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*J Immunol* published online 23 January 2013
http://www.jimmunol.org/content/early/2013/01/23/jimmunol.1202736

**Supplementary Material**
http://www.jimmunol.org/content/suppl/2013/01/24/jimmunol.1202736.DC1

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Sox4 Is Required for the Survival of Pro-B Cells

Baohua Sun,* Saradhi Mallampati,* Yun Gong, † Donghai Wang,‡ Véronique Lefebvre,§,* and Xiaoping Sun*†

The development of mature B cells from hematopoietic stem cells is a strictly orchestrated process involving multiple regulatory genes. The transcription factor Sox4 is required for this process, but its role has not been systematically studied, and the underlying mechanisms remain unknown. To determine when and how Sox4 functions in the stepwise process of B cell development, we used mice harboring conditional null alleles for Sox4 and a Cre transgene. Sox4 deletion in hematopoietic stem cells almost entirely eliminated pro-B cells in both fetal livers and adult bone marrow, resulting in a severe deficiency in later stage B cells, including circulating mature B cells. Sox4-deficient pro-B cells, particularly those expressing the stem cell factor receptor c-Kit, readily underwent apoptosis, and even more so when c-Kit activity was inhibited by imatinib. C-Kit–expressing pro-B cells showed decreased activation of the c-Kit downstream protein Src upon Sox4 deletion. Likewise, the level of the anti-apoptotic Bcl2 protein was decreased in residual pro-B cells, and its restoration using a Bcl2 transgene allowed not only partial rescue of pro-B cell survival but also B cell maturation in the absence of Sox4. Our findings indicate that Sox4 is required for the survival of pro-B cells and may functionally interact with c-Kit and Bcl2. The Journal of Immunology, 2013, 190: 000–000.

B cells play pivotal roles in humoral immunity and are a key component of the immune system. Like other blood cell types, mature B cells arise from self-renewing pluripotent hematopoietic stem cells (HSCs) through a stepwise process involving coordinated cell proliferation along with progressive lineage commitment and differentiation. In the B cell lineage, HSCs first develop into lymphoid-primed multipotent progenitors, which have lost their self-renewal ability but remain multipotent, and then into common lymphocyte progenitors (CLPs), which in turn develop into B cells, T cells, NK cells, and dendritic cells (1). The first B cell–specific progenitors arising from CLPs are pre-pro-B cells, which sequentially develop into pro-B, pre-B, immature, and ultimately mature B cells.

B cell development requires appropriate orchestration of a network of regulatory genes involved in cell survival, proliferation, and differentiation (2, 3). Particularly in early B cell development, several key transcription factors act in a hierarchical order to precisely control the expression of critical genes (4, 5). Pu.1 is involved in the hematopoietic lineage fate decision at the branch-point of myeloerythroid and myelolymphoid progenitor populations (6). Ikaros is a key factor in B lineage specification and commitment. Ikaros-deficient or hypomorphic mutant mice have severe defects in the development of the lymphoid system (7, 8). Ebf1 and Pu.1 activate IL-7R, the first gene that distinguishes CLPs from lymphoid-primed multipotent progenitors (9–11). E2A functions upstream of EBF, and the two transcription factors collaborate to activate Pax5 expression (12, 13). Pax5 is indispensable in B cell lineage commitment in that it activates B lineage–specific genes and represses lineage-inappropriate genes (14, 15).

Programmed cell death frequently occurs during B cell development, but despite intensive studies, the molecular mechanisms that control this process are still elusive. Accumulating evidence has demonstrated that members of the Bcl2 protein family, including Bcl2 (16), Bcl-xL (17, 18), Mcl1 (19), Bak (20), Bik (21), and Bim (22), are critical for B cell survival in vivo and in vitro. Other factors, such as Stat5 and Dicer1, are also involved in maintaining B cell survival. Stat5 has been shown to mediate the survival function of IL-7 signaling during B cell development (23, 24). Conditional deletion of Dicer1 in the earliest stage of B cell development results in an almost complete absence of pre-B cells, owing to a massive increase in apoptosis in pre-B cells (25, 26). In adult mice, signaling by stem cell factor and its receptor c-Kit is critical in pro-B and pro-T cell development (27). The downstream signaling pathways activated by c-Kit include activation of PI3K and Src kinase. However, c-Kit–mediated Src kinase activation, but not PI3K activation, is important for adult B cell development, likely through signaling events related to cell proliferation and cell survival (28).

