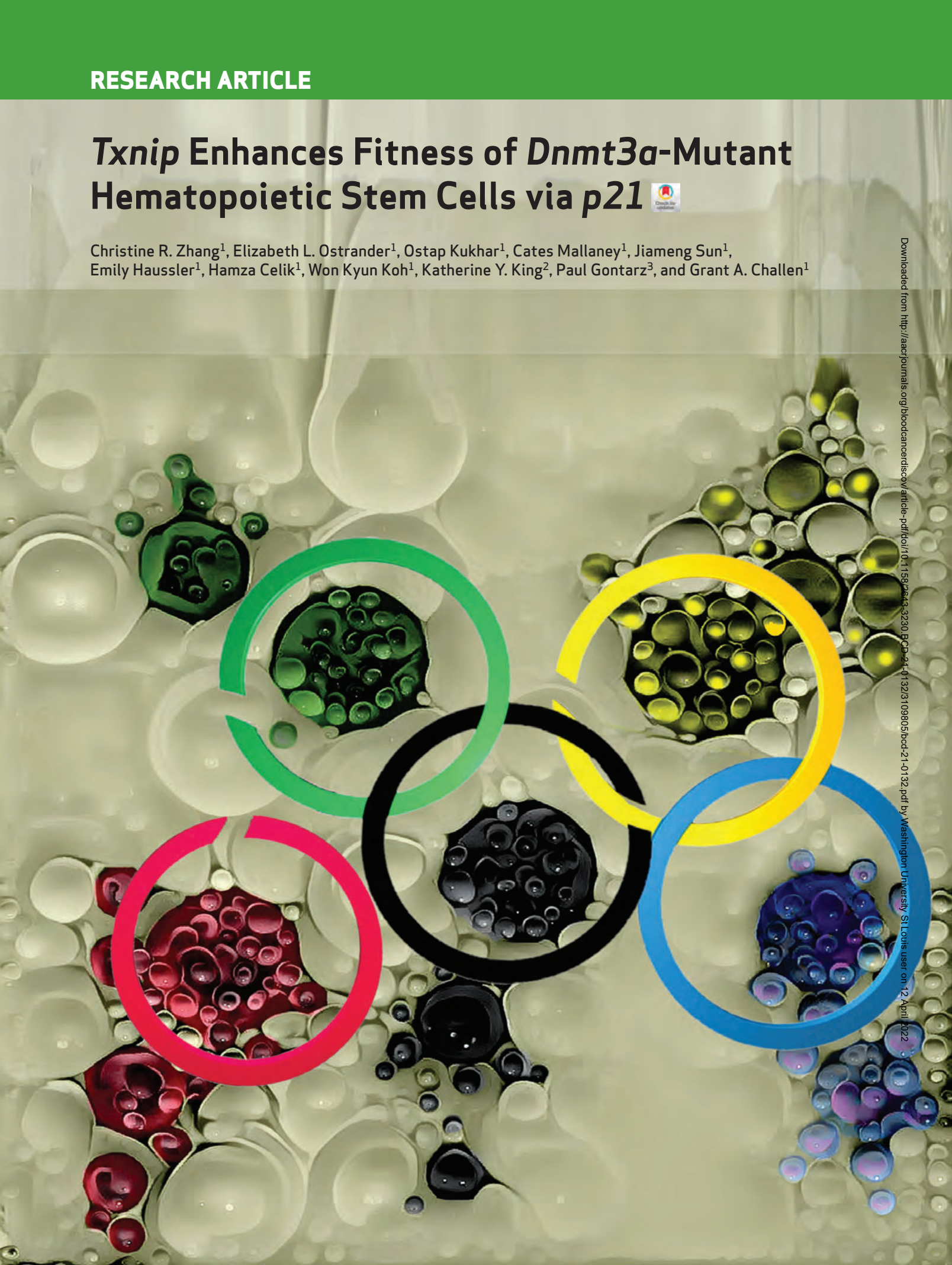


Txnip Enhances Fitness of *Dnmt3a*-Mutant Hematopoietic Stem Cells via *p21*

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ABSTRACT

Clonal hematopoiesis (CH) refers to the age-related expansion of specific clones in the blood system, and manifests from somatic mutations acquired in hematopoietic stem cells (HSCs). Most CH variants occur in the gene *DNMT3A*, but while *DNMT3A*-mutant CH becomes almost ubiquitous in aging humans, a unifying molecular mechanism to illuminate how *DNMT3A*-mutant HSCs outcompete their counterparts is lacking. Here, we used interferon gamma ($\text{IFN}\gamma$) as a model to study the mechanisms by which *Dnmt3a* mutations increase HSC fitness under hematopoietic stress. We found *Dnmt3a*-mutant HSCs resist $\text{IFN}\gamma$ -mediated depletion, and $\text{IFN}\gamma$ -signaling is required for clonal expansion of *Dnmt3a*-mutant HSCs *in vivo*. Mechanistically, DNA hypomethylation-associated overexpression of *Txnip* in *Dnmt3a*-mutant HSCs leads to p53 stabilization and upregulation of *p21*. This preserves the functional potential of *Dnmt3a*-mutant HSCs through increased quiescence and resistance to $\text{IFN}\gamma$ -induced apoptosis. These data identify a previously undescribed mechanism to explain increased fitness of *DNMT3A*-mutant clones under hematopoietic stress.

SIGNIFICANCE: *DNMT3A* mutations are common variants in clonal hematopoiesis, and recurrent events in blood cancers. Yet the mechanisms by which these mutations provide hematopoietic stem cells a competitive advantage as a precursor to malignant transformation remain unclear. Here, we use inflammatory stress to uncover molecular mechanisms leading to this fitness advantage.

See related article by De Dominici and James DeGregori.

INTRODUCTION

Hematopoietic stem cells (HSCs) randomly acquire somatic mutations with age (1). The vast majority of these mutations are inconsequential, but some variants provide a fitness advantage to HSC clones, allowing them to become disproportionately represented in the blood (2). This state of abnormal HSC expansion without overt hematologic disease is known as clonal hematopoiesis (CH), and is associated with increased risk of cardiovascular events and blood cancers (3–5). Approximately 50% of all genetic variants in CH occur in the gene *DNMT3A* (6), which encodes a *de novo* DNA methyltransferase enzyme responsible for establishing new DNA methylation patterns during development and stem cell fate decisions (7, 8). *DNMT3A* is also recurrently mutated in blood diseases such as acute myeloid leukemia (AML; ref. 9) and myelodysplastic syndromes (MDS; ref. 10) where it functions as an initiating mutation (11). Emerging evidence indicates somatic *DNMT3A* variants can be acquired early in life (12, 13), before becoming almost ubiquitous in the aging population (14).

Mouse models show *Dnmt3a* loss-of-function skews HSC fate decisions toward self-renewal, facilitating clonal

expansion in the bone marrow (BM; ref. 15, 16). However, analysis of DNA methylation and gene expression patterns in mouse (15, 17) and human (17, 18) systems have yet to identify a unifying molecular mechanism to explain the increased fitness of these mutant HSCs. The competitive advantage of *Dnmt3a*-mutant HSCs is exacerbated by hematopoietic stress such as serial transplantation (19) and microbial infection (20). In addition to genetics, environmental factors likely contribute to clonal selection in CH. In particular, chronic inflammation has been proposed as an environmental pressure that selects for certain clones in CH (21, 22). We recently showed ulcerative colitis patients have a higher incidence of *DNMT3A*-mutant CH associated with increased serum $\text{IFN}\gamma$ (23). Here, we used $\text{IFN}\gamma$ as a model of inflammatory stress to study the molecular mechanisms that empower *Dnmt3a*-mutant HSCs with a fitness advantage. We found *Dnmt3a*-mutant HSCs specifically resist $\text{IFN}\gamma$ -mediated depletion, and $\text{IFN}\gamma$ signaling is required for clonal expansion of *Dnmt3a*-mutant HSCs *in vivo*. Complementary genomic techniques identified a *Txnip*-*p53*-*p21* pathway that preserves the functional potential of *Dnmt3a*-mutant HSCs under conditions of inflammatory stress, which was validated by functional genetic rescue experiments *in vivo*. These findings highlight a novel mechanistic basis to explain the increased fitness of *Dnmt3a*-mutant HSCs, and supports rationale for developing interventions to mitigate expansion of premalignant clones as a method of blood cancer prevention.

RESULTS***DNMT3A*-Mutant HSCs Are Insensitive to the Deleterious Effects of $\text{IFN}\gamma$ *In Vivo***

Mouse genetic models were generated to represent the spectrum of *DNMT3A* mutations found in human CH; *Dnmt3a* heterozygous (*Vav-Cre;Dnmt3a^{fl/+} = Dnmt3a^{HET}*) and

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Note: Supplementary data for this article are available at Blood Cancer Discovery Online (<https://bloodcancerdiscov.aacrjournals.org/>).

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homozygous (Vav-Cre;*Dnmt3a*^{fl/fl} = *Dnmt3a*^{KO}) hematopoietic loss-of-function, and a knockin model (24) analogous to the point mutation most prevalent in AML (Vav-Cre;*Dnmt3a*^{R878H/+} = *Dnmt3a*^{R878}). Most DNMT3A variants in human CH are missense heterozygous loss-of-function (6). However, we and others have shown that *Dnmt3a*^{HET} HSCs produce only modest phenotypes in mice (25, 26), likely because these variants provide very minor fitness advantages to HSCs (2) that require the extended lifespan of humans to manifest measurable phenotypes. In mice, complete loss of *Dnmt3a* function is required to generate robust phenotypes. Inclusion of the *Dnmt3a*^{R878} model (although this variant is less prevalent in CH than AML) allows for the most complete analysis of *Dnmt3a* mutations in HSC function.

To examine how *Dnmt3a* mutations impact the HSC response to IFN γ , hematopoietic chimeras were created by transplanting recipient mice (CD45.1) with 5.0×10^5 BM cells from control (Vav-Cre;*Dnmt3a*^{+/+}), *Dnmt3a*^{HET}, *Dnmt3a*^{KO}, or *Dnmt3a*^{R878} mice in competition with 5.0×10^5 BM cells from congenic wild-type (CD45.1) mice. Recipients were treated with PBS (control) or recombinant mouse IFN γ (Fig. 1A). IFN γ treatment had minimal impact on peripheral blood engraftment (Supplementary Fig. S1A and S1B). Control HSCs (Supplementary Fig. S1C) were reduced following IFN γ treatment in relation to PBS-treated mice (Fig. 1B). In contrast, *Dnmt3a*^{HET}, *Dnmt3a*^{KO}, and *Dnmt3a*^{R878} HSCs (Supplementary Fig. S1D) were resistant to depletion in response to IFN γ (Fig. 1B; Supplementary Fig. S1E). To determine the functional consequences of prior IFN γ exposure on long-term HSC function, 200 donor-derived HSCs (CD45.2⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) were purified from primary transplants and transferred to secondary recipients along with 2.5×10^5 fresh BM competitor cells. The trend toward impaired peripheral blood engraftment of IFN γ -treated control HSCs was not observed from IFN γ -treated *Dnmt3a*-mutant HSCs (Supplementary Fig. S1F). While IFN γ was detrimental to the self-renewal of control HSCs, clonal expansion of *Dnmt3a*^{HET}, *Dnmt3a*^{KO}, and *Dnmt3a*^{R878} HSCs was unaffected (Fig. 1C).

To determine whether human HSCs showed a similar phenotype, cord blood CD34⁺ cells were nucleofected with Cas9-RNPs (27) complexed with guide RNA (gRNA) targeting DNMT3A or the inert AAVS1 locus (negative control). Edited CD34⁺ cells were transplanted into NSG mice, then randomized to receive either recombinant human (hIFN γ) or vehicle (PBS; Fig. 1D). As with mice, treatment with hIFN γ did not influence peripheral blood engraftment of human cells (Supplementary Fig. S2A and S2B). Analysis of the BM (Supplementary Fig. S2C) showed the increased abundance of DNMT3A-targeted HSCs was not influenced by IFN γ treatment (Fig. 1E). As the self-renewal phenotype of murine *Dnmt3a*-mutant HSCs is exacerbated by serial transplant, we attempted to perform secondary transfer of cord blood cells that has not previously been reported. We found transfer of 1×10^6 hCD34⁺ cells from primary recipients via intratibial injection was required for robust engraftment in secondary recipients. Again, while peripheral blood engraftment was not different between genotypes (Supplementary Fig. S2D), the trend toward reduced self-renewal of IFN γ -treated AAVS1-targeted HSCs was not observed in IFN γ -treated DNMT3A-targeted HSCs (Fig. 1F).

Analysis of CRISPR genome edit variant allele frequency (VAF) revealed that DNMT3A mutations maintained their clone size, whereas AAVS1 mutations were depleted over serial transplantation (Supplementary Fig. S2E) and by IFN γ treatment (Fig. 1G). Cumulatively, these data suggest that *Dnmt3a*-mutant HSCs are specifically resistant to IFN γ -mediated depletion.

DNMT3A-Mutant HSCs Are Resistant to IFN γ -Induced Apoptosis

One of the major mechanisms of inflammation-mediated attrition of HSCs is sensitization to stress-induced apoptosis (28). To examine this, mice were injected with two doses of recombinant cytokines 24 hours apart. HSCs were purified 2 hours after the second dose, then apoptosis was examined immediately after purification (fresh) or after a 16-hour culture period. All HSC genotypes express comparable levels of cell surface IFN γ R1 (Fig. 2A) and showed equivalent activation of the IFN γ target gene *Stat1* (Fig. 2B). Thus, any differences should not result from the inability of the mutant HSCs to sense or respond to IFN γ . There were no differences in apoptosis (cleaved Caspase 3/7 activity) between freshly isolated HSCs of any genotype (Fig. 2C). While IFN γ -treated control HSCs showed significantly elevated apoptosis after culture, this response was absent in *Dnmt3a*-mutant HSCs (Fig. 2D). Moreover, the apoptotic resistance phenotype of *Dnmt3a*-mutant HSCs to IFN γ was not observed after exposure to related proinflammatory cytokines (Fig. 2E). These results demonstrate that *Dnmt3a* mutations render HSCs specifically insensitive to IFN γ -induced apoptosis.

