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原田浩徳	MDS「分子病態」	臨床血液	52	1525-1534	2011
原田結花, 原田浩徳	MPNと遺伝	最新医学	66	2552-2557	2011
原田結花, 原田浩徳	放射線発がん（骨髄異形成症候群・白血病）の分子病態	血液フロンティア	21	1775-1781	2011
原田結花, 原田浩徳	造血器腫瘍におけるEZH2変異とその機能的意義	血液内科	64(2)	139-144	2012
今川 潤, 原田結花, 吉田徹巳, 樽谷美保, 木村昭郎, 松元加奈, 森田邦彦, 原田浩徳	ダサチニブ少量療法が有効であったイマチニブ不耐容慢性好酸球性白血病	臨床血液	52	546-550	2011

#### IV. 研究成果の刊行物・別刷

(主なもの)

# Evi1 is essential for hematopoietic stem cell self-renewal, and its expression marks hematopoietic cells with long-term multilineage repopulating activity

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Ecotropic viral integration site 1 (Evi1), a transcription factor of the SET/PR domain protein family, is essential for the maintenance of hematopoietic stem cells (HSCs) in mice and is overexpressed in several myeloid malignancies. Here, we generate reporter mice in which an internal ribosome entry site (IRES)-GFP cassette is knocked-in to the *Evi1* locus. Using these mice, we find that Evi1 is predominantly expressed in long-term HSCs (LT-HSCs) in adult bone marrow, and in the hematopoietic stem/progenitor fraction in the aorta-gonad-mesonephros, placenta, and fetal liver of embryos. In both fetal and adult hematopoietic systems, Evi1 expression marks cells with long-term multilineage repopulating activity. When combined with conventional HSC surface markers, sorting according to Evi1 expression markedly enhances purification of cells with HSC activity. Evi1 heterozygosity leads to marked impairment of the self-renewal capacity of LT-HSCs, whereas overexpression of Evi1 suppresses differentiation and boosts self-renewal activity. Reintroduction of Evi1, but not *Mds1-Evi1*, rescues the HSC defects caused by Evi1 heterozygosity. Thus, in addition to documenting a specific relationship between Evi1 expression and HSC self-renewal activity, these findings highlight the utility of Evi1-IRES-GFP reporter mice for the identification and sorting of functional HSCs.

Hematopoietic stem cells (HSCs) are distinguished by their inherent capacity to perpetuate themselves through self-renewal and to generate multiple blood cell lineages through differentiation. To maintain a steady-state pool of self-renewing HSCs and prevent HSC exhaustion, these defining properties of HSCs must be tightly regulated. Fine-tuning of stem cell properties requires stem cell-specific expression of their regulatory genes. To elucidate the stemness transcriptional profile, several gene expression microarray analyses have identified quite a few number of HSC-specific gene candidates (Ramalho-Santos et al., 2002; Akashi et al., 2003; Forsberg et al., 2010). However, most of the molecules established to be associated with the regulation of self-renewal capacity

in HSCs are widely expressed in the hematopoietic system, and their mutations in genetic models are exclusively accompanied with other hematological abnormalities. Thus, a bona fide stem cell-specific regulator of their function has not been identified, and the functional identification of HSCs based on their ability to self-renew remains difficult.

Ecotropic viral integration site 1 (Evi1) is an oncogenic transcription factor that belongs to the SET/PR domain protein family (Goyama and Kurokawa, 2009). We and others have reported that Evi1 accomplishes an important regulatory function in hematopoietic stem/progenitor

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Abbreviations used: AGM, aorta-gonad-mesonephros; AML, acute myeloid leukemia; BFU-E, burst-forming unit-erythrocyte; CFU-S, CFU-spleen; CLP, common lymphoid progenitor; CMP, common myeloid progenitor; CRA, competitive repopulation assay; EC, endothelial cell; ES, embryonic stem; Evi1, ecotropic viral integration site 1; FL, fetal liver; GEMM, granulocyte/erythrocyte/macrophage/megakaryocyte; GM, granulocyte/macrophage; GMP, GM progenitor; HSC, hematopoietic stem cell; HSPC, hematopoietic stem/progenitor cell; IRES, internal ribosome entry site; Lin, lineage; LSK, Lin<sup>-</sup>Sca-1<sup>+</sup>c-kit<sup>+</sup>; LT-HSC, long-term HSC; ME, Mds1-Evi1; MEP, megakaryocyte/erythrocyte progenitor; MPP, multipotent progenitor; MSC, mesenchymal stem cell; MSL, Mac-1<sup>+</sup>Sca-1<sup>+</sup>Lin<sup>-</sup>; OB, osteoblast; pA, polyadenylation; PB, peripheral blood; RQ-PCR, real-time quantitative PCR; SCF, stem cell factor; ST-HSC, short-term HSC; TPO, thrombopoietin.

K. Kataoka and T. Sato contributed equally to this paper.

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cells (HSPCs) during fetal and adult development. Evi1 expression is limited to HSPCs in the embryonic and adult hematopoietic systems. HSCs in *Evi1*<sup>-/-</sup> embryos are markedly decreased in numbers with defective repopulating capacity (Yuasa et al., 2005). Moreover, conditional deletion of Evi1 in adult mice revealed that Evi1 is essential for the maintenance of HSCs, but is dispensable for lineage commitment (Goyama et al., 2008). Besides the importance of Evi1 in normal hematopoiesis, dysregulation of Evi1 expression can have distinct oncogenic potential in various myeloid malignancies (Goyama and Kurokawa, 2009). Indeed, aberrant EVI1 expression defines a unique subset of acute myeloid leukemia (AML), and predicts adverse outcome in patients (Lugthart et al., 2008; Gröschel et al., 2010). Furthermore, Evi1 overexpression in hematopoietic cells leads to myelodysplasia in a murine BM transplant model (Buonamici et al., 2004).

In this study, using newly generated Evi1-GFP reporter mice, we demonstrate that Evi1 is preferentially expressed in LT-HSCs, and its expression can mark *in vivo* long-term multilineage repopulating HSCs and improve the conventional HSC isolation strategy in both adult BM and embryo,

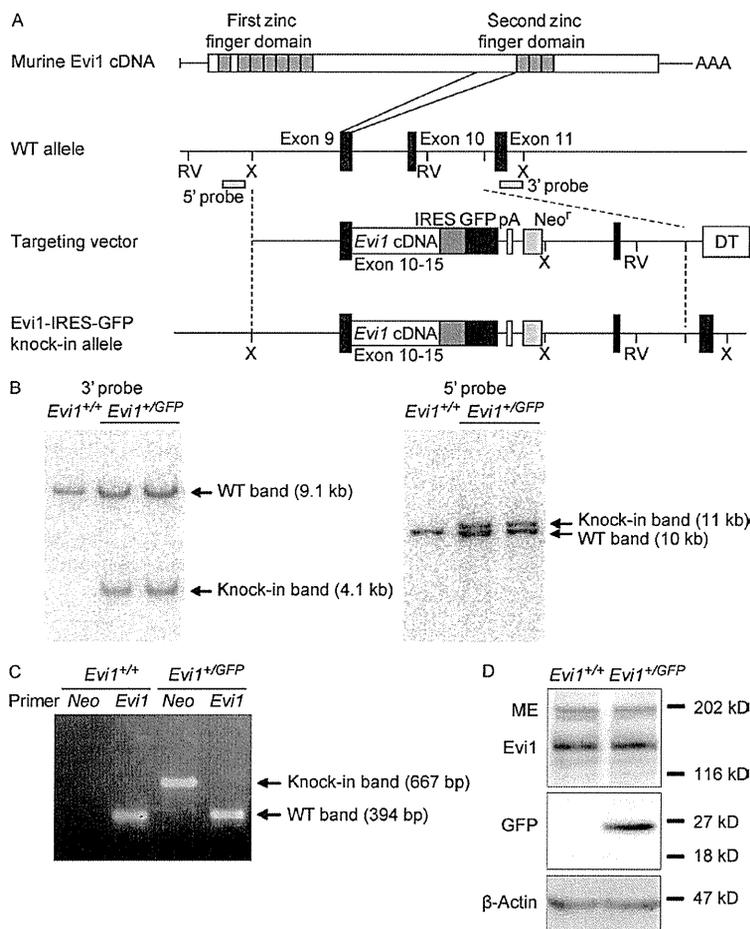
which suggests a distinctive relationship between Evi1 and HSC function. Consistent with this, heterozygosity of Evi1 causes a striking reduction in the number of LT-HSCs, with a specific defect of self-renewal capacity caused by accelerated differentiation. Our results point to a potential utility of an Evi1-GFP reporter mouse line for the functional identification of HSCs based on their self-renewal activity, and a central role of Evi1 in regulating the homeostasis of HSCs.

## RESULTS

### Evi1 is predominantly expressed in LT-HSCs in adult BM

To elucidate Evi1 expression within the hematopoietic system, we have generated gene-targeted mice in which an internal ribosome entry site (IRES)-GFP cassette is knocked-in to the *Evi1* locus by homologous recombination (Fig. 1 A). This knock-in allele functions in a bicistronic manner in that expression of both Evi1 and GFP is under the endogenous transcriptional regulatory elements of the *Evi1* gene, thus enabling us to track Evi1 expression on an individual cell basis. Appropriately targeted TT2 embryonic stem (ES) cell clones were identified by Southern blotting (Fig. 1 B). Mice heterozygous for the *Evi1*-IRES-GFP allele (*Evi1*<sup>+/<sup>GFP</sup>) were distinguished</sup>

from WT mice by genotyping PCR (Fig. 1 C). Western blot analysis showed the presence of GFP protein and comparable expression of Evi1 protein in embryonic fibroblast cells from *Evi1*<sup>+/<sup>GFP</sup> mice compared with WT mice (Fig. 1 D). *Evi1*<sup>+/<sup>GFP</sup> mice were phenotypically indistinguishable in survival, hematopoietic cellularity, and lineage composition from WT controls (unpublished data). Initial flow cytometric analysis of adult *Evi1*<sup>+/<sup>GFP</sup> mice revealed a small, but discrete, population of GFP<sup>+</sup> cells (0.15 ± 0.6%; Fig. 2 A), confirming the expression of the *Evi1*-IRES-GFP allele. To examine whether GFP expression levels correlated with those of endogenous *Evi1* mRNA expression, *Evi1* expression of sorted GFP<sup>-</sup> and GFP<sup>+</sup> cells from BM of *Evi1*<sup>+/<sup>GFP</sup> mice was analyzed by real-time quantitative PCR (RQ-PCR). *Evi1* mRNA was exclusively expressed in the GFP<sup>+</sup> cells, and almost no expression was found in the GFP<sup>-</sup> cells (Fig. 2 B), indicating that GFP expression in this mouse model faithfully marks cells with active Evi1 expression.</sup></sup></sup></sup>



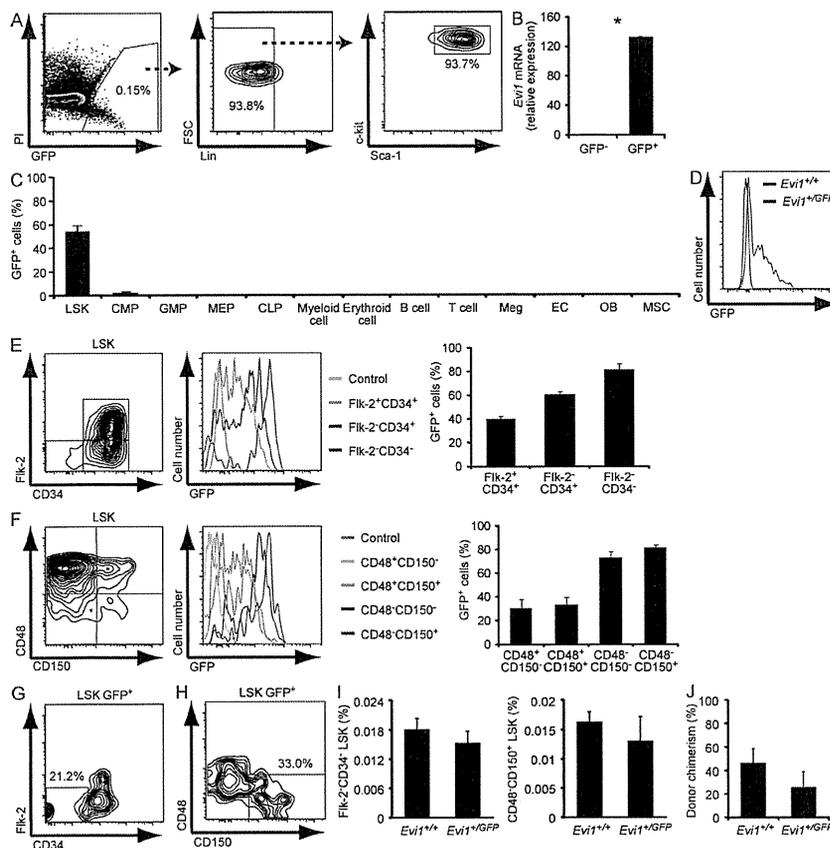
**Figure 1. Generation of Evi1-IRES-GFP knock-in mice.**

(A) The structure of murine *Evi1* and the targeted *Evi1*-IRES-GFP locus is shown. RV, EcoRV; X, XbaI. (B) Southern blot analysis of genomic DNA isolated from WT ES cells (*Evi1*<sup>+/+</sup>) and two independent clones of targeted ES cells (*Evi1*<sup>+/<sup>GFP</sup>). DNA was digested with XbaI (left) or EcoRV (right), and hybridized with the indicated probes. (C) Genotyping of *Evi1*<sup>+/<sup>GFP</sup> mice by PCR. (D) Western blot analysis for GFP and Evi1 in embryonic fibroblast cells from *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/<sup>GFP</sup> mice.  $\beta$ -Actin was used as a loading control. ME, Mds1-Evi1.</sup></sup></sup>

*Evi1* mRNA has been shown to be expressed at significantly higher levels in HSPCs ( $\text{Lin}^- \text{Sca-1}^+ \text{c-kit}^+$  [LSK]) and common lymphoid progenitors (CLPs) than in other hematopoietic cells (Yuasa et al., 2005; Chen et al., 2008). To gain insight into the biological function of *Evi1* through its cell type-specific expression pattern, the distribution of  $\text{GFP}^+$  cells was examined in adult BM from *Evi1*<sup>+/GFP</sup> mice. Beyond expectation, GFP expression was highly restricted to the LSK fraction (Fig. 2 A). To confirm stem/progenitor-specific expression of *Evi1*, we analyzed the GFP fluorescence of various hematopoietic cell populations from BM and spleen of *Evi1*<sup>+/GFP</sup> mice. We found a heterogeneous expression of GFP in the LSK fraction, in which about half of the cells were  $\text{GFP}^+$  (Fig. 2, C and D). Conversely, only 2.5% of common myeloid progenitors (CMPs) expressed GFP, and almost no expression was found in granulocyte/monocyte progenitors (GMPs) and megakaryocyte/erythrocyte progenitors (MEPs; Fig. 2 C). In contrast to the previous study (Chen et al., 2008), GFP was not expressed in CLPs (Fig. 2 C). In addition, no GFP expression was observed in mature hematopoietic lineages or nonhematopoietic cells in BM (Fig. 2 C). Together, these results suggest that *Evi1* is uniquely expressed in HSPCs, but its expression is sharply down-regulated along with differentiation.

