MYELOID NEOPLASIA

Fbxl10 overexpression in murine hematopoietic stem cells induces leukemia involving metabolic activation and upregulation of Nsg2

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Key Points

- Fbxl10 is a bona fide oncogene in vivo.
- Fbxl10 overexpression in HSCs induces mitochondrial metabolic activation and enhanced expression of Nsg2.

We previously reported that deficiency for *Samd9L*, which was cloned as a candidate gene for -7/7q- syndrome, accelerated leukemia cooperatively with enhanced expression of a histone demethylase: *F-box* and leucine-rich repeat protein 10 (*FbxI10*, also known as *Jhdm1b*, *Kdm2b*, and *Ndy1*). To further investigate the role of *FbxI10* in leukemogenesis, we generated transgenic (Tg) mice that overexpress *FbxI10* in hematopoietic stem cells (HSCs). Interestingly, *FbxI10* Tg mice developed myeloid or B-lymphoid leukemia with complete penetrance. HSCs from the Tg mice exhibited an accelerated G0/G1-to-S transition with a normal G0 to G1 entry, resulting in pleiotropic progenitor cell expansion. *FbxI10*Tg HSCs displayed enhanced expression of neuron-specific gene family member 2

(*Nsg2*), and forced expression of *Nsg2* in primary bone marrow cells resulted in expansion of immature cells. In addition, the genes involved in mitochondrial oxidative phosphorylation were markedly enriched in *Fbx110* Tg HSCs, coupled with increased cellular adenosine 5'-triphosphate levels. Moreover, chromatin immunoprecipitation followed by sequencing analysis demonstrated that Fbx110 directly binds to the regulatory regions of *Nsg2* and oxidative phosphorylation genes. These findings define *Fbx110* as a bona fide oncogene, whose deregulated expression contributes to the development of leukemia involving metabolic proliferative advantage and *Nsg2*-mediated impaired differentiation. (*Blood.* 2015;125(22):3437-3446)

Introduction

Appropriate patterns of epigenetic alterations in histone modifications are required to assure cell identity, and their deregulation can contribute to human diseases such as cancer.¹ We previously generated knockout mice for *Samd9L* (which was cloned as a candidate gene for the -7/7q- syndrome frequently that is observed in myelodysplastic syndrome and acute leukemia patients) and demonstrated that mice deficient in *Samd9L* developed leukemia after a long latent period.² In addition, retroviral insertional mutagenesis revealed that the onset of the disease was highly accelerated with upregulation of *F-box* and leucine-rich repeat protein 10 (*Fbx110*) or ectopic virus integration 1 (*Evi1*).² Clinically, an elevated expression of Fbx110 is observed in patients with acute myeloid leukemia (AML) or acute lymphoid leukemia (ALL), seminoma, and pancreatic ductal adenocarcinoma.³⁻⁶ These findings prompted us to further investigate the role of Fbx110 in leukemia development in vivo.

Fbx110 belongs to the JmjC domain-containing histone demethylases, and contains an N-terminal JmjC domain, followed by a CXXC zinc finger domain, a plant homeodomain finger, an F-box, and 8 leucine-rich repeats.⁷ Fbx110 preferentially demethylates the dimethylation of histone H3 at lysine 36 (H3K36me2), but not H3K36me1 and H3K36me3.^{6,8} Fbx110 was also recently reported to mediate the monoubiquitination of the histone H2A at lysine 119 as a component of noncanonical polycomb-repressive complex 1 in embryonic stem (ES) cells.^{9,10} Several lines of evidence have so far suggested that Fbx110 is a tumor accelerator. Overexpression of Fbx110 in fibroblasts inhibited cellular senescence by repressing p16ink4ap19Arf and p15ink4b loci in mouse as well as Rb and p53 loci in humans, resulting in cellular immortalization.^{6,11} In addition, wild-type (WT) Fbx110 but not a demethylase activity-deficient mutant, accelerated the progression of pancreatic cancer in a mouse allograft model.⁵ Fbx110 is highly expressed in lineage marker (Lin)⁻, Sca-1⁺, c-Kit⁺ (LSK) cells and Lin⁻ undifferentiated bone marrow (BM) cells, and forced expression of Fbx110 in hematopoietic stem cells (HSCs) retained high BM repopulation capacity.12 Furthermore, another study demonstrated that the demethylase enzymatic activity was required for leukemic transformation in a Hoxa9-Meis1-induced mouse BM transplantation model.¹³ These findings suggest that Fbx110 contributes to the development of tumor in vivo.