Sox4 is a member of the SOX (SRY-related HMG box) transcription factor family and has been shown to play important roles in embryonic development and differentiation (29). Germline deletion of Sox4 is embryonically lethal in mice (30). Embryos with this deletion died at day 14 of development owing to circu-
Sox4 in Regulation of B Cell Development

In our study, we used dancy with its close relatives Sox11 and Sox12 (31–35). In this Sox4 has since been shown to be critical for cell survival and what role Sox4 plays in adult B cell development remain unknown. Sox4 deficiency causes the fetal B cell developmental arrest and the pro-B cell stage. These findings indicated that Sox4 is indispensable for B cell development in the fetal liver. However, how Sox4 deficiency causes the fetal B cell developmental arrest and what role Sox4 plays in adult B cell development remain unknown. Sox4-deficient embryos failed to generate B cells in the presence of IL-7. Reconstitution of lethally irradiated adult mice with the Sox4+/− fetal liver cells showed a stringent arrest of donor B cell development at the pro-B cell stage. These findings indicated that Sox4 is indispensable for pro-B cell survival and might functionally interact with c-Kit and Bcl2.

Materials and Methods

Mice

Mice with loxP sites flanking the entire coding region of the Sox4 gene were described previously (36). Vav-Cre mice were provided by Dr. Dimitris Kiossis at the National Institute for Medical Research, The Ridgeway, London (37). H2K-Bc12 transgenic (Tg) mice were provided by Dr. Irving Weissman at Stanford University, Stanford, CA (38). Genotyping was performed by PCR, using genomic DNA extracted from mouse tails. All mice were bred and maintained in a specific pathogen-free animal facility at The University of Texas MD Anderson Cancer Center, Houston, TX. All mouse experiments were performed in accordance with federal laws as well as guidelines of the National Institutes of Health, and protocols were approved by the MD Anderson Animal Care and Use Committee.

Imatinib treatment

To inhibit the c-Kit signaling pathway, 4- to 5-wk-old mice were given i.p. injections of 100 mg/kg imatinib (LC Laboratories, Woburn, MA) twice daily in a volume of 100 μl PBS for 2, 3, or 7 consecutive days, as indicated. Mice were euthanized the day after the last injection, and samples were harvested for flow cytometric analysis.

Cell surface staining, flow cytometry, and cell sorting

Single-cell suspensions were prepared at the time of autopsy from fetal liver, bone marrow (femurs and tibias), and spleen. All samples were incubated with ice-cold RBC lysis buffer with ammonium chloride for 4 min in the dark before staining with a combination of mouse-specific Abs at 4°C for 15 min in staining solution (PBS supplemented with 1% BSA). Blood was collected from the hearts of the mice in 15-ml Falcon tubes (Becton-Dickinson Biosciences, San Jose, CA) containing 50 μl 0.5 M EDTA in PBS. After two rounds of RBC lysis, the WBCs were subjected to staining.

Ab use for flow cytometric analysis included the following: FITC–anti-CD45 (Jackson ImmunoResearch, West Grove, PA); FITC–anti-CD21 (SD9), PE–anti-CD11c (N418), PE–anti-CD25 (PC61,5), PE–anti-NK1.1 (PK136), allophycocyanin–anti-AA4.1 (AA4.1), PerCP-Cy5.5–anti-B220 (RA3–6B2), biotin–anti-β-globin (FG53.4), and eFluor 450–streptavidin, all from eBioscience (San Diego, CA); FITC–anti-CD24 (M1–69), FITC–anti-IgD (11–26c.2a), FITC–anti–CD43 (St7), PE–anti-Ter119 (Ter-119), PE–anti–c-kit (2B8), allophycocyanin–anti–CD19 (1D3), biotin–anti–Ccr4 (2B11), and biotin–anti–IL-7R (B12-1), all from Becton-Dickinson Biosciences. Biotinylated Abs were detected with FITC–allophycocyanin, PerCP-Cy5.5–, eFluor 450–, or PE–conjugated streptavidin (Becton-Dickinson Biosciences). Samples were acquired on a FACS Calibur or Influx high-speed sorter (BD), and data were analyzed with FlowJo software (Tree Star, Ashland, OR). The absolute number of each B cell population was derived from the total number of bone marrow cells per mouse and the percentage of the population in question.

