

# Bmi1 regulates memory CD4 T cell survival via repression of the *Noxa* gene

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The maintenance of memory T cells is central to the establishment of immunological memory, although molecular details of the process are poorly understood. In the absence of the polycomb group (PcG) gene *Bmi1*, the number of memory CD4<sup>+</sup> T helper (Th)1/Th2 cells was reduced significantly. Enhanced cell death of *Bmi1*<sup>-/-</sup> memory Th2 cells was observed both in vivo and in vitro. Among various proapoptotic genes that are regulated by *Bmi1*, the expression of proapoptotic BH3-only protein *Noxa* was increased in *Bmi1*<sup>-/-</sup> effector Th1/Th2 cells. The generation of memory Th2 cells was restored by the deletion of *Noxa*, but not by *Ink4a* and *Arf*. Direct binding of *Bmi1* to the *Noxa* gene locus was accompanied by histone H3-K27 methylation. The recruitment of other PcG gene products and *Dnmt1* to the *Noxa* gene was highly dependent on the expression of *Bmi1*. In addition, *Bmi1* was required for DNA CpG methylation of the *Noxa* gene. Moreover, memory Th2-dependent airway inflammation was attenuated substantially in the absence of *Bmi1*. Thus, *Bmi1* controls memory CD4<sup>+</sup> Th1/Th2 cell survival and function through the direct repression of the *Noxa* gene.

## CORRESPONDENCE

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Abbreviations used: 5-Aza, 5-Aza-2'-deoxycytidine; BAL, bronchioalveolar lavage; ChIP, chromatin immunoprecipitation; MeDIP, methylated DNA immunoprecipitation; PcG, polycomb group; PRC, polycomb repressive complex; RL, lung resistance; Tg, transgenic.

The quality of adaptive immune responses depends on the size of the antigen-specific memory T cell pool, which is regulated through specific homeostatic mechanisms controlling both cell survival and proliferation. Upon antigen recognition, naive CD4 T cells undergo a rapid clonal expansion, followed by the differentiation into functionally distinct Th cell subsets, such as effector Th1 and Th2 cells (1–3). These effector T cells undergo a dramatic contraction in numbers after antigen clearance, with 90–95% succumbing to apoptosis within weeks (4–6). However, some of the effector T cells are maintained as memory T cells for long periods in vivo (7, 8).

In contrast to CD8 memory T cells, CD4 memory T cells may not require the signals through common cytokine receptor  $\gamma$  chain (9, 10). However, critical regulatory roles of IL-7 in the generation and survival of memory CD4 T cells were reported recently (11, 12). In addition, the homeostasis of memory CD4 T cells is dependent on the signals through the TCR as well as the IL-7 receptor (13). *Bcl-2*

and *Mcl-1* are reported to be the downstream targets of the IL-7 receptor and promote T cell survival (14–16).

Several properties that distinguish memory T cells from naive T cells have been described, such as increased longevity and enhanced capacity for recall response to cognate antigen (17). Memory T cells have several features associated with stem cells (18), and the similarity of the gene expression pattern between memory T/B cells and long-term hematopoietic stem cells was reported (19). Similar to hematopoietic stem cells, memory T cells appear to possess the ability to proliferate in response to homeostatic signals. The homeostatic signals may drive self-renewal, whereas antigenic signals drive effector cell differentiation and function.

The polycomb group (PcG) gene *Bmi1* has recently been implicated in the maintenance of hematopoietic (20, 21), neural (22), and cancer stem cells (23). PcG gene products form multimeric complexes and maintain the early determined gene expression patterns of key developmental regulators, such as homeobox genes both in invertebrates and vertebrates (24, 25). *Bmi1*, *Mel-18*, *M33*, *Pc2*, *Rae-28/Mph1*,

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Mph2, and Ring1A/B are constituents of a multimeric protein complex similar to the *polycomb* repressive complex (PRC)1 identified in *Drosophila*. Another PcG complex, which contains Eed, Suz12, Ezh1, and Ezh2, is PRC2. PRC2 possesses an intrinsic histone H3-K27 methyltransferase activity (26–29), which implicates a likely mechanism for PcG-mediated gene silencing. Lack of individual components of PRC1 results in apoptosis and the loss of proliferative responses of immature lymphoid cells (30, 31). In mature lymphocytes, PcG gene products play several roles in differentiation and cell fate. Mel-18 controls Th2 cell differentiation through the regulation of GATA3 expression (32), and Bmi1 controls the stability of GAT3 protein in developing Th2 cells (33). Ezh2 is involved in B cell development and controls IgH V(D)J rearrangement (34).