Because LSK cells, a population which contains multipotent progenitors (MPPs), short-term HSCs (ST-HSCs), and long-term HSCs (LT-HSCs), include both a  $\text{GFP}^+$  fraction and a  $\text{GFP}^-$  fraction, we next resolved GFP expression within the LSK compartment for other markers characteristic of LT-HSCs. When LSK cells were subdivided according to CD34 and Flk-2 expression (Orford and Scadden, 2008), the  $\text{Flk-2}^- \text{CD34}^-$  LSK fraction, which is considered to contain most LT-HSC activity, had the highest expression of GFP, and its expression decreased with differentiation to hematopoietic progenitors (Fig. 2 E). In addition, further enrichment for LT-HSCs within the LSK fraction using SLAM family receptors (CD48 and CD150; Kiel et al., 2005) revealed that  $\text{GFP}^+$  cells were found in greatest abundance within  $\text{CD48}^- \text{CD150}^+$  LSK cells, in which LT-HSCs are highly enriched. In contrast, GFP expression was substantially down-regulated in  $\text{CD48}^+$  LSK cells, irrespective of CD150 expression (Fig. 2 F). When we examined how  $\text{GFP}^+$  cells were distributed within the LSK fraction, GFP expression was highly enriched in the  $\text{Flk-2}^- \text{CD34}^-$  LSK or  $\text{CD48}^- \text{CD150}^+$  LSK fractions (Fig. 2, G and H). Therefore, these results indicate that *Evi1* is dynamically regulated within HSPCs; its

**Figure 2. *Evi1* is predominantly expressed in LT-HSCs in adult BM.** (A) FACS analysis of expression of lineage markers (Lin), c-kit, and Sca-1 on  $\text{GFP}^+$  cells in adult BM from *Evi1*<sup>+/GFP</sup> mice. Data are representative of three independent experiments. PI, propidium iodide; FSC, forward scatter. (B) RQ-PCR analysis of the expression of *Evi1* mRNA in sorted  $\text{GFP}^-$  or  $\text{GFP}^+$  cells from BM of *Evi1*<sup>+/GFP</sup> mice, presented relative to GAPDH expression (\*,  $P < 0.0001$ ;  $n = 2$ ). (C) Frequency of  $\text{GFP}^+$  cells in indicated BM subpopulation and in splenic T cells from *Evi1*<sup>+/GFP</sup> mice ( $n = 3-5$ ). Meg, megakaryocyte; EC, endothelial cell. (D) FACS analysis of expression of GFP in LSK cells from *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/GFP</sup> mice. Data are representative of at least twenty independent experiments. (E and F) Frequency of  $\text{GFP}^+$  cells in subpopulations of LSK cells divided using Flk-2 and CD34 (E) or CD48 and CD150 (F) in *Evi1*<sup>+/GFP</sup> mice. (left) Representative plot is shown. (right) Bar graph represents mean  $\pm$  SD ( $n = 3-4$ ). (G-H) FACS analysis of expression of Flk-2 and CD34 (G) or CD48 and CD150 (H) on LSK  $\text{GFP}^+$  cells in BM from *Evi1*<sup>+/GFP</sup> mice. Data are representative of two independent experiments. (I) Frequency of  $\text{Flk-2}^- \text{CD34}^-$  LSK or  $\text{CD48}^- \text{CD150}^+$  LSK cells in BM from *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/GFP</sup> mice ( $n = 3-5$ ). (J) PB donor chimerism in CRAs, in which  $2 \times 10^5$  BM cells from *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/GFP</sup> mice (Ly5.1) were transplanted into lethally irradiated recipients (Ly5.2) together with  $2 \times 10^5$  competitor BM cells (Ly5.2). Percentages of donor-derived cells (Ly5.1) in PB 16 wk after transplantation are shown ( $P = 0.12$ ;  $n = 3$ ). Data represent mean  $\pm$  SD.



expression is predominantly enriched in LT-HSCs and rapidly extinguished during early stages of lineage commitment.

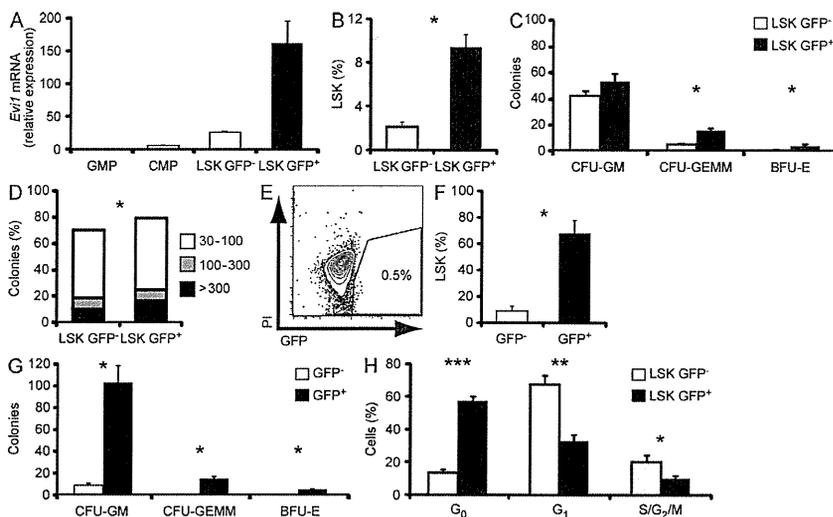
To reinforce *Evi1*-IRES-GFP knock-in mice as a faithful tool for investigating HSCs, we assessed the number and function of LT-HSCs in BM from *Evi1*<sup>+/GFP</sup> mice. Flow cytometric analysis revealed that the frequencies of Flk-2<sup>-</sup> CD34<sup>-</sup> LSK or CD48<sup>-</sup> CD150<sup>+</sup> LSK cells were comparable between *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/GFP</sup> mice (Fig. 2 I). In addition, a competitive repopulation assay (CRA) showed that *Evi1*<sup>+/GFP</sup> BM cells exhibited slightly less, but not significantly different, long-term reconstitution capacity (Fig. 2 J), indicating that the number and function of HSCs in *Evi1*<sup>+/GFP</sup> mice are similar to WT controls.

### ***Evi1* expression represents a functionally distinct population that remains in an undifferentiated and quiescent state within HSPCs**

As only a subset of LSK cells expressed GFP in *Evi1*<sup>+/GFP</sup> mice, we hypothesized that *Evi1* expression functionally divides the LSK population and marks a more undifferentiated and quiescent state with multipotent differentiation properties in this population. To test this idea, we separated the LSK population into LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells and compared their biological functions. Initially, we confirmed that LSK GFP<sup>+</sup> cells had a much higher level of *Evi1* transcripts than LSK GFP<sup>-</sup> cells by RQ-PCR analysis (Fig. 3 A). Interestingly, despite the negative GFP expression, LSK GFP<sup>-</sup> cells expressed *Evi1* mRNA at a higher level compared with CMPs and GMPs (Fig. 3 A), which also suggests that *Evi1* expression is inversely proportional to the differentiation status. To achieve an estimate of the differentiation stage of these

two populations, LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells were cultured in serum-free medium containing stem cell factor (SCF) and thrombopoietin (TPO). After 3 d of culture, the proportion that remained in the LSK fraction was significantly higher in LSK GFP<sup>+</sup> cells than in LSK GFP<sup>-</sup> cells (Fig. 3 B), suggesting that LSK GFP<sup>+</sup> cells are more primitive HSCs. Next, to evaluate the differentiation potential of LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells, we performed colony-forming assays in vitro. Although both populations generated an equivalent number of myeloid colonies CFU-granulocyte/macrophage [CFU-GM]), LSK GFP<sup>+</sup> cells gave rise to greater numbers of erythroid (burst-forming unit-erythrocyte [BFU-E]) and multipotential (CFU-granulocyte/erythrocyte/macrophage/megakaryocyte [CFU-GEMM]) colonies than LSK GFP<sup>-</sup> cells (Fig. 3 C). These data suggest that *Evi1* expression correlates with multipotent differentiation capacity. In addition, to assess the colony-forming capacity at the clonal level, single LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells were cultured in serum-free medium. LSK GFP<sup>-</sup> cells formed detectable colonies at a frequency comparable to LSK GFP<sup>+</sup> cells, but generated smaller numbers of highly proliferative colonies (>300 cells; Fig. 3 D), indicating that the LSK GFP<sup>+</sup> fraction comprises a higher proportion of HSPCs with enhanced proliferative capacity.

Our observations suggested that *Evi1* reporter activity is down-regulated as HSCs differentiate. To examine this issue, we forced LSK GFP<sup>+</sup> cells to differentiate in vitro in response to SCF, TPO, IL-3, and IL-6. These LSK GFP<sup>+</sup> cells predominantly generated GFP<sup>-</sup> cells (Fig. 3 E). After culture, the majority of cells that had become GFP<sup>-</sup> lost the LSK phenotype, whereas most cells that remained in GFP<sup>+</sup> continued to express



**Figure 3. *Evi1* expression represents a functionally distinct population that remains in an undifferentiated and quiescent state within HSPCs.** (A) RQ-PCR analysis of the expression of *Evi1* mRNA in sorted GMPs, CMPs, LSK GFP<sup>-</sup> cells, and LSK GFP<sup>+</sup> cells from *Evi1*<sup>+/GFP</sup> mice, presented relative to *GAPDH* expression ( $n = 2$ ). (B) LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells were cultured in serum-free medium with 20 ng/ml SCF and 20 ng/ml TPO for 3 d, and the percentage of the remaining LSK fraction was analyzed (\*,  $P < 0.001$ ;  $n = 3$ ). (C) Numbers of CFU-GM, CFU-GEMM, and BFU-E colonies derived from 100 sorted LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells (\*,  $P < 0.05$ ;  $n = 3$ ). (D) Single LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells from *Evi1*<sup>+/GFP</sup> mice were clonally sorted and cultured in serum-free medium. After 14 d of culture, cell numbers in each colony were analyzed. Their relative distribution is shown (\*,  $P < 0.05$ ;  $n = 192$  clones

from 2 independent experiments). (E) LSK GFP<sup>+</sup> cells were cultured in medium containing 10% serum with 50 ng/ml SCF, 50 ng/ml TPO, 10 ng/ml IL-3, and 10 ng/ml IL-6 for 5 d, and the percentage of the remaining GFP<sup>+</sup> fraction were analyzed. Data are representative of four independent experiments. (F) The percentages of the remaining LSK fraction in GFP<sup>-</sup> and GFP<sup>+</sup> cells after culture were analyzed (\*,  $P < 0.0001$ ;  $n = 4$ ). (G) Numbers of CFU-GM, CFU-GEMM, and BFU-E colonies derived from 200 GFP<sup>-</sup> and GFP<sup>+</sup> cells were analyzed (\*,  $P < 0.0001$ ;  $n = 4$ ). (H) Cell cycle status of LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells from *Evi1*<sup>+/GFP</sup> mice, analyzed by Hoechst 33342 and pronyon Y staining (\*,  $P < 0.05$ ; \*\*,  $P < 0.005$ ; \*\*\*,  $P < 0.0005$ ,  $n = 3$ ). Data represent mean  $\pm$  SD.

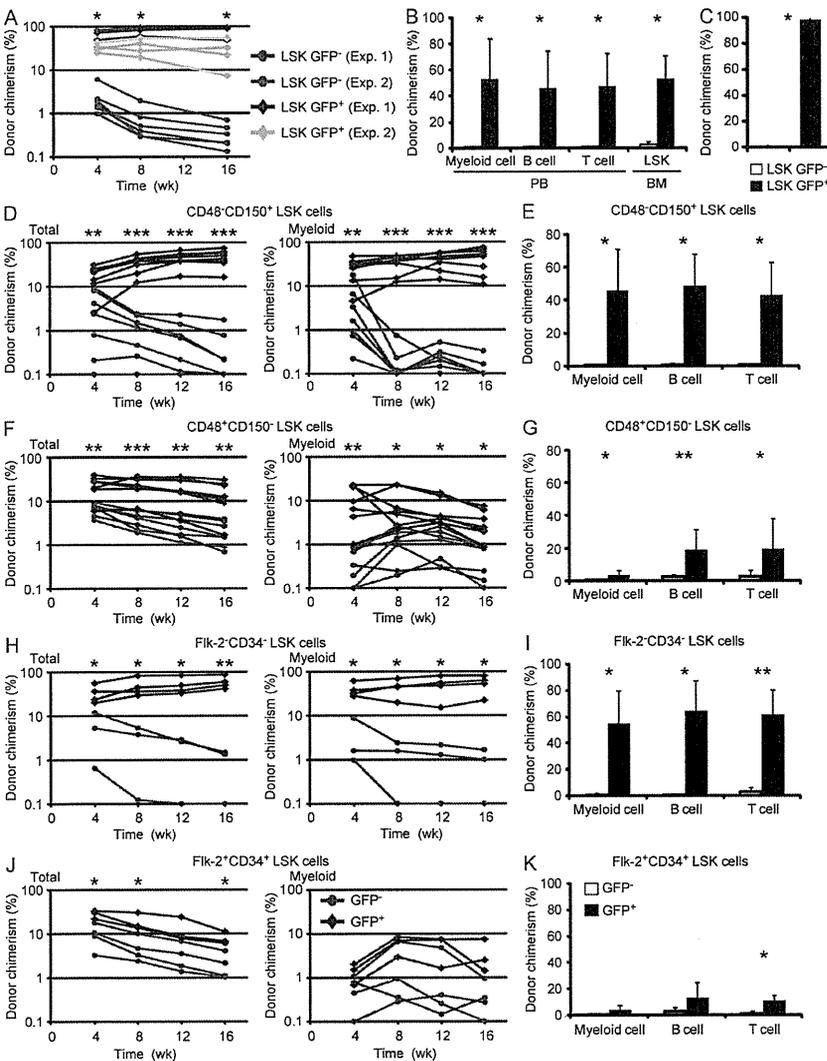
the LSK phenotype (Fig. 3 F), indicating that loss of GFP correlates with phenotypic differentiation. To confirm the differential phenotype of the GFP<sup>-</sup> and GFP<sup>+</sup> cells after culture reflected their functional status, we compared their ability to form colonies in methylcellulose. GFP<sup>+</sup> cells yielded significantly more colonies than GFP<sup>-</sup> cells (Fig. 3 G), suggesting that functionally primitive HSCs predominantly reside in the GFP<sup>+</sup> fraction. Collectively, these data indicate that in vitro culture of LSK GFP<sup>+</sup> cells leads to generation of GFP<sup>-</sup> cells that are more differentiated, and lend credence to the use of GFP as a fluorescent sensor for the differentiation state of hematopoietic cells.