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Figure 1. Transgenic expression of *Fbx110* **in mice develop leukemia.** (A) Kaplan-Meier survival plots for the *Fbx110* Tg (Tg) (n = 30) and control WT (Ctrl) (n = 20) mice. (B) Representative peripheral blood smear (left) and macroscopic appearance (right) of a leukemic Tg mouse. Proliferation of blast cells and massive hepatosplenomegaly are shown. (C) Flow cytometric profiles in the leukemic tissues of 2 Tg mice. TG-1 was positive for Mac1 and/or Gr1 positive but negative for Thy1.2 and B220 and diagnosed with AML, whereas TG-2 was positive for B220 but negative for Thy1.2, Gr1, and Mac1 and accordingly diagnosed with B-lymphocytic leukemia (B-ALL). (D) Survival curves of Tg, Tg/MOL4070LTR (Tg mice infected with MOL4070LTR), control (Ctrl), Ctrl/MOL4070LTR. MOL4070LTR infection induced acute leukemias in all Tg/MOL4070LTR mice showed significantly accelerated mortality compared with the original Tg mice, whereas WT Ctrl mice did not. The time of MOL4070LTR injection is indicated by an arrow. (E) Survival curves of Tg, Tg/ENU (Tg mice injected with ENU), Ctrl/ENU mice. ENU-induced acute leukemias in all Tg/MOL4070LTR injection is indicated by an arrow. For survival comparisons, the Tg and control cohorts shown in panel A were merged in panels D and E.

Genomic and functional studies have identified 2 broad classes of mutations that cooperate during the development of acute leukemia.^{14,15} Class I mutations confer a proliferative and/or survival advantage of HSCs and progenitor cells. Class II mutations are associated with impaired hematopoietic differentiation.^{14,15} As a simplified model, it has been proposed that growth advantage together with impaired differentiation is required for the development of the acute leukemia phenotype. In this study, we report that *Fbx110* transgenic (Tg) mice that overexpress *Fbx110* in HSCs spontaneously develop leukemia involving these class I/class II mutationmimetic properties.

Methods

Generation of Fbxl10 transgenic mice

Murine *Fbxl10* complementary DNA (cDNA) with a *Flag* tag at the 3' end was inserted at the *ClaI* site of the *Ly-6A* transgenic cassette (*pLy-6A14*) that encompasses the mouse *Sca-1* gene.^{16,17} A fragment containing the *Sca-1* promoter, *Fbxl10* cDNA with *Flag* tag and *Sca-1* 3' locus control regions was excised and microinjected into the pronuclei of C57BL/6N mice. All the mice were kept according to the guidelines of the Institute of Laboratory Animal Science, Hiroshima University.

Statistics

Mouse survival curves were constructed using the Kaplan-Meier methodology and compared by the log-rank test using the GraphPad Prism software. Other statistical analyses were performed using the Student t test, unless otherwise stated.

A complete and detailed description of methods can be found in the supplemental Methods (available on the *Blood* Web site).

Results

Fbxl10 transgenic mice spontaneously develop myeloid and B-lymphocytic leukemias

The *Sca-1* gene promoter is activated in functional repopulating adult HSCs in mice and has been, so far, widely available for Tg research.¹⁶⁻¹⁹ To achieve Tg expression of *Fbxl10* in mice, the *Flag*tagged mouse Fbxl10 cDNA was inserted into the cloning site of the Ly6A/Sca-1 Tg cassette that allows high expression in the HSC compartment¹⁷ (supplemental Figure 1A). We confirmed a >10-fold upregulation of *Fbxl10* messenger RNA (mRNA) in Sca-1⁺ cells in 2 independent lines (lines 3 and 4) (supplemental Figure 1B), both of which provided similar results in this study (thus, hereafter we refer to mice in both lines as Fbxl10 Tg mice). We found that the expression levels of Fbxl10 mRNA were higher in Tg cells than control cells in all of the compartments, especially in HSC-early progenitor fractions (supplemental Figure 1C). This is consistent with the result of the initial study reporting that the same Sca-1 promoter used in this study is vigorously active predominantly in murine HSCs.¹⁷ The overexpression of the Fbx110 protein was confirmed by immunoprecipitation followed by western blot using an anti-Flag antibody (supplemental Figure 1D). Immunoblot of extracted histones obtained from sorted Sca-1⁺ cells and differentiated Sca-1⁻ cells showed a strong reduction of dimethylated H3K36 (H3K36me2) levels in the Tg cells, indicating an increased catalytic function of Fbx110 in mice (supplemental Figure 1E). One possible explanation for decreased H3K36me2 levels in Sca-1 cells may be that Fbx110 protein was sustained in Sca-1⁻ progenitor and other differentiated populations.

