O-Fucose Modulates Notch-Controlled Blood Lineage Commitment

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Notch receptors are cell surface molecules essential for cell fate determination. Notch signaling is subject to tight regulation at multiple levels, including the posttranslational modification of Notch receptors by O-linked fucosylation, a reaction that is catalyzed by protein O-fucosyltransferase-1 (Pofut1). Our previous studies identified a myeloproliferative phenotype in mice conditionally deficient in cellular fucosylation that is attributable to a loss of Notch-dependent suppression of myelopoiesis. Here, we report that hematopoietic stem cells deficient in cellular fucosylation display decreased frequency and defective repopulating ability as well as decreased lymphoid but increased myeloid developmental potential. This phenotype may be attributed to suppressed Notch ligand binding and reduced downstream signaling of Notch activity in hematopoietic stem cells. Consistent with this finding, we further demonstrate that mouse embryonic stem cells deficient in Notch1 (*Notch1*^{-/-}) or Pofut1 (*Pofut1*^{-/-}) fail to generate T lymphocytes but differentiate into myeloid cells while coculturing with Notch ligand-expressing bone marrow stromal cells *in vitro*. Moreover, *in vivo* hematopoietic reconstitution of CD34⁺ progenitor cells derived from either *Notch1*^{-/-} or *Pofut1*^{-/-} embryonic stem cells show enhanced granulopoiesis with depressed lymphoid lineage development. Together, these results indicate that Notch signaling maintains hematopoietic lineage homeostasis by promoting lymphoid development and suppressing overt myelopoiesis, in part through processes controlled by O-linked fucosylation of Notch receptors. (Am J Pathol 2010, 176:2921–2934; DOI: 10.2353/ajpath.2010.090702)

Notch signaling is essential for T cell commitment and development.^{1,2} Intracellular signaling mediated by Notch also plays a key role in maintaining various stem cell systems.³ Studies have shown that Notch activation by the exogenous Notch ligands promotes hematopoietic stem cell (HSC) expansion *in vitro* and favors lymphoid over myeloid lineage outcome *in vivo*.^{4–7} More recently, Calvi et al demonstrated that Notch activation by Notch ligand-expressing osteoblasts in bone marrow promotes self-renewal of primitive hematopoietic cells.⁸ These observations identify Notch as an essential component of bone marrow niche that supports HSC function. In comparison, *Notch1* and/or Jagged1 loss-of-function studies using targeted gene ablation, or expression of a dominant negative inhibitor of canonical Notch activation, did not reveal an essential role of Notch in HSC self-renewal.^{9–11} To reconcile these two findings, we hypothesize that the interaction between other Notch receptor and Notch ligand pairs upstream of Notch intracellular signaling, or aspects of Notch pathways in the absence of fucose that are independent of the transcriptional control of Notch signaling mediated by Mastermind-like 1, may be critical for HSC biology. Further, the role of Notch and Notch ligand interaction or Notch signaling transduction in lymphoid versus myeloid fate commitment of HSC has not been defined.

Notch signaling is subject to multiple regulating events that modify Notch–Notch ligand interactions or intracellular signaling. O-linked fucose attached to the epidermal growth factor (EGF)-like repeats of Notch receptors can modulate Notch signaling. Additionally, Fringe-mediated extension of O-fucose can differentially influence Notch signaling in *Drosophila* development, in murine embryo-

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genesis, and T lymphopoiesis.^{12–14} Recently we reported that mice carrying a targeted deficiency in the *FX* locus, of which gene encodes an enzyme that converts GDP-mannose to GDP-fucose in the *de novo* fucose synthesis pathway, develop chronic myeloproliferation. This faulty myeloid development is a result of loss of controlled suppression of myelopoiesis exerted by Notch, and that protein O-fucosylation modulates these Notch-dependent processes by controlling the interactions between Notch and its ligands.¹⁵

Mice with fucose deficiency also exhibit thymic hypoplasia and shortened survival of unknown causes. Expansion of the myeloid compartment is accompanied by a decreased common lymphoid progenitor (CLP) compartment.¹⁵ Given the evidence that Notch is important for HSC function, we speculate that fucosylation-modulated Notch signaling does not only regulate myelopoiesis of committed myeloid progenitors but also may affect HSC biology, particularly during early myeloid versus lymphoid diversification. We now report that HSC from fucose-deficient mice display suppressed Notch signaling and decreased self-renewal. Marrow cells isolated from fucose-deprived mice are less effective than wild-type cells, or cells from fucose-repleted *FX*^{−/−} mice, in competitive repopulation transplant assays. These cells, however, exhibit an enhanced myelopoietic phenotype after engraftment. The enhanced myelopoiesis from marrow progenitors can be partially suppressed by the expression of the activated Notch1 intracellular domain. Further, using both *in vitro* mouse embryonic stem cell (ESC) hematopoietic differentiation assay and *in vivo* hematopoietic reconstitution, we observe that Notch activation promotes T lymphopoiesis while suppressing myelopoiesis. These processes of ES hematopoiesis appear to be dependent on both Notch1 activity and O-fucose modification, as absence of *Notch1* or loss of fucosylation leads to suppressed T lymphopoiesis and enhanced myelopoiesis *in vitro*, as well as an aberrant blood lineage development *in vivo* characterized by an expansion of the myeloid compartment and suppressed lymphoid reconstitution.

Materials and Methods

Animals and ES Cell Lines

The animal research related to this article was approved by Case Western Reserve University Institutional Animal Care and Use Committee. Mice used include 8- to 18-week-old wild-type and *FX*^{−/−} mice maintained and prepared as described.¹⁶ The Notch reporter transgenic (NTg) mouse was purchased from the Jackson Laboratory (stock # 005854, Bar Harbor, Maine). *FX*^{−/−}-NTg mouse was generated by crossing the NTg with the *FX*^{−/−} mouse. *Notch1*^{−/−} ES cell lines were kindly provided by Dr. Rafel Kopan (Washington University School of Medicine, St. Louis, MO). *Pofut1*^{−/−} ES cells were kindly provided by Dr. Pamela Stanley (Albert Einstein College of Medicine, New York, NY). The *Pofut2*^{−/−} mouse ES line was generated from blastocyst out-

growths of *Pofut2* heterozygotes crossed at the Case Transgenic & Targeting Facility (BayGenomics, ES cell line RST434). The genotype was confirmed by PCR.

Flow Cytometry Analysis and Cell Sorting

Flow cytometric analyses were performed as described.¹⁵ When isolating multipotent progenitor cells (MPP) (Lin[−]Scal⁺c-kit⁺Flt3⁺), lineage-depleted cells were further stained with streptavidin-allophycocyanin (APC)-Cy7, fluorescein isothiocyanate (FITC)-anti-Sca-1, phycoerythrin (PE)-anti-Flt3, and APC-anti-c-kit, and sorted using FACSaria (BD Biosciences, San Jose, CA).

CFU Assay and MPP in Vitro Culture

For single CFU assays, single Lin[−]Scal⁺c-kit⁺ (LSK) cells were sorted into 96-well plates containing methylcellulose (MethoCult M3434, StemCell Technologies, Vancouver, BC, Canada). For pre-B CFU assay, bone marrow (BM) cells were plated onto plates containing MethoCult M3630 (MethoCult M3630, StemCell Technologies, Vancouver, BC, Canada). In OP9 co-culture assays, MPPs from wild-type and *FX*^{−/−} mice were cultured for 28 days on OP9-control or OP9-Notch ligand expressing cells in the presence of rmlL-7 (5 ng/ml) and rmFlt3L (5 ng/ml; R&D Systems, Minneapolis, MN) and analyzed at the end of culture as described.¹⁵

Bone Marrow Transplantation and Competitive Repopulation Assays

Bone marrow transplant was performed as described.¹⁵ For competitive repopulation assays, BM nucleated cells from wild-type or *FX*^{−/−} mice (Ly5.2) were mixed with wild-type competitor cells (Ly5.1) in a ratio of 1:3, 1:1, and 3:1 (1.8 × 10⁶ total cells per mouse) and transplanted into lethal-irradiated wild-type recipients (Ly5.1). Peripheral blood was collected from the recipients at 12 to 16 weeks to investigate the percentage of test cells based on Ly5.2 expression and different lineage contributions.