IFN γ Signaling Is Required for Clonal Expansion of *Dnmt3a*-Mutant HSCs

To study the cell-intrinsic role of IFN γ -signaling in HSC function, *Dnmt3a*-mutant mice were crossed to mice lacking the IFN γ receptor (*IFN γ R1*^{-/-} = *IFN γ R1*^{KO}). In unperturbed adult (~12-week old) mice, genetic loss of *IFN γ R1* normalized the increased abundance of *Dnmt3a*-mutant HSCs in the BM (Supplementary Fig. S3A and S3B). HSC function was assessed by competitive transplantation of 200 phenotypically defined (Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) HSCs against 2.5×10^5 congenic BM cells (Fig. 3A). While loss of *IFN γ R1* had no effect on blood contribution from control HSCs, *Dnmt3a*^{KO} and *Dnmt3a*^{R878H} HSCs lacking *IFN γ R1* showed significantly reduced myeloid cell output (Fig. 3B). Loss of *IFN γ R1* also hindered lymphoid production from *Dnmt3a*^{R878H} HSCs (Fig. 3B). Analysis of recipient mice BM 18 weeks posttransplant showed loss of *IFN γ R1* ameliorated the clonal expansion of *Dnmt3a*^{KO} and *Dnmt3a*^{R878H} HSCs while having no effect on the self-renewal of control HSCs (Fig. 3C). These data show IFN γ signaling is required for the enhanced self-renewal of *Dnmt3a*-mutant HSCs. IFN γ signaling does not appear a universal CH selection mechanism as genetic deletion of *IFN γ R1* from Vav-Cre;*Tet2*^{fl/fl} (= *Tet2*^{KO}) HSCs had no effect on their competitive advantage in analogous transplantation experiments (Supplementary Fig. S3C and S3D).

If *Dnmt3a*-mutant HSCs require IFN γ signaling for clonal expansion, then reducing systemic IFN γ levels could represent a targetable approach for impeding DNMT3A-mutant

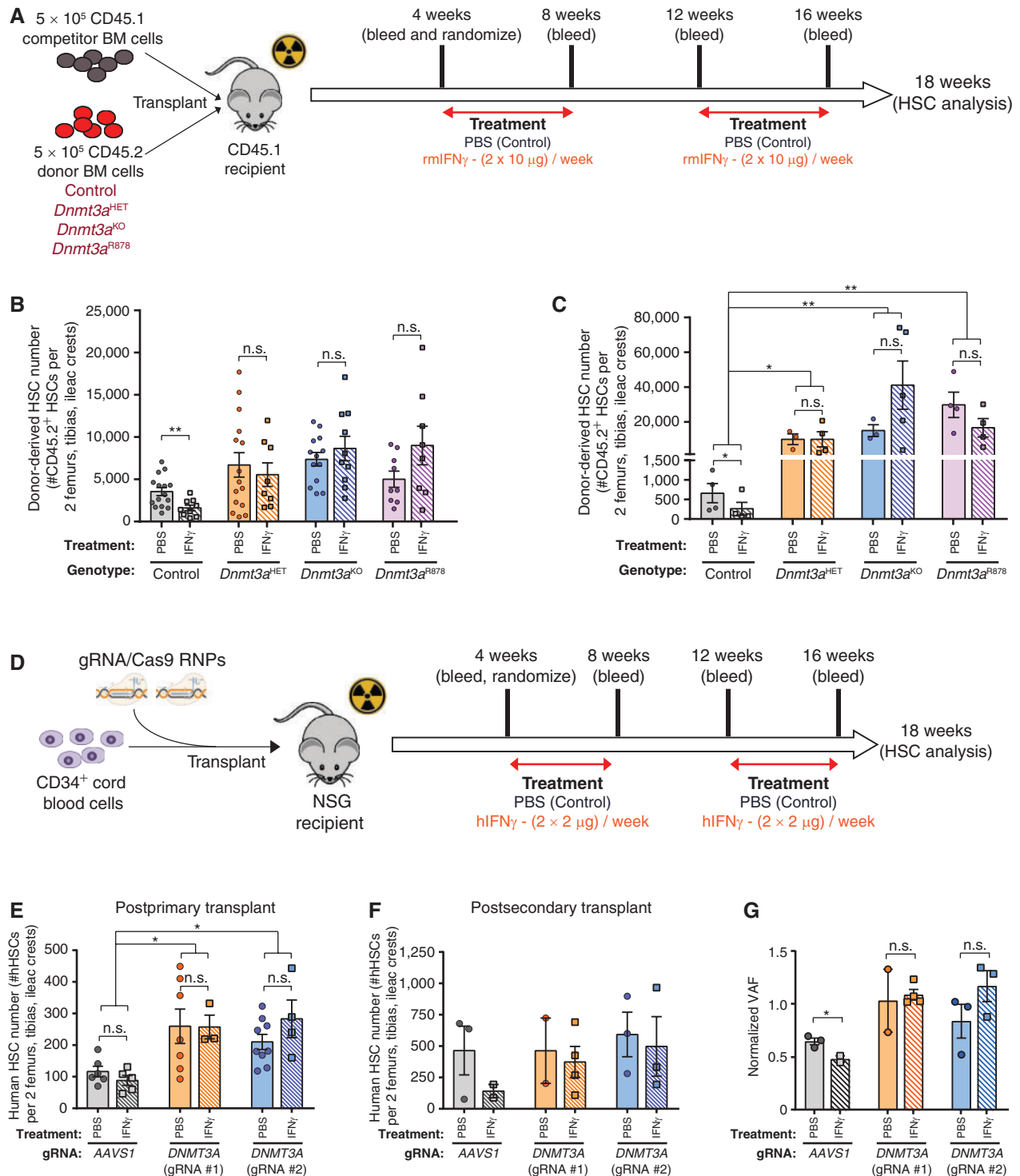


Figure 1. DNMT3A-mutant HSCs are insensitive to the deleterious effects of IFN γ *in vivo*. **A**, Schematic of competitive transplant where recipients were challenged with IFN γ . **B**, Number of donor-derived HSCs (CD45.2⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) in BM of recipients 18 weeks postprimary transplant ($n = 8-15$). **C**, Quantification of donor-derived HSCs in BM of recipients 18 weeks postsecondary transplant ($n = 3-5$). **D**, Schematic of transplantation of CRISPR-edited human CD34⁺ cord blood cells into NSG mice. **E**, Number of human HSCs (mCD45⁻hCD45⁺Lineage⁻CD38⁻CD34⁺CD45RA⁻CD90⁺CD49f⁺) in BM of NSG recipients 18 weeks posttransplant ($n = 5-9$). **F**, Number of human HSCs in BM of NSG recipients 18 weeks postsecondary transplant ($n = 2-4$). **G**, Relative clone size of CRISPR-edited human cells postsecondary transplant (hCD45⁺ BM). VAF postsecondary transplant was normalized to that of pretransplant CD34⁺ cells. One-way ANOVA; *, $P < 0.05$; **, $P < 0.01$. Data represent mean \pm SEM.

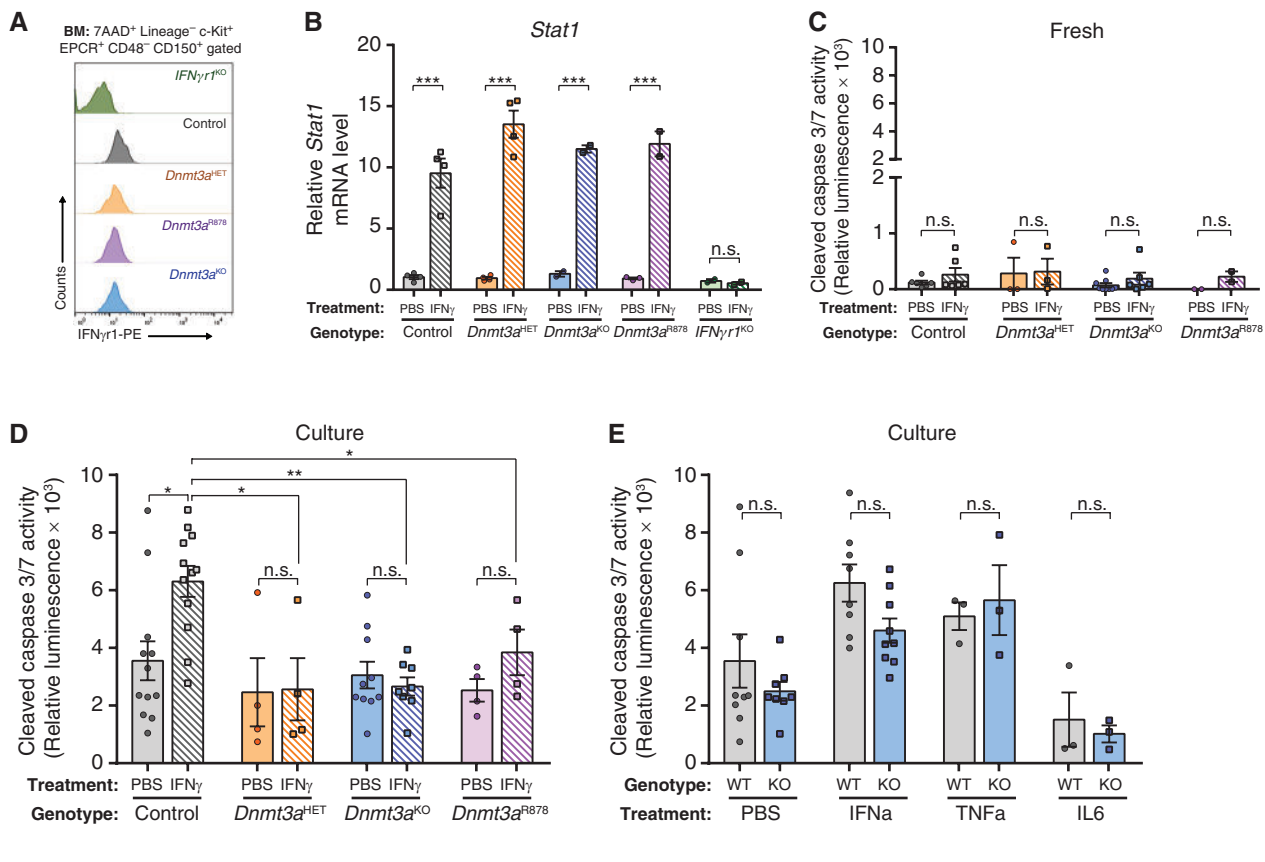


Figure 2. DNMT3A-mutant HSCs are resistant to IFN γ -induced apoptosis. **A**, Cell surface protein expression (MFI) of IFN γ 1 on HSCs (Lineage⁻ c-Kit⁺ EPCR⁺ CD48⁺ CD150⁺) measured by flow cytometry. **B**, Relative mRNA level of *Stat1* in HSCs 2-hours after *in vivo* exposure to IFN γ ($n = 2-4$) measured by qPCR. **C**, Cleaved caspase 3/7 activity of HSCs freshly purified from mice treated with PBS or IFN γ ($n = 2-9$). **D**, Cleaved caspase 3/7 activity in HSCs purified from mice treated with PBS or IFN γ after a 16-hour culture period ($n = 4-12$). **E**, Cleaved caspase 3/7 activity in HSCs (WT = Control; KO = *Dnmt3a*^{KO}) from mice treated with indicated cytokines or PBS after a 16-hour culture period ($n = 3-9$). One-way ANOVA; *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$. Data represent mean \pm SEM.

CH. To examine this, 200 control or *Dnmt3a*^{KO} HSCs were transplanted into wild-type (WT) or IFN γ -deficient (*IFN γ ^{-/-}*) mice with 2.5×10^5 BM cells matched to recipient genotype. Loss of systemic IFN γ reduced blood chimerism of both control and *Dnmt3a*^{KO} cells compared with transplantation into IFN γ -replete environments (Fig. 3D). BM analysis post-transplant showed comparable numbers of control HSCs in WT and *IFN γ ^{-/-}* recipients. Conversely, *Dnmt3a*^{KO} HSCs were significantly reduced in *IFN γ ^{-/-}* recipients (Fig. 3E), suggesting that both intrinsic IFN γ signaling and extrinsic IFN γ stimulation are required for clonal expansion of *Dnmt3a*-mutant HSCs.