We followed 30 Tg and 20 control littermates over a 1.5-year period. Interestingly, all of the Fbxl10 Tg mice developed acute leukemias and died during the observation period (median survival = 367 days), whereas no hematopoietic disease was observed in the WT mice



Figure 2. Ly6A/Sca-1 promoter-driven Fbx/10 overexpression induces cell-autonomous proliferation in HSCs. (A) Absolute numbers of HSCs (LT-HSC, ST-HSC, and MPP) and progenitor (CLP, CMP, GMP, and MEP) cells (n = 4-8, *P < .025). Surface markers to separate mouse stem and progenitor cell subsets are shown in supplemental Table 3. (B) Increased in vivo BrdU incorporation in LT-HSCs of Tg mice. BrdU-positive cells in LT-HSC and ST-HSC were analyzed by flow cytometry (n = 4 per group, *P = .024). (C) Summary of flow cytometric pyronin Y analysis of CD34⁻ LSK or CD34⁺ LSK fractions in control or Tg BM cells (n = 4 per group). Error bars, SD. (D) Percent chimerism of donor cells in the peripheral blood after serial BM transplantation. Upper panel shows a schematic diagram of the experimental strategies. Each dot shows the percent chimerism of donor cells in the peripheral blood of recipients 12 weeks after the secondary transplant. Bars indicate the average (n = 6 per group, *P < .05, **P < .01). CBC, complete blood cell count; MEP, megakaryocyte/erythroid progenitor; n.s., not significant; SD, standard deviation; WBC, white blood cell count.

(Figure 1A). All moribund Tg mice showed leukocytosis and numerous circulating leukemic blasts in the peripheral blood (Figure 1B, left) and frequently displayed gross hepatosplenomegaly (Figure 1B, right). Leukemic cells from 14 of the 25 tested mice were abundantly positive for both Gr1 and Mac1 and were accordingly diagnosed as AML (Tg-1 in Figure 1C and supplemental Table 1). Seven mice were diagnosed as B-cell ALL (B-ALL), in which B220⁺ cells dominantly proliferated (Tg-2 in Figure 1C and supplemental Table 1). Regarding the remaining 4 leukemic mice, although surface marker analysis was not applicable, tumor tissues exhibited germline patterns in the $TCR\beta$ and IgH loci (supplemental Table 1). Therefore, we did not find any mice that had an expansion of Thy1.2 T-cell antigen-positive cells and/ or rearrangements in $TCR\beta$ locus in the leukemic tissues, suggesting that Fbxl10 Tg mice exclusively developed myeloid and B-cell lineage leukemias. We intraperitoneally injected cells disaggregated from Tg leukemic spleens into immunodeficient NOD/SCID mice. The recipient mice exhibited splenomegaly as did the Fbxl10 Tg mice, indicating that leukemic cells were transplantable (supplemental Figure 2).

To further assess the tumor susceptibility caused by the expression of Fbx110, neonate mice were infected with replication-competent MOL4070LTR mouse leukemia retrovirus, which is capable of inducing both myeloid disease and lymphoid diseases.²⁰ The injection of MOL4070LTR profoundly accelerated the onset of the disease in the *Fbxl10* Tg group (median survival = 274 days, P < .0001) (Figure 1D). We also applied a chemically (N-ethyl-N-nitrosourea [ENU]) induced leukemia model to the same cohort, which enhanced leukemogenesis in Tg mice (median survival = 210 days, P < .0001) (Figure 1E). Consistent with the spontaneously developed leukemias, the injection of virus predisposed Tg mice to AML and B-ALL but not T-cell (T-ALL) (supplemental Figure 3). Thus, Tg expression of Fbx110 in HSCs drives myeloid and B-cell lineage leukemias with enhanced tumor susceptibility.

Transgenic expression of FbxI10 induces proliferation advantage in HSCs

We first analyzed BM cellularity at 8 weeks of age in control and Tg mice, and found similar absolute cell numbers of multiple lineages among these mice (supplemental Figure 4A-B). To characterize more primitive cells, we further compared the HSC compartment and progenitor populations in the BM of control and Tg mice. Flow cytometric analysis revealed that the populations of myeloid progenitor cells (common myeloid progenitor [CMP] and granulocyte/macrophage progenitor [GMP]) and common lymphoid progenitor (CLP) cells were

Refseq_name	Gene symbol	Ctrl	Tg	Tg/Ctrl	Description
Genes upregulated in Tg HSCs					
NM_008741	Nsg2	1.37	62.8	45.9	Neuron-specific gene family member 2
NM_013805	Cldn5	0.37	2.49	6.73	Claudin 5
NM_001018013	Izumo1	0.80	4.93	6.16	Izumo sperm-egg fusion 1
NM_178618	Fam83g	0.58	2.33	4.02	Family with sequence similarity 83, member G
NM_001033227	Slc5a10	1.32	5.10	3.86	Solute carrier family 5, member 10
NM_011627	Tpbg	0.59	2.19	3.71	Trophoblast glycoprotein
NM_010738	Ly6a	21.9	78.5	3.58	Lymphocyte antigen 6 complex, locus A
NM_010218	Fjx1	0.70	2.38	3.40	Four jointed box 1
NM_001159748	Serpinb8	0.90	3.02	3.36	Serine (or cysteine) peptidase inhibitor, clade B, member 8
NM_007388	Acp5	0.75	2.41	3.21	Acid phosphatase 5, tartrate resistant
Genes downregulated in Tg HSCs					
NM_181596	Retnlg	8.31	1.23	0.15	Resistin-like y
NM_008611	Mmp8	1.87	0.28	0.15	Matrix metallopeptidase 8
NM_008405	ltgb2l	2.97	0.55	0.19	Integrin β2-like
NM_145143	Mpp4	2.80	0.66	0.24	MAGUK p55 subfamily member 4
NM_021407	Trem3	4.36	1.07	0.25	Triggering receptor expressed on myeloid cells 3
NM_013599	Mmp9	4.20	1.04	0.25	Matrix metallopeptidase 9
NM_009639	Crisp3	2.51	0.63	0.25	Cysteine-rich secretory protein 3
NM_011410	Slfn4	1.08	0.28	0.26	Schlafen 4
NM_145389	BC016579	4.44	1.20	0.27	cDNA sequence, BC016579
NM_178885	AI182371	1.36	0.37	0.27	Expressed sequence AI182371