Annexin V and intracellular staining

For cell apoptosis analysis, the staining buffers containing surface Abs were carefully removed before biotin-conjugated Annexin V was applied according to the manufacturer’s instructions (Becton-Dickinson Biosciences). For intracellular detection of Bcl2 (2870), total Src (2109), or p-Src (2101; Cell Signaling Technology, Danvers, MA), cells were incubated with surface Abs as indicated. After being washed with staining solution, cells were incubated in fixation solution (00-8222-49; eBioscience) for 20 min and subsequently washed twice with permeabilization buffer (00-8333-56; eBioscience). Cells were then incubated with specific primary Abs for 30 min at room temperature and then incubated for 20 min on ice with PE–, allophycocyanin–, or FITC–conjugated anti-rabbit secondary Abs.

Flowsort Flow Cytometry

Flow cytometric analysis revealed that the number of pro-B cells, including fraction B (B220+CD43+CD24+B1) and fraction C–C' (B220–CD43+CD24+B1), was much lower in the bone marrow of adult Sox4fl/fl-Vav-Cre mice (Hardy’s nomenclature for subpopulations of developing B cells (41)). Whereas control littermates had 0.55 million pro-B cells, the Sox4fl/fl-Vav-Cre mice had only 0.04 million (p = 0.0019; Fig. 1A). The number of fraction A cells (i.e., prepro-B, B220+CD43+CD24+B1) was slightly decreased (0.14 million in control mice versus 0.07 million in Sox4fl/fl-Vav-Cre mice; p = 0.064). Fraction A cells are heterogeneous and can give rise to, in addition to B cells, NK and plasmacytoid dendritic cells (pDCs) (42–44). The AA4.1+ subset of fraction A cells contains B lineage precursors (45), the NK1.1+ subset produces NK cells, and the CD11c+ subset generates pDCs (46, 47). The numbers of AA4.1+, NK1.1+, and CD11c+ fraction A cells in Sox4fl/fl-Vav-Cre bone marrow were comparable to those in the corresponding subsets of control mice (Fig. 1B). These data indicate that deletion of Sox4 significantly affected pro-B cells but only minimally affected prepro-B cells. Moreover, a moderate decrease in the number of Lin–c-Kit+Sc-1+ primitive HSCs (36 × 103 in control mice versus 12 × 103 in Sox4fl/fl-Vav-Cre mice; p = 0.03; Fig. 1C) and a minimal increase in the number of CLPs (Lin–IL-7R–c-Kit–Sc-1+; 65 × 103 in control mice versus 78 × 103 in Sox4fl/fl-Vav-Cre mice; p = 0.18; Fig. 1D) suggested that the marked reduction in pro-B cells was unlikely to be caused by changes in earlier stage cells (HSCs, CLPs, and prepro-B cells). As a consequence of the pro-B cell depletion, bone marrow fraction D (pre-B, IgM B220+CD25+), fraction E (immature B, IgM+ B220–), and fraction F (mature B, IgM+ B220+) cells were almost completely eliminated (Fig. 1E). In the spleen, the percentages of immature (B220+AA4.1+, including transitional T1, T2, and T3) and mature (B220+AA4.1+, including marginal zone and follicular) B cells were severely reduced (7.12% and 61.3% in control mice versus 0.14% and 17.9% in Sox4fl/fl-Vav-Cre mice; p = 0.03; Fig. 1F).
FIGURE 1.  B cell development in the bone marrow of Sox4fl/flVav-Cre mice. Flow cytometric analyses were performed to assess B cells at different developmental stages in Sox4fl/flVav-Cre (fl/+ Cre) and Sox4fl/flVav-Cre (fl/fl Cre) mice. The numbers in the plots are the percentages of the indicated cell groups. Bar graphs show the total number of cells per mouse, presented as mean ± SD with p values indicated. (A) Bone marrow B220+CD43+ lymphocytes were analyzed for fraction (Fr.) A (prepro-B; B220+CD43+Cd24–), fraction B (early pro-B; B220+CD43+Cd24+BP1–), and fraction C–C9 (late pro-B; B220+CD43+CD24+BP1+). Note that the percentage of fraction A cells was 24.3% in Sox4fl/+Vav-Cre mice and 79.3% in Sox4fl/flVav-Cre mice (flow cytometric plot). This difference was mostly due to the decrease in fraction B–C9 cells; the total number of fraction A cells was 0.14 million in Sox4fl/+Vav-Cre mice and 0.07 million in Sox4fl/flVav-Cre mice (bar graph). (B) Bone marrow fraction A cells were analyzed for the expression of NK1.1, AA4.1, and CD11c. (C) Bone marrow Lin– cells were analyzed for the expression of c-Kit and Sca-1 to assess the number of hematopoietic stem cells. Lin, lineage markers, including CD11b, B220, CD4, CD8, Ter119, NK1.1, Gr-1, and CD19. (D) Bone marrow Lin– and IL-7R+ cells were analyzed for the expression of c-Kit and Sca-1 to assess the number of common lymphocyte progenitors. (E) Bone marrow lymphocytes were analyzed for fractions D (pre-B; IgM+B220+CD25+), E (immature B; IgM+B220low), and F (mature B; IgM+B220high) B cells (n = 3–5 mice per group).
versus 0.56% and 5.31% in Sox4fl/fl Vav-Cre mice, respectively; Fig. 2A, 2B). As shown in Fig. 2C and 2D, B220+CD19+ peripheral blood B cells were essentially absent. However, the peripheral blood residual B cells in Sox4fl/fl Vav-Cre mice expressed IgM and IgD on the surface, indicating that these cells were able to develop into mature B cells.