ES Cell in Vitro Hematopoietic Differentiation

ES cells (8 × 10³) were cultured on Iscove modified Dulbecco medium containing 1% methylcellulose (M03120, StemCell Technologies) and 40 ng/ml stem cell factor (R&D Systems, Minneapolis, MN) as described.¹⁷ After 7 to 10 days, the embryoid bodies were removed, and CD34⁺ cells were isolated by biotinylated rat anti-mouse CD34 and anti-biotin-beads (Miltenyi, Auburn, CA). These cells were then seeded onto OP9-control or OP9-Notch ligand-expressing cells in the presence of rmFlt3 ligand and rmlL-7.¹⁸ Cells were recovered from the day 15 culture and analyzed by FITC-anti-CD11b or CD25; PE-anti-Ter119 or CD44; APC-anti-B220 or CD4; APC-Cy7-anti-CD8; and PE-Cy7-anti-CD45 (BD Biosciences,

San Jose, CA). Cell surface expression of Notch receptors in CD34⁺ HPCs was characterized by staining cells with PE-conjugated anti-Notch1, Notch2, Notch3, and Notch4 antibodies (Biolegend, San Diego, CA).

In Vivo Assessment of ESC-Derived Hematopoietic Development by Intrafemoral Transplantation and Intrathymic Transplantation

ESCs were induced to differentiate toward hematopoiesis as described above. A single cell suspension of CD34⁺ cells (10⁵ cells per 20 μ l) was used for intrafemoral injection. Recipients (Ly5.1) were sublethally (5.5 Gy) or lethally irradiated (9.5 Gy) 24 hours before injection as described.¹⁹ The presence of donor-derived (ESC-Ly5.2) T, B, and granulocytes was determined by flow analysis of peripheral blood collected 2, 4, 8, 12 weeks, and monthly after injection. Of a total of 74 mice injected, only one mouse developed teratoma. No mice showed signs of graft-versus-host disease. For intrathymic transplantation, CD34⁺ cells (10⁵ cells/20 μ l) were injected into thymic lobes of anesthetized *FX*^{-/-} mice (maintained on fucose-supplemented chow until 8 week of age and then maintained on standard chow for 10 to 14 days). T lymphopoiesis was analyzed by flow cytometry on total thymocytes collected 2 weeks after transplantation.

Recombinant Mouse Notch Ligand Binding Assays

Recombinant Notch ligand comprising the extracellular domains of Dll1 or Dll4 fused with human IgG Fc were constructed as described.²⁰ Recombinant Notch ligand was prepared from HEK 293 T cells transfected with hlgG-Dll1, Dll4, or the vector and quantified by ELISA.¹⁵ Binding assay was performed by incubating LSK cells or ES cells with recombinant Notch ligand in Hanks balanced salt solutions supplemented with Ca²⁺ and analyzed by FACS using PE-anti-human IgG Fc.

Generation and Bone Marrow Infection of Activated Notch1-Expressing Retrovirus

To generate retrovirus containing constitutive active Notch1 molecules (ICN1), Phoenix E cells were transiently transfected with either pMig (kindly provided by Dr. Luk Van Parijs at Massachusetts Institute of Technology Center for Cancer Research, Cambridge, MA), or pMig-ICN1 using the Profection mammalian transfection system according to the manufacturer's specifications (Promega, Madison, WI). Notch1 intracellular domain (ICN1; 1735-2531) was inserted into pMig, a vector that allows coexpression of cloned cDNA and green fluorescent protein (GFP) from a single bicistronic message, to form pMig-ICN1-GFP. Viral supernatants were collected on three consecutive days 24 hours after the transfection. Infection of each viral vector on bone marrow progenitors was performed as described.²¹ Briefly, *FX*^{-/-} mice mar-

row cells were collected 4 days after 5-FU treatment and cultured in a 6-well plate in the growth medium (DMEM supplemented with 10% of FBS, 6 ng/ml of IL3, 10 ng/ml of IL6, and 10 ng/ml of stem cell factor) for 24 hours. Infections were performed on 1 $\times$ 10⁶ bone marrow progenitors per well in 1.5 ml of growth medium containing 4 μ g/ml of polybrene using 1.5 ml of each viral supernatant. A second infection was performed 24 hours after the initial infection. Forty-eight hours after the secondary infection, cells were harvested for bone marrow transplant. Approximately 2 $\times$ 10⁶ cells were injected into the tail veins of lethally irradiated syngeneic wild-type recipient mice. Recipients were sacrificed at the indicated time, and single-cell suspensions were prepared from peripheral blood, BM, and spleens.

Statistical Analysis

Data are presented as means $\pm$ SD, unless otherwise stated. Statistical significance was assessed by Student *t* test.

Results

*Decreased HSC Frequency and Lymphoid Progenitor Populations in *FX*^{-/-} Mice, Accompanied by Suppressed Notch Activation*

Notch signaling strength can be modified by O-fucose that is present on Notch EGF repeats.²²⁻²⁵ We recently identified that O-fucosylation of Notch receptors is required for the Notch receptor-ligand interaction on myeloid progenitor cells and thus suggests a requirement for O-fucosylation in normal control of myelopoiesis.¹⁵ To examine whether O-fucosylation of Notch modulates HSC biology, we first assessed the frequency of LSK (Lin⁻Sca-1⁺c-kit⁺) cells in *FX*^{-/-} mice that are deficient in GDP-fucose synthesis. Flow cytometric analysis indicates that there is a twofold decrease in the LSK percentage in the marrows of *FX*^{-/-} mice at steady state. A large proportion of this decrement was accounted for by a decrease in the percentage of MPPs (Flt3⁺LSK; Figure 1A). This change in the HSC population was confirmed by CFU analysis, which revealed that *FX*^{-/-} mice have a decreased number of granulocyte/erythrocyte/monocyte/megakaryocyte-CFU (mix-CFU; Figure 1B). To further examine the stem cell frequency in *FX*^{-/-} marrow, we performed limiting-dilution assays using a common wild-type marrow competitor (Ly5.1; Table 1). *FX*^{-/-} mice had a 13.7-fold reduction of stem cell frequency relative to wild-type mice after myeloablative transplantation. Individual LSKs from *FX*^{-/-} mice also displayed a decreased ability to form multilineage colonies but an increased frequency of single-lineage myeloid colonies when compared with wild-type LSKs (Figure 1C). Furthermore, marrow cells from *FX*^{-/-} mice formed a reduced number of pre-B cell colonies (Figure 1, C and D). This finding is consistent with our prior reports that *FX*^{-/-} mice have reduced CLP numbers.¹⁵ In comparison, *FX*^{-/-} mice maintained on