Dnmt3a Mutations Confer HSCs a Fitness Advantage under Repeated IFN γ Exposure

Although the competitive advantage of *Dnmt3a*-mutant HSCs during inflammation has been implied (20, 23), no direct experiments have been performed to quantify fitness between mutant and WT HSCs in response to IFN γ . We created a novel model to examine HSC competition under chronic IFN γ exposure. Mice bearing a doxycycline-inducible IFN γ allele were crossed to *Rosa26*^{trTA-M2} mice on an *IFN γ 1^{-/-}* background (*tetO-IFN γ* ; *Rosa26*^{trTA-M2};

IFN γ 1^{KO} = “TetON-IFN γ ”), so the resultant BM can over-express IFN γ in response to doxycycline, but remain insensitive to the effects of this IFN γ overexpression and serve as a consistent internal control. Titration of doxycycline concentration in chow (Supplementary Fig. S4A) identified a dose that approximated serum levels of IFN γ in mice infected with the *Mycobacterium avium* pathogen (29). To model a 10% VAF for CH variants, chimeras were created by transplanting a 1:1:8 ratio of test BM (1.5×10^5 CD45.2 - control, *Dnmt3a*^{HET}, *Dnmt3a*^{KO}, *Dnmt3a*^{R878}), wild-type competitor BM (1.5×10^5 CD45.1/2 *Ubc*-GFP), and TetON-IFN γ or control (*Rosa26*^{trTA-M2}, *IFN γ 1^{KO}* = “TetON”) BM (1.2×10^6 CD45.1). Four weeks posttransplant, recipients were placed on weekly cycles on 1,250 ppm doxycycline chow (Fig. 3F), mimicking episodic inflammation. Chronic IFN γ exposure did not significantly alter blood chimerism (Supplementary Fig. S4B) for each donor cell genotype relative to engraftment in TetON chimeras (Supplementary Fig. S4C). HSC fitness was calculated as the ratio of test CD45.2: CD45.1/2 *Ubc*-GFP HSCs per mouse (Supplementary Fig. S4D). A ratio of 1 indicates equal competition. Control test HSCs possessed an advantage without IFN γ due to the described fitness deficit of *Ubc*-GFP HSCs (30). However, this advantage

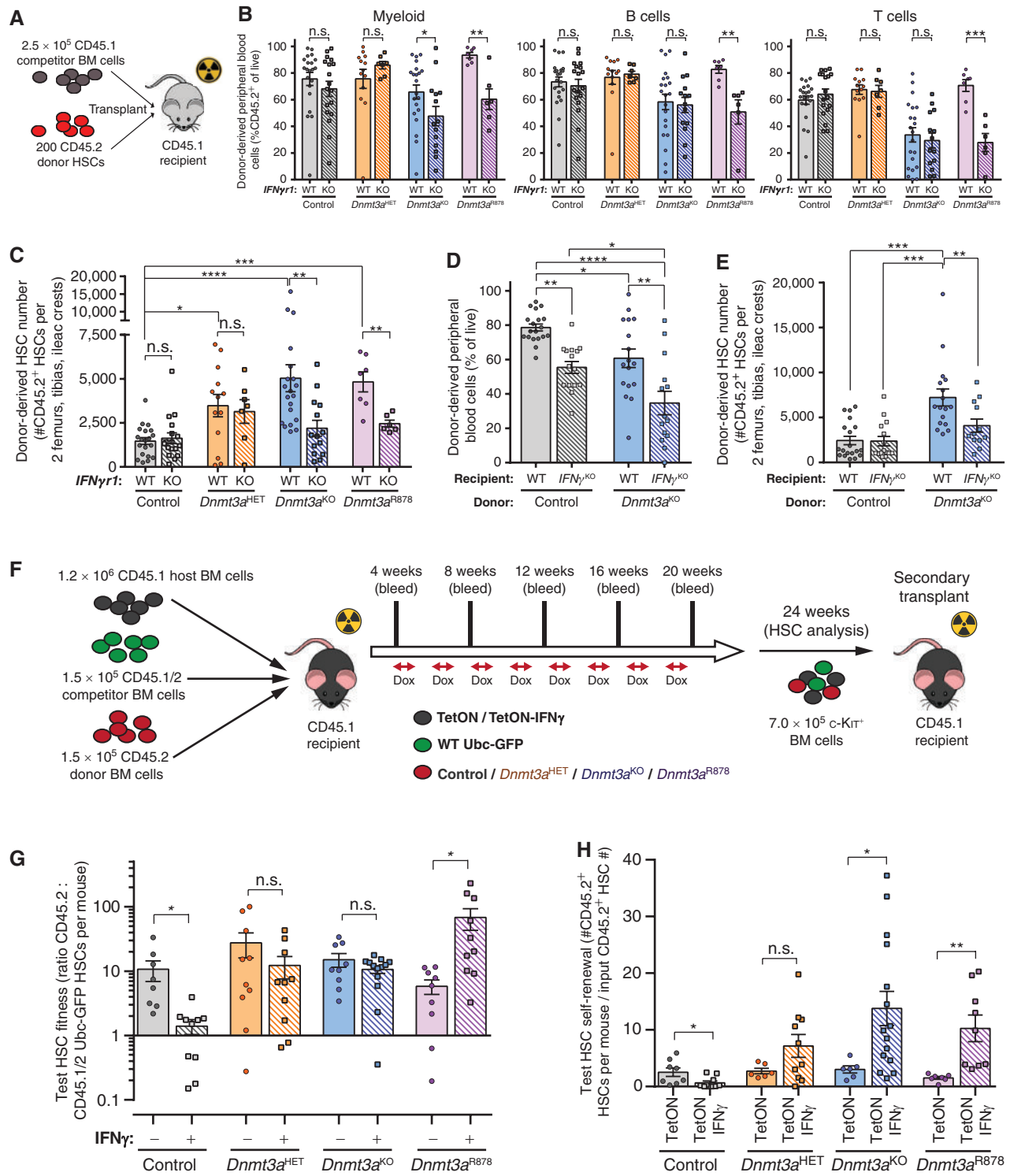


Figure 3. IFN γ signaling is required for clonal expansion of *Dnmt3a*-mutant HSCs. **A**, Schematic of competitive HSC transplantation. **B**, Donor-derived chimerism in peripheral blood lineages 16 weeks posttransplant ($n = 6-21$). **C**, Number of donor-derived HSCs (CD45.2⁺Lineage^{c-Kit}⁺EPCR⁺CD48⁻CD150⁺) in BM of recipients 18 weeks posttransplant ($n = 6-21$). **D**, Donor-derived chimerism in peripheral blood 16 weeks posttransplant in wild-type (WT) or *IFN γ* ^{KO} recipients ($n = 14-19$). **E**, Number of donor-derived HSCs in BM of recipients 18 weeks posttransplant in WT or *IFN γ* ^{KO} recipients ($n = 14-19$). **F**, Schematic showing CH competition model. **G**, Test HSC fitness ratio 24 weeks posttransplant in CH competition model ($n = 8-13$). **H**, Number of test HSCs (CD45.2⁺Lineage^{c-Kit}⁺EPCR⁺CD48⁻CD150⁺Ubc-GFP⁺) generated in BM of secondary recipients 18 weeks posttransplant per input HSC from primary transplant shown in **G** ($n = 7-15$). One-way ANOVA (**D** and **E**) or two-tailed *t* test (**B**, **C**, **G**, and **H**; data are compared for treatment/condition relative to parental genotype); *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$; ****, $P < 0.0001$. Data represent mean \pm SEM.

was diminished under chronic IFN γ (Fig. 3G). Conversely, fitness of *Dnmt3a*-mutant HSCs was generally unaffected by chronic IFN γ , and was in fact enhanced for *Dnmt3a*^{R878} HSCs (Fig. 3G). Serial transplantation was performed by transferring 7.0×10^5 c-Kit-enriched BM cells from primary recipients into secondary mice to preserve clonal composition (Fig. 3F). In secondary recipients, there was a marked reduction in blood output of control HSCs previously exposed to IFN γ (Supplementary Fig. S4E). In contrast, IFN γ -exposed *Dnmt3a*-mutant cells showed significantly increased blood engraftment in secondary transplant (Supplementary Fig. S4E), showing a fitness advantage over WT cells previously exposed to chronic IFN γ . Self-renewal of *Dnmt3a*-mutant HSCs with prior IFN γ treatment was enhanced postsecondary transplant (Fig. 3H; Supplementary Fig. S4F).

Dnmt3a-Mutant HSCs Retain Quiescence after IFN γ Challenge

To investigate the molecular mechanisms underlying the fitness advantage of *Dnmt3a*-mutant HSCs under IFN γ , RNA-seq was performed on HSCs from mice at baseline and after a 2-hour IFN γ treatment to examine gene expression. In unmanipulated mice, there were few significantly differentially expressed genes (DEG) between control and *Dnmt3a*^{KO} HSCs (Fig. 4A), consistent with prior studies (15). In support of the functional data, genetic deletion of *IFNgr1* in *Dnmt3a*^{KO} HSCs normalized gene expression, including the previously reported (15) upregulation of HSC “stemness” genes (Supplementary Fig. S5A). Control and *Dnmt3a*^{KO} HSCs displayed a similar transcriptional response to IFN γ (Supplementary Fig. S5B). In HSCs at baseline, gene-set enrichment analysis (GSEA) revealed suppression of cell-cycle networks (E2F targets, G₂M checkpoint) in *Dnmt3a*^{KO} HSCs, whereas p53 pathway targets were activated (Fig. 4B). Comparison of IFN γ -stimulated HSCs showed a more robust IFN response in *Dnmt3a*^{KO} HSCs, but cell cycle-related genesets remained suppressed (Fig. 4C), suggesting *Dnmt3a*^{KO} HSCs may have a different proliferative response to IFN γ exposure.

Loss of quiescence is a major factor leading to erosion of HSC self-renewal (31, 32). As IFN stimulation is known to induce HSC proliferation (29, 33), we hypothesized that *Dnmt3a*-mutant HSCs may resist the detrimental effects of IFN γ exposure by maintaining a quiescent state. To examine HSC proliferation *in vivo*, mice were labeled with BrdU and treated with PBS (control) or IFN γ (Fig. 4D). Control HSCs were stimulated to proliferate in response to IFN γ (Fig. 4E). Proliferation of *IFNgr1*^{KO} HSCs was not different than control HSCs in mice treated with PBS, and remained unchanged following IFN γ treatment. However, *Dnmt3a*^{KO} HSCs were more quiescent at baseline, and their proliferation was unaltered in response to IFN γ (Fig. 4E). This suggests an uncoupling of the molecular response of *Dnmt3a*^{KO} HSCs to IFN γ exposure with the normal functional effect of increased proliferation. Deletion of *IFNgr1* in *Dnmt3a*^{KO} HSCs restored their proliferation rate at baseline to that of control HSCs (Fig. 4E). *Dnmt3a*^{KO}*IFNgr1*^{KO} HSCs remained insensitive to IFN γ , suggesting the effect is cell intrinsic. *Dnmt3a*^{HET} HSCs were not more quiescent than

the control HSCs at baseline, but like *Dnmt3a*^{KO} HSCs were resistant to IFN γ -induced proliferation (Fig. 4E).

To track HSC division on a clonal level, a histone labeling strategy was used. Control (*Vav-Cre;Dnmt3a*^{+/+}; *Rosa26*^{rtTA-M2}; *Col1a1*^{tetO_H2B-GFP}) and *Dnmt3a*^{KO} (*Vav-Cre;Dnmt3a*^{fl/fl}; *Rosa26*^{rtTA-M2}; *Col1a1*^{tetO_H2B-GFP}) mice were administered doxycycline chow for three weeks to label cells with H2B-GFP, then withdrawn from doxycycline for one week prior to experimentation to “wash out” the H2B-GFP label. H2B-GFP is quickly lost from proliferative hematopoietic cells, while being retained by the quiescent HSCs. The purpose of using H2B-GFP labeling in this experiment was to aid microscopic visualization of single HSCs. H2B-GFP labeled mice were treated with PBS (control) or IFN γ , then individual H2B-GFP⁺ HSCs were purified and monitored *in vitro* (Fig. 4F). Time to first division was not different between PBS-treated control and *Dnmt3a*^{KO} HSCs. While IFN γ treatment accelerated time to first division for control HSCs, *Dnmt3a*^{KO} HSCs were unaffected (Fig. 4G), again indicating *Dnmt3a*-mutant HSCs resist proliferation following IFN γ exposure. To more specifically determine whether *Dnmt3a*^{KO} HSCs were restricted at defined stages in cell cycle, Ki67/DAPI staining was performed on BM from PBS- or IFN γ -treated mice. Analysis was performed on both freshly isolated BM (Supplementary Fig. S5C), and also after HSCs were purified and cultured for 26 hours to stimulate proliferation. While there were no significant cell-cycle differences in freshly analyzed HSCs (Supplementary Fig. S5D), *Dnmt3a*^{R878} and *Dnmt3a*^{KO} HSCs showed a delayed exit from G₀ after a 26-hour culture period with concomitant decreases of HSCs in G₁ and S/G₂-M (Fig. 4H). This suggests quiescence-enforcing mechanisms may delay the proliferative activation of *Dnmt3a*-mutant HSCs following IFN γ exposure.

Txnip Is a DNA Methylation-Sensitive Gene Involved in HSC Stress Response

The single-cell division assay also revealed another interesting observation: while virtually all IFN γ -treated control HSCs had divided by 72 hours in culture, approximately 30% of *Dnmt3a*^{KO} HSCs had not divided by this timepoint, suggesting some clonal heterogeneity even within this highly purified cell population. To investigate this, single-cell RNA-seq was performed on PBS- and IFN γ -treated control and *Dnmt3a*^{KO} Lineage⁻c-Kit⁺EPCR⁺ cells. UMAP clustering showed treatment was a stronger influence on transcriptome than genotype (Fig. 5A, left). Assignment of cell identities (Fig. 5B) resolved two clusters of HSCs, with both genotypes grouped together but separated by treatment (Fig. 5A, right). Similar to the bulk HSC RNA-seq data, analysis of single-cell transcriptomic data identified only approximately 50 DEGs in the HSC clusters between genotypes and treatments. GSEA comparison of PBS-treated HSCs replicated many of the findings of bulk RNA-seq such as downregulation of “E2F targets” and “MYC targets” in *Dnmt3a*^{KO} HSCs, whereas analysis of IFN γ -treated cells confirmed the increased IFN γ signaling in *Dnmt3a*^{KO} HSCs (Supplementary Fig. S5E). The functional differences of *Dnmt3a*^{KO} HSCs were not due to the inability to upregulate immediate early response genes and myeloid differentiation