To determine the cell-cycle distribution of LSK GFP<sup>-</sup> and LSK GFP<sup>+</sup> cells, we performed Hoechst 33342 and pyronin Y staining, which revealed that the majority of LSK GFP<sup>+</sup> cells were in G<sub>0</sub> phase, whereas a significant proportion of LSK GFP<sup>-</sup> cells were actively cycling (Fig. 3 H). These data indicate that, within HSPCs, Evi1 expression represents

a functionally distinct population that remains in an undifferentiated and quiescent state.

**Evi1 expression marks in vivo long-term multilineage repopulating HSCs in adult BM**

Based on the aforementioned data, we hypothesized that Evi1 expression would have the potential to effectively mark long-term multilineage repopulating HSCs. To examine this issue, we performed a CRA, in which 500 purified LSK GFP<sup>-</sup> or LSK GFP<sup>+</sup> cells were transplanted with 2 × 10<sup>5</sup> competitor BM cells into lethally irradiated recipients (Fig. S1 A). At 16 wk after transplantation, flow cytometric analysis of donor-derived cells revealed long-term reconstitution in all recipients transplanted with LSK GFP<sup>+</sup> cells (Fig. 4 A). Moreover, LSK GFP<sup>+</sup> cells displayed multilineage potential with robust contribution to myeloid, B, and T cells in peripheral blood (PB) as well as the LSK

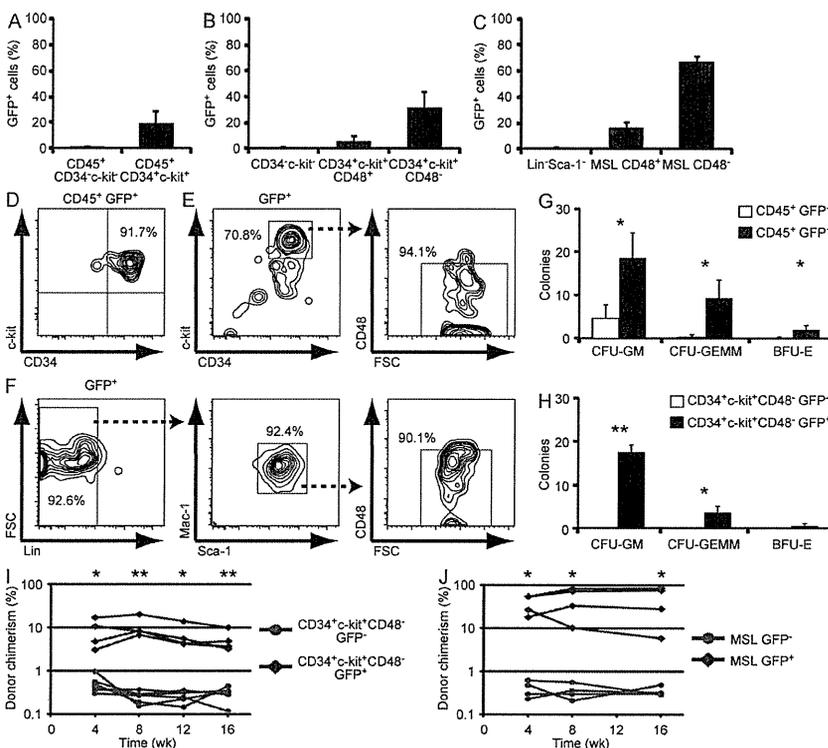


**Figure 4. Evi1 expression marks in vivo long-term multilineage repopulating HSCs in adult BM.** (A and B) PB donor chimerism in CRAs, in which 500 LSK GFP<sup>-</sup> or LSK GFP<sup>+</sup> cells sorted from *Evi1*<sup>+/GFP</sup> mice (Ly5.2) were transplanted into lethally irradiated recipients (Ly5.1) together with 2 × 10<sup>5</sup> competitor BM cells (Ly5.1 × Ly5.2). (A) Percentages of donor-derived cells (Ly5.2) in PB after transplantation are shown. Each dot indicates an individual recipient mouse (\*, P < 0.005; n = 6–7 from 2 independent experiments). (B) Percentages of donor-derived cells (Ly5.2) in myeloid, B, and T cells of PB and LSK cells of BM 16 wk after transplantation. Recipient mice in experiment 1 were used for the analysis of BM (\*, P < 0.01; n = 6–7 for PB and n = 3 for BM). (C) Percentages of donor-derived cells (Ly5.2) in PB of secondary recipient mice (Ly5.1) 16 wk after transplantation. Recipient mice in experiment 1 were used for secondary transplantation (\*, P < 0.0001, n = 3). (D–K) PB donor chimerism in CRAs, in which 100 CD48<sup>-</sup> CD150<sup>+</sup> LSK GFP<sup>-</sup> or CD48<sup>-</sup> CD150<sup>+</sup> LSK GFP<sup>+</sup> cells (D and E; n = 7–8), or 500 CD48<sup>-</sup> CD150<sup>-</sup> LSK GFP<sup>-</sup> or CD48<sup>-</sup> CD150<sup>-</sup> LSK GFP<sup>+</sup> cells (F and G; n = 7), or 100 Fik-2<sup>+</sup> CD34<sup>-</sup> LSK GFP<sup>-</sup> or Fik-2<sup>+</sup> CD34<sup>-</sup> LSK GFP<sup>+</sup> cells (H and I; n = 3–4), or 500 Fik-2<sup>+</sup> CD34<sup>+</sup> LSK GFP<sup>-</sup> or Fik-2<sup>+</sup> CD34<sup>+</sup> LSK GFP<sup>+</sup> cells (J and K; n = 3–4) sorted from *Evi1*<sup>+/GFP</sup> mice (Ly5.1) were transplanted into lethally irradiated recipients (Ly5.2) together with 2 × 10<sup>5</sup> competitor BM cells (Ly5.2). (D, F, H, and J) Percentages of donor-derived cells (Ly5.1) in total (left) and myeloid cells (right) of PB after transplantation are shown. Each dot indicates an individual recipient mouse (\*, P < 0.05; \*\*, P < 0.005; \*\*\*, P < 0.0005). (E, G, I, and K) Percentages of donor-derived cells (Ly5.1) in myeloid, B, and T cells of PB 16 wk after transplantation (\*, P < 0.05; \*\*, P < 0.005). Data represent mean ± SD.

fraction in BM (Fig. 4 B). In contrast, LSK GFP<sup>-</sup> cells yielded an almost total inability to generate long-term chimerism (Fig. 4, A and B), which suggests that this population is devoid of self-renewal activity. To confirm the *in vivo* repopulating capacity of LSK GFP<sup>+</sup> cells, we performed secondary transplantation. Similarly, LSK GFP<sup>+</sup> cells showed remarkable long-term reconstitution, whereas LSK GFP<sup>-</sup> cells consistently failed to produce detectable donor-derived cells (Fig. 4 C), demonstrating that *in vivo* long-term multilineage repopulating cells are exclusively enriched in the LSK GFP<sup>+</sup> fraction in adult BM.

To further refine our analysis designating Evi1 expression as a robust and reliable HSC marker, we compared the repopulating capacity of GFP<sup>-</sup> and GFP<sup>+</sup> cells within the CD48<sup>-</sup> CD150<sup>+</sup> LSK fraction, which is enriched for LT-HSCs (Fig. S1 B). Intriguingly, CD48<sup>-</sup> CD150<sup>+</sup> LSK GFP<sup>+</sup> cells exhibited long-term multilineage reconstitution, whereas no engraftment was observed in recipients of CD48<sup>-</sup> CD150<sup>+</sup> LSK GFP<sup>-</sup> cells (Fig. 4, D and E), suggesting that long-term repopulating HSCs predominantly reside in the GFP<sup>+</sup> fraction even within the highly subfractionated LT-HSC fraction. We then examined whether Evi1 expression is associated with repopulating capacity in the CD48<sup>+</sup> CD150<sup>-</sup> LSK fraction, which is enriched for ST-HSCs/MPPs with limited self-renewal activity (Fig. S1 B). Although CD48<sup>+</sup> CD150<sup>-</sup> LSK

GFP<sup>-</sup> cells provided only a transient reconstitution, CD48<sup>+</sup> CD150<sup>-</sup> LSK GFP<sup>+</sup> cells showed declining, but sustained engraftment 16 wk after transplantation (Fig. 4 F). In contrast to CD48<sup>-</sup> CD150<sup>+</sup> LSK GFP<sup>+</sup> cells, CD48<sup>+</sup> CD150<sup>-</sup> LSK GFP<sup>+</sup> cells mediated faint myeloid but superior lymphoid reconstitution (Fig. 4, F and G). Although it is controversial whether CD48<sup>+</sup> CD150<sup>-</sup> LSK cells are transiently reconstituting MPPs/ST-HSCs or lymphoid-biased LT-HSCs with limited long-term engraftment and strong predominance of lymphoid reconstitution (Kiel et al., 2005; Weksberg et al., 2008; Grassinger et al., 2010), Evi1-expressing cells possess higher repopulating capacity within this fraction. When we subfractionated the LSK fraction according to CD34 and Flk-2 expression, and compared the repopulating capacity of GFP<sup>-</sup> and GFP<sup>+</sup> cells within these subsets (Fig. S1 C), we obtained similar results to the aforementioned findings using SLAM markers (Fig. 4, H–K). These data reveal that, irrespective of the combination of HSC surface markers used, GFP<sup>+</sup> cells are the exclusive reservoir of HSC activity, with no reconstitution ability being observed in GFP<sup>-</sup> cells within the LT-HSC compartment. Altogether, our results demonstrate that Evi1 expression can further augment the conventional HSC purification strategy, and suggest that Evi1-IRES-GFP knock-in mice allow us to functionally identify HSCs on the ground of self-renewal capacity.



in CRAs, in which 500 MSL GFP<sup>-</sup> or MSL GFP<sup>+</sup> cells sorted from E14.5 FL of *Evi1*<sup>+/GFP</sup> embryos (Ly5.2) were transplanted into lethally irradiated recipients (Ly5.1 × Ly5.2) together with 2 × 10<sup>5</sup> competitor BM cells (Ly5.1 × Ly5.2). Percentages of donor-derived cells (Ly5.2) in PB weeks after transplantation are shown. Each dot indicates an individual recipient mouse (\*, *P* < 0.05; *n* = 4). Data represent mean ± SD.

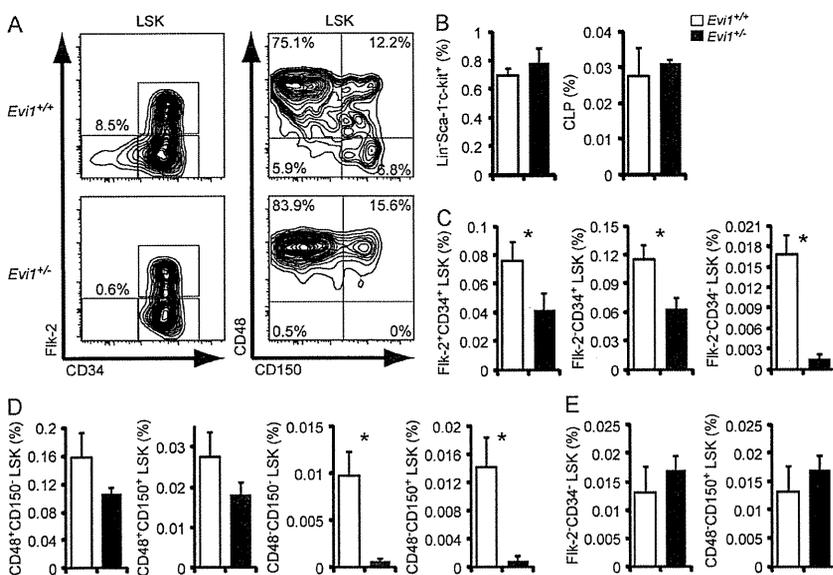
### Evi1 expression marks in vivo long-term multilineage repopulating HSCs in embryo

The formation of blood cells begins in the yolk sac of the embryo, and then shifts to the aorta-gonad-mesonephros (AGM) region, and then sequentially to the placenta, fetal liver (FL), and adult BM. There are several major phenotypic and functional differences between fetal and adult HSCs in surface marker profile, cell cycle status, self-renewal potential, gene expression profile, and regulatory mechanism (Mikkola and Orkin, 2006; Orkin and Zon, 2008). Fetal HSCs, in particular, divide rapidly and undergo massive expansion, whereas adult HSCs are mostly quiescent (Bowie et al., 2006). It is known that *Evi1* is highly expressed in the yolk sac, paraortic splanchnopleura, and HSPCs ( $CD45^+ CD34^+ c-kit^+$ ) in early embryo (Yuasa et al., 2005). Therefore, we sought to determine whether *Evi1* expression can mark fetal HSCs despite their distinct features from adult HSCs, and thus analyzed the expression pattern of GFP in *Evi1<sup>+/GFP</sup>* embryos. As expected, GFP expression was highly restricted to HSPCs in the embryonic tissues;  $CD45^+ CD34^+ c-kit^+$  cells in embryonic day 10.5 (E10.5) AGM,  $CD34^+ c-kit^+ CD48^-$  cells in E12.5 placenta, and  $Mac-1^+ Sca-1^+ Lin^-$  (MSL)  $CD48^-$  cells in E14.5 FL (Fig. 5, A-C; Takakura et al., 2000; Kim et al., 2006; McKinney-Freeman et al., 2009). When the distribution of GFP<sup>+</sup> cells in the fetal hematopoietic system was analyzed, most GFP<sup>+</sup> cells exhibited the HSPC-specific marker profile in all embryonic tissues examined (Fig. 5, D-F), indicating the predominant expression of *Evi1* in HSPCs during fetal hematopoiesis.