Figure 1. *FX*^{-/-} mice have decreased HSC and lymphoid progenitors and LSK have suppressed Notch activation. **A:** FACS analysis of the HSC compartment (LT & ST-HSC: Lin⁻c-kit⁺Sca-1⁺Flt3⁻; MPP: Lin⁻c-kit⁺Sca-1⁺Flt3⁺) in the bone marrow in 3- to 4-month-old control mice (wild-type: WT; *n* = 8). *FX*^{-/-} mice were reared on fucose-supplemented chow until 12 weeks of age and then on standard chow for at least four weeks (*n* = 8; *FX*^{-/-}, no fucose), and *FX*^{-/-} mice were reared on fucose-supplement chow until use (*n* = 8; *FX*^{-/-}, with fucose). Bar graphs indicate the average total number of LT & ST-HSC and MPP cells from two tibias and two femurs of each mouse. **B:** CFU-Mix numbers as colonies per 2 × 10⁵ BMNC (wild-type, *n* = 6; *FX*^{-/-}, no fucose, *n* = 6; *FX*^{-/-}, with fucose, *n* = 6). **C:** Multilineage or single myeloid lineage colonies formed by single LSK cells in 96-well plates (wild-type, *n* = 4; *FX*^{-/-}, no fucose, *n* = 4; *FX*^{-/-}, with fucose, *n* = 4). **D:** Number of CFU-preB colonies per 2 × 10⁵ BMNC (wild-type, *n* = 4; *FX*^{-/-}, no fucose, *n* = 4; *FX*^{-/-}, with fucose, *n* = 4). **E:** FACS analysis of bone marrow LSK cells displaying GFP signal in wild-type-NTg and *FX*^{-/-}-NTg mice. **F:** FACS analysis of binding of recombinant Notch ligands, Dll1 (1.3 μg/ml), Dll4 (0.5 μg/ml), and control (human IgG1 Fc, 1 μg/ml), to LSK cells isolated from wild-type mice, from *FX*^{-/-} mice reared in the presence of fucose, or from *FX*^{-/-} mice reared in the absence of fucose. The bar graph shown is one representative of the mean fluorescence intensity (MFI) of the binding of LSKs with control and Notch ligands (Dll1 and Dll4) from three independent experiments. **P* < 0.05, ***P* < 0.01.

fucose-supplemented chow had normal frequencies of LSK subpopulations and pre-B cells. These findings indicate that fucose-deficiency results in marrow myeloproliferation that is accompanied by a suppressed HSC compartment and a severely suppressed lymphoid progenitor compartment.

To correlate Notch activation in fucose-deficient HSC with HSC numbers, we used a Notch reporter transgenic (NTg) mouse to examine whether fucosylation-deficient progenitors display a down-regulated Notch signaling phenotype. The NTg mouse carries a hemizygous allele of Notch-signaling CBF-1 response element and a minimal SV40 promoter driven enhanced green fluorescent protein (eGFP). This mouse has been used to define

active Notch signaling in HSCs *in vivo*.²⁶ We found that 22% of wild-type LSKs were GFP⁺, whereas only 11.6% of fucose-depleted LSKs from *FX*^{-/-}-NTg mice were GFP⁺. The number of GFP⁺ LSKs was restored to the wild-type level when mice were reared on a fucose-supplemented diet (Figure 1E). These observations are consistent with our prior observations that Notch signaling is suppressed in HSCs and progenitors as a result of fucosylation deficiency.¹⁵ We then determined whether there is a similar fucosylation-dependent requirement for ligand binding to HSCs as for myeloid progenitors. We found that recombinant Dll4 and Dll1 ligands bound to wild-type LSKs with mean fluorescence intensity (MFI) of 1401 and 1728, respectively, whereas the MFI of human

Table 1. Competitive Limiting Dilutions of Wild-Type and *FX*^{-/-} Marrow HSC

Donor HSC	Cell dose injected	Calculated number of LSK cells injected	Number of mice engrafted/ number of mice injected	HSC frequency (± 95% confidence interval)
Wild-type	5 × 10 ⁵	500	6/6	1/17,631 (9660 to 32,179)
	5 × 10 ⁴	50	7/7	
	2.5 × 10 ⁴	25	6/8	
	5 × 10 ³	5	1/8	
<i>FX</i> ^{-/-}	5 × 10 ⁵	260	6/6	1/240,741 (96,151 to 602,764)
	5 × 10 ⁴	26	0/5	
	2.5 × 10 ⁴	13	1/9	
	5 × 10 ³	3	0/13	

Bone marrow cells were isolated from *FX*^{-/-} and wild-type mice and were transplanted into lethally irradiated CD45.1 recipient mice along with 5 × 10⁵ CD45.1 competitor cells. Numbers of LSK cells injected are calculated based on their frequency in the marrow of *FX*^{-/-} and wild-type mice. Shown are cumulative data from three separate transplant experiments. Donor engraftment was determined by FACS analysis on peripheral blood leukocytes. Donor engraftment was defined as >5% in myeloid Gr-1⁺ and lymphoid B220⁺ lineages. The cell dose at which 37% of mice tested were negative for donor multi-lineage engraftment at 12 weeks was used to calculate the frequency of HSC based on Poisson statistics using L-Cal software (StemCell Technologies). *P* < 0.01 by two-tailed Student *t* test.

Figure 2. Enhanced myelopoiesis and suppressed lymphopoiesis from MPP cells from *FX*^{-/-} mice. **A:** A representative flow cytometry dot plot analysis on day 21 of culture of bone marrow-derived MPP cells, cocultured with OP9-control cells (no ligand) or OP9-Dll1, for Gr-1 and CD11b expression. **B:** Data shown are mean values of at least four determinations. The bar graph indicates percentage of myeloid cells (expressing Gr-1 or CD11b) among total viable cells. **C–E:** Quantitative RT-PCR analysis of *Flt3* expression in total *Flt3*⁻LSK and *Flt3*⁺LSK cells (**C**), *GATA-3* expression (**D**), and *PU.1* and *C/EBPα* expression (**E**) on day 21 cultured MPP cells. Results of *Flt3* (mean ± SD; *n* = 3) or *GATA-3/PU.1/C/EBPα* (one representative of three similar experiments) have been standardized for GAPDH levels and were expressed as fold expression change relative to the levels detected in wild-type *Flt3*⁻LSK cells (**C**) or MPPs in OP9-Ret10 cultures (**D** and **E**; set at 1). **P* < 0.05 versus wild-type cell numbers or expression level.

IgG Fc staining was 218 (Figure 1F). LSKs from fucose-repleted *FX*^{-/-} mice exhibited a wild-type binding phenotype. In contrast, we observed an approximately 50% reduction in the binding of these recombinant ligands to fucose-depleted LSKs. These observations indicate that the decreased Notch signaling in fucose-depleted LSKs is accompanied by a loss of effective binding of Notch ligands to these cells.

Fucose-Deficiency Promotes Multipotent Progenitors to Adopt Myeloid Fate at the Expense of Lymphoid Specification

It has been shown that the up-regulation of *Flt3* confers a primed lymphoid potential in marrow MPPs (*Flt3*⁺LSK),^{27,28} and the suppression of myeloid differentiation potential is required for T cell commitment.²⁹ To explore the underlying mechanism of the increase in myeloid progenitors and the decrease in lymphoid progenitors in fucose-deficient mice, we sought to examine the myeloid versus lymphoid development potential of MPPs in response to Notch activation.