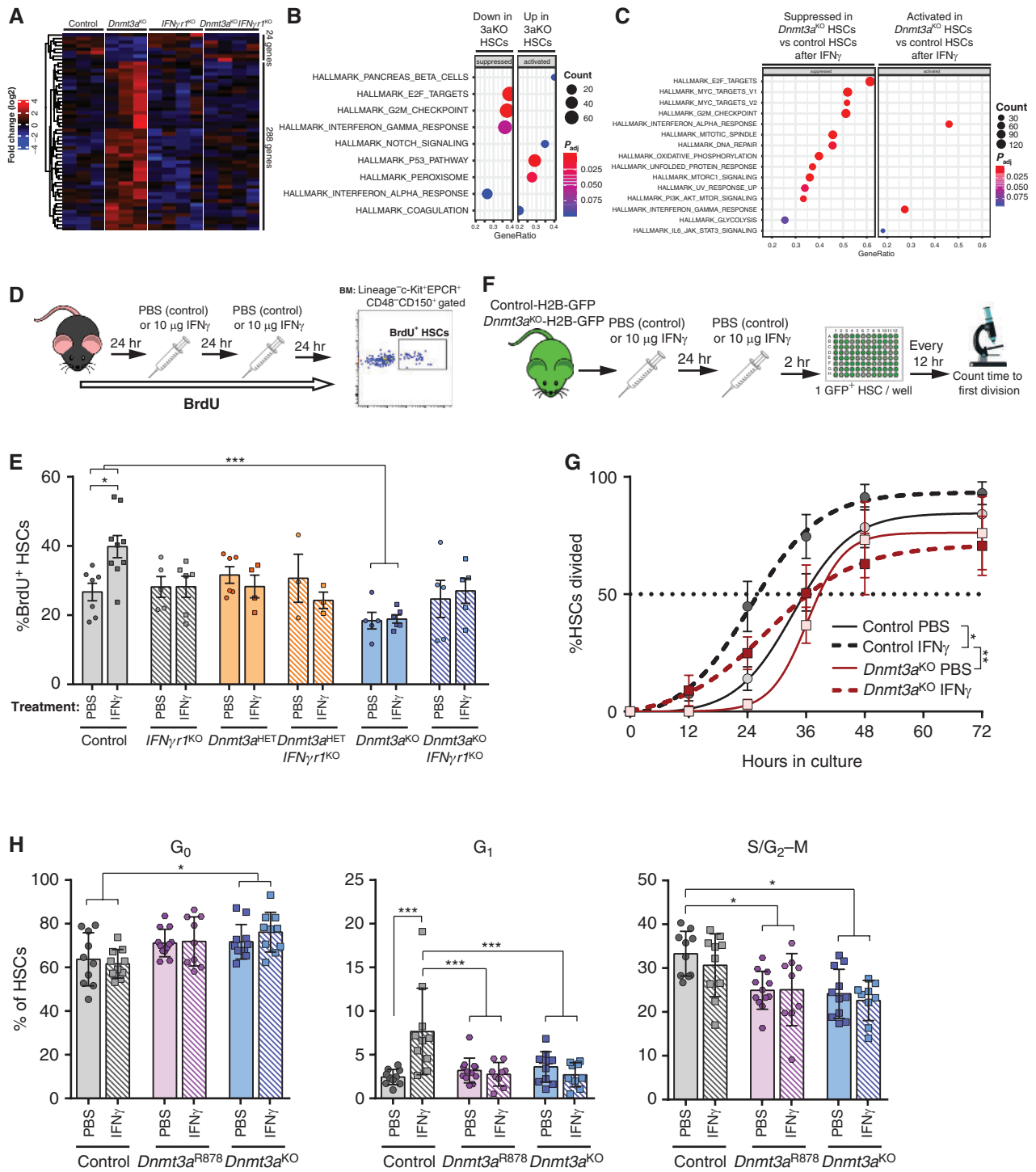
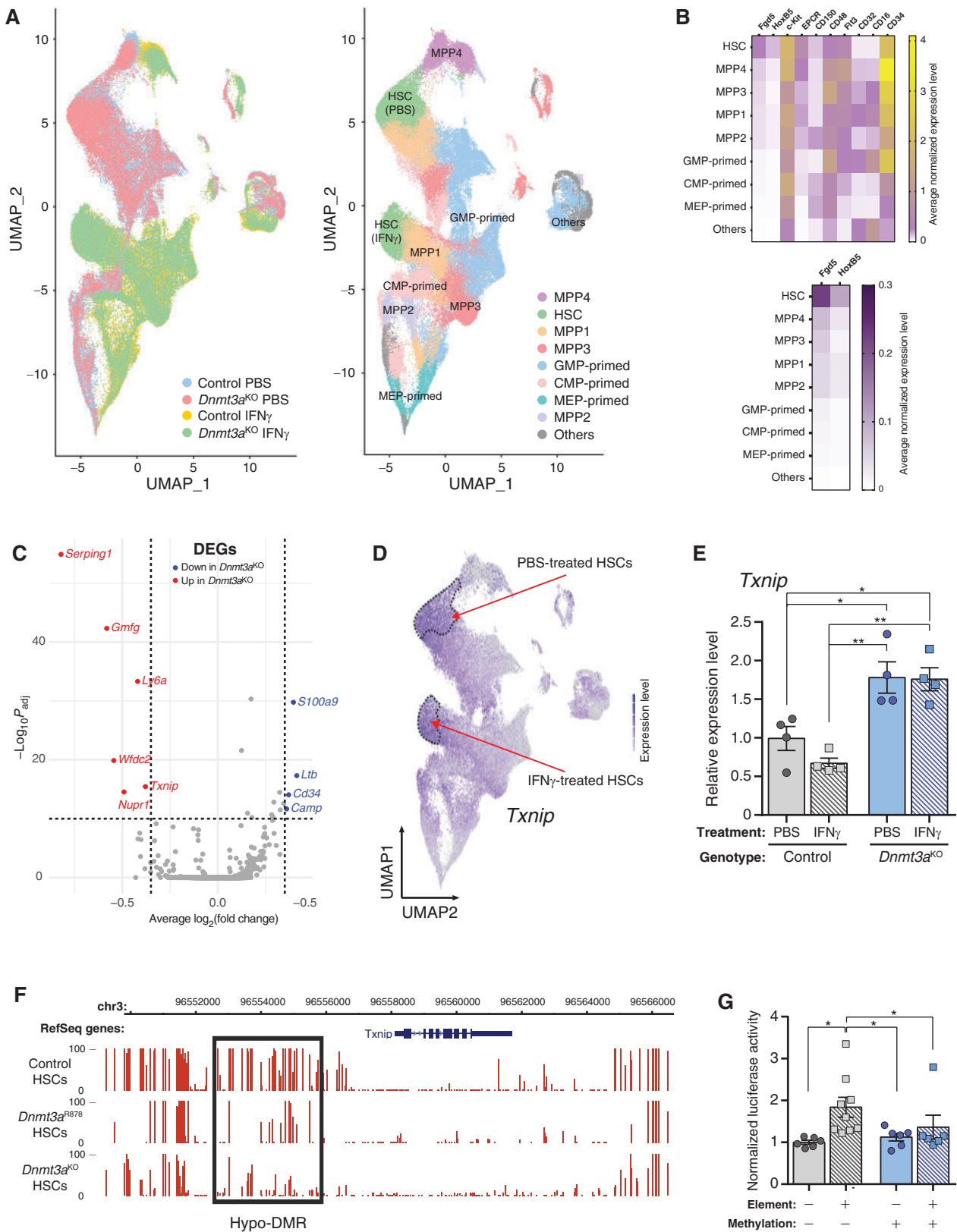


Figure 4. *Dnmt3a*-mutant HSCs retain quiescence after IFN γ challenge. **A**, Expression pattern of differentially expressed genes (DEGs; $P < 0.05$, fold-change > 2) between control and *Dnmt3a*^{KO} HSCs across *IFN γ r1*^{KO} and *Dnmt3a*^{KO}/*IFN γ r1*^{KO} HSCs. **B**, Gene-set enrichment analysis (GSEA) comparison of control and *Dnmt3a*^{KO} HSCs at baseline showing suppression of cell cycle-related genesets and activation of p53 targets. **C**, GSEA comparison of control and *Dnmt3a*^{KO} HSC after acute IFN γ challenge. **D**, Schematic of BrdU incorporation assay to assess HSC proliferative responses. **E**, Percentage of BrdU⁺ HSCs (Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) from indicated genotypes after 72-hour timecourse and treatment with either PBS or IFN γ (n = 3-9). **F**, Experimental workflow for single HSC division assays using H2B-GFP-labeled HSCs. **G**, Time to first division of HSCs (GFP⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) in response to PBS or IFN γ treatment (mean \pm SEM plus line of best fit, n = 5-6). **H**, Flow cytometric Ki67/7AAD cell-cycle analysis of HSCs (Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) from PBS- or IFN γ -treated mice following 26-hour culture to stimulate proliferation (n = 9-12). One-way ANOVA (**E** and **H**) or two-way ANOVA (**G**); *, $P < 0.05$; ***, $P < 0.001$. Data represent mean \pm SEM.



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factors in response to IFN γ (Supplementary Fig. S5F), but the single-cell RNA-seq was not able to resolve discrete *Dnmt3a*^{KO} HSC subpopulations which may be primed for differential IFN γ responses as suggested from the single HSC division assay (Fig. 4G).

Thioredoxin interacting protein (*Txnip*) was one of the few genes upregulated specifically in *Dnmt3a*^{KO} HSCs (Fig. 5C), and its expression pattern was HSC-predominant (Fig. 5D). This gene was of interest due to its described role in HSC function (34), cell-cycle regulation (35), and stress response (36). Targeted qPCR confirmed the upregulation of *Txnip* in *Dnmt3a*^{KO} HSCs at baseline, and while there was a trend for reduced *Txnip* in IFN γ -treated control HSCs, there was no diminution of the levels in IFN γ -treated *Dnmt3a*^{KO} HSCs (Fig. 5E). To determine whether the gene expression changes of *Dnmt3a*-mutant HSCs under IFN γ stress were associated with DNA methylation differences, whole genome bisulfite sequencing (WGBS) was performed on cells from TetON-IFN γ chimeric mice postprimary transplant. WGBS was performed on donor-derived (CD45.2⁺) BM cells as we have previously shown that the DNA methylation changes in *Dnmt3a*-mutant HSCs manifest more prominently in their differentiated progeny (15). *Dnmt3a*^{KO} cells showed global hypomethylation, whereas the overall genomic DNA methylation profile of *Dnmt3a*^{HET} cells was virtually indistinguishable from control cells (Supplementary Fig. S6A). Exposure to chronic IFN γ did not dramatically alter the methylome of any genotype (Supplementary Fig. S6B). Identification of differentially methylated regions (DMR) showed virtually all differences in DNA methylation were driven by genotype, rather than treatment (Supplementary Fig. S6C–S6E). This suggests the differences of *Dnmt3a*-mutant HSCs in response to IFN γ were not associated with DNA methylome remodeling. However, it has been shown that focal changes in DNA methylation in *DNMT3A*-mutant samples may have important biological consequences (18, 37). *Dnmt3a*-mutant cells showed a hypomethylated DMR in the *Txnip* regulatory region (Supplementary Fig. S6F), although the DNA methylation level was not altered by IFN γ treatment, suggesting this loss of methylation was a canonical consequence to *Dnmt3a* loss-of-function. The same region was also hypomethylated specifically in *Dnmt3a*^{KO} and *Dnmt3a*^{R878} HSCs (Fig. 5F). As this genomic region overlapped with ENCODE active enhancer histone marks, the DMR was cloned into a methylation-sensitive reporter assay. In the absence of DNA methylation, the DMR functioned as an active regulatory element (Fig. 5G). Artificial methylation of the reporter plasmid with the bacterial CpG methyltransferase *M.SssI* ablated this regulatory element activity (Fig. 5G). These observations suggest *Txnip* may be

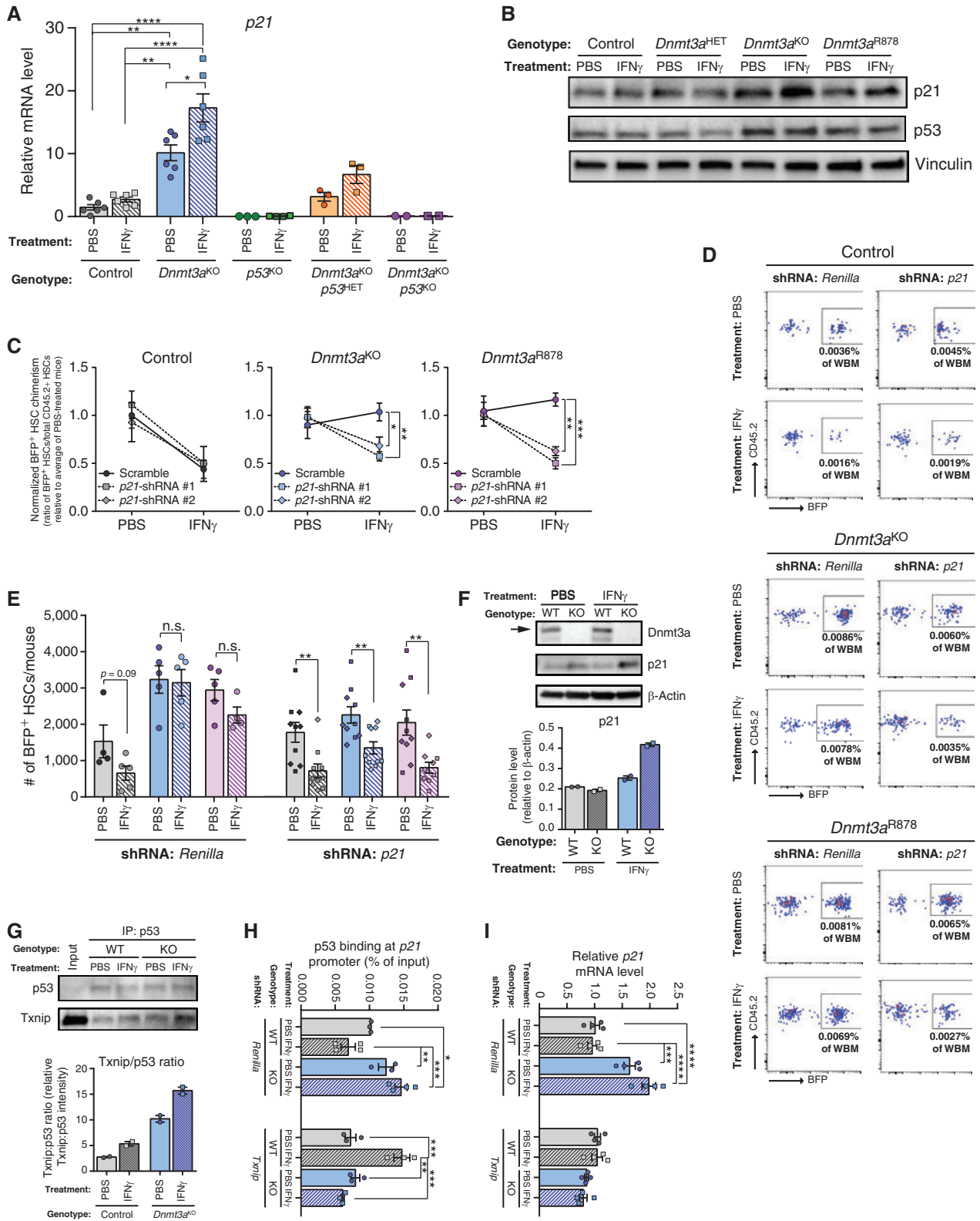
a DNA methylation-sensitive gene that primes the stress response of *Dnmt3a*-mutant HSCs.

