirmed equal, widespread expression of Cre in the control and Rb mutant BAT (third panel). H+E staining of BAT (bottom panel).