By using a coculture system, Schmitt et al has shown that Notch ligand-expressing OP9 stromal cells support a robust generation of CD4⁺CD8⁺ (DP) T cells from hematopoietic progenitors.³⁰ Further, the T lymphopoietic potential of LSKs is fucosylation dependent, as LSKs from *FX*^{-/-} marrow lost their T cell differentiation ability.³¹ Sim-

ilarly, we observed that when cultured on OP9s expressing Notch ligand (Jagged2, Dll1, or Dll4), wild-type MPPs generated robust DP T cells (data not shown) but minimal granulocytes. In contrast, when culturing with Notch-ligand bearing OP9s, *FX*^{-/-} MPPs differentiated into those primarily of the myeloid lineage (>50% express Gr-1 or CD11b; see Figure 2, A and B), whereas others expressed early immature T cell markers at DN1 stage (data not shown).

To further explore the underlying mechanism of the increased myeloid development potential of fucosylation-deficient MPPs, we examined the expression of *Flt3* receptor,²⁸ and *GATA-3*, a T-cell transcription factor that is known to be regulated by Notch,³⁰ in both freshly-isolated and cultured MPPs after their exposure to Notch ligands. We observed that the expression of *Flt3* mRNA in freshly-isolated *Flt3*⁺LSK and *Flt3*⁻LSK from *FX*^{-/-} mice was only 40% and 50%, respectively of that seen in wild-type subpopulations. The expression of *Flt3* was not significantly changed in fucose-repleted LSKs (Figure 2C). Wild-type cells taken from day 10 OP9 cultures in the presence of Jagged2 and Dll4 exhibited 18.4- and 27.9-fold increases in *GATA-3* expression, respectively (Figure 2D). In comparison, cultured MPPs taken from fucose-depleted *FX*^{-/-} mice showed minimal increases in *GATA-3* expression when interacting with Dll4 and Jagged 2 (3.7- and 2-fold increases, respectively). This

Figure 3. $FX^{-/-}$ HSC have impaired repopulating ability and enhanced myelopoiesis and suppressed lymphopoiesis *in vivo*. **A:** Schematic representation of the competitive transplantation protocol. **B:** Reconstitution of peripheral blood by Ly5.2 cells 12 weeks after transplantation. Reconstitution is calculated as the percentage of testing cells (Ly5.2) of total peripheral blood mononuclear cells (Ly5.2 + Ly5.1) after red cell lysis. **C:** Absolute numbers of LSK cells derived from testing cells (Ly5.2) in the recipient bone marrow of each mouse receiving 1:1 of testing (wild-type or $FX^{-/-}$) versus competing cells (wild-type, $n = 4$; $FX^{-/-}$, $n = 4$). **D:** Percentage of neutrophils (Gr-1⁺) and lymphocytes (B220⁺ and CD3e⁺) in Ly5.2 cells by FACS 12 weeks after competitive transplantation in recipient mice. Data shown are mean values $\pm$ SD. A Student *t* test was performed to compare the percentage of neutrophils or lymphocytes in transplant setting receiving $FX^{-/-}$ cells with those of wild-type cells. * $P < 0.05$, ** $P < 0.01$.

phenotype was rescued when fucose-repleted MPPs were cultured in the presence of exogenous fucose where GATA-3 expression was increased by 13.9- to 14.9-fold. Furthermore, we observed that the expression of myeloid transcription factors, PU.1 and C/EBP α , remained unchanged or was suppressed on Notch ligand stimulation in wild-type or fucose-repleted MPP cells, whereas their expressions were increased four- to eight-fold in fucose-depleted MPPs when compared with that in cells from cultures without Notch ligand (Figure 2E). Together, these observations suggest that the reduced T lymphoid and increased myeloid potential associated in fucose-depleted LSKs is partly attributable to a suppressed expression of lymphoid-priming transcription factors that are normally up-regulated, and a sustained expression of myeloid transcription factors that are normally down-regulated, during the HSC lymphoid lineage specification.

Fucose-Depleted HSC Exhibit Enhanced Myelopoiesis and Suppressed Lymphopoiesis in Vivo that is Partially Reversed by Activated Notch1 Intracellular Domain

Although $FX^{-/-}$ mice displayed a twofold decrease in the frequency of LSK cells by immunophenotyping, they revealed a 13.7-fold reduction in HSC frequency by the functional analysis. Thus a 6.8-fold discrepancy of stem cell frequency determined by limiting dilution transplantation and by immunophenotyping suggests a functional defect of fucose-deficient HSC in marrow engraftment or

self-renewal. To further assess the effect of fucose deficiency on marrow HSC reconstitution and blood lineage development, we performed competitive repopulating analyses to compare the contribution of wild-type or $FX^{-/-}$ testing cells (Ly5.2) to that of competing cells (Ly5.1) after transplantation into lethally irradiated wild-type mice (Figure 3A).

The contribution of $FX^{-/-}$ but not wild-type cells (percentage of Ly5.2) was substantially decreased when testing cells were mixed with competing cells at 1:1 and 3:1 ratios (Figure 3B). The contribution of $FX^{-/-}$ cells was almost undetectable when one part of the $FX^{-/-}$ cells was injected with three parts of the competing cells. In comparison, the contribution of wild-type cells injected at the same ratio was present in proportion to the number of the cells injected. Analysis of the bone marrow at 12 weeks after transplanting 1:1 mixture of testing and competing cells revealed that $FX^{-/-}$ LSK cells decreased to about one-third of the wild-type counterparts. Thus, the general decline in numbers observed in the contribution of $FX^{-/-}$ cells to the bone marrow reflects a depletion of the fucose-deficient HSCs (Figure 3C).

The impact of fucose deficiency on blood lineage distribution was further examined in competitive repopulation assays. We found that $FX^{-/-}$ cells displayed an increased granulocytic but a decreased lymphoid reconstitution, relative to the wild-type control, when $FX^{-/-}$ cells were mixed with wild-type competitors in a ratio of 1:1 or 3:1 (Figure 3D). These observations indicate that fucosylation-deficient marrow cells have enhanced myeloid but decreased lymphoid developmental potential. The absolute neutrophil numbers were unchanged be-

Figure 4. Expression of activated Notch1 (ICN1) partially suppresses the neutrophilia and marrow myeloid expansion induced by the $FX^{-/-}$ progenitors. **A:** Analysis of peripheral neutrophils (Gr-1⁺) by FACS in wild-type mice 4 and 6 weeks after receiving syngeneic $FX^{-/-}$ BM progenitors transduced with ICN1 or Mig vector (ICN1-wild-type, Mig-wild-type), and in $FX^{-/-}$ mice receiving $FX^{-/-}$ BM progenitors transduced with Mig vector (Mig-FX). **B:** Flow cytometric analysis of peripheral T cells (CD4⁺ or CD8⁺), B cells (B220⁺), and granulocytes (Gr1⁺) in the GFP⁺ populations 6 weeks after transplantation. **C:** Flow cytometric analysis of marrow and spleen T cells, B cells, and granulocytes in the GFP⁺ populations six weeks after transplantation. Data shown (**A–C**) are one representative of two to three mice in each group, with GFP⁺ populations ranging from 5% to 12% in all mice groups. **D:** CFU analysis of marrow or spleen progenitors six weeks after transplantation. Bar graphs indicate average numbers of myeloid colonies (CFU-G, CFU-M, and CFU-GM) per 2×10^4 total marrow nucleated cells, or per 2×10^5 splenic cells, from duplicates of two to three mice in each group.

cause of the decreased chimerism established by the $FX^{-/-}$ cells.