It was shown previously that Sox4null fetal liver hematopoietic cells fail to develop into mature B cells in recipient adult mice after transplantation (30). However, it was not known which early stage B cells were absent in the fetal livers of the Sox4-null embryos. We found that pro-B cells (B220+CD43+CD24+CD19+) were almost undetectable in the fetal livers of day 13.5 Sox4null embryos.

FIGURE 2. B cell development in the spleen, peripheral blood, and fetal liver of Sox4fl/fl Vav-Cre mice. (A) Splenocytes were analyzed for immature (I, B220AA4.1+) and mature (M, B220AA4.1) B cells. The immature B cells were further analyzed for transitional T1 (B220AA4.1IgM+CD23–), T2 (B220AA4.1IgM–CD23+), and T3 (B220AA4.1IgM–D23+) cells, and the mature B cells were further analyzed for marginal zone (MZ, B220AA4.1IgM+CD23–) and follicular (FO, B220AA4.1IgM–CD23+) B cells. (B) The number of individual cells per million spleen lymphocytes. (C) Peripheral blood B cells (B220CD19+) were analyzed for the expression of IgM and IgD. (D) Percentage of B cells in peripheral blood. (E) Fetal liver cells from day E13.5 embryos were analyzed for B220CD43CD24CD19 pro-B cells (data are representative of two independent experiments). Gating strategies and abbreviations are the same as in Fig. 1, and n = 3–5 mice per group unless otherwise specified.

FIGURE 3. Sox4 deletion in B cells at various stages in Sox4fl/fl Vav-Cre mice. (A and B) Total RNA prepared from bone marrow FACS-sorted prepro-B cells (A) and pro-B cells (B) was subjected to RT-PCR to assess Sox4 expression. The cDNA of pro-B cells was 5X serially diluted for semiquantitative RT-PCR (B). (C) Quantitative RT-PCR analysis of Sox4 mRNA expression in pro-B cells of Sox4fl/+Vav-Cre and Sox4fl/fl Vav-Cre mice, with or without Bcl2 Tg overexpression. (D and E) Fractions A–F (D) and peripheral blood B cells [B220CD19+ (E)] from Sox4fl/Vav-CreRosa26EYFP (blue line) and Sox4fl/fl Vav-CreRosa26EYFP (red line) mice were subjected to flow cytometric analysis to assess EYFP expression, which serves as an indicator for Sox4 deletion.
FIGURE 4. Effect of Sox4 deletion and c-Kit inhibition on the apoptosis of B cells. (A) Apoptotic frequency measured as the percentage of Annexin V–positive cells in fractions A–F in Sox4fl/+Vav-Cre (blue line) and Sox4fl/flVav-Cre (red line) mice. The histogram shows one of the representative experiments, whereas the bar graph shows results (mean ± SD with p values indicated) from five experiments. (B) The expression of critical B cell developmental genes in pro-B cells was determined by semiquantitative RT-PCR. Also included were two antiapoptotic genes, Bcl2 and Mcl1. The actin cDNA was used as control. Three cDNA concentrations with 5× serial dilution were used for each gene. The size of the PCR products is in- (Figure legend continues)
Vav-Cre embryos (Fig. 2E). This result suggested that deletion of the Sox4 gene in embryonic HSCs resulted in the abrogation of fetal liver pro-B cells and that this abrogation was responsible for the B cell deficiency in fetal liver–reconstituted adult bone marrow and in vitro culture.