The values indicate reads per kilobase of transcript per million mapped reads and the Tg vs control.

significantly higher in Tg mice than in the controls, whereas the HSC subpopulations (long-term [LT] HSC, short-term [ST] HSC, and multipotent progenitor [MPP]) were apparently not affected by the overexpression of *Fbx110* (Figure 2A).

Subsequently, to assess the specific activation of the cell cycle of HSCs, we examined the ratio of 5-bromo 2'-deoxyuridine (BrdU)positive cells in the BM after the injection of BrdU into mice combined with pyronin Y staining. The ratio of BrdU-incorporated LT-HSCs in Tg mice was higher than that in the controls (Figure 2B), whereas pyronin Y analysis demonstrated no significant difference of G0/G1 ratio between Tg and control mice (Figure 2C). Thus, the transgenic expression of Fbxl10 in HSCs accelerated the progression of cell cycle accompanying a normal G0 to G1 entry, resulting in pleiotropic progenitor expansion. In addition, peripheral blood chimerism in competitive and serial transplantation analyses showed that whereas no difference was observed in the primary transplantation, the secondary transplantation exhibited a significant increase in the long-term repopulating activity of Fbxl10 Tg BM cells as compared with control BM cells (Figure 2D). Tg mouse-derived transplanted cells strongly contributed to myeloid and B-cell lineages but not to T-cell lineage (Figure 2D), which is in good agreement with the lineage specificity of Fbxl10-induced leukemia (Figure 1C; supplemental Table 1). The percentage of donor-derived cells was slightly higher in the BM recipients that received Fbxl10 Tg cells compared with those that received control cells, although the difference was not statistically significant at the time point (supplemental Figure 4C). These results suggest that Fbx110 enhances self-renewal activity and a long-term BM repopulation capacity with a lasting myeloid and B-lymphoid expansion in the peripheral blood in a cell-autonomous manner.

Nsg2, a neuron-specific gene aberrantly induced by FbxI10, impairs hematopoietic differentiation in vitro

To elucidate the mechanistic link between the overexpression of *Fbxl10* and development of leukemia, we analyzed the transcriptional

profiles of IL-7R α^{-} LSK HSCs in nonleukemic young Tg mice (8 weeks of age) together with the corresponding controls (Table 1). A remarkable observation was the markedly increased expression of Nsg2 (also known as HMP19), encoding a 19-kDa protein localized in the Golgi apparatus,²¹ in Tg cells (Figure 3A; Table 1). We found that the levels of Nsg2 mRNA were low in control HSCs (Table 1), whereas a 40-fold higher expression was observed in the Tg HSCs (Figure 3B). A significantly higher expression of Nsg2 was also detected in most of the Fbxl10 Tg leukemic tissues of myeloid lineage (Figure 3C), suggesting that Nsg2 may be implicated in the pathogenesis of leukemia. To assess this possibility, we performed small interfering RNA (siRNA) knockdown in cells prepared from a Tg leukemic spleen diagnosed as AML exhibiting high Nsg2 expression (mouse no. 8 in Figure 3C and supplemental Table 1) (supplemental Figure 5A). siRNA oligos were successfully introduced into these cultured leukemic cells by electroporation (supplemental Figure 5B), and efficient Nsg2 knockdown was verified by quantitative real-time polymerase chain reaction (PCR) (quantitative PCR [qPCR]) (supplemental Figure 5C). As a result, siRNA-mediated Nsg2 knockdown in the leukemic cells significantly decreased the colony number and cell viability in methylcellulose culture (supplemental Figure 5D-E).

Next, to evaluate the impact of the aberrant expression of *Nsg2* on normal hematopoiesis, WT BM cells were infected with retroviruses expressing *Nsg2*. The overexpression of *Nsg2* in murine primary WT c-Kit⁺ BM cells increases the proportion of c-Kit⁺ hematopoietic stem/progenitor cells and LSK cell fraction compared with the cells expressing empty vector in short-term ex vivo liquid culture (Figure 3D-E). In addition, colony numbers of *Nsg2*-transduced control cells were significantly higher than those of vector-transduced control cells through 3 rounds of plating, and were almost comparable to those of vector-transduced Tg cells after the first and second platings (Figure 3F). The *Nsg2*-transduced cells formed fewer colonies than vector-transduced Tg cells at the third round, but the colonies retained their concentric and compact morphology (Figure 3F-G). These results suggest that *Nsg2* potentially