To determine whether the aberrant lineage homeostasis developed from $FX^{-/-}$ marrow cells can be attributed to defective Notch1 signaling activation, we performed bone marrow transplantation where donor $FX^{-/-}$ progenitors were transduced with either constitutively active Notch1-expressing retroviral vectors (ICN1) or vector controls (pMig). Previous studies have shown that expression of activated Notch1 resulted in the development of T cell leukemia/lymphoma 2 months after transplantation, whereas mice receiving activated Notch1 marrow progenitors 3 weeks after transplantation contained immature CD4⁺CD8⁺ T cells while their early B cells progenitor development was blocked.^{21,32} Here, we found that one month after transplantation, neutrophil counts were not elevated in wild-type recipients receiving ICN1-expressing $FX^{-/-}$ progenitors, when compared with the neutrophil numbers in wild-type recipients receiving wild-type marrow progenitors (data not shown). The neutrophil counts, however, were elevated 1.8-fold and 7.5-fold in wild-type and $FX^{-/-}$ recipients receiving $FX^{-/-}$ progenitors transduced with pMig, respectively, when compared with wild-type recipients receiving ICN1 (Figure 4A). Six weeks after transplantation, neutrophils continued to mildly increase in wild-type mice receiving either pMig or ICN1-expressing $FX^{-/-}$ progenitors. In $FX^{-/-}$ recipients that received pMig $FX^{-/-}$ progenitors, their neutrophil counts remained markedly elevated (Figure 4A). Analysis of peripheral T, B, and granulocytes in the GFP⁺ cells revealed that, when compared with wild-type mice receiving pMig, wild-type mice receiving ICN1 displayed an increase in frequency of CD4⁺CD8⁺ T cells (73.4% versus 37.1%), and a decrease in frequency of Gr-1⁺

cells (6.1% vs 33%). $FX^{-/-}$ mice receiving pMig had decreased lymphocytes and drastically elevated granulocytes (Figure 4B). Together, it appears that activated Notch1 partially corrects the mild neutrophilia conferred from the $FX^{-/-}$ progenitors in a wild-type recipient background. The effect of activated Notch1 on the severe neutrophilia contributed by the fucose-deficient stromal environment, however, cannot be assessed as all of the $FX^{-/-}$ mice receiving ICN1-expressing $FX^{-/-}$ progenitors did not survive beyond 2 weeks after transplantation. The effect of Notch1 activation in $FX^{-/-}$ progenitors on bone marrow and spleen lineage homeostasis was assessed by flow cytometry, using GFP expression to identify transduced cells. Six weeks after transplantation, the frequency of mature Gr1⁺ granulocytes was 13.2% of GFP⁺ cells in the marrow of wild-type recipients receiving ICN1-expressing $FX^{-/-}$ progenitors, 42.9% and 48.7% of GFP⁺ cells in wild-type and $FX^{-/-}$ recipients receiving pMig, respectively. The frequency of mature granulocytes in the spleen was 20.1%, 12.7%, and 23.4% of GFP⁺ cells in wild-type mice receiving ICN1, wild-type, and $FX^{-/-}$ mice receiving pMig, respectively (Figure 4C). The frequencies of mature granulocytes were in the range of 45% to 63% of GFP⁺ cells in the marrow of all three groups of mice, with the highest frequency found in $FX^{-/-}$ recipients receiving pMig (data not shown). This observation is consistent with our previous findings of the stromal environment-dependent control of myeloproliferation in $FX^{-/-}$ mice. Similarly, the frequency of GFP⁺ granulocytes in the spleen was in the range of 3.7% to 10.8% in all three groups of mice, with the highest frequency found in $FX^{-/-}$ recipients receiving pMig (data not shown). Analysis of marrow and spleen progenitors by methylcellulose culture revealed that single or double myeloid colony num-

bers were increased in the marrow and the spleen of wild-type mice receiving pMig and were further increased in $FX^{-/-}$ mice receiving pMig. However, the myeloid colony numbers appeared to be decreased in the marrow and remained unchanged in the spleen of the wild-type mice receiving $FX^{-/-}$ progenitors expressing ICN1 (Figure 4D). Similar to a previous report,³¹ we identified a population of immature $CD4^+CD8^+$ T cells in the marrow and the spleen of wild-type mice receiving ICN1. This phenotype was further accompanied by a block of B cell development (Figure 4C). Considered together, these data suggest that activated Notch1 partially suppressed the expansion of granulocyte derived from the $FX^{-/-}$ progenitors in the marrow. Moreover, expression of constitutively active Notch1 did not suppress the splenic myeloproliferation induced by fucose-deficiency.

ES Cells Deficient in Notch1 Receptor or Pofut1 Exhibit Suppressed Lymphopoiesis but Enhanced Myeloopoiesis in Vitro

The observations from $FX^{-/-}$ mice are consistent with the hypothesis that enhanced myeloopoiesis and suppressed lymphopoiesis are accounted for by loss of Notch-dependent control on lineage homeostasis as a consequence of loss of Notch O-fucosylation. To confirm this hypothesis, we sought to assess the role played by the O-fucosyltransferase Pofut1 in the control of myeloopoiesis and lymphopoiesis because Pofut1 is responsible for adding fucose molecules to consensus EGF repeats on Notch receptors.^{23,33} The embryonic lethal phenotypes of *Notch1*-null and *Pofut1*-null mice prohibit the use of their hematopoietic progenitors to study the effect of Notch1 or Pofut1 on myeloopoiesis.^{34–36} Instead, to determine whether the aberrant myeloid reconstitution by $FX^{-/-}$ cells is attributable specifically to suppressed Notch signaling as a consequence of either loss of Notch1 receptor or loss of O-linked fucose on Notch receptors, we analyzed the blood lineage specification from mouse ESCs carrying either a *Notch1*-null (*Notch1*^{-/-}) or *Pofut1*-null allele (*Pofut1*^{-/-}) using a modified *in vitro* culture system (Figure 5A). ESCs deficient in Pofut2, which catalyzes the addition of O-linked fucose to thrombospondin type 1 repeats (TSRs)^{37,38} were studied in parallel to confirm that it does not modulate Notch-dependent signaling in hematopoiesis. In this system, ESCs were initially differentiated into $CD34^+$ cells in methylcellulose before coculturing with either OP9 cells bearing different Notch ligands or control OP9 cells (Figure 5A).^{30,39}

After a 15-day culture with control OP9 cells, wild-type ESC-derived $CD34^+$ cells had differentiated into myeloid cells expressing Gr-1 (Figure 5B) and B lymphoid cells (data not shown). In contrast, these cells differentiated into DP/CD8⁺ T lymphocytes (69.3%, 73.4%, and 70.9%, respectively) when cocultured with OP9s bearing Jagged2, Dll1, or Dll4. The induction of T lymphopoiesis by the Notch ligand-expressing OP9 is consistent with previously published observations of marrow LSK cell differentiation.³⁰ The development of mature T cells in culture correlates with the up-regulation of Notch-specific

transcripts, including Hes1 and Deltex1 (Figure 5C). Wild-type ESC-derived progenitor cells also displayed efficient Notch ligand (Dll1 and Dll4) binding (Figure 5D). When $CD34^+$ *Pofut2*^{-/-} progenitors were cultured under similar conditions, they showed comparable T lymphopoiesis to wild-type cells. These cells also exhibited similar levels of Notch activation and Notch ligand binding as observed in wild-type ESCs (Figure 5, B–D). Together, these latter observations indicate that Notch activation and its role in inducing T lymphoid development proceeds normally in the absence of O-fucosylation of TSR-1 repeats.