To exclude the possibility of incomplete or escape of Sox4 deletion in the residual B cells in Sox4<sup>−/−</sup>Vav-Cre mice, we sorted out the minute populations of prepro-B cells and pro-B cells and analyzed the Sox4 mRNA levels. Sox4 mRNA was not detectable in bone marrow fraction A cells (Fig. 3A) or residual pro-B cells (Fig. 3B) in Sox4<sup>−/−</sup>Vav-Cre mice by RT-PCR. Real-time RT-PCR showed that Sox4 mRNA in pro-B cells was decreased 133-fold in Sox4<sup>−/−</sup>Vav-Cre mice (Fig. 3C, left). We also crossed the Sox4<sup>−/−</sup>Vav-Cre mice with Rosa26–enhanced yellow fluorescent protein (EYFP) Cre reporter mice to assess Cre activity in residual B cells. It was found that >90% of bone marrow prepro-B (fraction A) cells and >95% of peripheral blood residual B220<sup>−</sup>CD19<sup>+</sup> B cells were EYFP positive (Fig. 3D, 3E), implying that the floxed Sox4 allele had been deleted in these cells. The relative low percentages of EYFP<sup>+</sup> cells in pro-B (fractions B–C) and pre-B (fraction D) cells of Sox4<sup>−/−</sup>Vav-Cre mice compared with control mice were expected because of the high frequency of apoptosis in these cells (see below); EYFP<sup>+</sup> (or GFP<sup>+</sup>) cells become EYFP<sup>−</sup> (or GFP<sup>−</sup>) cells when undergoing apoptosis (48). The presence of EYFP<sup>+</sup> B cells in peripheral blood suggested that a minor portion of pro-B cells overcame the effect of Sox4 deficiency and continued to develop into mature B cells.

**Sox4 is vital for the survival of pro-B and pre-B cells**

To measure apoptosis in minute populations of B cells, particularly in mice with B cell deficiency caused by Sox4 deletion, we used highly sensitive multicolor flow cytometry with Annexin V staining (49, 50). Annexin V staining may vary in B cells with a different status of activation (51), but is a reliable marker for apoptosis when comparing cells at similar stages in development (52, 53). As shown in Fig. 4A, a minimal increase in the frequency of apoptosis in prepro-B (fraction A) cells was detected in Sox4<sup>−/−</sup> Vav-Cre mice (3.55% versus 6.44%; p = 0.39), whereas the frequency of apoptosis in pro-B (fraction B–C) and pre-B (fraction D) cells increased dramatically (respectively, 5.61% versus 33.6%; p < 0.001) and 1.76% versus 10.1%; p = 0.014). In contrast, bone marrow immature and mature (fractions E–F) B cells of Sox4<sup>−/−</sup>Vav-Cre mice had a frequency of apoptosis comparable to that in controls. These results suggested that Sox4 was crucial for maintaining the survival of pro-B and pre-B cells but not required for the survival of later stage B cells.

**Sox4 cooperates with the c-Kit signaling pathway to regulate pro-B cell survival**

To find out whether Sox4 exerts its function by regulating the genes known to be critical in early B cell development, we sorted out the small number of residual pro-B cells and characterized the expression of these genes by semiquantitative RT-PCR. Of the genes studied, including IL-7r, Cxcr4, E2a, Ebf1, Pax5, Foxp1, Stat5a, Ezh2, and Rag1, none showed a significant difference in expression between Sox4<sup>−/−</sup>Vav-Cre mice and their control littermates (Fig. 4B). Because Sox4 deficiency caused pro-B cell apoptosis, we also determined the mRNA levels of two antiapoptotic genes—Bcl2 and McI1, which were also known to be involved in early B cell development—and found no significant difference in them, either (Fig. 4B). However, the possibility still existed that Sox4 regulates some of these gene products at the posttranscriptional level.