Figure 3. *Nsg2* is highly upregulated in HSCs and leukemic tissues of Tg mice and impairs hematopoietic differentiation. (A) Scatter plot comparing the transcriptomes of control and Tg IL-7R α^- LSK HSCs. Transcription levels are shown on a logarithmic scale. A linear regression curve and correlation coefficient are indicated. (B) *Nsg2* mRNA in IL-7R α^- LSK HSCs was measured by real-time quantitative RT-PCR, and is presented as the relative expression level in arbitrary units where the mRNA level in the control was defined as 1 unit. (C) Comparison of *Nsg2* mRNA levels in IL-7R α^- LSK HSCs from the control and Tg BM, and in the control normal spleen as well as leukemic spleens from Tg mice. *Nsg2* mRNA was measured and shown as in panel B. Among leukemic spleens, gray and black bars indicate the leukemic tissues that were diagnosed with AML and B-ALL, respectively. *Unclassified samples. (D) Murine primary c-Kit⁺ BM cells were retrovirally transduced with pMYs-IRES-EGFP empty vector or the vector expressing *Nsg2*. Cells were grown in liquid culture with SCF, IL-3, IL-6, GM-CSF, TPO, and FIt3L for 5 days and assessed for the percentage of c-Kit⁺ population in EGFP^{high} cells. (Left) Flow cytometric profiles. (Right) The ratio of c-Kit⁺ cells (percentage of EGFP⁺ cells). (E) Cells treated as in panel D were assessed using flow cytometry for the percentage of LSK in total Lin⁻EGFP^{high} cells. (Left) Flow cytometric profiles. (Right) The ratio of LSK cells (percentage of EGFP⁺ cells). All results are shown as the mean \pm SD of independent experiments; ***P* < .005. (F) c-Kit⁺ BM cells from the control and Tg littermates were retrovirally transduced with pMYs-IRES-EGFP empty vector. c-Kit⁺ BM cells from control mice were retrovirally transduced with the vector expressing *Nsg2*. EGFP⁺ cells were sorted and then cultured in methylcellulose media as described in "Methods." Colonies were counted and replated at 2 × 10⁴ cells every 7 days. Error bars, SD. ***P* < .005. (G) Representative

possesses undifferentiated properties and may contribute to leukemogenesis in a manner similar to canonical class II mutations. We further transplanted the WT c-Kit⁺ BM progenitor cells overexpressing *Nsg2* into irradiated mice. These recipients, however, did not show leukemia at least during the observation period of 8 months.



Figure 4. *Fbx110* overexpression leads to mitochondrial metabolic activation characterized by increased oxidative phosphorylation. (A) GSEA enrichment plot of KEGG OXPHOS gene set in nonleukemic (preleukemic) BM HSCs (HSCs from Tg mice vs HSCs from controls). Genes involved in OXPHOS were significantly enriched in Tg HSCs. Pooled RNAs from L-7Ra⁻ LSK HSCs in Tg and control individuals were used for the analysis. (B) GSEA enrichment plot of KEGG OXPHOS gene set in leukemic spleens (Tg leukemic spleens vs control spleens). Genes involved in OXPHOS were significantly enriched in Tg leukemic spleens vs control spleens). Genes involved in OXPHOS were significantly enriched in Tg leukemic spleens. (C) Heat map of OXPHOS genes. Genes involved in OXPHOS are profoundly highly expressed in both pooled Tg HSCs and individual Tg leukemic spleens compared with the corresponding WT control HSCs and spleens. (D) Relative ATP levels in IL-7Ra⁻ LSK HSCs of control and Tg mice at 20 weeks of age (n = 6 per group). Horizontal line, mean of ATP levels. Tg cells produced significantly more ATP than control cells (*P = .032). FDR, false discovery rate; NES, normalized enrichment score.

Fbxl10-overexpressing HSCs are metabolically active and characterized by an increased oxidative phosphorylation

Previous studies have reported that Fbx110 suppresses senescenceassociated cyclin-dependent kinase inhibitors (CDKIs) such as *p15ink4b*, *p16ink4a*, *p18ink4c*, and *p57kip2* in murine blood cells.^{12,13} Therefore, we investigated the possibility that suppression of these CDKI genes might account for the survival/proliferation advantage in Tg HSCs, which is reminiscent of the canonical class I mutations. The former 2 genes, *p15ink4b* and *p16ink4a*, were very weakly expressed in primary control HSCs, as reported in a previous transcriptome analysis²²; no expressional change of these genes was observed in Tg HSCs. The latter 2 genes, *p18ink4c* and *p57kip2*, were expressed in both genotypes and slightly suppressed by 20% to 45% in Tg HSCs as compared with wild-type cells (supplemental Table 2). Although the suppression of *p57kip2* was most remarkable, this would not account for the enhanced self-renewal activity of *Fbx110* Tg HSCs because a previous report showed that *p57kip2* was required for HSC quiescence and its deficiency abrogated the self-renewal capacity of HSCs in mice.²³

Table 2. Gene sets enriched by FbxI10 in HSCs on GSEA (NES > 2.0)

Pathway	Size*	NES	Nominal P	FDR q value
Upregulated gene set in Tg HSCs				
Oxidative phosphorylation	98	2.50	.000	0.000
Ribosome	56	2.39	.000	0.000
Proteasome	22	2.13	.000	0.001
Downregulated gene set in Tg HSCs				
Net explicable				

Not applicable

FDR, false discovery rate; NES, normalized enrichment score. *Number of genes in each set.