To determine whether *Evi1* expression is associated with hematopoietic activity in the embryonic tissues, we performed colony-forming assays in vitro using  $CD45^+ GFP^-$  and  $CD45^+ GFP^+$  cells from E10.5 AGM, and found that

$CD45^+ GFP^+$  cells contained almost all colony-forming cells, with few detectable hematopoietic colonies in  $CD45^+ GFP^-$  cells (Fig. 5 G). In the same manner, within the  $CD34^+ c-kit^+ CD48^-$  fraction from E12.5 placenta of *Evi1<sup>+/GFP</sup>* embryos, colony-forming activity was exclusively present in GFP<sup>+</sup> cells, regardless of colony type (Fig. 5 H). These data suggest that clonogenic hematopoietic progenitors predominantly reside in the *Evi1*-expressing fraction in fetal hematopoiesis.

To examine whether *Evi1* expression would have the potential to effectively mark long-term repopulating HSCs in embryo, we performed a CRA using sorted  $CD34^+ c-kit^+ CD48^- GFP^-$  and  $CD34^+ c-kit^+ CD48^- GFP^+$  cells from E12.5 placenta of *Evi1<sup>+/GFP</sup>* embryos (Fig. S2 A). It was obvious that  $CD34^+ c-kit^+ CD48^- GFP^+$  cells contributed to the long-term reconstitution of irradiated recipients, whereas donor chimerism was almost undetectable in mice transplanted with  $CD34^+ c-kit^+ CD48^- GFP^-$  cells (Fig. 5 I), which is in agreement with the results obtained with their adult counterparts. To further assess whether *Evi1* expression can enrich long-term repopulating HSCs in embryo, we performed a CRA using purified MSL GFP<sup>-</sup> and MSL GFP<sup>+</sup> cells from E14.5 FL (Fig. S2 B). Along with cells in E12.5 placenta, MSL GFP<sup>+</sup> cells gave rise to long-term multilineage reconstitution, whereas no engraftment was observed in recipients of MSL GFP<sup>-</sup> cells (Fig. 5 J). These results indicate that fetal HSCs with active *Evi1* transcription exclusively harbor stem cell activity. Collectively, despite the functional differences between fetal and adult HSCs, *Evi1* expression marks long-term multilineage repopulating HSCs throughout ontogeny, suggesting a specific relationship between *Evi1* expression and HSC self-renewal capacity.



**Figure 6. *Evi1* heterozygosity leads to an almost complete loss of LT-HSCs in a cell-autonomous manner.** (A) FACS analysis of expression of Flk-2 and CD34 or CD48 and CD150 on LSK cells in BM from *Evi1<sup>+/+</sup>* and *Evi1<sup>+/-</sup>* mice. Data are representative of at least three independent experiments. (B) Frequency of myeloid (Lin<sup>-</sup> Sca-1<sup>-</sup> c-kit<sup>+</sup>) and lymphoid progenitors (CLPs) in BM from *Evi1<sup>+/+</sup>* and *Evi1<sup>+/-</sup>* mice ( $n = 3$ ). (C) Frequency of Flk-2<sup>+</sup> CD34<sup>-</sup>, Flk-2<sup>-</sup> CD34<sup>+</sup>, and Flk-2<sup>-</sup> CD34<sup>-</sup> subsets within LSK cells in BM from *Evi1<sup>+/+</sup>* and *Evi1<sup>+/-</sup>* mice (\*,  $P < 0.0001$ ;  $n = 8$ ). (D) Frequency of CD48<sup>+</sup> CD150<sup>-</sup>, CD48<sup>-</sup> CD150<sup>+</sup>, and CD48<sup>-</sup> CD150<sup>-</sup> subsets within LSK cells in BM from *Evi1<sup>+/+</sup>* and *Evi1<sup>+/-</sup>* mice (\*,  $P < 0.01$ ;  $n = 3$ ). (E) Reciprocal transplantation assay was performed by transplantation of  $2 \times 10^5$  WT BM cells (Ly5.1) into lethally irradiated *Evi1<sup>+/+</sup>* or *Evi1<sup>+/-</sup>* mice (Ly5.2). Frequency of Flk-2<sup>-</sup> CD34<sup>-</sup> LSK or CD48<sup>-</sup> CD150<sup>+</sup> LSK cells in BM of recipients 16 wk after transplantation is shown ( $n = 4$ ). Data represent mean  $\pm$  SD.

### Evi1 heterozygosity leads to an almost complete loss of LT-HSCs in a cell-autonomous manner

The aforementioned observations led us to predict that Evi1 plays a functional role specifically in LT-HSCs. To clarify this issue, we analyzed heterozygous *Evi1* KO mice (*Evi1*<sup>+/-</sup>). We previously showed that heterozygosity of Evi1 leads to decreased numbers of LSK and CD34<sup>-</sup> LSK cells, as well as impaired long-term repopulating activity (Goyama et al., 2008). In the current study, although Flk-2<sup>+</sup> CD34<sup>+</sup> and Flk-2<sup>-</sup> CD34<sup>+</sup> LSK cells were moderately decreased, Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells from *Evi1*<sup>+/-</sup> mice exhibited a marked reduction in frequency compared with WT controls (Fig. 6, A and C). Likewise, when LSK cells were subdivided according to SLAM markers, we observed substantial decreases in CD48<sup>-</sup> CD150<sup>-</sup> and CD48<sup>+</sup> CD150<sup>+</sup> LSK subsets (Fig. 6, A and D). Therefore, the number of each subpopulation within the LSK fraction in *Evi1*<sup>+/-</sup> mice was declined in proportion to their expression level of Evi1, indicating that Evi1 has a dominating effect on the maintenance of LT-HSCs. In contrast, there were no significant differences in BM cellularity and the frequencies of lymphoid and myeloid progenitors, and mature blood cells between *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> mice (Fig. 6 B and not depicted), indicating that the differentiation potential to all mature lineages and committed progenitors is normal in *Evi1*<sup>+/-</sup> mice. Collectively, these observations suggest that Evi1 serves as a specific regulator in the earliest stage of adult hematopoietic development.

To exclude the possibility that a defect of BM micro-environment could be responsible for the observed hematopoietic abnormalities in *Evi1*<sup>+/-</sup> mice, we performed reciprocal transplantation experiments, in which WT BM cells were transplanted into lethally irradiated *Evi1*<sup>+/+</sup> or *Evi1*<sup>+/-</sup> mice. At 16 wk after transplantation, flow cytometric analysis showed no differences in the percentages of Flk-2<sup>-</sup> CD34<sup>-</sup> LSK or CD48<sup>-</sup> CD150<sup>+</sup> LSK cells in both groups of recipient mice (Fig. 6 E), demonstrating that the profound loss of LT-HSCs in *Evi1*<sup>+/-</sup> mice is attributed to cell-intrinsic mechanisms.

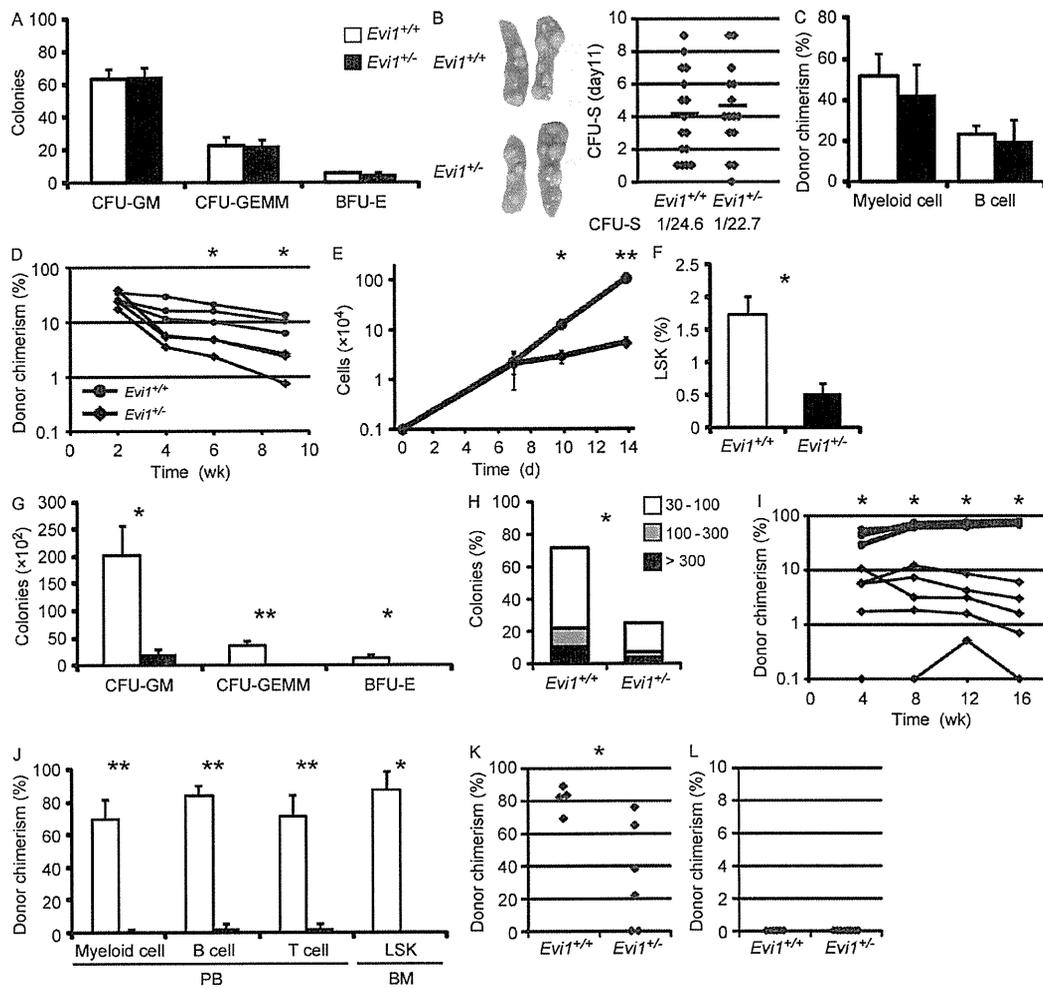
### Evi1 heterozygosity causes specific abrogation of self-renewal capacity in ST- and LT-HSCs

To further characterize which subpopulation in HSPCs is most dependent on Evi1, we purified CD34<sup>+</sup> and Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells from *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> mice and compared their differentiation and self-renewal capacity in vitro and in vivo. First, to assess the effect of Evi1 heterozygosity on the biological functions of ST-HSCs/MPPs, we performed colony-forming assays in vitro using *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> CD34<sup>+</sup> LSK cells, which demonstrated no significant differences in the number and type of colonies (Fig. 7 A). Similarly, we found the capacity of *Evi1*<sup>+/-</sup> CD34<sup>+</sup> LSK cells to form colonies in the spleen 11 d after transplantation (CFU-spleen [CFU-S]) was also equivalent to that of WT littermates (Fig. 7 B), indicating Evi1 is dispensable for the regulation of the differentiation and proliferation capacity in ST-HSCs/MPPs. Moreover, to investigate the self-renewal ability of ST-HSCs/MPPs in vivo, we evaluated the short-term

repopulating capacity of purified CD34<sup>+</sup> LSK cells using a CRA. At 2 wk after transplantation, we detected comparable frequencies of *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> CD34<sup>+</sup> LSK cell-derived myeloid and B cells (Fig. 7 C), suggesting that heterozygosity of Evi1 does not affect the engraftment and differentiation potential of ST-HSCs/MPPs in vivo. However, at later time points in the experiment, we found a moderate but significant decline in the percentage of donor-derived cells from *Evi1*<sup>+/-</sup> mice (Fig. 7 D). These data indicate that heterozygosity of Evi1 attenuates the self-renewal capacity of ST-HSCs/MPPs, but is not accompanied by any specific differentiation defects in them.

To assess whether Evi1 is required for the functions of LT-HSCs, we compared the self-renewal and proliferation capacity of *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells when cultured in serum-free medium. *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells showed comparable proliferation with WT cells for the first week of culture, but thereafter they exhibited pronouncedly impaired growth (Fig. 7 E). After incubation, a significantly lower proportion of cultured *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells remained in the LSK fraction than those from *Evi1*<sup>+/+</sup> mice (Fig. 7 F). In addition, we observed a prominent reduction of hematopoietic colonies contained in cultured *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells (Fig. 7 G). Besides, most of the colonies generated from cultured *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells consisted of only CFU-GM. These data indicate that heterozygosity of Evi1 results in accelerated loss of HSPCs, leading to the inefficient expansion of their progeny. To evaluate the colony-forming capacity at the single cell level, *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells were clonally sorted and cultured in serum-free medium. Evi1 heterozygosity diminished the colony-forming efficiency of clone-sorted Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells, and single *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells generated smaller colonies compared with control cells (Fig. 7 H), which indicates that the disruption of Evi1 gene not only decreases the number of clonogenic HSCs but also impairs the functional output per cell.

To assess the repopulating capacity of *Evi1*<sup>+/-</sup> LT-HSCs in vivo, we performed a CRA using purified Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells from *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> mice. Notably, *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells were almost unable to efficiently repopulate all mature lineages as well as stem and progenitor cells 16 wk after transplantation (Fig. 7, I and J), suggesting that LT-HSC function is critically dependent on Evi1 gene dosage. In a noncompetitive setting, although recipients of *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells had similar survival after transplantation (not depicted), *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells showed impaired engraftment (Fig. 7 K), suggesting that *Evi1*<sup>+/-</sup> HSCs were outcompeted by residual host HSCs. However, some of those recipients exhibited long-term multilineage reconstitution (Fig. 7 K and not depicted), confirming that the multipotent differentiation capacity is not abrogated in *Evi1*<sup>+/-</sup> mice. To further explore the competitive disadvantage of *Evi1*<sup>+/-</sup> HSCs, we transplanted WT BM cells into unirradiated *Evi1*<sup>+/+</sup> or *Evi1*<sup>+/-</sup> mice.