We analyzed apoptosis in residual c-Kit<sup>+</sup> pro-B cells from the bone marrow of the Sox4<sup>−/−</sup>Vav-Cre mice. Lack of Sox4 expression significantly increased cell death in c-Kit<sup>+</sup> pro-B cells (IgM<sup>−</sup>B220<sup>−</sup>c-Kit<sup>−</sup>CD19<sup>+</sup>; 6.95% versus 36.2%; p < 0.001) but not in c-Kit<sup>−</sup> non–pro-B cells (IgM<sup>−</sup>B220<sup>−</sup>c-Kit<sup>−</sup>CD19<sup>+</sup>; 11.5% versus 11.1%; p = 0.9; Fig. 4C). Although deletion of one copy of the Sox4 gene (Sox4<sup>+/−</sup>Vav-Cre) resulted in only minimal apoptosis of c-Kit<sup>+</sup> pro-B cells (Fig. 4D, left), it caused nearly three times the frequency of apoptosis in pre-B cells compared with control (3.36% versus 9.65%; Fig. 4D, right), which did not express c-Kit. We hypothesized that c-Kit compensates for the deleterious effect of Sox4 deletion in pro-B cells. We administered 100 μg of the c-Kit inhibitor imatinib by i.p. injection twice daily for 2, 3, or 7 d. Whereas the frequency of apoptosis in c-Kit<sup>−</sup> pro-B cells was comparable between Sox4<sup>+/−</sup>Vav-Cre and Sox4<sup>+/−</sup>Vav-Cre mice after 0 and 2 d of injection, the difference in the frequency of apoptosis was 3-fold (29.1% versus 10.9%) and 1.5-fold (46.9% versus 30.2%) after 3 and 7 d of injection, respectively (Fig. 4E). Because Src kinase is downstream of c-Kit and plays a critical role in pro-B cell development (28, 54), we determined the levels of total Src and phosphorylated Src (p-Y416) in pro-B cells from Sox4<sup>−/−</sup>Vav-Cre and Sox4<sup>+/−</sup>Vav-Cre mice by flow cytometric analysis. We found no overt difference in total Src, but phosphorylated Src (p-Y416) was significantly reduced upon Sox4 deletion (median fluorescence intensity [MFI] 29.4 versus 13.2; Fig. 4F).

**B cell development in Sox4<sup>+/−</sup>Vav-Cre mice can be partially rescued by Bcl2**

Because the frequency of apoptosis in Sox4-deficient pro-B cells was significantly increased and lack of the antiapoptosis protein Bcl2 was reported to increase the susceptibility of pro-B cells to proapoptotic agents (55, 56), we analyzed the level of Bcl2 protein in Sox4-deficient pro-B cells. Although Bcl2 mRNA did not show significant change (Fig. 4B), Bcl2 protein was reduced in pro-B cells (MFI 261.3 versus 96.7), but not in prepro-B cells (MFI 509.1 versus 633.3; Fig. 5A, left) in the Sox4<sup>−/−</sup>Vav-Cre mice, according to flow cytometric analysis. A reduction by 66% in Bcl2 protein was detected by Western blot in B220<sup>+</sup> bone marrow B cells (Fig. 5A, right). These data indicated that the proapoptotic effect of Sox4 deficiency in pro-B cells was associated with reduction of Bcl2 at the protein level.

Forced expression of Bcl2 in Bcl2-Tg mice has been shown to restore hematopoietic cell development in a number of gene
FIGURE 5. Effect of Bcl2 on B cell apoptosis induced by Sox4 deletion. (A) Left panel, Flow cytometric analysis of Bcl2 expression in prepro-B (B220\(^+\)CD43\(^+\)CD24\(^-\)CD19\(^-\)) and pro-B cells (B220\(^+\)CD43\(^-\)CD24\(^+\)CD19\(^+\)) from Sox4\(^{fl/fl}\) Vav-Cre (blue line) and Sox4\(^{fl/+}\) Vav-Cre (red line) mice. The numbers represent MFI. Data represent three independent experiments. Right panel, Western blot results of the B220\(^+\) bone marrow cells from these mice. (B–D) Frequency of apoptosis measured as the percentage of Annexin V–positive cells in bone marrow c-Kit\(^+\) pro-B cells (B), pre-B cells (C), and IgM\(^+\) (immature and mature) B cells (D), with or without forced Bcl2 expression. Data represent three mice per group. Gating strategies and abbreviations are the same as in Figs. 1 and 2. (E) Semiquantitative RT-PCR of Sox4 expression in prepro-B (B220\(^+\)CD43\(^+\)CD24\(^-\)CD19\(^-\)) and pro-B (B220\(^+\)CD43\(^-\)CD24\(^+\)CD19\(^+\)) cells of Sox4\(^{fl/Cre}\) and Sox4\(^{fl/+}\) Vav-Cre mice with forced Bcl2 expression.
deficient mice (25). We crossed Sox4fl/flVav-Cre mice with H2K-Bcl2-Tg mice (38) to test whether forced Bcl2 expression could rescue the B cell deficiency caused by Sox4 deletion. Flow cytometric analysis showed that the number of B cells of various stages was increased in Sox4fl/flVav-Cre Bcl2-Tg mice compared with Sox4fl/flVav-Cre mice (Table I). Comparing the frequency of apoptosis in B cells of various stages from Sox4fl/flVav-Cre and Sox4fl/flVav-Cre mice, with or without Bcl2-Tg, showed that the increase in the number of B cells in Bcl2-Tg mice was associated with a reduction in the frequency of apoptosis in pro-B cells (Fig. 5B) and pre-B cells (Fig. 5C), whereas the frequency of apoptosis in immature and mature B cells remained essentially the same (Fig. 5D). The rescued B cells did not have detectable Sox4 mRNA expression by semiquantitative RT-PCR (Fig. 5E), and real-time RT-PCR showed a 2632-fold reduction in Sox4 mRNA levels in pro-B cells of Sox4fl/flVav-Cre mice (Fig. 3C, right), yet the rescued pro-B cells retained the normal expression pattern of critical B cell development genes listed in Fig. 4B. These results suggested that forced Bcl2 expression could partially restore B cell development by reducing apoptosis caused by Sox4 deficiency. Noticeably, bone marrow IgM+ late stage B cells (fractions E and F) and peripheral blood B220+CD19+ B cells were increased in number as well (Fig. 5D, Table I), suggesting that the rescued pro-B cells could pass through the checkpoint from pre-BCR”pre-B cells to BCR”immature and mature B cells without the function of Sox4.