Although *Notch1*^{-/-} ESCs displayed a normal myeloid and B lymphoid differentiation pattern when cultured on OP9-controls, they failed to generate T lymphocytes on Notch ligand-expressing OP9s. Instead, these cultures contained a robust population of myeloid cells, which accounted for between 39.4% and 45.3% of the total hematopoietic cells in the culture (Figure 5B). The absence of T lymphopoiesis in these cultures correlated with the basal levels of Hes1 and Deltex1 expression and the background level of Notch ligand binding, all of which indicate complete abrogation of Notch activation (Figure 5, B–D). Regardless of whether Notch ligand was present in the culture, when *Pofut1*^{-/-} ESCs were assayed, they generated a pattern of increased myeloopoiesis and absence of T lymphopoiesis that was essentially identical to that of *Notch1*^{-/-} ESCs. In contrast to the complete abrogation of Notch activation and loss of Notch ligand binding observed in *Notch1*^{-/-} ESCs, we found that the expression of Notch-targeted transcripts was diminished (but not absent) whereas Notch ligand binding was markedly decreased (but not absent), relative to wild-type cells (Figure 5, B–D). These observations indicate that the partial suppression of Notch activation in the absence of O-linked fucosylation on Notch yields an ESC hematopoietic phenotype similar to that observed in *Notch1*^{-/-} cells. Further analysis of these ESCs revealed that the surface expression of Notch1 was absent in *Notch1*^{-/-} HPCs but mildly decreased in *Pofut1*^{-/-} HPCs. The expression levels of other Notch receptors (Notch2, Notch3, and Notch4) in *Pofut1*^{-/-} HPCs were not up-regulated when compared with the expression of these receptors in *Notch1*^{-/-} cells (see Supplemental Figure S1 at <http://ajp.amjpathol.org>). Together, these findings suggest that a defective Notch-ligand binding mediated by Notch1 could be the primary cause for the phenotype seen in *Pofut1*^{-/-} cells.

Aberrant in Vivo Hematopoiesis from ES Cells Deficient in Notch1 Receptor or Notch O-Fucose Modification

To confirm that the anomalous *in vitro* lineage developmental potential of *Notch1*^{-/-} and *Pofut1*^{-/-} ESCs is reflective of their *in vivo* features, we performed intrafemoral transplantation of ESC-derived $CD34^+$, $Ly5.2^+$ hematopoietic progenitors (HPCs) into nonlethally irradiated recipient mice ($Ly5.1^+$). ESCs were first differentiated into hematopoietic progenitors in methylcellulose me-

Figure 5. Notch1^{-/-} or Pofut1^{-/-} ESCs have suppressed lymphopoiesis but enhanced myeloopoiesis *in vitro*. **A:** Schematic representation of the *in vitro* coculture of ESCs (wild-type, Notch1^{-/-}, Pofut1^{-/-}, and Pofut2^{-/-}) with OP9 cells expressing Notch ligands in the presence of IL7 and Flt3 ligand. **B:** FACS analysis of CD8 and Gr-1 expression on four lines of ESC-derived hematopoietic cells after coculture with control OP9 (Ret10) or OP9 expressing Jagged2, DII1, or DII4. **C:** Expression of *Hes1* and *Deltex1* was analyzed by quantitative RT-PCR on ESC-derived hematopoietic cells relative to the levels detected in OP9-Ret10 (set at 1). Data shown are one representative of three similar experiments. **D:** One representative flow cytometry analysis of binding of recombinant Notch ligands DII1, DII4, and control (human IgG1 Fc) to ESC-derived hematopoietic cells. Bar graph indicates MFI.

dium (Figure 6A). The percentage of CD34⁺ HPCs in these differentiated ESCs is consistent with other reports ranging from 8 to 14%.¹⁷

The earliest hematopoietic reconstitution from ESC-derived HPCs was observed 2 weeks after the injection of CD34⁺ cells. By 12 weeks, the proportion of ESC-derived Ly5.2⁺ cells in the peripheral blood ranged from 52 ± 25% to 79 ± 13% for all types of cells transplanted. The fraction of Ly5.2⁺ cells remained steady through 16 to 20

weeks after transplantation. Flow cytometric analysis of peripheral blood revealed that the population of wild-type HPC-derived T cells, B cells, and granulocytes comprising 25.1 ± 8.4%, 54.1 ± 9.2%, and 12.3 ± 1.4%, Ly5.2⁺ cells, respectively. A similar lineage distribution was found in mice transplanted with Pofut2^{-/-} HPCs (Figure 6, B and C). At 12 weeks, mature T and B cells derived from Notch1^{-/-} HPCs represented only 8.8 ± 2.5% and 16.9 ± 3.3% of the transplanted cells, respectively. In

Figure 6. *Notch1*^{-/-} or *Pofut1*^{-/-} HPCs have aberrant *in vivo* hematopoiesis. **A:** Schematic representation of HPC intrafemoral injection and *in vivo* hematopoietic reconstitution in recipient mice. **B:** FACS analysis of peripheral blood reconstitution by antibodies against T cells (CD3), B cells (B220), and neutrophils (Gr-1) after gating on ESC-derived cells (Ly5.2). **C:** Line graphs of peripheral blood lymphoid and myeloid lineage reconstitution by ESC-derived cells (Ly5.2) at 2, 4, 8, and 12 weeks after intrafemoral transplantation. **D:** Comparison of total thymocyte numbers and the numbers of thymocytes contributed by the host (Ly5.1) and HPCs (Ly5.2). FACS analysis was performed with anti-CD4 and anti-CD8 after gating on Ly5.2 or Ly5.1 cells. Data shown are one representative of three similar experiments. **E:** Thymocyte development after intrathymic transplantation of HPCs (Ly5.2) into *FX*^{-/-} mice (Ly5.1) maintained on standard chow. Data shown are one representative of three similar experiments.

comparison, the proportion of the granulocytes from *Notch1*^{-/-} HPCs reached $57.7 \pm 9.6\%$. The hematopoietic reconstitution of the *Pofut1*^{-/-} HPC donor had a pattern of lineage distribution similar to that of *Notch1*^{-/-} HPC-reconstituted mice: decreased T and B lymphoid compartments, increased granulocytic chimerism (Figure 6, B and C) and absolute neutrophilia (see Supplemental Table 1 at <http://ajp.amjpathol.org>). As summarized in Table 2, 12 weeks after transplantation, hematopoietic reconstitution by *Pofut2*^{-/-} HPCs resulted in a lineage distribution that was essentially identical to mice reconstituted with wild-type HPCs. In contrast, *Notch1*^{-/-} and *Pofut1*^{-/-} HPCs yielded a lymphoid-suppressed but

myeloid-expanded phenotype. The absolute number of B cells derived from donors carrying *Notch1*^{-/-} or *Pofut1*^{-/-} alleles remained similar because of increased WBC numbers (see Supplemental Table 1 at <http://ajp.amjpathol.org>). The proportion and absolute numbers of T cells, B cells, and granulocytes generated from hosts (Ly5.1⁺) were similar among mice receiving all four ESC lines, suggesting that the lymphoid-suppressive and myeloid-enhancing phenotype in recipients transplanted with either *Notch1*^{-/-} or *Pofut1*^{-/-} HPC is cell-autonomous.