Txnip Upregulates *p21* in *Dnmt3a*-Mutant HSCs in Response to IFN γ

Txnip is a thiol-oxidoreductase that controls cellular redox signaling to protect cells from oxidative stress, and regulates HSC quiescence under stress conditions (34) by stabilizing p53 protein via direct interaction (38). H₂DCFDA staining was performed in HSCs to correlate *Txnip* upregulation in *Dnmt3a*^{KO} with reactive oxygen species (ROS; Supplementary Fig. S7A). While IFN γ -treatment increased ROS in control HSCs, this response was absent in *Dnmt3a*^{HET} and *Dnmt3a*^{KO} HSCs (Supplementary Fig. S7B). Given the activated p53 gene expression signature in naïve *Dnmt3a*-mutant HSCs (Fig. 4B), we hypothesized *Txnip* may function in the response of *Dnmt3a*-mutant HSCs to IFN γ by regulating p53 and its targets. p53 has diverse functions in HSC responses to inflammation (28), which can be broadly classified as survival, apoptosis, and cell-cycle depending on the context. To examine p53 target genes that might explain the functional differences of *Dnmt3a*-mutant HSCs to IFN γ , gene expression analysis was performed on HSCs from mice treated with IFN γ . There were no IFN γ -induced differences in transcript levels of p53 itself, or major p53 targets for prosurvival (*Bcl2*, *Mcl1*) or proapoptosis (*Bax*, *Puma*) across HSC genotypes (Supplementary Fig. S7C). However, the p53 target gene *Cdkn1a* (*p21*) was significantly upregulated in *Dnmt3a*^{KO} HSCs at baseline and further increased after IFN γ treatment (Fig. 6A). A similar expression profile was observed in *DNMT3A*-edited cord blood CD34⁺ cells after transplant (Supplementary Fig. S7D). Hematopoietic progenitor-enriched BM cells (Lineage⁻c-Kit⁺) from *Dnmt3a*-mutant mice also showed upregulation of p21 associated with p53 stabilization following IFN γ (Fig. 6B; Supplementary Fig. S7E). *Dnmt3a*^{KO} mice were crossed to p53^{-/-} mice and the progeny treated with PBS or IFN γ . Gene expression analysis showed the upregulation of p21 in *Dnmt3a*^{KO} HSCs was p53-dependent (Fig. 6A).

p21 is a potent cell-cycle inhibitor (39), thus we hypothesized that p21 upregulation may be the downstream effector that preserves the quiescence of *Dnmt3a*-mutant HSCs under IFN γ challenge. To test this, c-Kit-enriched BM from control, *Dnmt3a*^{KO} and *Dnmt3a*^{R878} (the *Dnmt3a*-mutant strains with robust HSC phenotypes) mice was transduced with lentiviruses expressing one of two shRNAs targeting p21 (Supplementary Fig. S7F) or a nontargeting *Renilla* control shRNA. 1.5 × 10⁵ transduced c-Kit⁺ cells were transplanted in competition with congenic 2.5 × 10⁵ BM cells, and then the recipients were treated with PBS (control) or IFN γ (Fig. 1A). In terms of HSC clonal expansion, the reduction of *Renilla* shRNA-transduced control HSCs following IFN γ treatment

Figure 5. *Txnip* is a DNA methylation-sensitive gene involved in HSC stress response. **A**, UMAP clustering of single-cell RNA-seq data of control and *Dnmt3a*^{KO} Lineage⁻c-Kit⁺EPCR⁺ hematopoietic cells after treatment with PBS or IFN γ . Left, clustering of cells by genotype and treatment. Right, assignment of cell identities to cell clusters. Two clusters of HSCs are identified, which represents HSCs from both genotypes separated by treatment. **B**, Marker gene expression used to assign identities to cell clusters. **C**, Volcano plot of differentially expressed genes (DEG) between IFN γ -treated control and *Dnmt3a*^{KO} HSCs. **D**, *Txnip* expression in UMAP-clustered scRNA-seq data. Dashed lines outline HSC clusters. **E**, *Txnip* expression in HSCs purified from mice treated with PBS or IFN γ (n = 4). **F**, DNA methylation profile of *Txnip* locus in HSCs determined by WGBS. Height of red bar indicates methylation level of individual CpG. Black box denotes hypomethylated DMR in *Dnmt3a*-mutant HSCs. **G**, Luciferase activity of *Txnip* regulatory element in methylation-sensitive reporter assay. Firefly luciferase activity was measured 48 hours posttransfection and normalized to *Renilla* luciferase levels (n = 6–9). One-way ANOVA; *, P < 0.05; **, P < 0.01. Data represent mean ± SEM.



was completely paralleled in the same HSCs transduced with *p21*-targeting shRNAs (Fig. 6C and D). That is, *p21* knockdown had no influence on the response of normal HSCs to IFN γ . In contrast, the increased chimerism of *Dnmt3a*^{KO} and *Dnmt3a*^{R878} HSCs after IFN γ (Fig. 6C) was completely reversed following *p21* knockdown (Fig. 6C-E). These data suggest that *p21* has genotype-specific functions in the stress response of HSCs to inflammation.

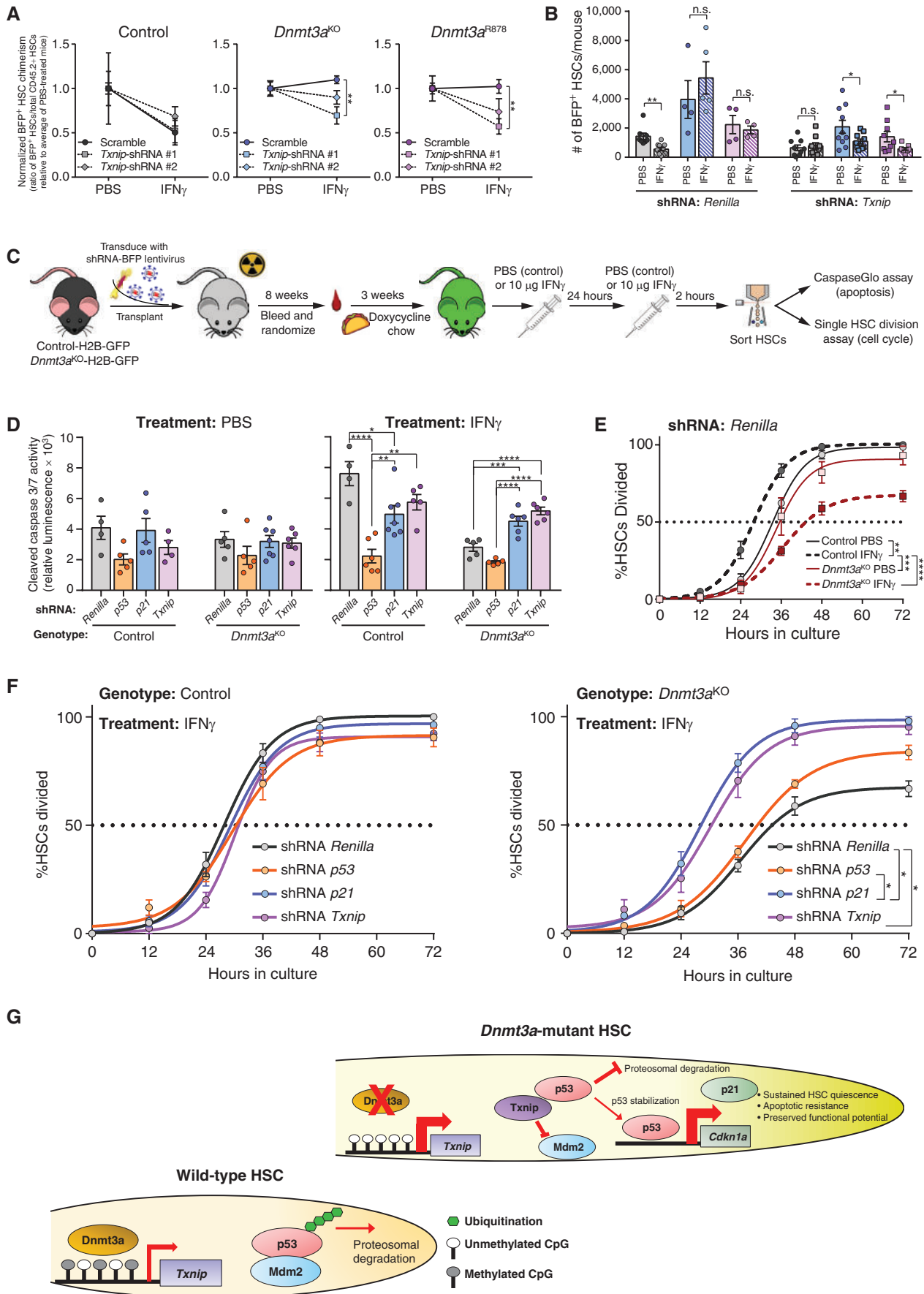
Biochemical studies were performed to examine the role of *Txnip* and *p53* in the regulation of *p21* in *Dnmt3a*-mutant cells. For experiments requiring greater input than can be obtained from primary HSCs, we used the mouse myeloid progenitor 32D cell line as a surrogate model. After CRISPR/Cas9-mediated *Dnmt3a* deletion, we confirmed these cells replicated the IFN γ -dependent increase in *p21* seen in *Dnmt3a*-mutant BM cells (Fig. 6F). As *Txnip* has been reported to stabilize *p53* protein in hematopoietic stress conditions, protein interaction between *Txnip* and *p53* was examined by immunoprecipitation. This showed an IFN γ -dependent increase in binding of *Txnip* to *p53* specifically in *Dnmt3a*-null 32D cells (Fig. 6G). As *p53* regulates *p21* expression by transcriptional activation, ChIP-qPCR was performed for *p53* occupancy at the well-annotated binding site in the *p21* promoter. WT and *Dnmt3a*-null 32D cells were transduced with lentiviral shRNAs targeting *Txnip* (Supplementary Fig. S7F) or *Renilla* control, then treated with either PBS (control) or IFN γ . In *Renilla*-transduced cells, there was increased binding of *p53* to the *p21* promoter in *Dnmt3a*-null 32D cells at baseline, which increased further after IFN γ treatment (Fig. 6H). In contrast, there was reduced *p53* binding at the *p21* promoter in WT 32D cells treated with IFN γ . Knockdown of *Txnip* normalized binding of *p53* to the *p21* promoter in *Dnmt3a*-null 32D cells to approximate WT 32D cells, and was unchanged in these cells after IFN γ (Fig. 6H). This suggests that in response to IFN γ , *Txnip* binds *p53* in *Dnmt3a*-mutant cells, leading to *p53* stabilization and increased binding to the *p21* promoter. The subsequent increased *p21* expression (Fig. 6I) hinders proliferation of *Dnmt3a*-mutant cells to preserve functional potential. We did note that *p53* binding was significantly increased at the *p21* promoter in WT 32D cells following *Txnip* knockdown which was the opposite response to *Renilla*-transduced cells (Fig. 6H). However, this did not translate to increased *p21* transcription (Fig. 6I) and implies a fundamental difference in the molecular response to IFN γ between control and *Dnmt3a*-mutant cells.

Txnip Preserves Functional Integrity of *Dnmt3a*-Mutant HSCs via *p21*

To validate *Txnip* as the upstream regulator of *p21* in the response of *Dnmt3a*-mutant HSCs to IFN γ , shRNA transplants were performed to reduce expression of *Txnip* as previously described for *p21*. Similar to inhibition of *p21*, *Txnip* knockdown did not influence the impaired self-renewal of control HSCs exposed to IFN γ (Fig. 7A). However, *Txnip* shRNA-transduced *Dnmt3a*^{KO} or *Dnmt3a*^{R878} HSCs phenocopied *p21* knockdown whereby the competitive advantage and clonal expansion of these HSCs under IFN γ challenge was mitigated by *Txnip* inhibition (Fig. 7B). *p21* expression was reduced in *Txnip* shRNA-transduced *Dnmt3a*^{KO} or *Dnmt3a*^{R878} HSCs (Supplementary Fig. S7G), implicating *Txnip* as an important regulator of *p21* in these mutant HSCs under IFN γ challenge.

To further the mechanistic connection between *Txnip*-*p53*-*p21* in response to IFN γ , a genetic experiment was performed by transducing H2B-GFP labeled control and *Dnmt3a*^{KO} HSCs with lentiviral shRNAs (Supplementary Fig. S7F), followed by BM transplantation. After stable engraftment, recipients were acutely treated with either PBS or IFN γ (Fig. 7C). HSC functional responses to IFN γ in terms of apoptosis and cell cycle were examined by cleaved caspase 3/7 activity and single HSC division assay, respectively. There were no significant differences in apoptosis between any shRNA-transduced HSCs from PBS-treated mice after culture (Fig. 7D), similar to previous observations (Fig. 2D). HSCs transduced with the control *Renilla* shRNA recapitulated the native setting with *Dnmt3a*^{KO} HSCs resistant to IFN γ -induced apoptosis (Fig. 7D). Knockdown of *p53* reduced apoptosis in both IFN γ -treated control and *Dnmt3a*^{KO} HSCs, likely because the genetic inhibition non-specifically altered a myriad of *p53*-related cellular functions, including induction of apoptosis (40). More specific effects were observed from genetic inhibition of *p21* and *Txnip*. While *p21* and *Txnip* inhibition was protective for control HSCs following IFN γ exposure, knockdown of these genes in *Dnmt3a*^{KO} HSCs increased IFN γ -induced apoptosis to approximate the levels observed in control HSCs (Fig. 7D). These data again highlight genotype-specific HSC responses and implicate a protective function for *Txnip* and *p21* in preventing stress-induced apoptosis in *Dnmt3a*-mutant HSCs in response to IFN γ .