We herein investigated the role of Bmi1 in the generation and maintenance of memory CD4<sup>+</sup> Th1/Th2 cells. In the absence of Bmi1, the generation of both Th1 and Th2 memory cells was impaired as a consequence of increased Noxa expression. Because our previous study indicated that the expression of Bmi1 is higher in Th2 cells than Th1 cells, we sought to investigate the molecular mechanisms underlying the Bmi1/Noxa-mediated regulation of memory T cell generation by focusing primarily on memory Th2 cells. Our results indicate that Bmi1 controls memory CD4 T cell survival and function through the direct repression of the *Noxa* gene.

## RESULTS

### Generation of memory Th1/Th2 cells was impaired in the absence of Bmi1

In *Bmi1*<sup>-/-</sup> mice, although a reduction of cell numbers in the lymphoid organs was seen, substantial numbers of CD4 and CD8 T cells developed with normal expression of developmental cell surface antigens (not depicted). After in vitro antigenic stimulation, *Bmi1*<sup>-/-</sup> CD4 T cells were shown to proliferate well (Fig. S1 A, available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>) and differentiate normally into IL-4–producing effector Th2 cells under Th2 conditions or IFN- $\gamma$ –producing effector Th1 cells under Th1 conditions (33). Under IL-4–limiting conditions, Th2 cell differentiation was impaired significantly in the absence of Bmi1, and the Bmi1-mediated regulation of the stability of GATA3 protein was demonstrated (33).

To investigate the role of *Bmi1* in the generation and maintenance of memory Th1/Th2 cells, we used a “memory Th1/Th2 mouse” system (35, 36) in which OVA-specific  $\alpha\beta$ TCR transgenic (Tg) CD4 T cells from *Bmi1*<sup>+/+</sup>, *Bmi1*<sup>+/-</sup>, or *Bmi1*<sup>-/-</sup> mice were stimulated with antigenic OVA peptide and wild-type antigen-presenting cells under Th2 or Th1 culture conditions in vitro for 6 d, and these effector Th cells were then adoptively transferred into syngeneic BALB/c *nu/nu* mice (memory Th1/Th2 mice). 5 wk after cell transfer, the numbers of donor-derived KJ1<sup>+</sup> memory Th1/Th2 cells were assessed in various organs. As shown in Fig. 1 A, a *Bmi1* gene dose-dependent affect on the numbers of memory Th2 cells was observed in all tissues tested (spleen, liver, lung,