A decreased T lymphocyte population in the periphery of recipients receiving *Notch1*^{-/-} or *Pofut1*^{-/-} HPCs is a result of suppressed thymic T lymphopoiesis in these

Table 2. Lineage Distributions of ESC-Derived Ly5.2 and Host-Derived Ly5.1 Cells in Nonlethally Irradiated Recipient Mice Transplanted with CD34⁺ ESC-HPCs (12 Weeks after Injection)

	Peripheral blood					
	Ly5.2+ Cells (n = 4)			Ly5.1+ Cells (n = 4)		
	T (%)	B (%)	G (%)	T (%)	B (%)	G (%)
Wild-type	25.1 ± 8.4	54.1 ± 9.2	12.3 ± 1.4	44.8 ± 9.8	25.2 ± 9.4	16.7 ± 7.0
Pofut2 ^{-/-}	21.0 ± 2.5	60.9 ± 6.6	11.6 ± 6.2	43.5 ± 6.5	24.4 ± 10.6	23.0 ± 12.5
Notch1 ^{-/-}	8.8 ± 2.5**	16.9 ± 3.2*	57.7 ± 9.6**	32.3 ± 2.0	34.3 ± 3.9	23.8 ± 1.5
Pofut1 ^{-/-}	8.0 ± 3.4**	23.3 ± 8.6*	51.6 ± 7.6**	36.7 ± 9.4	30.5 ± 6.8	17.2 ± 7.2

*P < 0.01; **P < 0.0001.

mice. In one representative experiment, the absolute cell count of the donor-derived thymocytes (Ly5.2⁺) were only 0.3×10^6 and 0.04×10^6 in each mouse receiving *Notch1*^{-/-} or *Pofut1*^{-/-} HPCs, respectively, whereas the donor thymocyte numbers in each mouse receiving wild-type or *Pofut2*^{-/-} HPCs were 34.4×10^6 and 29.1×10^6 , respectively. Both single positive (SP) and double positive (DP) T cells derived from either *Notch1*^{-/-} or *Pofut1*^{-/-} HPCs were significantly decreased (Figure 6D). In comparison, the development of mature thymocytes generated from the hosts (Ly5.1⁺) was not disturbed. To exclude the possibility that defective thymopoiesis is attributable to a thymus-homing defect of HPCs, CD34⁺ cells were directly injected into the thymus of fucose-depleted *FX*^{-/-} mice (Ly5.1) whose ability to support the host thymocyte development is retarded (unpublished data). Essentially no SP or DP mature T cells were found derived from the injected *Notch1*^{-/-} or *Pofut1*^{-/-} HPCs, whereas robust T cells were identified from wild-type and *Pofut2*^{-/-} HPCs (Figure 6E).

The increase in granulocytes in mice receiving *Notch1*^{-/-} or *Pofut1*^{-/-} HPCs was accompanied by marrow myeloproliferation and extramedullary hematopoiesis. The spleens of these mice were mildly enlarged, of which the red pulps were expanded because of myeloid hyperplasia (see Supplemental Fig S2, A and B at <http://ajp.amjpathol.org>). The B lymphocyte progenitors were mildly decreased but did not display any obvious developmental blockade in the marrow (data not shown). There is a decrease of IgM^{hi}IgD^{lo} B cells in spleens of mice receiving *Pofut1*^{-/-} HPCs, as was found in *FX*^{-/-} mice spleens (Figure 7). The splenic IgM^{hi}IgD^{lo} B cells comprise type1 transitional (T1) and marginal zone B cells (MZB), which are decreased in *Notch2* conditional

knock-out mice.⁴⁰ Mice receiving either *Notch1*^{-/-} or *Pofut1*^{-/-} HPCs also displayed infiltrations of neutrophils and myeloid progenitors to the peripheral organs such as lymph nodes (see Supplemental Fig S2C at <http://ajp.amjpathol.org>). However, these mice displayed a normal peripheral blood platelet count (see Supplemental Fig S3 at <http://ajp.amjpathol.org>). These findings further support that loss of *Notch1* or *Notch* O-fucosylation leads to chronic granulomonocytic proliferation.

Discussion

In this study, we identified a role of Notch receptor O-fucose modification in regulating blood lineage homeostasis. We confirmed our previous findings that a myeloproliferation in mice with fucosylation deficiency is attributable to an enhanced myeloid developmental potential of HSCs. The data presented in our current study indicate that Notch activation modulates lineage commitment and development from HSCs. This regulation exerted by Notch is controlled by O-fucosylation of Notch receptors. We identified that the absence of O-fucose on Notch receptors results in decreased lymphoid priming but enhanced myeloid potential. This phenotype is accompanied by suppressed Notch ligand binding and Notch signaling activation in HSCs.

Commitment of HSCs to either a myeloid or lymphoid lineage is believed to be regulated by the activation of several key transcription factors, which establish the profile of genes characteristic of that lineage.²⁹ The tyrosine kinase receptor Flt3 and the transmembrane protein Notch1 receptor are important proteins for lymphoid lineage initiation,^{41,42} whereas PU.1 and C/EBP α are transcription factors that control myeloid lineage development.⁴³ It is known, and confirmed by our observations in this study, that Notch1 promotes T lymphoid commitment by stimulating T cell-specific transcription factors while down-regulating myeloid transcription factors.^{29,44} We identified an enhanced myeloid but a suppressed lymphoid potential of HSCs in the absence of fucose. This was demonstrated that fucose-deficient HSCs form more myeloid progenitor colonies and MPPs form *FX*^{-/-} mice failed to respond to Notch ligand-induced T lineage commitment while exhibiting a sustained expression of myeloid transcriptional activity *in vitro*. The suppressed lymphoid potential of fucose-deficient hematopoietic progenitors is also suggested by the reduction of Flt3 tran-

Figure 7. A Decreased MZB cell compartment in the spleens of *FX*^{-/-} mice and mice receiving *Pofut1*^{-/-} HPCs. A representative FACS analysis of T1 B cells and MZB cells (IgM^{hi}IgD^{lo}) in the spleens of wild-type and *FX*^{-/-} mice (**top**; n = 4) and mice receiving ES-derived HPCs (**bottom**; n = 4). The numbers denote the percentage of cells in the indicated squares.

scripts in lymphoid-primed MPPs. It remains unclear why the expression of myeloid transcription factors found in fucose-deficient progenitors in response to Notch ligand stimulation is increased. We speculate that weak Notch activation may promote myeloid-specific events in fucose-deficient progenitors. In comparison, strong Notch signaling may promote the elevation of lymphoid transcription factors as seen in wild-type or fucose-repleted cells. Alternatively, altered Notch signaling pathways attributable to O-fucosylation deficiency, such as misfolding of Notch receptors in the ER,²² decreased Notch receptor or Notch ligand endocytosis,^{45,46} may cause enhanced myeloid commitment from MPPs through unidentified molecular mechanisms.

Our data from *in vitro* hematopoietic differentiation of ESC is consistent with the hypothesis that Notch activation supports T lymphopoiesis and inhibits myelopoiesis. Consistent with this finding, we observed that a transient and balanced leukocyte lineage development established by wild-type ESCs, whereas loss of Notch1 results in a diminished lymphoid but enhanced myeloid lineage outcome. We also observed that the balanced lymphoid versus myeloid lineage outcome exerted by Notch activation is controlled by the modification of O-fucose on Notch. Both *in vitro* and *in vivo* hematopoiesis generated by ESCs display enhanced myeloid but inhibited lymphoid development in the absence of Pofut1. Indeed, mice reconstituted with *Notch1*^{-/-} or *Pofut1*^{-/-} progenitors displayed both neutrophilia and myelohyperplasia. This finding is consistent with the myeloproliferation observed in the *FX*^{-/-} mice and the enhanced granulopoietic reconstitution produced by fucose-deficient progenitor cells in wild-type recipients. Furthermore, both myeloproliferation and suppression of T lymphopoiesis are observed in conditional *Pofut1* knock-out mice (manuscript in preparation), suggesting that the myeloproliferative phenotype observed in *FX*^{-/-} mice can be largely attributed to an abnormal Notch signaling control of myelopoiesis in the absence of O-linked fucose on Notch receptors. These findings strongly support the hypothesis that Notch signaling is essential for blood lineage homeostasis by promoting lymphoid development while suppressing overt myelopoiesis. Further studies at the single HSC level are required to define whether such control mechanism is exerted by Notch at the earliest diversification of lymphoid and myeloid progenitor during HSC lineage commitment. Because we have shown that this control mechanism exerted by Notch is dependent on the O-fucose modification of Notch EGF repeats, future studies will examine how each informative EGF-like repeat domain regulates the activity of Notch signaling.