**Figure 7. Evi1 heterozygosity causes specific abrogation of self-renewal capacity in ST- and LT-HSCs.** (A) Numbers of CFU-GM, CFU-GEMM, and BFU-E colonies derived from 100 sorted *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> CD34<sup>+</sup> LSK cells (*n* = 3). (B) Appearance and number of CFU-S colonies in the spleen 11 d after injection of 100 sorted *Evi1*<sup>+/+</sup> or *Evi1*<sup>+/-</sup> CD34<sup>+</sup> LSK cells into lethally irradiated recipients. (left) Representative appearance is shown. (right) Data are shown as a dot plot and each bar represents mean (*n* = 15–16 from 3 independent experiments). (C and D) Short-term in vivo repopulating assay, in which 500 sorted *Evi1*<sup>+/+</sup> or *Evi1*<sup>+/-</sup> CD34<sup>+</sup> LSK cells (Ly5.2) were transplanted into lethally irradiated recipients (Ly5.1) together with  $2 \times 10^5$  competitor BM cells (Ly5.1). (C) Percentages of donor-derived myeloid and B cells (Ly5.2) in PB 2 wk after transplantation are shown (*n* = 3). (D) Short-term kinetics of the percentages of donor-derived cells (Ly5.2) in PB. Each dot indicates an individual recipient mouse (\*, *P* < 0.05; *n* = 3). (E) Proliferation of 1,000 sorted *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells cultured in serum-free medium supplemented with 20 ng/ml SCF and 20 ng/ml TPO for 14 d (\*, *P* < 0.01; \*\* *P* < 0.001; *n* = 3–4). (F) After 7 d of culture, the percentage of the remaining LSK fraction in cultured *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells was analyzed (\*, *P* < 0.005; *n* = 3). (G) In vitro colony-forming assay was performed to assess the numbers of CFU-GM, CFU-GEMM, and BFU-E colonies after 1,000 *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells were cultured for 14 d (\*, *P* < 0.01; \*\* *P* < 0.001; *n* = 3–4). (H) Single *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells were clone-sorted and cultured in serum-free medium. After 14 d of culture, cell numbers in each colony were analyzed. Their relative distribution is shown (\*, *P* < 0.0001; *n* = 192 clones from 2 independent experiments). (I–J) Long-term in vivo repopulating assay, in which 200 sorted *Evi1*<sup>+/+</sup> or *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells (Ly5.2) were transplanted into lethally irradiated recipients (Ly5.1) together with  $2 \times 10^5$  competitor BM cells (Ly5.1). (I) Percentages of donor-derived cells (Ly5.2) in PB after transplantation are shown. Each dot indicates an individual recipient mouse (\*, *P* < 0.0001, *n* = 5–6). (J) Percentages of donor-derived cells (Ly5.2) in myeloid, B, and T cells of PB and LSK cells of BM 16 wk after transplantation (\*, *P* < 0.001; \*\* *P* < 0.0001; *n* = 5–6 for PB and *n* = 3 for BM). (K) Noncompetitive repopulating assay, in which 200 sorted *Evi1*<sup>+/+</sup> or *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells (Ly5.2) were transplanted into lethally irradiated recipients (Ly5.1) without competitor. Percentages of donor-derived cells (Ly5.2) in PB of recipient mice that survived 12 wk after transplantation are shown (\*, *P* < 0.05, *n* = 4–6). (L) Reciprocal transplantation assay was performed by transplantation of  $2 \times 10^5$  WT BM cells (Ly5.1) into unirradiated *Evi1*<sup>+/+</sup> or *Evi1*<sup>+/-</sup> mice (Ly5.2). Percentages of donor-derived cells (Ly5.1) in PB 16 wk after transplantation are shown (*n* = 6–8). Data represent mean  $\pm$  SD.

We found no engraftment in both mice (Fig. 7 L), which indicates that the resistance to the donor HSC engraftment during steady-state hematopoiesis is maintained in *Evi1*<sup>+/-</sup> mice. Collectively, these data suggest that *Evi1* is dispensable for the regulation of proliferative and differentiation capacity of ST-HSCs/MPPs, but is strictly required for the maintenance of LT-HSC activity.

To investigate the mechanism behind the impaired HSC activity, we performed cell-cycle and apoptosis assays, but found no differences in the cell-cycle profile or apoptotic rates of Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells between *Evi1*<sup>+/+</sup> and *Evi1*<sup>+/-</sup> mice (unpublished data). Collectively, in consideration of the accelerated loss of LT-HSC activity in *Evi1*<sup>+/-</sup> mice, it is supposed that *Evi1* heterozygosity directs LT-HSCs from self-renewal toward differentiation to generate more committed progenitors, which is uncoupled from cell-cycle progression or apoptosis.

#### Forced expression of *Evi1* prevents HSPC differentiation and enhances their expansion

The findings noted above led us to hypothesize *Evi1* has the potential to inhibit differentiation and enhance HSC self-renewal independent of cell-cycle progression. To clarify this, we adopted a gain-of-function approach, in which WT LSK cells were transduced with *Evi1*, and then incubated in serum-free medium. Although forced expression of *Evi1* gave no apparent growth advantage for the first 10 d of culture, *Evi1*-transduced LSK cells subsequently manifested a mild but significant increase in proliferation rate (Fig. 8 A). Moreover, we found a substantial increase in the frequency of the remaining LSK fraction in cultured *Evi1*-transduced cells compared with control cells (Fig. 8 B). In parallel, the number of colonies derived from cultured *Evi1*-transduced LSK cell was drastically increased (Fig. 8 C). These results suggest that *Evi1* activation restricts lineage differentiation and enhances self-renewal activity of HSPCs. Collectively, our data provide compelling evidence that *Evi1* regulates the developmental transition from HSPCs to more committed progenitors, suggesting a crucial role of *Evi1* in controlling the balance between self-renewal and differentiation.

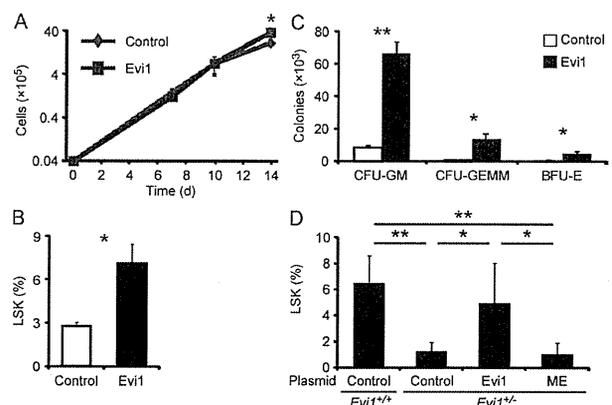
A recent work suggests that the longer, PR domain-containing isoform Mds1-*Evi1* (ME) deficiency alone causes a reduction in the number of HSCs with a loss of long-term repopulation capacity (Zhang et al., 2011). Because both ME and *Evi1* are inactivated in our *Evi1* KO model (Goyama et al., 2008), we attempted to genetically dissect the relative roles of ME and *Evi1* in maintaining LT-HSCs. For this purpose, we transduced *Evi1* or ME into *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells and examined whether they could maintain stem cell phenotype after in vitro culture. Reintroduction of *Evi1* led to a significant increase in the proportion that remained in the LSK fraction, similar to observations made in *Evi1*<sup>+/+</sup> cells (Fig. 8 D). However, retroviral transfer of ME was unable to normalize the frequency of the remaining LSK fraction (Fig. 8 D), indicating that *Evi1* preferentially rescues *Evi1*<sup>+/-</sup> LT-HSC defects. Given that ME has broader effects

on the hematopoietic system than *Evi1* and acts in part by maintaining HSC quiescence through up-regulation of *Cdkn1c* transcription (Zhang et al., 2011), *Evi1* and ME may exert their functions in regulating hematopoiesis at different stages and by different mechanisms.

#### DISCUSSION

In this study, we show that the amount of *Evi1* transcript can be indicative of an undifferentiated state with multipotent differentiation capacity within HSPCs. In both the fetal and adult hematopoietic systems, *Evi1* expression can mark long-term multilineage repopulating HSCs, and enhance HSC purification with a combination of other surface markers, suggesting a specific relationship between HSC activity and *Evi1* expression throughout ontogeny. This stem cell-specific expression pattern of *Evi1* allows us to functionally identify self-renewing HSCs by using *Evi1*-IRES-GFP knock-in mice, and suggests the relevance of *Evi1* in fine-tuning of stem cell properties. Indeed, we provide the genetic evidence confirming that *Evi1* has a predominant effect on LT-HSCs by specifically regulating their self-renewal capacity.

The prospective isolation of HSCs is the most important step to dissect their function. The strategy commonly used for HSC isolation is purification based on the expression of a combination of cell surface markers. However, some of these parameters differ between strains of mice, change dramatically during development, and are expressed on many non-HSCs.



**Figure 8. Forced expression of *Evi1* prevents HSPC differentiation and promotes their expansion.** (A) Proliferation of 3,000 control- or *Evi1*-transduced LSK cells cultured in serum-free medium with 20 ng/ml SCF and 20 ng/ml TPO for 14 d (\*,  $P < 0.05$ ;  $n = 3$ ). (B) After 7 d of culture, the percentage of the remaining LSK fraction in cultured control- or *Evi1*-transduced LSK cells was analyzed (\*,  $P < 0.01$ ;  $n = 3$ ). (C) In vitro colony-forming assay was performed to assess the numbers of CFU-GM, CFU-GEMM, and BFU-E colonies after 3,000 control- or *Evi1*-transduced LSK cells were cultured for 14 d (\*,  $P < 0.005$ ; \*\*,  $P < 0.0005$ ;  $n = 3$ ). (D) Control-, *Evi1*<sup>-</sup>, or ME-transduced *Evi1*<sup>+/+</sup> or *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells were cultured in medium containing 10% serum with 50 ng/ml SCF, 50 ng/ml TPO, 10 ng/ml IL-3, and 10 ng/ml IL-6 for 5 d, and the percentages of the remaining LSK fraction were analyzed (\*,  $P < 0.05$ ; \*\*,  $P < 0.0005$ ;  $n = 5-6$ ). Data represent mean  $\pm$  SD.

Here, we reveal that *Evi1* expression specifically correlates with functional HSCs, whereas lack of *Evi1* expression exclusively identifies cells without functional HSC activity in both the fetal and adult hematopoietic systems. In particular, *Evi1* expression can segregate long-term repopulating HSCs from cells without self-renewal potential even within the highly subfractionated Flk-2<sup>-</sup> CD34<sup>-</sup> LSK or CD48<sup>-</sup> CD150<sup>+</sup> LSK compartments. In addition, *Evi1*-IRES-GFP knock-in mice offer advantages over the conventional HSC surface markers, as the GFP<sup>-</sup> and GFP<sup>+</sup> subfractions of the Flk-2<sup>-</sup> CD34<sup>-</sup> LSK or CD48<sup>-</sup> CD150<sup>+</sup> LSK cells show a quite similar distribution of these markers (unpublished data). Moreover, our findings that *Evi1* specifically regulates the self-renewal capacity of HSCs guarantee the potential utility of *Evi1* expression as an indicator of HSC activity. Therefore, the *Evi1*-IRES-GFP knock-in mouse line provides a powerful approach for the functional identification of self-renewing HSCs *in vivo*, thus opening a new avenue for investigating HSC biology.

Although functional HSCs exclusively reside in the GFP<sup>+</sup> population, a proportion of GFP<sup>+</sup> cells lack HSC function. As GFP protein is quite stable and degraded more slowly than *Evi1* protein (unpublished data), these observations may reflect a remnant of GFP expression from cells that have just differentiated from GFP<sup>+</sup> HSCs. However, it is possible that *Evi1* expression distinguishes self-renewing ST-HSCs from cells with no self-renewal activity in the ST-HSC/MPP fraction, as *Evi1* heterozygosity affects the short-term repopulating capacity of CD34<sup>+</sup> LSK cells.

The present findings in the hematopoietic system encourage us to examine the possibility that *Evi1* expression serves as a selective marker for stem cells in other tissues or in cancer systems. We show that mesenchymal stem cells (MSCs), one of the few tissue stem cell types that have been established to self-renew *in vivo* (Morikawa et al., 2009), do not express *Evi1*. However, a mouse gene expression atlas and prior studies examining the expression pattern of *Evi1* in various tissues have reported *Evi1* expression in the kidney, ovary, uterus, intestine, stomach, lung, trachea, and nasal cavity in the adult mouse (Morishita et al., 1990; Perkins et al., 1991; Su et al., 2004). It will be interesting to determine, using *Evi1*-IRES-GFP knock-in mice, whether *Evi1*-expressing cells in these organs are enriched with tissue stem cells.

Our data suggest a unique association between *Evi1* expression and HSC self-renewal activity throughout hematopoietic ontogeny. Along with this stem cell-specific expression pattern of *Evi1*, the fact that the disruption of a single allele of *Evi1* leads to a near total loss of self-renewing HSCs implicates *Evi1* as a central regulator in HSC self-renewal. In addition, a recent gene expression profile analysis showed that *Evi1* binding sites are enriched in the upstream region of genes expressed selectively in LT-HSCs (Forsberg et al., 2010). In fact, several molecules involved in the regulation of HSC self-renewal have been identified as downstream targets or interacting proteins of *Evi1*, including *Gata2* (Sato et al., 2008; Yuasa et al., 2005), *Pbx1* (Shimabe et al., 2009), *Runx1*

(Senyuk et al., 2007), and TGF- $\beta$  (Kurokawa et al., 1998). Together with these findings, our data strongly support a model in which *Evi1* gene dosage is a critical determinant of HSC self-renewal potential.