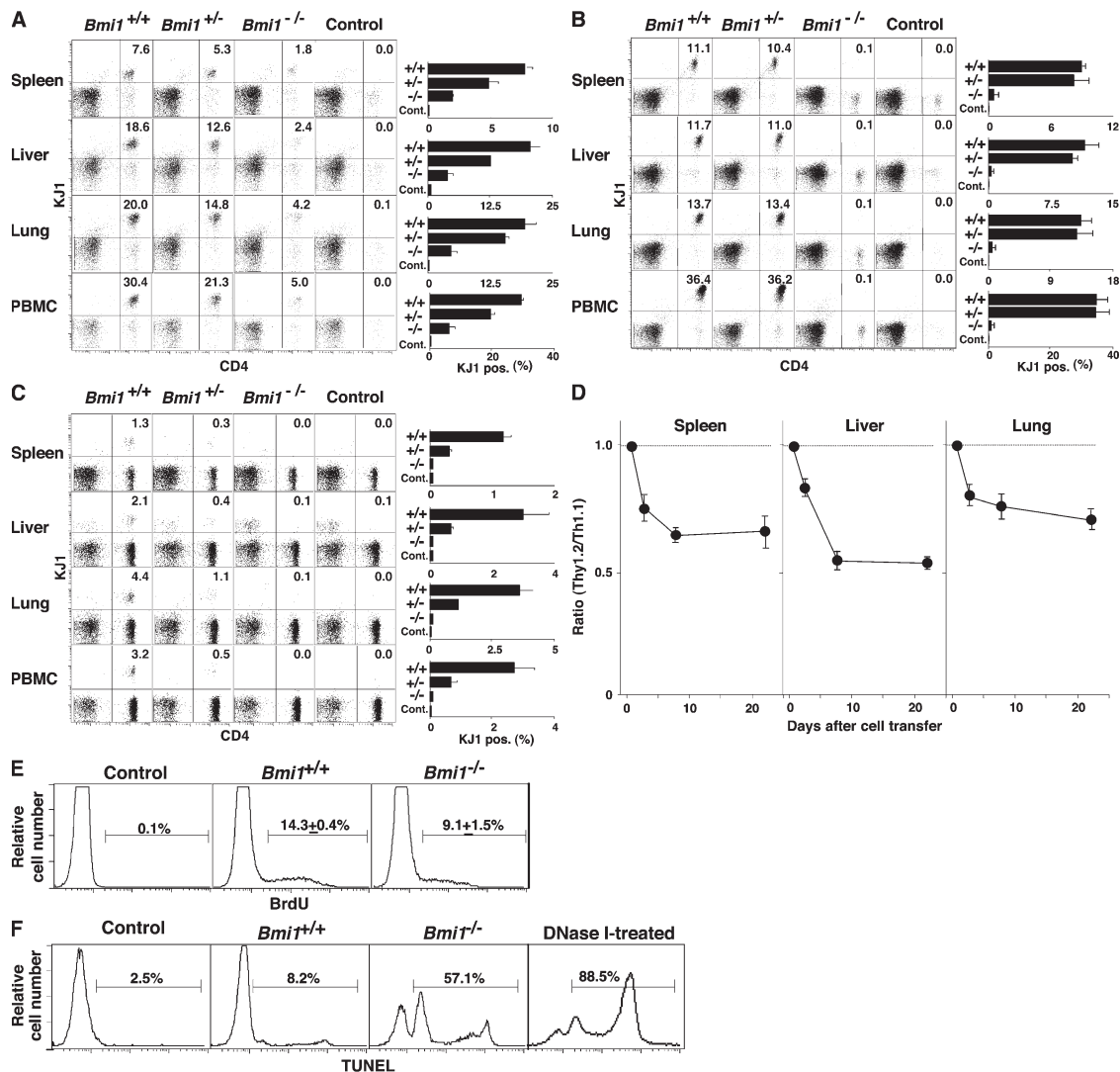
and PBMCs). It is worth noting that even heterozygous *Bmi1*<sup>+/-</sup> groups showed marked effects. Although the expression levels of IL-7R $\alpha$  and IL-2R $\beta$  were slightly lower on *Bmi1*<sup>-/-</sup> memory Th2 cells as compared with wild-type cells, other surface marker antigens were expressed normally (Fig. S1 B). Next, KJ1<sup>+</sup> effector Th1 cells were transferred into BALB/c *nu/nu* mice. The memory Th1 cell generation from *Bmi1*<sup>-/-</sup> effector Th1 cells was also impaired in all tissues tested (Fig. 1 B). The reduction in the *Bmi1*<sup>+/-</sup> group was not obvious. A similar loss of memory Th cell generation was observed in both Th2 (Fig. 1 C) and Th1 cells (Fig. S2, available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>) when *Bmi1*<sup>-/-</sup> effector cells were transferred into nonlymphopenic normal BALB/c mice. These results indicate that the generation of memory Th1/Th2 cells was impaired in the absence of Bmi1.