Discussion
In this study, we investigated the role of Sox4 in B cell development by conditionally knocking out the Sox4 gene in HSCs. We found that deletion of Sox4 markedly reduced the number of pro-B cells associated with an increased frequency of apoptosis, indicating that Sox4 was essential for the survival of pro-B cells. Our data also suggested that Sox4 functionally interacted with c-Kit and Bcl2 in protecting pro-B cells from apoptosis.

In the stepwise analysis of B cell development, depletion of fraction B pro-B cells is the earliest major change caused by Sox4 deficiency. In Sox4fl/flVav-Cre mice, although the number of Lin–c-Kit+Sca-1+ HSC-enriched cells was moderately decreased, no significant change was noted in the number of CLPs and prepro-B cells. Thus, the nearly complete absence of pro-B cells was unlikely to have resulted from a decrease in the earlier stage cells. Nevertheless, to confirm that heterogeneity in the fraction A population did not obscure the B lineage subset, we measured the expression of various markers representing individual subsets within it, including AA4.1 for B cells (45), NK1.1 for NK cells, and CD11c for pDCs (46, 47). We found that the size of each individual subset was not significantly altered in the absence of Sox4. Therefore, marked reduction in the number of pro-B cells is the major cause for deficiency of downstream B cells. Our data also showed that deletion of Sox4 in embryonic HSCs caused fetal liver pro-B cell deficiency in Sox4fl/flVav-Cre mice and that deletion of Sox4 in adult bone marrow resulted in pro-B cell deficiency in Sox4fl/flVav-Cre mice. We conclude, therefore, that Sox4 is required for B cell development in both the fetal liver and adult bone marrow and that its deficiency causes pro-B cell abrogation.

Although controversy exists in the literature, Sox4 appears to be a cell survival factor. This function has been postulated for neural cells during spinal cord development (35) and for the development of late sympathetic ganglia (31, 33). In the current study, a significant increase in the frequency of apoptosis was observed in residual pro-B and pre-B cells in Sox4fl/flVav-Cre mice. However, the frequency of apoptosis in prepro-B cells in Sox4fl/flVav-Cre mice was not significantly different from that in control mice. Likewise, the status of apoptosis of immature and mature B cells was not affected by Sox4 deletion. Thus, during B cell development, Sox4 is required for the survival of pro-B and pre-B cells, but unlikely for B cells in other stages.

As a transcription factor, Sox4 is expected to be part of the hierarchy of multiple transcription factors that control early B cell development, including Ebf1, E2A, and Pax5. However, the mRNA levels of these transcription factors and several other proteins such as CXCR4 and IL-7R that are critical in early B cell development were not changed in Sox4-deficient pro-B cells, suggesting that the expression of these genes was not directly regulated by Sox4 at the transcriptional level. Nevertheless, it could not be ruled out that Sox4 might indirectly regulate these gene products at the protein level. Although the Sox4 transcriptional program remains to be elucidated, our data show that c-Kit signaling and Bcl2 function are important in the control of pro-B cell survival supported by Sox4.

