Supporting Information

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Fig. S1. Lysyl oxidase (LOX) is required for MEFs to differentiation into adipocytes. (*A*) MEFs were plated at 30% confluency and 24 h later transfected with LOX Stealth RNAi, cell extracts were prepared at postconfluence, and the expression of LOX was measured. (*B*) The effect of LOX knockdown on adipocyte commitment and terminal differentiation assessed by the expression of an adipocyte-specific marker (422/aP2) and by the accumulation of cytoplasmic triglyceride.



Fig. S2. Overexpression of dominant-negative bone morphogenetic protein receptor (DN-BMPr)1A prevents the induction of LOX by BMP2. C3H10T1/2 stem cells were infected with retrovirus harboring constitutively active (CA)-BMPr1A or DN-BMPr1A and treated with/out BMP2. (A) One day after infection, total RNA was isolated and subjected to RT-PCR to confirm the expression of CA-BMPr1A or DN-BMPr1A; β -actin was used loading control. (B) Whole-cell lysates were prepared, and the effects of overexpressing CA-BMPr1A or DN-BMPr1A on the induction of LOX by BMP2 were assessed by immunoblotting; β -actin was used loading control.

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