Inappropriate expression of *EVI1* confers poor prognosis in patients with AML (Lugthart et al., 2008; Gröschel et al., 2010), and therefore improvement of the therapeutic outcome of leukemia with high *EVI1* expression is needed. In this study, we reveal that *Evi1* overexpression blocks differentiation and induces HSPC expansion. Our data fit with other studies showing that retroviral integration at the *Evi1* locus can be associated with long-term *in vivo* clonal dominance, occasionally leading to leukemic transformation (Stein et al., 2010). The genetic events underlying AML pathogenesis fall into two groups: (1) mutations that enhance proliferation and survival of hematopoietic progenitors, or (2) mutations that result in impaired differentiation or aberrant acquisition of self-renewal properties of HSPCs (Fröhling et al., 2005). Our data indicate that *Evi1* activation can function as the latter mutation and confer enhanced self-renewal capacity in myeloid neoplasms. In addition, we demonstrate that retroviral transfer of *Evi1*, but not ME, can ameliorate the self-renewal defects in *Evi1*<sup>+/-</sup> HSCs, highlighting a distinct role of *Evi1* in HSC self-renewal. These findings may explain the underlying mechanisms of the clinical observations that, irrespective of ME expression, aberrant *EVI1* expression carries an adverse prognostic value in AML (Lugthart et al., 2008, 2010). As it is becoming evident that leukemic stem cells share self-renewal machinery with normal HSCs, the elucidation of how *Evi1* controls HSC self-renewal may provide biological insight into the pathogenesis of *Evi1*-related leukemia.

## MATERIALS AND METHODS

**Generation of *Evi1*-IRES-GFP knock-in mice.** The targeting construct was assembled in the plasmid vector pBluescript KS. The 5' arm of the targeting vector consists of a 5.1-kb fragment of BAC clone RP24-481A14 and the 3' arm consists of a 3.1-kb fragment. The 5' arm contains *Evi1* intron 8 and exon 9, and the 3' arm contains intron 9, exon 10, and intron 10. Both arms were obtained by PCR using BAC clone RP24-481A14 as a template, all sequenced, and then inserted into pBluescript KS. Mouse *Evi1* cDNA was isolated from murine embryo cDNA by PCR, with an EcoRI site at the 5' end and a BamHI site at the 3' end, which was cloned into pBluescript KS. A 1.3-kb IRES-GFP cassette derived from pGCDNsam-IRES-GFP retroviral vector was inserted downstream of the aforementioned *Evi1* fragment. A polyadenylation (pA) cassette was then ligated 3' to the IRES-GFP cassette. A neomycin-positive selection (Neo<sup>r</sup>) cassette, expressed under control of the PGK promoter, was inserted downstream of the pA cassette. Both pA cassette and loxP-flanked neomycin-positive selection cassette were derived from DT-A/AFP(EGFP)/Neo vector (a gift from the Institute of Physical and Chemical Research Center for Developmental Biology, Kobe, Japan). The partial *Evi1* cDNA-IRES-GFP-pA-loxP-neo was released intact by digestion with *Sse8387I* and cloned into a unique *Sse8387I* site in exon 9 of BAC arm. A diphtheria toxin-negative selection cassette was cloned into pBluescript KS, 3' to the targeting construct. The targeting construct was linearized by *SacII* and transfected into TT2 ES cells by electroporation. Homologous recombinant clones were identified by Southern blot analysis of genomic DNA isolated from individual G418/FIAU-resistant ES cell colonies. The DNA was digested with *XbaI*, blotted to nylon membranes, and

hybridized with a 3' external *Evi1* probe. Confirmatory Southern blotting could detect a 9.1-kb WT *Evi1* allele band and a 4.1-kb correctly targeted *Evi1-IRES-GFP* allele band with this 3' probe. In EcoRV-digested genomic DNA from positive ES cell clones, a 5' external probe detected 10- and 11-kb bands from the WT and targeted alleles, respectively. Next, appropriately targeted ES clones were aggregated with 8-cell stage of ICR embryo, and resultant blastocysts were injected into pseudopregnant foster ICR mothers. Chimeric males, which gave germ-line transmission, were crossed with C57BL/6 females. Blastocyst injection and breeding of chimeras were performed in the Animal Center for Biomedical Research, University of Tokyo, Tokyo, Japan.

**Mice.** *Evi1-IRES-GFP* knock-in mice were backcrossed onto a C57BL/6 background (Ly5.1 or Ly5.2) for at least four generations (Sankyo-Laboratory Service). Heterozygous *Evi1* KO mice (*Evi1*<sup>-/-</sup> mice) were previously described (Goyama et al., 2008). C57BL/6-Ly5.1 mice were crossed with Ly5.2 mice to obtain Ly5.1/Ly5.2 mice. Littermates were used as controls in all experiments. All animal experiments were approved by the University of Tokyo Ethics Committee for Animal Experiments and strictly adhered to the guidelines for animal experiments of the University of Tokyo.

**Genotype analysis.** *Evi1*<sup>+/-</sup> mice were genotyped by PCR as previously described (Goyama et al., 2008). *Evi1*<sup>+/*GFP*</sup> mice were genotyped using a multiplex PCR to detect both WT and *Evi1-IRES-GFP* alleles. Genomic DNA was isolated from tail biopsies and subjected to PCR using *Neo* and *Evi1* primers. PCR with *Neo* primers detects the knock-in allele, and that with *Evi1* primers detects the WT allele. The PCR samples were denatured at 94°C for 2 min, subjected to 30 cycles of amplification (94°C for 30 s, 65°C for 1 min, and 72°C for 1 min), and followed by a final extension step at 72°C. PCR products were resolved by agarose gel electrophoresis. PCR primers are listed below: *Neo* primer forward, 5'-AGGGGATCCGCTG-TAAGTCT-3', reverse, 5'-GCACTGACTGCTCATCCAAA-3'; *Evi1* primer forward, 5'-ATGTCAGCAATTGAGAACATGG-3', reverse, 5'-ATCCAAAGGTCCTGAGTTCAAA-3'.

**Flow cytometry.** A list of antibodies is provided in Table S1. Stained cells were sorted with a FACSAriaII, and analysis was performed on LSRII (both from BD). A mixture of antibodies recognizing CD3, CD4, CD8, B220, TER-119, Mac-1, or Gr-1 was used to identify Lin<sup>+</sup> cells. Anti-CD127 antibody was added to the lineage mixture, except for the analysis of CLPs. The data analyses were performed with FlowJo software (Tree Star). In experiments with the *Evi1-IRES-GFP* knock-in mouse, a "fluorescence minus one" littermate control was analyzed in parallel to set GFP gates.

**Cell-cycle analyses.** For Hoechst 33342 and pyronin Y staining, cells were incubated with 5 ng/ml Hoechst 33342 (Invitrogen) and 25 μg/ml verapamil at 37°C for 45 min. Next, pyronin Y (Sigma-Aldrich) was added to 1 μg/ml, and the cells were incubated for an additional 15 min.

**In vitro culture.** For in vitro serum-free culture, cells were cultured in StemSpan SFEM (StemCell Technologies) supplemented with 20 ng/ml mouse SCF and 20 ng/ml human TPO, and subsequently subjected to flow cytometry or colony-forming assay at the indicated day after incubation. For colony-forming assay, cells were seeded in duplicate and cultured in cytokine-supplemented methylcellulose medium (MethoCult GF M3434; Stem Cell Technologies). Subsequently, colonies were counted and identified based on morphological examination on day 12. For in vitro differentiation, LSK GFP<sup>+</sup> cells were cultured in RPMI-1640 medium (Wako) containing 10% serum with 50 ng/ml mouse SCF, 50 ng/ml human TPO, 10 ng/ml mouse IL-3, and 10 ng/ml human IL-6, and subjected to flow cytometry or colony-forming assay after 5 d of incubation.

**Single-cell culture.** Cells were clone-sorted into 96-well plates using FACS-based automated cell deposition unit and cultured in StemSpan SFEM supplemented with 20 ng/ml mouse SCF and 20 ng/ml human TPO. After 14 d of culture, cell numbers in each colony were analyzed.

**In vivo transplantation assay.** Transplantation assays were performed using the Ly5 congenic mouse system. In CRAs, lethally irradiated (9.5 Gy) mice were reconstituted with the indicated subsets from *Evi1*<sup>+/-</sup>, *Evi1*<sup>+/*GFP*</sup>, or *Evi1*<sup>-/-</sup> mice, in competition with 2 × 10<sup>5</sup> unfractionated BM cells from congenic mice. For second BM transplantation, BM cells (1/2 femur equivalent) were obtained from recipient mice 16 wk after transplantation, and transplanted into a second set of lethally irradiated (9.5 Gy) mice. For reciprocal transplantation assays, lethally irradiated (9.5 Gy) or unirradiated *Evi1*<sup>+/-</sup> and *Evi1*<sup>-/-</sup> mice were transplanted with 2 × 10<sup>5</sup> WT BM cells without competitor cells. In non-CRAs, lethally irradiated (9.5 Gy) mice were reconstituted with *Evi1*<sup>+/-</sup> and *Evi1*<sup>+/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells without competitor cells. Reconstitution of donor-derived cells was monitored by staining PB cells with antibodies against Ly5.1, Ly5.2, CD3, CD4, CD8, B220, Mac-1, and Gr-1. When CD48<sup>+</sup> CD150<sup>-</sup> LSK cells were transplanted, nonblocking anti-CD48 antibody (MRC OX78 clone) was used (Grassinger et al., 2010).

**CFU-S assay.** For CFU-S assay, 100 CD34<sup>+</sup> LSK cells were injected into lethally irradiated (9.5 Gy) mice. Spleens in transplanted mice were isolated 11 d later, and visually inspected for the presence of macroscopic colonies after fixation in Tellyesniczky's solution.

**RQ-PCR.** Total RNA was prepared using RNeasy Mini kit (QIAGEN), then cDNA was synthesized with a QuantiTect Reverse Transcription kit (QIAGEN), and used for RQ-PCR with FastStart SYBR Green Master and LightCycler 480 System (Roche Applied Science) according to the manufacturer's instructions. All assays were performed in triplicate and relative expression was normalized to the internal control *GAPDH*. PCR primers are listed below: *GAPDH* primer forward, 5'-CCATCACCATCTTC-CAGGAG-3', reverse, 5'-CCTGCTTCACCACCTCTTG-3'; *Evi1* primer forward, 5'-ATCGGAAGATCTTAGATGAGTTTTG-3', reverse, 5'-CTTCTACATCTGGTTGACTGG-3'.

**Western blotting.** Western blotting was performed as previously described (Goyama et al., 2008). In brief, mouse embryo fibroblast cells were lysed in TNE buffer, subjected to 7% SDS-PAGE, and transferred to a PVDF membrane (Millipore). The blot was incubated with an Evi-1 (C50E12) rabbit monoclonal antibody (Cell Signaling Technology), and visualized by ECL Plus (GE Healthcare).

**AGM and placental cell preparation.** The day of vaginal plug observation was considered as day 0.5 postcoitum (E0.5). E10.5 AGM region or E12.5 placenta were carefully dissected from embryos, dissociated by incubation with 250 U/ml dispase (Godo Shusei) for 20 min and cell dissociation buffer (Invitrogen) for 20 min at 37°C, and followed by passages through 18–25 G needles. Single cell suspensions were filtered through 70-μm cell strainer and analyzed by flow cytometry.

**Endothelial cell (EC), osteoblast (OB), and MSC preparation.** After BM cells were flushed out, bones were crushed with a pestle and mortar. Then bone fragments were incubated with a Collagenase/Dispase (1 mg/ml; Roche Applied Science) in MEM α (Wako) with 20% serum and gently agitated for 90 min at 37°C. The dissociated cells were collected, and bone-associated mononuclear cells were isolated with the use of density centrifugation with Histopaque-1083 (Sigma-Aldrich). ECs were defined as CD31<sup>+</sup> TER-119<sup>-</sup> CD45<sup>-</sup>, OBs were defined as CD31<sup>-</sup> TER-119<sup>-</sup> CD45<sup>-</sup> Sca-1<sup>-</sup> ALCAM<sup>+</sup> cells (Nakamura et al., 2010), and MSCs were defined as CD31<sup>-</sup> TER-119<sup>-</sup> CD45<sup>-</sup> Sca-1<sup>+</sup> PDGFRα<sup>+</sup> cells (Morikawa et al., 2009).

**Plasmid construct and retroviral transduction of LSK cells.** The murine *Evi1* or *ME* cDNA were inserted into a site upstream of an IRES-EGFP cassette in the retroviral vector pGCDNsam. To produce *Evi1-GFP*-expressing retrovirus, Plat-E packaging cells were transiently transfected with retroviral constructs by using FuGENE 6 transfection reagent (Roche). LSK cells were purified and incubated in StemSpan SFEM medium and cytokines (100 ng/ml mouse SCF and 100 ng/ml human TPO) for 24 h.

Next, cultured LSK cells were infected with Evi1-GFP retrovirus in the presence of RetroNectin (Takara Bio Inc.). The infected LSK cells were harvested 36 h after retrovirus infection, and GFP<sup>+</sup> cells were sorted and subjected to *in vitro* culture. For retroviral transduction of Evi1 or ME into Evi1<sup>+/+</sup> and Evi1<sup>-/-</sup> Flk-2<sup>-</sup> CD34<sup>-</sup> LSK cells, those cells were sorted and immediately infected with Evi1- or ME-GFP retroviruses in the presence of RetroNectin. These cells were incubated in RPMI-1640 medium containing 10% serum and cytokines (50 ng/ml mouse SCF, 50 ng/ml human TPO, 10 ng/ml mouse IL-3, and 10 ng/ml human IL-6). After 5 d of culture, the percentage of the remaining LSK fraction in GFP<sup>+</sup> cells was analyzed by flow cytometry.

**Statistical analysis.** Statistical significance of differences between parameters was assessed using a two-tailed unpaired Student's *t* test or Wilcoxon rank sum test.

**Online supplemental material.** Fig. S1 shows FACS gating strategy used to identify GFP<sup>+</sup> population from adult BM of Evi1<sup>+/GFP</sup> mice. Fig. S2 shows FACS gating strategy used to identify GFP<sup>+</sup> population from E12.5 placenta or E14.5 FL of Evi1<sup>+/GFP</sup> embryos. Table S1 lists the antibodies used for flow cytometry. Online supplemental material is available at <http://www.jem.org/cgi/content/full/jem.20110447/DC1>.