Figure 6. *Txnip* upregulates *p21* in *Dnmt3a*-mutant HSCs in response to IFN γ . **A**, *p21* mRNA levels in HSCs (Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) from mice treated with PBS or IFN γ (two doses, cells purified 2 hours after the second dose) measured by qPCR ($n = 2-6$). **B**, Protein levels measured by Western blot in hematopoietic progenitor cells (Lineage⁻c-Kit⁺) from mice treated with PBS or IFN γ (two doses, cells purified 2 hours after the second dose). Representative of two independent experiments. **C**, Normalized level of BFP⁺ cell chimerism in donor-derived HSC (CD45.2⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) pool 18 weeks posttransplant. Ratio for each mouse is normalized to average BFP⁺ HSC chimerism in PBS treatment for individual shRNAs for each genotype to account for differences in transduction efficiency ($n = 4-5$ per group). **D**, Representative flow cytometry plots from mice transplanted with shRNA-transduced hematopoietic progenitor cells. Plots show BFP⁺ HSC chimerism for each genotype and treatment from Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺-gated BM. **E**, Number of BFP⁺ donor-derived HSCs (BFP⁺CD45.2⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁻CD150⁺) in BM of recipients 18 weeks posttransplant. Data for two independent shRNAs targeting *p21* are compiled, denoted by square (*p21*-shRNA #1) or diamond (*p21*-shRNA #2) shapes. **F**, Protein levels measured by western blot in 32D cells (WT = control; KO = *Dnmt3a*^{KO}) following 24-hour IFN γ (10 ng/mL) or PBS treatment. Arrow indicates full-length *Dnmt3a*. Representative of two independent experiments, quantification shown underneath. **G**, Western blot analysis of 32D cell lysate (treated with PBS or IFN γ for 24h; WT = control, KO = *Dnmt3a*^{KO}) following immunoprecipitation with *p53* antibody. Representative of two independent experiments, quantification shown underneath. **H**, Chromatin-immunoprecipitation qPCR analysis of *p53* enrichment at *p21* promoter in 32D cells (WT = control, KO = *Dnmt3a*^{KO}) transduced with shRNA targeting *Renilla* (control) or *Txnip*. Chromatin was isolated following a 24-hour exposure of IFN γ (10 ng/mL) or PBS ($n = 3-4$). **I**, qPCR for *p21* expression levels in the same cells used in **H**. One-way ANOVA (**A**, **C**, **H**, **I**) or two-tailed *t* test (**E**, data are compared for IFN γ relative to PBS within each genotype/shRNA); *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$; ****, $P < 0.0001$. Data represent mean \pm SEM.



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In terms of cell cycle, IFN γ treatment of control HSCs transduced with *Renilla* shRNA reduced time to first division, unlike *Dnmt3a*^{KO} HSCs (Fig. 7E), analogous to the native setting (Fig. 4G). In PBS-treated mice, *Txnip* inhibition had minimal effect on time to first division in both HSC genotypes compared with respective *Renilla* controls (Supplementary Fig. S7H). In IFN γ -treated mice, while *Txnip* inhibition in control HSCs had no effect compared with *Renilla*-shRNA, *Txnip* inhibition in *Dnmt3a*^{KO} HSCs restored their IFN γ -induced proliferative response (Fig. 7F). Analogous results were observed from *p21*-shRNA-expressing IFN γ -treated *Dnmt3a*^{KO} HSCs (Fig. 7F). Cumulatively, these data support a model whereby loss of DNA methylation at *Txnip* promoter in *Dnmt3a*-mutant HSCs, leads to increased expression and primes an inflammatory stress response. When *Dnmt3a*-mutant HSCs are challenged with IFN γ , *Txnip* stabilizes p53 through direct protein interaction, leading to upregulation of *p21* which helps prevent IFN γ -induced proliferation and apoptosis (Fig. 7G), providing the fitness advantage required to establish CH.

DISCUSSION

Because the identification of DNMT3A mutations in blood malignancies, a unifying mechanism of how these variants provide HSCs with a competitive advantage has proved elusive. Here, we used IFN γ as a selective pressure to identify molecular pathways that increase fitness of *Dnmt3a*-mutant HSCs. IFN γ signaling is required for clonal expansion of *Dnmt3a*-mutant HSCs *in vivo*, and chronic IFN γ provides an environment that selects these HSCs. Functionally, *Dnmt3a*-mutant HSCs resist IFN γ -induced apoptosis and preserve quiescence, allowing them to gain clonal dominance.

This study identified a *Txnip*-p53-p21 mechanism that protects *Dnmt3a*-mutant HSCs from the deleterious effects of IFN γ . While chronic IFN γ exposure did not dramatically alter the DNA methylome, *Txnip* upregulation in *Dnmt3a*-mutant HSCs was associated with a hypomethylated regulatory element. This reinforces previous findings that DNMT3A loss-of-function does not lead to global DNA methylation changes (15, 16, 19), yet focal methylation alterations appear physiologically relevant (17, 18, 37, 41). While there were few adaptive molecular changes in *Dnmt3a*-mutant HSCs in response to IFN γ , hypomethylation and upregulation of *Txnip* may prime *Dnmt3a*-mutant HSCs to respond differently to inflammation. Our data show under IFN γ challenge, upregulation of *Txnip* leads to p53

stabilization via direct interaction, with consequent upregulation of *p21* preserving functional integrity of *Dnmt3a*-mutant HSCs. Furthermore, dissecting each component of this *Txnip*-p53-p21 pathway revealed the relative importance and functional contribution of each factor. Although p53 and p21 have a plethora of functions in HSCs, we demonstrate a genotype-specific effect in preserving the functional integrity of *Dnmt3a*-mutant HSCs in response to IFN γ . But while the importance of the *Txnip*-p53-p21 mechanism was functionally validated *in vivo*, this pathway cannot be the only molecular mechanism that increases the fitness of *Dnmt3a*-mutant HSCs. p53 has a multitude of context-dependent activities in HSCs which may not be inextricably linked to the presented mechanism. Indeed, constitutive p53 activity impairs hematopoietic homeostasis and causes HSC depletion (42). In this study, while *Txnip* and *p21* shRNAs rescued the apoptosis and cell-cycle phenotypes of *Dnmt3a*-mutant HSCs in response to IFN γ , *p53* shRNAs largely did not. Moreover, only a fraction of *Txnip* appears to bind p53, suggesting *Txnip* has p53-independent actions. These data highlight the complexity of the mechanistic regulation of the HSC response to inflammation and how CH mutations can corrupt normal pathways.

Recent studies have identified a role for DNMT3A in the response of HSCs to chronic inflammation. The fitness advantage of *Dnmt3a*-mutant HSCs was quantified here using a novel transgenic IFN γ overexpression mouse model that enabled direct comparison between HSC genotypes. The IFN γ exposure regimen used in this study revealed an enforced quiescence phenotype in *Dnmt3a*-mutant HSCs by both *in vivo* BrdU labeling, cell-cycle analysis and *in vitro* single-cell division assay. We propose that enhanced quiescence-enforcing mechanisms, such as upregulation of *p21*, in *Dnmt3a*-mutant HSCs protect them from extrinsic forces that normally stimulate HSC proliferation and lead to functional decline. Cell-cycle analysis showed *Dnmt3a*-mutant HSCs show a delayed exit from G₀ in response to IFN γ . While p21 is more typically thought to function as a regulator of the G₁-S phase of cell cycle, p21 also causes cell-cycle arrest in G₀ in different settings (43, 44), again reinforcing the idea that these mechanisms are highly context-dependent. Moreover, there are likely many other molecular barriers that impede cell-cycle entry of *Dnmt3a*-mutant HSCs, one possibility being reduced metabolic activity suggested by GSEA of IFN γ -treated HSCs (Fig. 4C). Importantly, this study demonstrates the first *in vivo* evidence that human DNMT3A-mutant HSCs are sustained

Figure 7. *Txnip* preserves functional integrity of *Dnmt3a*-mutant HSCs via p21. **A**, Normalized level of BFP⁺ cell chimerism in donor-derived HSC (CD45.2⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁺CD150⁺) pool 18 weeks posttransplant. Ratio for each mouse is normalized to average BFP⁺ HSC chimerism in PBS treatment for individual shRNAs for each genotype to account for differences in transduction efficiency ($n = 4-5$ per group). **B**, Number of BFP⁺ donor-derived HSCs (BFP⁺CD45.2⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁺CD150⁺) in BM of recipients 18 weeks posttransplant. Data for two independent shRNAs targeting *Txnip* are compiled, denoted by square (*Txnip*-shRNA #1) or diamond (*Txnip*-shRNA #2) shapes. **C**, Schematic of functional genetic rescue by lentiviral shRNA transduction in H2B-GFP-labeled HSCs. **D**, Cleaved caspase 3/7 activity of shRNA-transduced HSCs (GFP⁺BFP⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁺CD150⁺) purified from mice treated with PBS or IFN γ (two doses, cells purified 2 hours after the second dose). Apoptosis was quantified following a 16-hour culture period ($n = 4-7$). Data for two independent shRNAs targeting *p53*, *p21* and *Txnip* are compiled. **E**, Time to first division of H2B-GFP-labeled HSCs (GFP⁺BFP⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁺CD150⁺) of indicated genotypes transduced with *Renilla*-targeting shRNA (control) from mice treated with PBS or IFN γ ($n = 4$). **F**, Time to first division of H2B-GFP-labeled control or *Dnmt3a*^{KO} HSCs (GFP⁺BFP⁺Lineage⁻c-Kit⁺EPCR⁺CD48⁺CD150⁺) transduced with shRNA targeting indicated genes from mice treated with IFN γ ($n = 4$). **G**, Schematic illustrating *Txnip*-p53-p21 axis in the preservation of *Dnmt3a*-mutant HSCs. One-way ANOVA (**A**), two-tailed t test (**B**, data are compared for IFN γ relative to PBS within each genotype/shRNA), two-way ANOVA (**D** and **E**) or one-way ANOVA with Tukey multiple test correction (**F**); *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$; ****, $P < 0.0001$. Data represent mean \pm SEM.

in response to episodic IFN γ exposure. The relatively small survival benefit observed in this model supports the premise that the fitness advantage conveyed by *DNMT3A* mutations in HSCs requires cumulative selection pressure over decades to manifest human CH. One caveat to these studies that must be acknowledged is the use of irradiation to condition the hosts, a process that induces inflammation, but appropriate controls were included in each experiment.

Despite tremendous advances in understanding the genetic basis of MDS and AML, cure rates have not substantially improved in the last 40 years (45). To our knowledge, this is one of the first studies to link changes in an environmental selection pressure with a fitness advantage for HSCs bearing a specific mutation. Targeting the mechanisms that allow *DNMT3A*-mutant HSCs to gain clonal dominance may highlight interventions that can selectively inhibit their growth and diminish future risk of malignant transformation.

METHODS

Mice and Transplantation

The Institutional Animal Care and Use Committee at Washington University School of Medicine approved all animal procedures. Mice were housed in specific pathogen-free conditions at Washington University School of Medicine on a 12:12-hour light:dark cycle in temperature- and humidity-controlled rooms. Donor mice were typically 12 weeks old for experimentation. Both male and female mice were used. All mice were from C57Bl/6 background. *Dnmt3a^{fl/fl}*, *Dnmt3a^{fl/R878}*, and *Tet2^{fl/fl}* mice were crossed to *Vav-Cre* strain. Recipient mice (C57Bl/6 CD45.1, The Jackson Laboratory strain #002014) mice were approximately 8 weeks old and transplanted by retro-orbital injection after a split dose (4 hours apart) of 10.5 Gy irradiation (Cesium-137). For general HSC-competitive transplants, 200 phenotypically defined HSCs (donors, CD45.2) were transplanted along with 2.5×10^5 whole BM (WBM) cells from wild-type (CD45.1) mice. For the competitive transplants challenged with stressors, 5×10^5 WBM donor cells (CD45.2) were cotransplanted with 5×10^5 WBM competitors (CD45.1). Recipients were randomized 4 weeks posttransplant and treated with PBS (control) or two consecutive doses of 10 μ g recombinant mouse IFN γ (mIFN γ ; retro-orbital injection). Recipients were sacrificed 18-weeks posttransplant for HSC analysis. For CH competition model, 1.2×10^6 WBM (CD45.1) from *Rosa26^{M2rtTA}*; *IFN γ ^{1KO}* (= TetOn; control) or tetO-*IFN γ* ; *Rosa26^{rtTA-M2}*; *IFN γ ^{1KO}* (= TetOn-IFN γ) were cotransplanted with 1.5×10^5 WBM cells from CD45.2 (donors) and 1.5×10^5 CD45.1/2 Ubc-GFP cells (competitors) into CD45.1 recipients. This models human CH with a 10% VAF of the test genotype. Recipients were fed 1,250 ppm doxycycline chow (TestDiet #1818439-203) on alternate weeks during 5 to 20 weeks posttransplant. This level of doxycycline chow was titrated to induce a physiologically relevant serum IFN γ level (~200 pg/mL). HSC quantification was performed 24 weeks posttransplant, and 7×10^5 c-Kit-enriched chimeric BM was transplanted into secondary recipients via retro-orbital injection. HSC quantification of secondary transplants was analyzed 18 weeks posttransplant. For all mouse transplants, experiments were repeated across at least two independent cohorts. Final graphs represent compiled data across multiple cohorts to control for experimental variability. The following mouse strains were used - *Vav-Cre* (46), *Dnmt3a^{fl/fl}* (47), *Dnmt3a^{fl/R878}* (24), *Tet2^{fl/fl}* (48), *p53^{-/-}* (49), Ubc-GFP (The Jackson Laboratory strain #004353), tetO-*IFN γ* (The Jackson Laboratory strain #009344), *Rosa26^{rtTA-M2}* (The Jackson Laboratory strain #006965); *IFN γ ^{1-/-}* (= *IFN γ ^{1KO}*); The Jackson Laboratory strain #003288), *IFN γ ^{1-/-}* (= *IFN γ ^{KO}*); The Jackson Laboratory strain # 002287), and *Col1a1^{tetO-H2B-GFP}* (= H2B-GFP; The Jackson Laboratory strain #016836). For studies requiring single

HSC division assays, H2B-GFP labeling of HSCs was induced with a 3-week course of 10,000 ppm doxycycline chow, followed by a 1-week chase period prior to isolation of HSCs. The purpose of H2B-GFP in this experiment was simply a label to more easily visualize HSCs during microscopic evaluation.