O-fucose moieties are found on other proteins that have EGF-like repeat motifs, including Cripto and coagulation factors,^{47–50} as well as on proteins containing TSR repeat domains.^{51,52} Unlike Pofut1, which is responsible for the addition of fucose to EGF-like repeats, Pofut2 modifies TSR repeats, which was first described in thrombospondin-1.⁵³ Our findings of normal blood lineage development using *Pofut2*^{-/-} ESC, both *in vitro* and *in vivo*, indicate that Pofut2 is not essential for the development of hematopoietic cells. In contrast, Pofut1 is required for

blood lineage homeostasis as its deficiency results in enhanced granulocytic potential and suppressed lymphoid potential. This effect is likely caused by decreased binding of *Pofut1*^{-/-} ESC-derived hematopoietic cells with Notch ligand. The faulty ligand-receptor interaction crippled the ability of hematopoietic progenitors to transcribe Notch-specific targets. This reduced ligand-receptor binding phenotype may also stem from the decreased cell surface expression of Notch1 in *Pofut1*^{-/-} HPCs. Interestingly, the cell surface expressions of Notch2, Notch3, and Notch4 are not changed in *Pofut1*^{-/-} HPCs when compared with that in *Notch1*^{-/-} HPCs. This is consistent with the observations made by Stahl et al, which indicate that *Pofut1*^{-/-} ES cells exhibit suppressed Notch signaling using a reporter assay.⁵⁴ Although Pofut1 also transfers fucose molecules to EGF repeat domains of Cripto, a coreceptor for Nodal, it is unlikely that the phenotype observed with *Pofut1*^{-/-} ESC is caused by a faulty Nodal-mediated signaling as it has been shown that Nodal signaling is not disturbed in *Pofut1*^{-/-} ESCs.⁵⁵ Further, the normal blood lineage distribution established by wild-type host cells in chimeric mice receiving *Pofut1*^{-/-} progenitors argues against that secreted plasma proteins, such as coagulation factors derived from *Pofut1*^{-/-} ESCs, may contribute to the aberrant phenotypes observed.

All four Notch receptors are expressed in HSCs, each of which has consensus sequences known to be modified by O-fucose.⁵⁶ The similarity of aberrant hematopoietic reconstitution between *Notch1*^{-/-} and *Pofut1*^{-/-} progenitors suggests that among the members of the Notch family, Notch1 probably plays the primary role in blood lineage maintenance. This notion is supported by the findings that the expression of a constitutively activated Notch1 partially suppressed myeloproliferation conferred by the *FX*^{-/-} marrow progenitors. However, we cannot exclude the possibility that the suppression of myeloproliferation in the marrow is only secondary to the expansion of immature T cells induced by activated Notch1. In contrast, Notch1 activation seems to play no role in splenic myelohyperplasia induced by fucose deficiency, suggesting that defective signaling of other Notch receptors may play a more important role in extramedullary myeloproliferation. Our observation that Notch1 deficiency leads to an enhanced myelopoiesis and suppressed T lymphopoiesis, however, differs from the findings by Radtke et al who previously demonstrated that conditional inactivation of Notch1 arrested T cell development whereas the differentiations of other lineages are unaffected.⁹ The discrepancy of these observations suggests that other Notch receptors may have redundant functions in myelopoietic suppression but not in ES-derived hematopoiesis in mice. Alternatively, an enhanced myeloid reconstitution by *Notch1*^{-/-} progenitors may reflect a developmental stage-specific effect of ES-derived progenitors on hematopoietic tissues, whereas such an effect is diminished or bypassed in Cre-LoxP inactivation system that targets the adult hematopoietic tissues and cells. Our findings of the suppressed T lymphopoiesis and the enhanced myelopoiesis with a rather normal peripheral B lymphoid compartment in *FX*^{-/-} mice and the *in*

vivo reconstitution of ESC may suggest a role of Notch signaling at the common progenitor of T/myeloid branching point, as recently demonstrated by Bell et al that early thymic progenitors possess T and myeloid potentials.⁵⁷ Alternatively, a rather normal B lymphoid compartment may reflect a combined effect of Notch signaling at two branching points (i.e., suppressed Notch activation decreases CLP numbers at the CLP versus CMP restriction while increasing B lymphocyte development at the expense of T lymphocytes at the T/B restriction). Despite a normal B cell compartment in the periphery, we observed a decrease of T1 B cells and MZB cells in *FX*^{-/-} mice and mice reconstituted with *Pofut1*^{-/-} HPCs. This finding suggests a role of fucose-dependent Notch2 signaling in MZB cell development. In contrast to the recent report that Notch signaling promotes megakaryocyte development,⁵⁸ we did not observe a significant reduction of platelet counts in *FX*^{-/-} mice or in mice receiving *Notch1*^{-/-} or *Pofut1*^{-/-} HPCs, suggesting O-fucose modification of Notch is not essential for megakaryocytopoiesis.

FX^{-/-} mice display a mild decreased concentration of LSKs under steady state. These mice have a profound decrease of HSC frequency and an aberrant repopulating function after myeloablative transplantation. The less competitive repopulating ability of fucose-deficient HSCs cannot be fully explained by an approximately twofold dilutional effect on the stem cell compartment attributable to myeloproliferation. It is, however, accompanied by a diminished HSC compartment in recipients of these cells after transplantation, which is likely attributable to the increased differentiation of HSC toward myeloid lineage.¹⁵ Alternatively, this observation is also possibly attributable to an defective stem cell engraftment or reduced HSC self-renewal, as demonstrated by a decreased stem cell frequency in the limiting dilution assay. Suppressed Notch activation in fucose-depleted LSKs, however, may not be relevant to the observed changes in the HSC compartment of *FX*^{-/-} mice, as suggested by a recent report that HSC self-renewal is not affected by a dominant negative inhibitor of canonical Notch signaling pathway.¹¹ In fact, we found that the expression of activated Notch1 is associated with a decrease of the LSK population (data not shown). Whether such aberrancies in HSC function observed in *FX*^{-/-} mice reflects an anomaly in Notch pathways upstream of Notch intracellular signaling (e.g., the interaction between Notch-expressing HSCs with the ligand-expressing osteoblasts, which dictates the physical association of HSC with marrow microenvironment that is critical for HSC biology) is unclear.⁴⁶ Further, other secreted or membrane-associated fucosylated stromal components may also work in concert with Notch directly, or indirectly, to maintain HSC function. In fact, only relatively enhanced myeloid potential is developed from fucose-deficient progenitors in a wild-type environment, whereas an absolute neutrophilia is induced by *Notch1*^{-/-} and *Pofut1*^{-/-} HPCs, which is likely caused by an altered microenvironment created by *Notch1*^{-/-} and *Pofut1*^{-/-} ES-derived cells. Combined with the finding that a more severe myeloproliferative phenotype can be induced by *FX*^{-/-} progenitors only in the *FX*^{-/-} marrow, our study suggests that fucose-defi-

cient marrow environment potentiates myeloproliferation induced by fucose-deficient marrow cells. Together these observations support the notion that marrow stem cell niche is maintained by complex interaction of HSCs with stromal cells and signaling molecules in a specialized microenvironment.^{59,60}

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