A kinetics study showed that the decrease in *Bmi1*<sup>-/-</sup> Th2 cells was observed from 2 d after cell transfer (Fig. S3, available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>). Consequently, we assessed the semi-acute survival of *Bmi1*<sup>+/-</sup> effector Th2 cells by a competition analysis using *Bmi1*<sup>+/+</sup> (Thy1.1) and *Bmi1*<sup>+/-</sup> (Thy1.2) C57BL/6 Ly5.2 effector Th2 cells and Ly5.1 host animals with a C57BL/6 background. Typical staining patterns of mixed cells for cell transfer and of transferred donor-derived cells (Ly5.2) from the indicated tissues are shown in Fig. S4. Interestingly, a significant decrease in the ratio of *Bmi1*<sup>+/-</sup>/*Bmi1*<sup>+/+</sup> (Thy1.2/Thy1.1) cells was already observed 2 d after cell transfer, and this decline continues through days 7–21 (Fig. 1 D). The kinetics of the preferential reduction of *Bmi1*<sup>+/-</sup> cells was similar in all tissues tested. The defect in semi-acute survival in *Bmi1*<sup>-/-</sup> Th2 cells was also observed under lymphopenic conditions (Fig. S5).

The homeostatic proliferation as measured by BrdU incorporation in vivo was decreased slightly in *Bmi1*<sup>-/-</sup> memory Th2 cells as compared with wild type ( $14.3 \pm 0.4\%$  vs.  $9.1 \pm 1.5\%$ ) (Fig. 1 E). Next, we assessed the extent of ongoing apoptotic cell death in vivo by a TUNEL assay. In *Bmi1*<sup>-/-</sup> memory Th2 cells, ongoing apoptosis was enhanced dramatically as compared with wild type (8.2 vs. 57.1%) (Fig. 1 F). In addition, increased annexin V<sup>+</sup>PI<sup>+</sup> cells were detected in *Bmi1*<sup>-/-</sup> memory Th2 cells after restimulation in vitro, indicating that these memory cells are highly prone to cell death (Fig. S1 C).

### Deletion of *p16*<sup>ink4a</sup> and *p19*<sup>arf</sup> genes failed to restore the decreased memory Th2 cell generation in the absence of Bmi1