Human Cord Blood Transplantation

Deidentified cord blood specimens were obtained from the St. Louis Cord Blood Bank. Because samples were anonymized, no member of the study team had access to information that allowed the specimens to be linked to identifiable individuals, and the provider of the specimens was not involved in the design or conduct of the research, these experiments were deemed “nonhuman studies” by the Washington University Human Research Protection Office (HRPO). Mononuclear cells were isolated by Ficoll gradient according to standard procedures. CD34⁺ cells were isolated using magnetic enrichment (Miltenyi Biotec #130-100-453) and cultured overnight in SFEMII media (StemCell Technologies #09605) supplemented with 50 U/mL penicillin-streptomycin (Fisher Scientific #MT30002CI), 50 ng/mL human stem cell factor (SCF; Miltenyi Biotec #130-096-695), 50 ng/mL human thrombopoietin (TPO; Miltenyi Biotec #130-094-013), and 50 ng/mL human FLT3 L (Miltenyi Biotec #130-096-479). Enrichment efficiency was confirmed by flow cytometry.

For each sublethally irradiated (200 rads) NSG (The Jackson Laboratory strain #005557) recipient, 7×10^4 live cells were transplanted via intratibial injection. Engraftment was determined 4 weeks posttransplantation, and recipients were randomized for IFN γ treatment. Two consecutive doses (24 hours apart) of recombinant human IFN γ (hIFN γ , 2 μ g) were retro-orbitally injected per week during two treatment blocks (week 5-8 and week 13-16). Volume-equivalent PBS was injected as a control. Recipients were sacrificed 18 weeks posttransplant for HSC analysis. Approximately 3×10^4 hCD45⁺ cells were sorted from each recipient's BM to quantify posttransplant VAF. For secondary transplantation, hCD34⁺ cells were enriched using anti-human CD34 antibody from pooled samples per group. 1×10^6 hCD34⁺ cells were transplanted into secondary recipients via intratibial injection. Recipients were sacrificed 18 weeks posttransplantation for HSC analysis and VAF quantification.

CRISPR and VAF Determination

Single guide RNAs (gRNA) targeting *DNMT3A* (exon 12 and exon 14) or *AAVS1* were designed using the UCSC Genome Browser (<https://genome.ucsc.edu>) with the following sequences - *AAVS1*: GGGGCCACTAGGGACAGGAT; *DNMT3A* exon 12 (1012): TCCC CAGCATCGGACCCAC; *DNMT3A* exon 14 (1014): CAGGCGTGG TAGCCACAGTG. Gene-knockout was performed with the CRISPR-RNP system (IDT) via nucleoporation using Neon Transfection Kit following the manufacturer's instructions. Briefly, 1.5 μ g of CRISPR-Cas9 (NLS, 2, IDT) and 1 μ g of gRNA were nucleofected into 2×10^5 CD34⁺ cells. Cells were incubated at 37°C with 5% CO₂ overnight postnucleofection and recovered for an additional 24 hours in fresh media. 5×10^4 edited cells were collected to determine CRISPR/Cas9 targeting efficiency using PCR amplicon-based deep sequencing. CRISPresso 2.0 was used to identify mutations and quantify VAF. The following primers were used to generate amplicons from genomic DNA for VAF quantification - *AAVS1*-forward: CACTCTTTCCCTACACGACGCTCTTC CGATCTACAGGAGGTGGGGTTAGAC with *AAVS1*-reverse: GTGA CTGGAGTTCAGACGTGTGCTCTTCCGATCTCCCCTATGTC CACTTCAGGA; *DNMT3A*-1012-forward: CACTCTTTCCCTACACG ACGCTCTCCGATCTCACCTCGTACTCTGGCTCGT with *DNMT3A*-1012-reverse: GTGACTCGAGTTCAGACGTGTGCTCTTCCGATC TCAGGAATGAATGCTGTGGAA; and *DNMT3A*-1014-forward: CACTCTTTCCCTACACGACGCTCTTCCGATCTCACCTCG

TACTCTGGCTCGT with *DNMT3A*-1014-reverse: GTGACTG GAGTTCAGACGTGTGCTCTTCCGATCTCAGGAATGAATGCT GTGGAA.

Cell Purification and Flow Cytometry

BM cells were isolated from iliac crests, femurs, and tibias of mice. Cells were stained in Hanks Balanced Salt Solution (HBSS, Corning #21021CV) containing 100 Units/mL penicillin/streptomycin (Fisher Scientific #MT30002CI), 10 μ mol/L HEPES (Life Technologies #15630080) and 2% Serum Plus II (Sigma #14009C) at a density of 1.0×10^8 /mL. Staining was performed for >20 minutes at 4°C with desired antibodies. For cell sorting, BM was enriched with anti-mouse CD117-conjugated microbeads (Miltenyi Biotec, #130-0910224) using AutoMacs Pro Separator (Miltenyi Biotec), then stained with appropriate antibody cocktails. For typical mouse HSC sorting and analysis, staining panel included: CD45.1-FITC (clone A20; BioLegend #110706), CD45.2-BV421 (clone 104; BioLegend #109831), B220-APCCy7 (clone RA3-6B2; BioLegend #103224), Gr-1-APCCy7 (clone RB6-8C5; BioLegend #108424), Mac-1-APCCy7 (clone M1/70; BioLegend #101226), CD3e-APCCy7 (clone 145-2C11; BioLegend #100330), Ter119-APCCy7 (clone TER-119; BioLegend #116223), CD48-PECy7 (clone HM48-1; BioLegend #103424), CD150-PE (clone TC15-12F12.2; BioLegend #115904), c-Kit-BV605 (clone 2B8; BioLegend #105847), EPCR-APC (clone RCR-16; BioLegend #141506). For BrdU analysis, mice were injected intraperitoneally with a single dose of 3.33 mg BrdU (Sigma #B5002) in 500 μ L PBS, then supplemented with 0.8 mg/mL BrdU in drinking water over a 72-hour timecourse. At 24 and 48 hours, mice were given either PBS (control) or 10 μ g recombinant mouse IFN γ (rmIFN γ) by retro-orbital injection. At 72 hours, mice were sacrificed for BrdU analysis. BM from each mouse was enriched with anti-CD117 microbeads as above, then stained with the following antibodies: B220-APCCy7 (clone RA3-6B2; BioLegend #103224), Gr-1-APCCy7 (clone RB6-8C5; BioLegend #108424), Mac-1-APCCy7 (clone M1/70; BioLegend #101226), CD3e-APCCy7 (clone 145-2C11; BioLegend #100330), Ter119-APCCy7 (clone TER-119; BioLegend #116223), CD48-PECy7 (clone HM48-1; BioLegend #103424), CD150-BV421 (clone TC15-12F12.2; BioLegend #115925), c-Kit-FITC (clone 2B8; BD #561680), EPCR-PE (clone 1560; BD #566337). After antibody staining, samples were processed with the BrdU-APC flow kit (BD #552598) as per manufacturer instructions. For measurement of ROS levels in HSCs, 10 million BM cells or 500 cultured HSCs stained with a HSC flow panel were incubated with 5 μ mol/L CM-H2DCFDA (Thermo Fisher) at 37°C for 30 minutes, and mean fluorescence intensity (MFI) was quantified. For determination of cell-cycle stages in HSCs, 10 million BM cells or 500 cultured HSCs stained with a HSC flow panel were fixed, permeabilized, and stained with anti-Ki67-BV605 (clone 16A8; BioLegend #652413; 1:50 dilution) at 4°C for 1 hour. DNA content was determined by 7AAD (BioLegend #420404, at 1:10 dilution).

For mouse peripheral blood analysis, red blood cells were lysed then samples were stained with the following antibodies: CD45.1-FITC (clone A20; BioLegend #110706), CD45.2-BV421 (clone 104; BioLegend #109831), B220-PECy7 (clone RA3-6B2; BioLegend #103222), B220-APC (clone RA3-6B2; BioLegend #103212), Gr-1-PECy7 (clone RB6-8C5; BioLegend #108416), Mac-1-PECy7 (clone M1/70; BioLegend #101216), CD3e-APC (clone 145-2C11; BioLegend #100312).

For peripheral blood analysis from NSG transplants, blood was collected as above then samples were stained with the following antibodies: anti-mouse CD45-BV605 (clone 30-F11; BioLegend #103139), anti-human CD45-APC (clone 2D1; BioLegend #368512), anti-human CD3-FITC (clone OKT3; BioLegend #317306), anti-human CD19-PECy7 (clone HIB19; BioLegend #302215), anti-human CD33-BV421 (clone WM53; BD #565949). For BM analysis of cord blood-transplanted NSG mice, BM was collected as above then samples were stained with the following antibodies - anti-mouse

CD45-BV605 (clone 30-F11; BioLegend #103139), anti-human CD45-APC (clone 2D1; BioLegend #368512), anti-human CD34-PE (clone 561; BioLegend #343606), anti-human CD90-PECy7 (clone 5E10; BioLegend #328124), anti-human CD45RA-BV421 (clone HI100; BioLegend #304130), anti-human CD38-FITC (clone HB7; Invitrogen #11-0388-42), anti-human CD49f-APCCy7 (clone GoH3; BioLegend #313628), and anti-human Lineage cocktail-BV510 (BioLegend #348807).

All antibodies were used at 1:100 dilution unless stated otherwise. Dead cells were excluded with 7AAD (BioLegend #420404, used at 1:100 dilution). Cell sorting was performed using MoFlo (Beckman Coulter) and FACS Aria II (BD) flow cytometers. Flow cytometric cell analysis was performed using Attune NxT (Thermo Fisher Scientific) and FACS Aria II (BD) flow cytometers. Acquired flow cytometry data were analyzed with FlowJo software (Tree Star).

Lentiviral Production and Transduction

293T cells were cotransfected with the packaging plasmids pMD.G, psPAX2 and with an shRNA-containing plasmid (BFP-(miR-E)-PGK-Puro) using PEI-based transfection protocol (Polysciences #23966-1). shRNAs were designed according to published protocols and the following 22-mer target sequences were used for final experimentation - *p21* shRNA#1: TTAAATAACTTTAAGTTTGGGA; *p21* shRNA#2: TTAAATAACTTTAAGTTTGGAG; *p53* shRNA#1: TTACA CATGTACTTGTAGTGGGA; *p53* shRNA#2: TAAAATAGGAAATTGATATATA; *Txnip* shRNA#1: TAATCATACAAAAGATACACA; *Txnip* shRNA#2: TTAATCATACAAAAGATACAC. The negative control shRNA targeting *Renilla* luciferase was subcloned from Addgene plasmid #111170. Transfections for lentiviral production were performed in 150 \times 25 mm tissue culture dishes (Falcon #353025) when 293T cells reached >85% confluency. Forty-eight hours posttransfection, 293T cell supernatants were collected and concentrated by centrifugation at 76,000 \times g for 1.5 hours at 4°C. For lentiviral transduction, c-Kit⁺ BM cells were purified using magnetic enrichment and adjusted to 5 \times 10⁵ cells/100 mL in Stempro-34 medium (Gibco #10639011) supplemented with 100 Units/mL penicillin/streptomycin (Fisher Scientific #MT30002CI), 2 mmol/L L-glutamine (Gibco #25030-081), 100 ng/mL murine stem cell factor (SCF; BioLegend #579704), 100 ng/mL murine thrombopoietin (TPO; PeproTech, #315-14), 50 ng/mL murine Flt3 L (PeproTech, #250-31L), 5 ng/mL murine IL3 (Miltenyi Biotec #130-099-510), 4 μ g/mL polybrene (Sigma; #TR-1003-G), and spin-infected with lentivirus preparations at 250 \times g for 2 hours in a 96-well plate. Twenty-four hours posttransduction, 1.6 \times 10⁵ transduced cells were transplanted into lethally irradiated mice (following a split dose of 10.5 Gy of irradiation) by retro-orbital injection.