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## AML1/RUNX1 functions as a cytoplasmic attenuator of NF- $\kappa$ B signaling in the repression of myeloid tumors

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Functional deregulation of transcription factors has been found in many types of tumors. Transcription factor AML1/RUNX1 is one of the most frequent targets of chromosomal abnormalities in human leukemia and altered function of AML1 is closely associated with malignant transformation of hematopoietic cells. However, the molecular basis and therapeutic targets of AML1-related leukemia are still elusive. Here, we explored immediate tar-

get pathways of AML1 by in vitro synchronous inactivation in hematopoietic cells. We found that AML1 inhibits NF- $\kappa$ B signaling through interaction with I $\kappa$ B kinase complex in the cytoplasm. Remarkably, AML1 mutants found in myeloid tumors lack the ability to inhibit NF- $\kappa$ B signaling, and human cases with AML1-related leukemia exhibits distinctly activated NF- $\kappa$ B signaling. Furthermore, inhibition of NF- $\kappa$ B signaling in leukemic cells with

mutated AML1 efficiently blocks their growth and development of leukemia. These findings reveal a novel role for AML1 as a cytoplasmic attenuator of NF- $\kappa$ B signaling and indicate that NF- $\kappa$ B signaling is one of the promising therapeutic targets of hematologic malignancies with AML1 abnormality. (*Blood*. 2011; 118(25):6626-6637)

### Introduction

Functional disruption of tumor-suppressive transcription factors, such as p53, has been widely found in many types of tumors. Although great efforts have been made to reactivate wild-type p53,<sup>1</sup> therapeutic interventions to impaired transcription factors still need innovative strategy. To overcome this difficulty, we need to seek the treatable and pathogenetic targets for deregulated transcription factors.

Transcription factor AML1, also known as RUNX1, is one of the most frequent targets of chromosomal abnormalities in human leukemia.<sup>2</sup> Functional impairment of AML1 caused by point mutation also is reported in patients with leukemia or myelodysplastic syndrome (MDS).<sup>3-6</sup> Patients who have *AML1* mutations are reported to be accompanied with poor prognosis.<sup>5,6</sup> Genetic disruptions of *AML1* also are known to cause familial platelet disorder, with predisposition to acute myelogenous leukemia (AML).<sup>7</sup> AML1 is a critical regulator in hematopoiesis and has an essential function in the establishment of definitive hematopoiesis, differentiation of lymphocytes, maturation of megakaryocytes, and regulation of hematopoietic stem cells (HSCs).<sup>8-11</sup> Point mutations of *AML1* have been found throughout the length of this gene in myeloid tumors such as AML or MDS; most of the mutants lose the potential to activate gene transcription, whereas some mutants show a dominant-negative effect over AML1 function.<sup>4,12</sup> The types of mutations are similar between AML and MDS. Significantly, it was recently reported that AML1 mutants cause MDS/AML in a mouse bone marrow transplantation (BMT) model.<sup>13</sup> To date, many target genes of AML1 have been reported.<sup>14,15</sup> However,

the signaling pathways involved in the pathogenesis of AML1-related leukemia are still elusive.

t(8;21) produces the chimeric protein AML1/ETO and is one of the most frequent chromosomal translocations found in AML. AML1/ETO is constituted of the N-terminal AML1-derived part and the C-terminal ETO part that contain a DNA-binding domain and a corepressor-binding domain, respectively. Besides the defect of trans-activation potential, AML1/ETO is a potent repressor of gene transcription and acts as a dominant-negative mutant of AML1.

Nuclear factor- $\kappa$ B is a dimeric complex of transcription factors mainly consisting of p65 (RelA)/p50 (NF $\kappa$ B1) or RelB/p52 (NF $\kappa$ B2). There are 2 major pathways in NF- $\kappa$ B signaling: the canonical pathway that broadly modulates cell proliferation, survival, or inflammation; and the noncanonical pathway that mainly controls lymphogenesis or B-cell maturation.<sup>16</sup> In the canonical pathway, p65 and p50 (NF $\kappa$ B1) constitute NF- $\kappa$ B complex and are localized in the cytoplasm with I $\kappa$ B in a steady state. Once inflammation is induced, TNF- $\alpha$  stimulates its receptor that in turn activates the I $\kappa$ B kinase (IKK) complex. Then, I $\kappa$ B is phosphorylated by the activated IKK complex and subsequently degraded through the ubiquitin-proteasome pathway, resulting in nuclear translocation of p65/p50 and transactivation of NF- $\kappa$ B target genes. In the noncanonical pathway, RelB stably binds to p100 (NF $\kappa$ B2) in the cytoplasm. During the maturation of B cells, B cell-activating factor of the TNF family (BAFF) stimulates its receptor and activates IKK complex that phosphorylates and processes p100 to p52, resulting in nuclear translocation.<sup>17</sup> Although these

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2 pathways usually act in distinct physiologic conditions, their regulations are closely related, as indicated by the fact that transcriptional control of NF $\kappa$ B2 is directly regulated by p65.<sup>18</sup> Recently, deregulated activation of NF- $\kappa$ B signaling has been reported to be responsible for many types of tumors, including hematologic malignancies.<sup>19-22</sup>

In this study, we revealed that AML1 represses NF- $\kappa$ B signaling in the hematopoietic cells. This repression is achieved by inhibition of the kinase activity of IKK through physical interaction between AML1 and IKK. Remarkably, aberrant activation of NF- $\kappa$ B signaling is induced not only by targeted disruption but also by leukemia-related gene alteration of *AML1*. Furthermore, we demonstrate that blocking the NF- $\kappa$ B signaling efficiently represses proliferation of leukemic cells with mutant AML1, including chimeric proteins, and development of AML1-related leukemia in mice.

## Methods

### Mice

AML1 cKO mice were described previously.<sup>11</sup> Homozygous AML1 floxed mice (AML1 *f/f*) and Mx-Cre-expressing homozygous AML1 floxed mice [AML1 *f/f* Mx (+)] were kept at the Animal Center for Biomedical Research (University of Tokyo, Tokyo, Japan), according to the institutional guidelines. NOD/Shi-scid, IL-2 $\gamma$  null (NOG) mice were purchased from the Central Institute for Experimental Animals (Kawasaki, Japan).

### Retrovirus production, BM harvest, and serial replating assays

To produce oncoprotein-expressing retrovirus, Plat-E packaging cells<sup>23</sup> were transiently transfected with retroviral constructs, as described previously.<sup>24,25</sup> To produce green fluorescent protein (GFP)- or Cre recombinase fused to the estrogen receptor (CreER)-GFP-expressing retrovirus, we used MP34 packaging cells transduced with pGCDNsam-enhanced (e)GFP (a gift from H. Nakauchi and M. Onodera, University of Tokyo) or pGCDNsam-eGFP-CreER. The viral supernatant was collected from the transfected Plat-E cells, filtered, and used for BM progenitor infection. Six- to 8-week-old female C57/B6 mice were primed with 3 mg of 5-fluorouracil 5 days before BM harvest (day 1). BM cells flushed from the tibia and femur were isolated by density centrifugation over Histopaque-1083 (Sigma-Aldrich) and were allowed to rest overnight in RPMI-1640 with 10% FCS, 1% penicillin streptomycin (PS), and cytokines (50 ng/mL SCF, 50 ng/mL megakaryocyte growth and development factor, 50 ng/mL fusion protein of IL-6R and IL-6, and 50 ng/mL Flt-3 ligand). The following day (day 2), the BM cells were combined with retroviral supernatant and RetroNectin (Takara). On day 4, the infected BM were harvested and resuspended in cold PBS (final concentration,  $1 \times 10^4/\mu\text{L}$ ). Serial replating assays were performed at a density of  $10^4/\text{mL}$  replated in duplicate every 7 days in Methocult 3434 (StemCell Technologies). BMS-345541 (Calbiochem) was reconstituted with DMSO (Sigma-Aldrich) and added to methylcellulose of the third generation colony-forming cells.

### Microarray analysis

These procedures were performed as described previously.<sup>25</sup> Lineage-, Sca-1+, c-Kit+ (LSK) cells derived from AML1 *f/f* mice were transduced with GFP or CreER-GFP-expressing retroviruses. GFP-positive cells were sorted by an FACS Aria cell sorter (BD Biosciences). Total RNA was prepared from sorted GFP+ cells after 24 or 48 hours from adding 4-hydroxytamoxifen (4-OHT; Sigma-Aldrich) using the RNeasy micro kit (QIAGEN). Amplification and biotin labeling of fragmented cDNA was carried out using the NuGen Ovation Biotin labeling system (NuGEN Technologies) in duplicate. Labeled probes

were hybridized to the Mouse Genome 430 2.0 Array (Affymetrix) and scanned with a GeneChip Scanner 3000 7G (Affymetrix). Expression data were extracted from image files produced on GeneChip Operating software 1.0 (Affymetrix). Normalization and expression value calculation were performed using DNA-Chip Analyzer ([www.dchip.org](http://www.dchip.org)).<sup>26</sup> Normalized data were filtered for minimal expression and then tested for gene set enrichment using gene set enrichment analysis (GSEA) Version 2.0 ([www.broad.mit.edu/gsea/](http://www.broad.mit.edu/gsea/)).<sup>27</sup> GSEA enrichment results were filtered for statistical significance using a nominal *P* value threshold of .05. We also used gene expression data of 285 individuals with AML ([www.ncbi.nlm.nih.gov/geo/](http://www.ncbi.nlm.nih.gov/geo/); accession GSE1159 [NCBI GEO])<sup>28</sup> and expression data of AML1/ETO transduced human cord blood cells ([www.ncbi.nlm.nih.gov/geo/](http://www.ncbi.nlm.nih.gov/geo/); accessions GSE8023, GSE7011 [NCBI GEO]).<sup>29,30</sup>

### Nuclear translocation assay

The HEK293T cells were cultured on glass coverslips and transfected with Lipofectamine (Invitrogen) according to the manufacturer's instructions. Thirty-six hours later, conditioning medium was replaced with starvation medium containing 0.5% FCS medium. Twelve hours later, 100 ng/mL TNF- $\alpha$  (Sigma-Aldrich) was added to the cells for 5 to 20 minutes. The cells were washed 3 times with PBS and fixed in 3.7% formaldehyde in PBS for 20 minutes. After washing with PBS, the cell membrane was permeabilized by treatment with 0.2% Triton in PBS for 10 minutes, and the cell membrane was blocked by treatment with 1% BSA in PBS for 40 minutes. The cells were treated with mouse anti-FLAG monoclonal antibody (dilution, 1:200; Sigma-Aldrich) and rabbit anti-p65 monoclonal antibody (dilution, 1:100; Santa Cruz Biotechnology) for 3 hours. After washing with PBS, the cells were stained with Alexa Fluor 555 goat anti-mouse IgG (dilution, 1:500; Invitrogen), Alexa Fluor 488 goat anti-rabbit IgG (dilution, 1:500; Invitrogen), and TO-PRO3 (dilution, 1:1000; Invitrogen) for 1 hour. The cells were washed and treated with ProLong Gold antifade reagent (Invitrogen), and the proteins were visualized using a confocal microscope (63 $\times$ /1.4 NA oil objective, TCS-LS, Leica). FLAG-tagged AML1 mutants were inserted into pME18S expression vector.<sup>24</sup>

### Quantification of nuclear p65

Nuclear intensity of p65 was quantified with ImageJ Version 1.41o software (National Institutes of Health).<sup>31</sup> Mean nuclear intensities of 10 cells from 2 pictures were quantified. Intensities were normalized by background subtraction.

### Quantitative real-time PCR

Real-time PCR was performed using the ABI PRISM 7000 Sequence Detection System (Applied Biosystems) or LightCycler 480 (Roche Diagnostics) according to the manufacturers' instructions. Results were normalized to GAPDH levels. PCR primers used for quantitative PCR were as follows: Nfkb2f, ACCAAGCTCCATGCTAATGTGAAT; Nfkb2r, GGGTGTGTTTCCAGCAAAGGT; GAPDHf, TGGTGAAGCAGGCATCTGAG; GAPDHr, TGCTGTTGAAGTCGCAGGAG; Cd74f, GCAGTGGCTCTT-GTTTGAAG; Cd74r, TTCTGGCACTTGGTCAGTA; Nfkb1af, CGAGGAGTACGAGCAAATGG; Nfkb1ar, TGATTGCCAAGTGCAGGA; Nfkb1ef, CGACTCTCTGCTGCTGAATG; Nfkb1er, GGTCATCGAAGG-GCAAATAA; Plekf, CCGGCTACCTGCACACTA; Plekr, CAGCCTT-CAAGTGAAGTGC; Tnfaip3f, AAGCTCGTGGCTCTGAAAAC; Tnfaip3r, CCCCACATGTACTGACAAGC; Birc3f, AGAGAGGAGCAGATG-GAGCA; Birc3r, TTTGTTCTTCCGGATTAGTGC; Il2rgf, AGCGGAGCT-GTACAGAAAGCTA; Il2rgf, CTGGGATTCACATTGCT; Relbf, GC-CTTGGGTTCCAGTGAC; and Relbr, TGTATTCTCGATGATTCCAA.

### Flow cytometric analysis

Cells were sorted with an FACS Aria cell sorter, and analysis was performed on an FACS LSRII flow cytometer (BD Biosciences). Intracellular staining of phospho-p65 was done according to the manufacturer's protocol (Cell

Signaling Technology). To gate AML1-transduced cells, we used Pcdcf3-IRES-GFP or Pcdcf3-IRES-GFP-AML1 vector.

### Immunoprecipitation and Western blotting

For immunoprecipitation, cell lysates were incubated with the anti-FLAG M2 monoclonal antibody (Sigma-Aldrich), anti-AML1 monoclonal antibody (Cell Signaling Technology), normal rabbit IgG (Santa Cruz Biotechnology) for 1 hour at 4°C. Then, the samples were incubated with protein G-Sepharose (GE Healthcare) for 1 hour at 4°C. The precipitates were washed twice with the TNE buffer<sup>32</sup>, twice with high salt-containing wash buffer (50mM Tris-HCl, pH 7.5, 500mM NaCl, 0.1% Nonidet P-40, and 0.05% sodium deoxycholate), and once with low salt-containing buffer (50mM Tris-HCl, pH 7.5, 0.1% Nonidet P-40, and 0.05% sodium deoxycholate), and they were subjected to sodium dodecylsulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and analysis by Western blotting. Antibodies used in immunoblotting were as follows: anti-FLAG (Sigma-Aldrich), anti-hemagglutinin (HA; Roche Diagnostics), anti-AML1 (Cell Signaling Technology), anti-IKK $\alpha$  (Cell Signaling Technology), anti-IKK $\beta$  (Cell Signaling Technology), anti-I $\kappa$ B $\alpha$  (Cell Signaling Technology), anti-p65 (Cell Signaling Technology), anti-NF- $\kappa$ B2 p100/p52 (Cell Signaling Technology), and anti- $\beta$ -actin (Cell Signaling Technology). ECL detection (GE Healthcare) was carried out according to the manufacturer's recommendations. Protein levels were quantified with ImageJ Version 1.41o software.<sup>31</sup> Nuclear fraction or cytoplasmic fraction was prepared using a nuclear extract kit (Active Motif) according to the manufacturer's instructions. Splenic B cells were separated by an auto-MACS separation column (Miltenyi Biotec) according to the manufacturer's protocol. These B cells were stimulated with 200 ng/mL recombinant human BAFF (BioSource) for 8 hours. HA-tagged AML1 mutants were inserted into pME18S expression vector.<sup>24</sup> FLAG-tagged IKK $\alpha$  and IKK $\beta$  were gifts from Dr T. D. Gilmore (Boston University, Boston, MA).