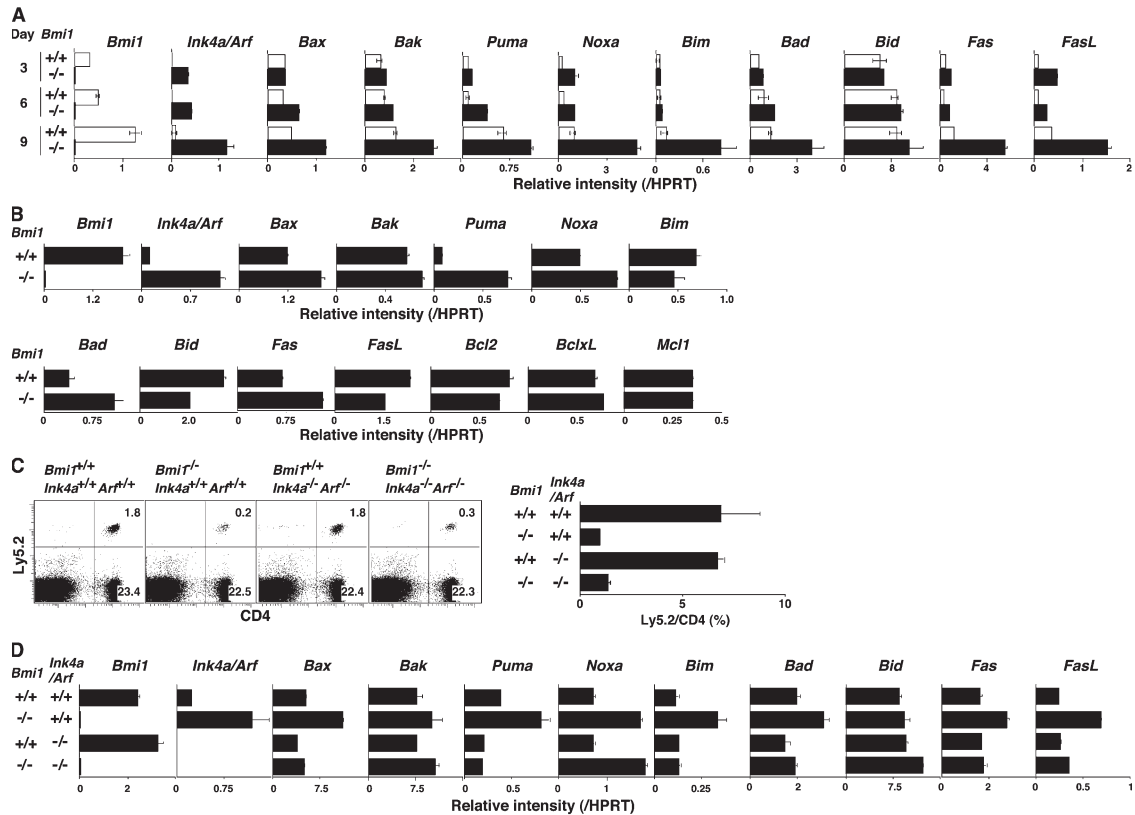
Bmi1 has been reported to promote cell proliferation, cell survival, and stem cell self-renewal by repressing the *Ink4a/Arf* locus (37). This locus codes for two proteins, p16<sup>ink4a</sup> and p19<sup>arf</sup> (Ink4a and Arf), through the use of alternative reading frames. Ink4a is a cyclin D–dependent kinase inhibitor that promotes cell cycle arrest after Rb activation. Arf induces p53 activation and p53-mediated cell death (38). In the *Bmi1*<sup>-/-</sup> effector Th2 cells, we confirmed the increased expression of



**Figure 1. Impaired generation of memory Th1/Th2 cells from *Bmi1*<sup>-/-</sup> effector Th2 cells.** (A and B) *Bmi1*<sup>+/+</sup>, *Bmi1*<sup>+/-</sup>, or *Bmi1*<sup>-/-</sup> effector Th2 (A) or Th1 (B) cells with DO11.10 Tg background were intravenously transferred into BALB/c *nu/nu* mice ( $n = 5$ ). 5 wk later, the number of KJ1<sup>+</sup> memory Th2 cells was determined by flow cytometry. Typical staining patterns of KJ1/CD4 and percentages of KJ1<sup>+</sup> cells are shown. Four independent experiments were performed with similar results. (C) *Bmi1*<sup>+/+</sup>, *Bmi1*<sup>+/-</sup>, or *Bmi1*<sup>-/-</sup> effector Th2 cells with DO11.10 Tg background were intravenously transferred into BALB/c mice ( $n = 5$ ). (D) Impaired semi-acute survival of *Bmi1*<sup>+/-</sup> Th2 cells. Thy1.1 *Bmi1*<sup>+/+</sup> (Ly5.2 background) and Thy1.2 *Bmi1*<sup>+/-</sup> (Ly5.2 background) effector Th2 cells were mixed (1:1) and transferred ( $3 \times 10^7$  cells/mouse) into syngeneic C57BL/6 mice with a Ly5.1 background. The ratio of Thy1.2<sup>+</sup>/Thy1.1<sup>+</sup> cells in Ly5.2<sup>+</sup> cells is shown ( $n = 3$ ). (E) Homeostatic proliferation of the splenic memory Th2 cells was assessed by the BrdU incorporation in vivo. Representative BrdU staining profiles of KJ1<sup>+</sup> memory Th2 cells prepared 5 wk after cell transfer are shown. The percentages of BrdU<sup>+</sup> cells with standard deviation are shown ( $n = 5$ ). Three independent experiments were performed with similar results. (F) Apoptotic cell death of freshly prepared memory Th2 cells from the spleen was measured by a TUNEL assay. As a positive control, cells were treated with DNase I. Representative TUNEL profiles are shown with percentages of TUNEL<sup>+</sup> cells. Two independent experiments were performed with similar results.