Single-Cell Division Assay

Control-H2B-GFP (*Vav-Cre; Dnmt3a^{+/+}; Rosa26^{M2rtTA}; Col1a1^{tetO-H2B-GFP}*) and *Dnmt3a^{KO}*-H2B-GFP (*Vav-Cre; Dnmt3a^{fl/fl}; Rosa26^{M2rtTA}; Col1a1^{tetO-H2B-GFP}*) mice were fed 10,000 ppm doxycycline chow (TestDiet #1815461-203) for 21-days. A 7-day recovery period was given prior to two consecutive doses of rmIFN γ (10 μ g) or PBS (24 hours apart). Mice were sacrificed 2 hours post second dose. Single immunophenotypically-defined HSC (Lineage⁻ c-Kit⁺ EPCR⁺ CD150⁺ CD48⁻ GFP^{hi}) were sorted into individual wells of 96-well plates containing complete Stempro-34 media described above. For single-cell division assay post-shRNA transductions, c-Kit⁺ cells were enriched with AutoMacs Pro Separator from three Control-H2B-GFP and three *Dnmt3a^{KO}*-H2B-GFP mice and transduced with shRNA-containing viral particles. Twenty-four hours posttransduction, 1.6 \times 10⁵ transduced cells (pooled by genotype) were transplanted into recipients. When stable engraftment was achieved at 8 weeks posttransplant, recipients were fed 10,000 ppm doxycycline chow to label HSCs with GFP. After a 7-day recovery period, two doses of rmIFN γ (10 μ g)

or PBS were injected into recipients and 2 hours posttreatment, single immunophenotypically defined HSCs were sorted into 96-well plates. Single HSCs were cultured in Stempro-34 medium and cell division behaviors were monitored. H2B-GFP allowed accurate identification of HSCs under a fluorescent microscope. Cell division was monitored at 12-hour intervals (60 hours postsorting was not evaluated). The latency to first division was determined as the time when 50% of trackable cells achieved their first division. Trackable cells were defined as cells that died postdivision and cells that survived throughout the 72-hour period.

CaspaseGlo Assay

Five-hundred HSCs were purified from mice treated with two consecutive doses of PBS or rmIFN γ (10 μ g) and sorted into white-walled 96-well plates (Corning #3610) containing complete Stempro-34 media. For measuring apoptosis in freshly purified HSCs, cleaved caspase 3/7 activity (CaspaseGlo 3/7 assay; Promega # G8090) was measured immediately postsorting. For measuring apoptosis in stressed HSCs, the CaspaseGlo assay was employed after a 16-hour culture period. CaspaseGlo assay was carried out according to manufacturer instructions. Cleaved caspases 3/7 activity was measured by luminescence intensity using BioTek Synergy H1 Hybrid Plate Reader.

Serum IFN γ Determination

Sera were isolated from peripheral blood using microtainer tubes with serum separator additive (BD #365967). Serum IFN γ levels were quantified using the Mouse IFN γ ELISA Kit II (BD #558258).

Quantitative Real-time PCR

Total RNA was extracted using the Nucleospin RNA XS kit (Macherey-Nagel #740902-250) and reverse-transcribed with the SuperScript VIL0 kit (Invitrogen #11754-050). Quantification was performed with TaqMan Master Mix (Applied Biosystems #4304437), the desired gene-specific probe (FAM-MGB; Applied Biosystems) and an 18S probe (VIC-MGB; Applied Biosystems). Transcript expression levels were normalized to endogenous 18S, and fold-changes were determined using the $\Delta\Delta C_t$ method.

RNA Sequencing

HSCs were purified from biological replicates of mice (pooled WBM from two male and two female 12-week-old mice) treated twice with either PBS or recombinant mouse IFN γ (rmIFN γ) 24 hours apart. HSCs were purified 2-hours after the second dose. A NucleoSpin RNA XS kit (Macherey-Nagel #740902.250) was used to isolate RNA. Library preparation, sequencing, and alignment were performed by the Genome Technology Access Center (Washington University). The SMARTer Ultra Low RNA kit (ClonTech) was used to prepare the libraries from 3 to 5 ng of total RNA. Sequencing was performed with an Illumina NextSeq 3000. Reads were aligned with STAR version 2.5.4b with Gencode release M20 (GRCm38.p6) genome assembly. Unambiguous read counts were calculated by HTSEQ-count version 0.6.0 with mode "intersection-strict." Expression data were imported into Noiseq v2.28.0 for differential gene expression analysis with TMM normalization and batch correction. Generalized linear models were then created to test for gene/transcript level differential expression. Differentially expressed genes (DEG) were then filtered for unadjusted *P* values <0.05. Gene-set enrichment analysis (GSEA) was performed with fGSEA v1.10.0.

Single-Cell RNA Sequencing

Two male and two female mice per genotype (control, *Dnmt3a*^{KO}) were treated with two doses of PBS (control) or rmIFN γ (10 μ g) 24 hours apart, then sacrificed 2 hours after the second injection. BM was isolated from individual mice and 17,000 Lineage- c-Kit⁺

EPCR⁺ BM cells per mouse were purified by flow cytometry. Cells were resuspended at 1,200 cells per μ L in PBS + 0.04% BSA, with cell concentration verified by flow cytometric counting. cDNA was prepared after the GEM generation and barcoding, followed by the GEM-RT reaction and bead cleanup steps. Purified cDNA was amplified for 11 to 13 cycles before being cleaned up using SPRIselect beads. Samples were then run on a Bioanalyzer to determine the cDNA concentration. GEX libraries were prepared as recommended by the 10x Genomics Chromium Single Cell 3' Reagent Kits v3 user guide with appropriate modifications to the PCR cycles based on the calculated cDNA concentration. For sample preparation on the 10x Genomics platform, the Chromium Single Cell 3' GEM, Library and Gel Bead Kit v3 (PN-1000075), Chromium Chip B Single Cell Kit, 48 rxns (10x Genomics PN-10000153), and Chromium Dual Index Kit TT Set A (PN-1000215) were used. The concentration of each library was accurately determined through qPCR utilizing the KAPA library Quantification Kit according to the manufacturer's protocol (KAPA Biosystems/Roche) to produce cluster counts appropriate for the Illumina NovaSeq6000 instrument. Normalized libraries were sequenced on a NovaSeq6000 S4 Flow Cell using the XP workflow and a 28 \times 10 \times 10 \times 150 sequencing recipe according to manufacturer protocol. A median sequencing depth of 50,000 reads per cell was targeted for each library. Single-cell RNA-seq data were demultiplexed and aligned to the Genome Reference Consortium mouse genome (mm10). The aligned data were then annotated and UMI-collapsed using Cellranger (v3.1.0, 10x Genomics). Cells with more than 10% mitochondrial gene expression and with top 5% unique feature counts were excluded. This resulted in the following number of unique cells that passed quality control and were examined in downstream analysis: Control-PBS: 26,199; Control-IFN γ : 19,829; *Dnmt3a*^{KO}-PBS: 11,804; *Dnmt3a*^{KO}-IFN γ : 20,396. Principal component analysis was performed to reduce data using Seurat 3.0. Clusters were identified applying the dimensional reduction techniques (tSNE and UMAP) using the same package. Expression levels of conventional cell surface markers were examined to annotate clusters. Differentially expressed genes were identified within the cluster of interest.

Whole Genome-Wide Bisulfite Sequencing

At the conclusion of the CH competition model transplant with chronic IFN γ (primary transplant), 3 \times 10⁵ donor-derived (CD45.2⁺) BM cells per genotype per treatment were sorted from pooled recipient BM cells. Genomic DNA (gDNA) was extracted using PureLink Genomic DNA Mini Kit (Invitrogen). 250 ng of gDNA for each sample, spiked in with lambda control, was fragmented using Covaris. The fragmented DNA was then subjected to an overnight cytosine-to-thymine conversion using EZ DNA Methylation-Gold Kit (Zymo Research). The CT converted samples were used for library construction with Accel-NGS Methyl-Seq DNA Library Prep Kit, indexed with Methyl-Seq Dual Indexing kit. The sequencing was performed on NovaSeqS4 300XP. Raw reads were quality and adapter trimmed using cutadapt (v2.4) with parameters -q 15,10 -minimum-length 36. Trimmed reads were aligned using bismark (v0.19.0) with bowtie2 and options -X 1000 -score_min L,0,-0.6, -N 0. After removal of PCR duplicates from aligned reads, methylation levels were calculated using bismark_methylation_extractor. CpG methylation files were loaded in R (v 3.6.3) and transformed into a BS object using bsseq. Differentially methylated loci (DML) were determined using the DMLtest function of DSS after with option smoothing = TRUE. Differentially methylated regions (DMR) were calculated from the result of DMLtest using the callDMR function of DSS with option p.threshold = 1e-5. Resultant DMRs were further filtered to remove any DMRs with fewer than 4 CGs or a methylation difference between conditions of less than 20%. Bed formatted coordinates of DMRs were extracted from filtered lists of DMRs. Methylation levels were transformed into bigwig format and plotted at DMRs using

deepools. Average methylation levels in all DMR comparisons were determined using the bedtools map function with options -o mean and compiled into a table. The table was plotted as a heatmap using the pheatmap package (v1.0.12) in R with clustering across the DMRs using the default for the hclust function.

In Vitro Methylation-Sensitive Reporter Assay

The hypomethylated DMR upstream region of *Txnip* (-3283 to -1793bp) was amplified from wild-type mouse gDNA using primers Upstream_Txnip_F (AAATTTAAAGGTACCGGTAGTCTCAAATCCCACAGACTCA) and Upstream_Txnip_R (TTTAAATTTT GAGCTCCCAGAAAGTTAACTAAGGCGTCA) and ligated to the pGL4.23 [Luc2] vector (Promega) using *KpnI* and *SacI* restriction enzyme sites. The bacterial CpG methyltransferase *M.SssI* (New England Biolabs) was used to methylate the element-containing construct and vector only according to manufacturer's instruction. *In vitro* methylation efficiency was determined by restriction enzyme digestion using BstUI that only recognize unmethylated CpG but is not able to cut methylated CpG sites. pGL4.23-based plasmids and pGL4.75 [hRluc] (Promega) (9:1) were transiently transfected into 3T3 cells using PEI-based transfection protocol (Polysciences #23966-1). pGL4.74 (hRluc/TK) contains *Renilla* luciferase and was used to control for transfection efficiency. Transfections were performed in triplicates. Luciferase assays were carried out 48 hours posttransfection using the Dual-Glo luciferase assay system (Promega) according to manufacturer's instructions.

In Vitro IFN γ Exposure

32D cells were cultured in RPMI1640 medium supplemented with 2 mmol/L L-glutamine, 10 mmol/L HEPES and 100 U/mL Penicillin-streptomycin and incubated at 37°C with 5% CO₂ until reaching 100% confluency. Thirty-six hours postconfluency, 2.6×10^6 cells were seeded into a new flask at the density of 2.6×10^5 cells/ml with 10 ng/mL recombinant mouse IFN γ or volume-equivalent PBS. Cells were collected 24 hours postexposure for protein or chromatin harvest.

Chromatin Immunoprecipitation and Real-time PCR

Chromatin was extracted, sheared and isolated from 5×10^6 32D cells using SimpleChIP Enzymatic Chromatin IP Kit (Cell Signaling Technology) following manufacturer instruction. Chromatin shearing efficiency was examined by agarose gel electrophoresis. Ten micrograms of prepared chromatin was used to precipitate p53-bound gDNA using p53 antibody (Cell Signaling Technology #32532). An IgG control antibody was also applied to each sample as isotype control. The precipitated gDNA was then eluted and subjected to qPCR using gene-specific primers for *p21* promoter (forward: CTTTCTG-GCCTTCAGGAACA; reverse: GCTGGTAGTTGGGTATCATCAG). For positive controls, all samples were immunoprecipitated with H3 antibody (Cell Signaling Technology #4620) and amplified with Mouse RPL30 Intron 2 Primers (Cell Signaling Technology #7015) to ensure the quality of chromatin was comparable. The enrichment of p53 was determined as $2^{(Ct_{input} - Ct_{sample})}$.

Immunoprecipitation and Western Blotting

Thirty to 50 μ g of protein samples were separated in precasted 4%–15% gradient SDS gels (Bio-Rad, #456-1084) and transferred to polyvinylidene difluoride membranes (Millipore #IPVH00010). Membranes were subsequently probed with antibodies to detect p21 (Santa Cruz Biotechnology, # sc-6264), p53 (Cell Signaling Technology #32532), Txnip (Cell Signaling Technology #32532), β -Actin (Santa Cruz Biotechnology, #SC-47778) and Vinculin (Santa Cruz Biotechnology #sc-73614). Detection was performed using horseradish peroxidase-conjugated secondary mouse or rabbit antibody and chemiluminescence HRP substrate (Millipore #WBKLS0100). For

protein coimmunoprecipitation, total protein was extracted from 10×10^6 32D cells using Cell Lysis Buffer (Cell Signaling Technology #9803) following manufacturer's instructions. Four-hundred micrograms of freshly prepared total protein was then incubated with p53 antibody (1:50 dilution; Cell Signaling Technology #32532) at 4°C overnight. Antibody-bound lysates were captured by protein G beads. The lysates were then eluted and prepared for Western blot analysis. Forty micrograms of pooled total protein was also loaded on Western blot as 10% input control. p53-bound proteins were examined to detect p53 and Txnip.

Statistical Analysis

Student *t* test and one-way ANOVA were used for statistical comparison where appropriate. Significance is indicated using the following convention: *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$; ****, $P < 0.0001$. All graphs represent mean \pm SEM, unless otherwise indicated.

Data Reporting

No statistical methods were used to predetermine sample size. Group sizes were based on historical data outcomes from our laboratories. The experiments were randomized where stated. Investigators were not blinded to allocation during experiments and outcome assessment.

Data Availability

Raw data of bulk RNA-seq, single-cell RNA-seq and whole-genome bisulfite-sequencing are available in the Gene Expression Omnibus (GEO) repository under accession number GSE168807 (<http://www.ncbi.nlm.nih.gov/geo/>). Previously generated WGBS data (19) used to generate Fig. 5F are available under accession number GSE98191. There are no restrictions on data availability.

Authors' Disclosures

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Authors' Contributions

C.R. Zhang: Conceptualization, resources, data curation, software, formal analysis, validation, investigation, visualization, methodology. **E.L. Ostrander:** Data curation, validation, investigation, methodology. **O. Kukhar:** Investigation. **C. Mallaney:** Investigation. **J. Sun:** Investigation. **E. Haussler:** Investigation. **H. Celik:** Investigation. **W.K. Koh:** Investigation. **K.Y. King:** Conceptualization, resources. **P. Gontarz:** Software, formal analysis, methodology. **G.A. Challen:** Conceptualization, formal analysis, supervision, funding acquisition, writing-original draft, project administration, writing-review and editing.

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