To identify the differentially regulated functional networks accounting for growth advantage of HSCs in Tg mice, we applied gene set enrichment analysis (GSEA) to the Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway, a database that represents molecular interaction networks, including metabolic pathways, regulatory pathways, and molecular complexes. This analysis revealed that in nonleukemic Tg HSCs, several gene sets were significantly upregulated with "oxidative phosphorylation (OXPHOS)" being the highest (Figure 4A; Table 2), whereas no gene sets were significantly downregulated, as compared with the control HSCs. The genes involved in OXPHOS were also most markedly enriched in the leukemic tissues compared with the corresponding controls (Figure 4B-C; Table 3). Adenosine 5'-triphosphate (ATP) is the central cellular metabolic marker and bioenergetic source.^{24,25} We observed a significant increase in basal intracellular ATP content in IL-7R α ⁻ LSK HSCs purified from nonleukemic Fbxl10 Tg mice compared with the corresponding controls (Figure 4D). These results strongly suggest that Fbx110 metabolically activates HSCs by increasing OXPHOS, which is linked to the progression of cell cycle and promotes the development of leukemia.

Fbxl10 has direct and indirect effects in proleukemogenic gene expression profiles

To investigate the regulatory mechanism of the proleukemogenic transcription signature caused by the overexpression of Fbx110, we performed chromatin immunoprecipitation sequencing (ChIP-seq) in Tg and control HSCs (LSKs) using a ChIP-grade polyclonal antibody for Fbx110.9 Recent studies demonstrated that Fbx110 preferentially binds to the regions flanking transcription start sites (TSSs) and containing unmethylated CpG sequences at actively transcribed genes but not only repressive genes in ES cells.⁹ Tag density of ChIP-seq data revealed 3 distinct regions flanking the TSS of the Nsg2 gene, in which additional bindings of Fbx110 were detected in Tg HSCs (boxed in Figure 5A). CpG sequences associated with an Fbxl10 binding region (+1991 to +2190 of Nsg2 gene locus) were detected using the sequence manipulation suite program²⁶ (Figure 5B). We also identified increased or additional peaks on OXPHOS-related Ndufs6, Ndufb2, and Uqcrfs1 gene regions flanking the TSS in Tg HSCs (Figure 5C). However, the bindings of Fbx110 were not pronouncedly increased at the TSS and promoter regions of most other OXPHOS genes that were upregulated in Tg HSCs. The activation of OXPHOS genes in Tg HSCs may involve overall effects triggered by the transgenic expression of Fbxl10. These results indicate direct and indirect regulation of proleukemic gene expression signature mediated by Fbx110. Because Fbx110 is an H3K36me2 demethylase, we also performed the ChIP-seq for HSCs using an anti-histone H3K36me2 antibody shown to be applicable for the ChIP-seq.²⁷ The result revealed that Fbx110 overexpression in HSCs reduced H3K36me2 levels more around TSSs (supplemental Figure 6A-B). The H3K36me2 ChIP-seq reads were, however, detected at low levels on Fbx110-targeted regions of Figure 5,

panels A and C, in control HSCs, and therefore, it was unable to clarify the corresponding decrease in reads in Tg HSCs.

Next, to assess the regulatory status in transformed cells, we performed ChIP-PCR to analyze the recruitment of Fbx110 to the *Nsg2* gene regions in proliferating myeloid leukemic cells in culture from a Tg spleen, using anti-Fbx110 and anti-Flag Abs (primer sets, P1-P3 in supplemental Figure 7A). A primer set P3 detected a significant binding of Fbx110 on the *Nsg2* gene region, corresponding to regions that were identified in Tg LSK by the ChIP-seq with Fbx110 Ab (supplemental Figure 7B). In addition, the primer set was able to detect reduced DNA binding for H3K36me2 in this tumor line (supplemental Figure 7C). These results suggest that the ectopic expression of *Nsg2* in Tg HSCs is targeted by the overexpression of Fbx110 and that the recruitment to Nsg2 gene locus is partly sustained in the resulting leukemic cells.