the *Ink4a/Arf* gene after anti-TCR stimulation (Fig. 2 A). An increased mRNA expression of both *Ink4a* and *Arf* was confirmed (not depicted). In addition, various proapoptotic genes, which are known to be targets for p53, such as *Bax*, *Puma*, *Noxa*, *Bim*, *Bad*, *Fas*, and *Fas ligand*, were increased in *Bmi1*<sup>-/-</sup> effector Th2 cells (Fig. 2 A). In *Bmi1*<sup>-/-</sup> memory Th2 cells, increased expression of mRNA was observed in the *Ink4a/Arf*, *Bax*, *Puma*, *Noxa*, *Bad*, and *Fas* genes, whereas expression of antiapoptotic genes *Bcl2*, *BclxL*, and *Md1* was

unchanged (Fig. 2 B). To examine whether the increased expression of *Ink4a/Arf* in *Bmi1*<sup>-/-</sup> Th2 cells plays a role in the *Bmi1*-mediated maintenance of memory Th2 cells, *Bmi1*<sup>-/-</sup> mice were crossed with *Ink4a*<sup>-/-</sup>/*Arf*<sup>-/-</sup> mice, and effector Th2 cells generated from *Bmi1*<sup>-/-</sup>/*Ink4a*<sup>-/-</sup>/*Arf*<sup>-/-</sup> with Ly5.2 background were transferred into Ly5.1 mice. 5 wk later, the number of Ly5.2<sup>+</sup> memory Th2 cells was assessed. Depletion of *Ink4a* and *Arf* genes itself had no effect on the generation of memory T cells (Fig. 2 C, panels 1 and 3) and failed to



**Figure 2. Deletion of the *p16<sup>Ink4a</sup>* and *p19<sup>Arf</sup>* genes failed to rescue the impaired generation of memory Th2 cells in the absence of *Bmi1*.** (A) mRNA expression of *Ink4a/Arf* and p53-related proapoptotic genes in *Bmi1*<sup>-/-</sup> effector Th2 cells was determined by quantitative RT-PCR. The relative intensity (/HPRT; mean of three samples) is shown with standard deviations. Three independent experiments were performed with similar results. (B) mRNA expression of *Ink4a/Arf*, p53-related proapoptotic genes, and antiapoptotic genes in *Bmi1*<sup>-/-</sup> memory Th2 cells was analyzed. The relative intensity (/HPRT; mean of three samples) is shown with standard deviations. Two independent experiments were performed with similar results. (C) Effects on *p16<sup>Ink4a</sup>* and *p19<sup>Arf</sup>* deficiency on the memory Th2 cell generation. The effector Th2 cells from the indicated mice (Ly5.2 background) were transferred into Ly5.1 host mice. 5 wk after cell transfer, the number of Ly5.2<sup>+</sup> memory Th2 cells was determined. The mean values are shown with standard deviations (*n* = 5; right). The experiments were performed twice with similar results. (D) mRNA levels of proapoptotic genes in *Bmi1*<sup>-/-</sup>/*Ink4a*<sup>-/-</sup>/*Arf*<sup>-/-</sup> effector Th2 cells were determined by quantitative RT-PCR. The relative intensity (/HPRT; mean of three samples) is shown with standard deviations. The experiments were performed twice with similar results.