The stem cell factor/c-Kit axis, through activation of Src, PI3K, and JAK/STAT, provides proliferation and survival signals to pro-B cells to ensure their proper development (57). C-Kit is a pro-B cell marker, and c-Kit–positive pro-B cells were diminished upon Sox4 deletion. We demonstrated that treatment with the tyrosine kinase inhibitor imatinib could sensitize the apoptotic effect of Sox4 deficiency in pro-B cells in vivo. Given the role of c-Kit in early B cell development, this sensitization most likely had resulted from the inhibition of c-Kit by imatinib and reflected functional cooperation of Sox4 and c-Kit in this process. This idea is supported by our findings that the level of Src phosphorylation (Y416) was lower in pro-B cells of Sox4fl/flVav-Cre mice than of Sox4fl/flVav-Cre mice. Src kinase is a downstream signaling component of the c-Kit pathway, and its signaling is critical in pro-B cell development (28). Src phosphorylation (Y416) has been reported to provide survival signals, and targeting Src activation could enhance apoptosis in multiple cell lines treated with proapoptotic agents (58, 59). In our study, the lower Src phosphorylation (Y416) in the Sox4fl/flVav-Cre mice indicates that c-Kit/Src signaling is weakened in the absence of Sox4.

Bcl2 was reported to be highly expressed in normal bone marrow pro-B cells (17). Deficiency in B cell development caused by disruption of a number of genes can be completely or partially rescued by a Bcl2 transgene (24). We also detected a high level of

Table I. Effect of forced Bcl2 expression on subpopulations of B cells

<table>
<thead>
<tr>
<th>Bone Marrow B Cells (millions)</th>
<th>CD19+ Pro-B</th>
<th>c-Kit+ Pro-B</th>
<th>Pre-B</th>
<th>Immature + Mature</th>
<th>Peripheral Blood B Cells (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>f/+  Vav-Cre</td>
<td>0.912 ± 0.087</td>
<td>0.354 ± 0.045</td>
<td>3.544 ± 0.285</td>
<td>2.576 ± 0.336</td>
<td>45.85 ± 9.67</td>
</tr>
<tr>
<td>f/+  Vav-Cre</td>
<td>0.006 ± 0.001</td>
<td>0.085 ± 0.013</td>
<td>0.173 ± 0.028</td>
<td>0.314 ± 0.041</td>
<td>1.81 ± 0.29</td>
</tr>
<tr>
<td>f/+  Vav-Cre Bcl2</td>
<td>0.766 ± 0.103</td>
<td>0.823 ± 0.147</td>
<td>2.256 ± 0.198</td>
<td>8.674 ± 1.201</td>
<td>59.36 ± 10.83</td>
</tr>
<tr>
<td>f/+  Vav-Cre Bcl2</td>
<td>0.054 ± 0.008</td>
<td>0.311 ± 0.096</td>
<td>1.203 ± 0.017</td>
<td>5.198 ± 1.172</td>
<td>17.56 ± 3.43</td>
</tr>
</tbody>
</table>

The numbers for bone marrow B cells are the total cells of each subpopulation from two femurs and two tibias. The numbers for peripheral blood B cells are the percentages of cells determined by flow cytometric analysis. f/+: Sox4fl/fl; f/+: Sox4fl/fl. Data represent the mean ± SD of three mice.
Bcl2 expression in normal prepro-B and pro-B cells by flow cytometric analysis, and the Bcl2 level was reduced in pro-B cells, but not in prepro-B cells, in Sox4fl/fl Vav-Cre mice. Because pro-B cells in the Sox4fl/fl Vav-Cre mice had an increased frequency of apoptosis, it is likely that Sox4 deletion caused pro-B cell apoptosis through reduction of the Bcl2 protein. We found that Bcl2 could partially restore B cell development in Sox4fl/fl Vav-Cre mice without affecting the mRNA expression of genes critical for this process. It is noteworthy that the numbers not only of pro-B cells but also of the later stage B cells, including pre-B and immature B cells, were higher in Sox4fl/fl Vav-Cre mice in the presence of the Bcl2 transgene. This observation indicated that the rescued pro-B cells had undergone full differentiation and given rise to functional later stage B cells, suggesting that Bcl2, beyond its inhibition of apoptosis, compensates Sox4 in an as-yet-unknown way in B cell development.

To conclude, we report that deletion of the Sox4 gene in HSCs causes severe deficiencies of pro-B and later stage B cells. Sox4 promotes survival during early B cell development in bone marrow and may functionally interact with c-Kit and Bcl2 in this process. Inhibition of c-Kit with imatinib sensitizes the apoptotic effect of Sox4 deficiency, and forced Bcl2 expression could partially rescue B cell development in the absence of Sox4.

Acknowledgments
We thank Drs. Dimitris Kioussi and Irving Weissman for providing the Vav-Cre mice and H2K-Bc12 Tg mice, respectively.

Disclosures
The authors have no financial interests of interest.

References