A subpopulation of AML patients exhibits high expression of both *HMP19* and *FBXL10*

To investigate the potential relevance of the findings to human disease, we investigated the expression levels of *HMP19* (human homolog of *Nsg2*) and *FBXL10* on The Cancer Genome Atlas (TCGA) database.²⁸ We found that *HMP19* mRNA was strongly elevated in 21 cases (12.8%) by more than fivefold (RPKM > 0.05) of the whole AML median value (0.01), whereas 82 of 179 cases (45.8%) showed no expressions of the gene (Figure 6A). Eight cases (4.5%) exhibited even more than 10-fold upregulation (RPKM > 0.1) of the *HMP19* gene expression. Of note, 6 cases (3.4%) that exhibited a higher expression of *FBXL10* over the median value were coupled with more than fivefold upregulation of the *HMP19* gene expression (Figure 6). These data may support that our mouse model is clinically relevant with a specific population of leukemia patients.

Discussion

Genetic and functional studies have so far suggested that there are at least 2 classes of myeloid leukemia disease alleles that contribute to hematopoietic transformation: class I alleles, which confer a growth advantage, and class II alleles, which impair the differentiation.²⁹ However, recent studies have demonstrated that mutations of single genes in epigenetic modifiers can contribute to myeloid malignancies, suggesting another class of somatic functional mutations or altered gene expression besides the canonically categorized class I/class II mutations (*TET2, ASXL1, DNMT3A, EZH2, MLL*, etc).²⁹ In this study, we used a genetically engineered mouse system to show that the Tg expression of epigenetic regulator Fbx110 induced acute leukemias (Figure 1).

Mechanistically, Fbx110 confers HSCs on the progression of cell cycle through the metabolic activation, whereas Fbx110 activated the expression of *Nsg2*, which predisposes hematopoietic stem/early

Table 3. Gene sets enriched by Fbxl10 in leukemic tissues on GSEA (NES > 2.5)

Pathway	Size	NES	Nominal P	FDR q value
Upregulated gene set in Tg I	leukemic	tissues		
Oxidative phosphorylation	94	3.56	.000	0.000
Ribosome	56	2.56	.000	0.000
Proteasome	22	2.50	.000	0.000
Downregulated gene set in 1	ſg leuken	nic tissue	S	
Not applicable				

Size, number of genes in each set.

FDR, false discovery rate; NES, normalized enrichment score.



Figure 5. Nsg2 and mitochondrial genes are directly targeted by FbxI10. (A) FbxI10 ChIP-seq signal at Nsg2 gene regions flanking promoter and TSS in HSCs. Rectangles indicate regions where FbxI10 bindings are higher in Tg HSCs than in control HSCs (left) and the bar graph demonstrates relative ChIP/Input ratio [ChIP DNA (ppm) divided by Input DNA (ppm)] in the indicated regions (right). (B) CpG sequences detected in Nsg2 gene regions flanking promoter and TSS, which was sought using sequence manipulation suite program.26 Fbx110 was highly recruited to the region +1991 to +2190 of Nsg2 gene in Tg HSCs, as shown in panel A. (C) Fbxl10 ChIP-seg signal at OXPHOS genes (Ndufs6, Ndufb2, and Uqcrfs1) regions flanking promoter and TSS. Rectangles indicate regions where Fbxl10 bindings are higher in Tg HSCs than in control HSCs (left) and the bar graphs demonstrate relative ChIP/Input ratio in the indicated regions (right)

progenitor cells to an undifferentiated state. Thus, elevated levels of Fbx110 involve both class I– and class II–like properties (Figure 7). However, the mice did not show an evident HSC differentiation block at a steady state, and the development of leukemia in Tg mice required long latency. Although these properties may underlie tumor susceptibility, additional genetic event(s) and/or microenvironmental effects may be required for full transformation (Figure 7). This idea was validated by the results that leukemogenesis in Tg mice is drastically accelerated by retrovirus insertional mutagenesis or a carcinogenic mutagen (Figure 1D-E).

Although the biological function of Nsg2 has not been fully understood so far, structurally similar transmembrane molecules NEEP21 (also known as Nsg1) and Caly (also known as Nsg3) are crucial for glutamate and/or dopamine receptor sorting through endosomes and their recruitment to the plasma membrane in neural cells.³⁰⁻³² Because the expression of Nsg2 is severely repressed in normal HSCs, its ectopic upregulation might deregulate cytokine receptor recycling/intracellular sorting, thereby resulting in impaired differentiation of HSCs (Figure 3D-F). Wu et al described Fbx110 binding loci in WT and Fbx110-overexpressed ES cells.9 Although the binding loci of Fbx110 were entirely shared between these 2 cell types, Fbx110 was bound to the Nsg2 locus specifically in the overexpressed condition.9 Likewise, our ChIP-seq for HSCs and ChIP-PCR for the Tg leukemic cells demonstrated that Nsg2 was targeted by the overexpression of Fbxl10 (Figure 5A; supplemental Figure 7). To examine the oncogenic activity of Nsg2, we transplanted BM c-Kit⁺ cells retrovirally overexpressing Nsg2 into irradiated mice. However, these recipients did not develop leukemia during the limited period of the 8 months, suggesting that overexpression of Nsg2 alone is not leukemogenic.