### Leukemogenesis assays in vivo

These procedures were performed as described previously.<sup>13</sup> In brief, BM mononuclear cells were prestimulated and infected for 60 hours with the retroviruses harboring AML1 S291fsX300-IRES-GFP. Then,  $0.2$  to  $1.2 \times 10^6$  infected BM cells (Ly5.1) were injected through the tail vein into C57/B6 (Ly5.2) recipient mice (8 weeks of age) that had been administered a sublethal dose of 5.25 Gy of total body  $\gamma$ -irradiation. In survival assays,  $2 \times 10^6$  S291fsX300-induced leukemic spleen cells or  $1 \times 10^5$  MLL/ENL-induced leukemic spleen cells were injected into sublethally irradiated (7.5 Gy) C57/B6 (Ly5.2) mice. Bortezomib (LC Laboratories) was administered at doses of 1.0 mg/kg, given by intraperitoneal bolus injection twice weekly.

### Orthotopic cell line model and in vivo bioluminescence imaging

These procedures were performed as described previously.<sup>33</sup> In brief, 1 million SKNO-1 cells were injected into sublethally irradiated (2.0 Gy) NOG mice via tail vein. One week after injection, bortezomib was administered at doses of 1.0 mg/kg, given by intraperitoneal bolus injection twice weekly. Total body bioluminescence was determined by in vivo bioluminescence imaging (IVIS Lumina2; Caliper Life Sciences) and quantitated using Living Image 2.60 software (Caliper Life Sciences). Luciferase positive SKNO-1 cells were kindly provided by Dr Andrew L. Kung (Dana-Farber Cancer Institute, Boston, MA).

### In vitro kinase assay

These procedures were performed as described previously.<sup>34</sup> In brief, HEK293T cells transiently expressing either AML1 or mock vector were treated with 100 ng/mL TNF- $\alpha$  (Sigma-Aldrich) for the indicated times and then harvested. Cell extracts were precipitated with anti-IKK antibody (Cell Signaling Technology) for 1 hour at 4°C, and protein G-Sepharose was added for 1 hour at 4°C. Immune complexes were washed twice with lysis buffer and then washed twice with kinase buffer (20mM HEPES, pH 7.4,

1mM MnCl<sub>2</sub>, 5mM MgCl<sub>2</sub>, 10mM  $\beta$ -glycerol phosphate, 0.1mM sodium orthovanadate, 2mM NaF, and 1mM dithiothreitol). Kinase assays were performed for 30 minutes at 30°C in 20  $\mu$ L of kinase buffer containing 5  $\mu$ Ci of [ $\gamma$ -<sup>32</sup>P]ATP (GE healthcare) and 1  $\mu$ g of GST-I $\kappa$ B $\alpha$  (Millipore) as a substrate. The reaction mixtures were resolved by SDS-PAGE and then detected by autoradiography and quantified with ImageJ Version 1.41o software.<sup>31</sup>

### Statistical analysis

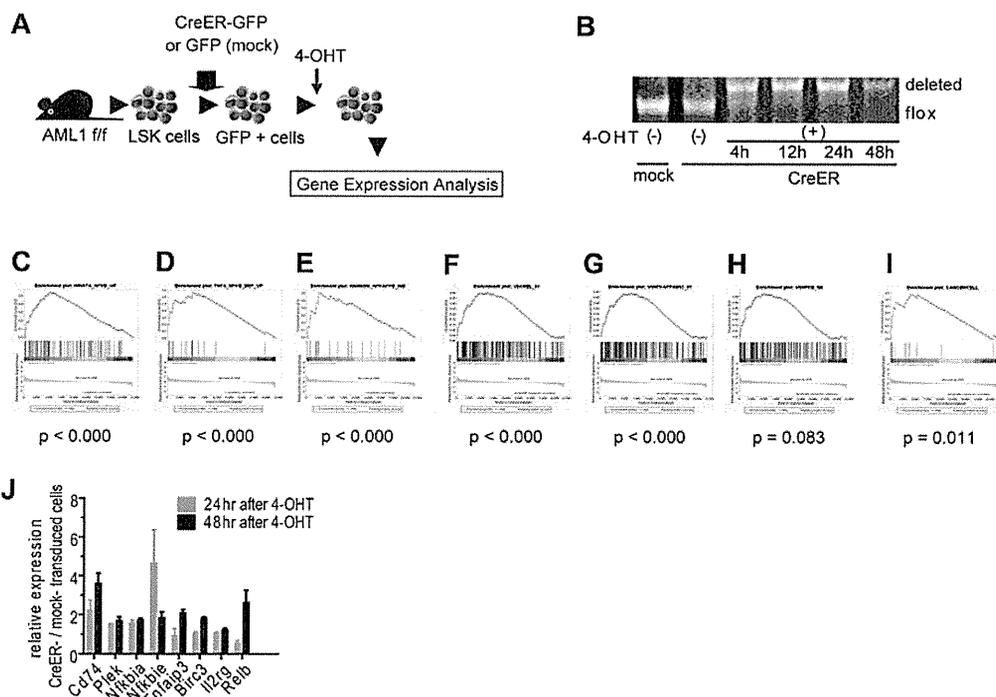
To compare data between groups, unpaired Student *t* test was used when equal variance were met by F-test. When unequal variances were detected, the Welch *t* test was used. Differences were considered statistically significant at a *P* value of less than .05. To analyze the survival curve, log-rank (Mantel-Cox) test was used. These analysis were done using Prism 5 (GraphPad Software).

## Results

### NF- $\kappa$ B signaling is activated in AML1-deficient cells

To seek the therapeutic target in AML1-related leukemia, we investigated the deregulated signaling pathway induced by loss of AML1. We analyzed the change of gene expression profiles in immature hematopoietic cells induced by AML1 inactivation. We purified LSK cells as hematopoietic stem and progenitor cells from AML1 conditional knockout (cKO) mice and retrovirally transduced CreER into them.<sup>35</sup> Immediately after deleting AML1 by the activation of CreER with 4-OHT, we collected them and analyzed their gene expression profiles with oligonucleotide microarray (Figure 1A, and supplemental Figure 1A, available on the *Blood* Web site; see the Supplemental Materials link at the top of the online article). As a control, we used LSK cells infected with an empty retrovirus and treated them with 4-OHT. We confirmed that AML1 was efficiently deleted after the addition of 4-OHT (Figure 1B). To analyze their gene expression profiles, we used GSEA with molecular signature database.<sup>27,36</sup> First, we analyzed the expression profiles with all curated gene sets (C2) and found that sets of NF- $\kappa$ B-related genes are significantly enriched in AML1-deficient cells (AML1 flox/flox, CreER+, 24 and 48 hours after adding 4-OHT; Figure 1C-E and supplemental Figure 1B-D).<sup>37-39</sup> Next, we analyzed them with all motif gene sets (C3), and again we found that NF- $\kappa$ B-regulated genes are significantly enriched in the AML1-deficient cells (Figure 1F-H and supplemental Figure 1E-G). In addition, the target genes of NF- $\kappa$ B signaling detected in hematopoietic malignant cells were distinctly enriched in AML1-deficient cells (Figure 1I and supplemental Figure 1H).<sup>21</sup> We confirmed the elevated expression of NF- $\kappa$ B target genes by real-time PCR (Figure 1J). Because we added 4-OHT both to mock- and CreER-transduced cells, the effects of 4-OHT to our gene expression analysis are unlikely.

To confirm that NF- $\kappa$ B signaling is activated in AML1-deficient cells, we performed immunofluorescent staining of p65, a major component of NF- $\kappa$ B in the canonical pathway, in AML1-deficient hematopoietic progenitor cells. As shown in Figure 2A, nuclear localized p65 was increased in AML1-deficient cells (AML1 flox/flox, CreER). This increase was attenuated by BMS-345541, an inhibitor of IKK. We also found that nuclear p65 protein was increased in AML1-deficient hematopoietic progenitor cells by Western blotting (Figure 2B). In addition, transcription level of NFKB2, a major component of noncanonical NF- $\kappa$ B pathway, was increased in AML1-deficient cells (Figure 2C), compatible with the transcription of NFKB2 being regulated by p65.<sup>18</sup> In the noncanonical NF- $\kappa$ B pathway, cytokines such as BAFF promote processing



**Figure 1. NF- $\kappa$ B signaling is activated in AML1-deficient cells.** (A) Schematic representation of the gene expression analysis. LSK cells from AML1 cKO mice (AML1 *f/f*) were transduced CreER-GFP or GFP expressing retroviruses. GFP+ cells were sorted and were subjected to gene expression analysis 24 or 48 hours after adding 4-OHT. (B) PCR genotyping of BM cells from AML1 cKO mice after adding 4-OHT. (C-I) NF- $\kappa$ B-related gene sets enriched in AML1-deficient cells (AML1 *flx/flx*, CreER+ 24 and 48 hours after adding 4-OHT) by GSEA Version 2.0. (C) A gene set that includes genes up-regulated by NF- $\kappa$ B (HINATA\_NFKB\_UP). (D) A gene set that includes genes which are up-regulated TNF- $\alpha$  treatment (TNFA\_NFKB\_DEP\_UP). (E) A gene set that includes Ras-inducible, NF- $\kappa$ B-regulated genes (HANSON\_NFKAPPB\_IND). (F) A gene set that includes genes with promoter regions containing REL motif (V\$CREL\_01); (G) a gene set which includes Genes with promoter regions containing p65 motif (V\$NFKAPPAB65\_01). (H) A gene set that includes genes with promoter regions containing NF- $\kappa$ B motif (SNFKB\_Q6). (I) Target genes of NF- $\kappa$ B.<sup>21</sup> Nominal *P* value was estimated by GSEA software. (J) mRNA expression for NF- $\kappa$ B target genes in LSK cells transduced with CreER or mock from AML1 *f/f* mice 24 or 48 hours after 4-OHT addition. Error bars show mean  $\pm$  SEM.

of p52 from p100 (NFKB2), protein level of which reflects activating status of the pathway. As shown in Figure 2D, both of basal and BAFF-mediated processing of p52 are augmented in AML1-deficient splenic B cells (AML1 *flx/flx* Mx+) compared with control cells (AML1 *flx/flx* Mx-) that express intact AML1. These data indicate that both canonical and noncanonical pathways of NF- $\kappa$ B signaling are aberrantly activated in AML1-deficient murine hematopoietic cells.

#### AML1 attenuates NF- $\kappa$ B signaling through interaction with IKK complex

Because the NF- $\kappa$ B signaling is activated by the loss of AML1, we investigated whether AML1 represses NF- $\kappa$ B signaling. We performed nuclear translocation assays using HEK293T cells. When HEK293T cells are starved with culture medium containing 0.5% FCS, p65 stays in the cytoplasmic fraction (Figure 3A top left panel). Twenty minutes after adding TNF- $\alpha$ , p65 translocates to the nuclear fraction (Figure 3A top right panel). In AML1-transduced cells, p65 does not translocate to nuclear fraction after adding TNF- $\alpha$  (Figure 3A bottom right panel). These data show that AML1 inhibits nuclear translocation of p65 when expressed in HEK293T cells (Figure 3A). TNF- $\alpha$ -induced phosphorylation of p65 at serine 536, which is important for its activation, also was inhibited by AML1 in those cells (Figure 3B).

To seek the underlying mechanism for inhibition of NF- $\kappa$ B signaling by AML1, we assessed physical and functional interaction between AML1 and signaling components of NF- $\kappa$ B signaling. IKK is a multimeric complex consisting of IKK $\alpha$ ,

IKK $\beta$ , and IKK $\gamma$ . The kinase activity of IKK is essential for I $\kappa$ B degradation and nuclear translocation of p65 and is exerted by IKK $\alpha$  and IKK $\beta$ . As shown in Figure 4A-B, AML1 was coprecipitated with IKK $\alpha$  and IKK $\beta$  when they are expressed in HEK293T cells. These results indicate that AML1 physically interacts with IKK $\alpha$  and IKK $\beta$ . Using Jurkat cells, we confirmed that endogenous AML1 forms a complex with IKK $\alpha$  and IKK $\beta$  (Figure 4C). Because AML1 is a DNA-binding protein and IKK complex is thought to act mainly in the cytoplasm, we examined where they interact with each other. As shown in Figure 4D, interaction between AML1 and IKK complex was mainly detected in the cytoplasmic fraction in HEK293T cells. Interestingly, protein level of AML1 was increased in the cytoplasmic fraction when IKK $\alpha$  was overexpressed with AML1. We confirmed that endogenous AML1 physically interacts with IKK complex mainly in the cytoplasmic fraction in Jurkat cells (Figure 4E). We also confirmed that AML1 physically interacts with IKK complex in U937 cells, a human myeloid cell line (Figure 4F). Next, we tested whether AML1 affects the kinase activity of IKK complex. Using the *in vitro* kinase assay, we found that AML1 significantly suppresses the kinase activity of IKK $\alpha$  and IKK $\beta$  (Figure 4G-H). In accordance with these findings, AML1 significantly attenuated TNF- $\alpha$ -induced degradation of I $\kappa$ B $\alpha$  in HEK293T cells (Figure 4I).

Taken together, AML1 acts as a cytoplasmic attenuator of IKK complex, thus accounting for a mechanistic basis for the inhibition of NF- $\kappa$ B signaling by AML1. Suppression of IKK activity by