Another remarkable characteristic of Tg HSCs is metabolic proliferation advantage coupled with mitochondrial energy activation (Figures 2B and 4A,C-D). This feature is also retained in leukemic tissues (Figure 4B), indicating that this change plays a critical role in the leukemogenic process. Accumulating evidence has shown that tumor cells increase their metabolic requirements for cell survival and proliferation.³³ In addition, clinical studies also have provided evidence of a close relationship between altered cellular metabolism and the



Figure. 6. A subpopulation of AML patients exhibits high expression of both *HMP19* and *FBXL10*. RNA-Seq data of 179 AML cases available in the TCGA database were analyzed. Expression levels of *HMP19* (A) and *FBXL10* (B) mRNAs are shown as RPKM. Six that expressed HMP19 mRNA more than fivefold over the average and *FBXL10* mRNA over the median value are indicated by red squares (TCGA-AB-2807, 2832, 2892, 2943, 2982, and 3009). RPKM, reads per kilobase per million mapped reads.

Figure 7. Schematic illustration of Fbx110-mediated leukemogenic processes. Fbx110 induces leukemia involving metabolic proliferation advantage and impaired differentiation mediated by Nsg2. Fbx110 confers HSCs on cell cycle progression through metabolic activation whereas Fbx110 upregulates the expression of the *Nsg2*, impairing the differentiation of HSCs. Thus, Fbx110 profoundly enhances leukemogenic potential/tumor susceptibility and leads to acute leukemia in mice after a long latency. Additional gene alterations and/or microenvironmental effects possibly intensify the proliferative potential and accelerate leukemic transformation mediated by Fbx110 overexpression.



cancer outcome.³⁴ Reactive oxygen species (ROS), which are generated by mitochondrial oxidative phosphorylation, was reported to limit the life span of HSCs.³⁵ However, mitochondrial ROS levels in HSCs were not increased before the development of disease by the overexpression of Fbxl10, as examined by flow cytometry using a fluorogenic molecular probe (supplemental Figure 8). Consistent with these findings, cell cycle analysis demonstrated that Tg HSCs had a growth advantage without exhausting the HSC pool (Figure 2B-D). In mouse embryonic fibroblasts, the overexpression of Fbxl10 was reported to detoxify ROS by directly upregulating antioxidative genes and contributes to the inhibition of cellular senescence.³⁶ Our results suggest that Fbx110 has a substantially similar effect on HSCs, although the gene expression of individual canonical ROS scavenger was not significantly altered.³⁷ Therefore, we cannot unequivocally state the mechanism underlying the lack of ROS increase in Tg HSCs at this point, and the possible involvement of Fbxl10 to maintain redox homeostasis in the HSC unique context remains to be defined. A recent study demonstrated that leukemic stem cells show relatively low levels of ROS and depend on oxidative respiration rather than glycolysis for energy generation, which may support our results.³⁸ In addition, GSEA revealed that genes involved in ribosomal biogenesis were also enriched in both Tg HSCs and leukemic tissues (Tables 2-3), which would provide a direct connection between metabolic activation and the development of tumor because increased protein synthesis is a highly energy demanding process.

Fbx110 was recently reported to promote *K-RAS*–mutated pancreatic cancer through upregulating a module of metabolic genes involved in OXPHOS in a mouse allograft model.⁵ In addition, *IMP2*-positive glioma and *PPARGC1A* (*PGC1a*)–positive melanoma showed tumor progression associated with mitochondrial energy activation in the absence and presence of enhanced ROS levels, respectively.^{39,40} Thus, despite the general enhancement of glycolytic bioenergetic requirements in many cancers (so called as "Warburg effect"), several recent studies have revealed unanticipated tumor metabolism by Warburg, that is, the bioenergetic type of tumors can vary widely from glycolysis to OXPHOS.⁴¹⁻⁴⁴ Our findings suggest that OXPHOS-mediated metabolic activation provides a bioenergetic source to promote leukemogenesis in *Fbxl10*-upregulated HSCs. However, a number of questions remain to be answered, including the lineage specificity of leukemia and the possible involvement of other factors in *Fbxl10*-induced leukemogenesis. The search for these effectors will no doubt contribute to elucidating the mechanism by which overexpressed Fbxl10 leads to leukemia.

In summary, we established *Fbxl10* as a bona fide oncogene in vivo and provided a novel mechanistic insight into leukemogenesis. Taken together with the upregulation of *FBXL10* in human leukemia and solid tumors,³⁻⁶ selective inhibition of FBXL10 may be a therapeutic strategy to achieve improvement in these diseases.

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Authorship

Contribution: T.U., H.H., Z.-i.H., T.S., T.I., and A.I. designed the research; T.U., A.N., K.T., N.Y., Y.N., K.I., H.M., A.K., T.K., H.O., and H.H. performed the research; L.W., X.W., and K.H. contributed new reagents or analytic tools; T.U., K.T., H.H., H.M., A.K., T.K., and A.I. analyzed the data; and T.U. and H.H wrote the manuscript.

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