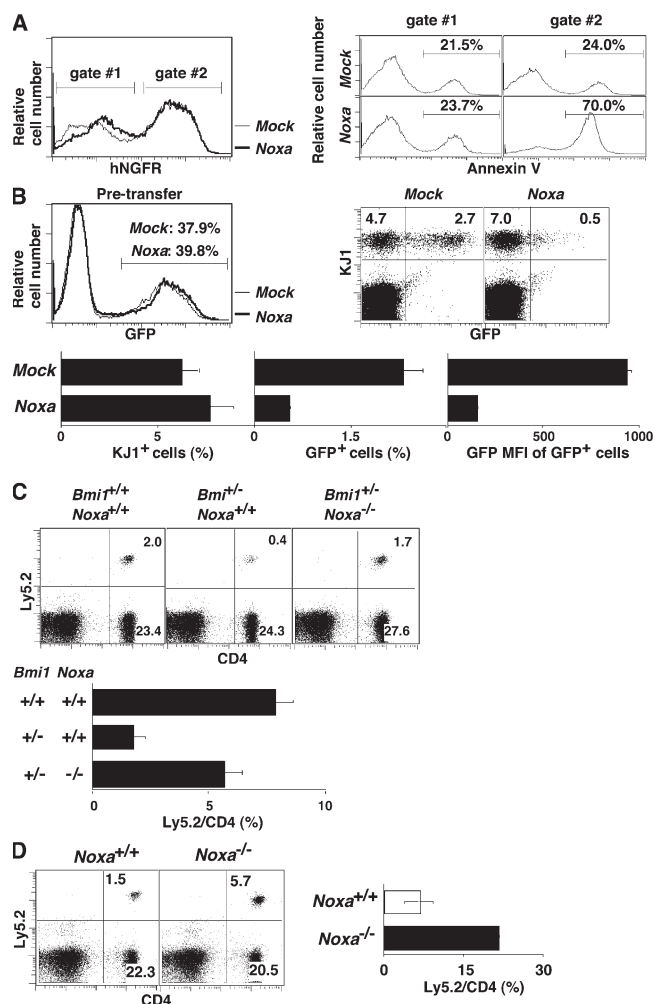
restore memory Th2 cell generation in *Bmi1*<sup>-/-</sup> background (Fig. 2 C, panels 2 and 4). Furthermore, the deletion of the *p53* gene failed to rescue the *Bmi1*<sup>+/-</sup> memory Th2 cell numbers (Fig. S6, available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>). These results suggest that the *Ink4/Arf* and *p53*-dependent pathways are not key pathways responsible for the reduced memory Th2 cell generation in *Bmi1*<sup>-/-</sup> memory mice. To identify any possible proapoptotic genes whose expression is regulated by *Bmi1* but not regulated by the *Ink4/Arf* and *p53*-dependent pathways, the expression levels of these proapoptotic genes were assessed in the *Bmi1*<sup>-/-</sup>/*Ink4a*<sup>-/-</sup>/*Arf*<sup>-/-</sup> Th2 cells. Among these proapoptotic genes, the level of *Noxa* mRNA increased in *Bmi1*<sup>-/-</sup> Th2 cells and remained high in *Bmi1*<sup>-/-</sup>/*Ink4a*<sup>-/-</sup>/*Arf*<sup>-/-</sup> background, whereas other targets such as *Bax*, *Puma*, *Bim*, *Bad*, *Fas*, and *Fas ligand* were at the normal expression levels (Fig. 2 D). Increased expression of *Noxa* mRNA was observed also in effector Th1, Tc1, and Tc2 cells in the absence of *Bmi1* (Fig. S7). These results indicate that the

expression of *Noxa* is suppressed by the expression of *Bmi1*, but it is independent from the control of the *Ink4/Arf* and *p53* pathways. The results thus far obtained suggest that *Bmi1* controls the generation of both Th1 and Th2 memory cells accompanied with an increased expression of *Noxa*. Our previous study indicated that the expression of *Bmi1* is higher in Th2 cells than Th1 cells, and *Bmi1* controls the generation of Th2 cells more profoundly as compared with Th1 cells (33). Therefore, to better understand the molecular mechanisms underlying the *Bmi1/Noxa*-mediated regulation of memory CD4 T cell generation, we focused much of our studies on memory Th2 cells.

### The expression of *Noxa* controls the generation of memory Th2 cells

Consequently, to test whether the changes in the level of *Noxa* affect the induction of cell death, effector Th2 cells were infected with a retrovirus containing the human *NGFR* and *Noxa*, and the number of annexin V<sup>+</sup> cells was assessed





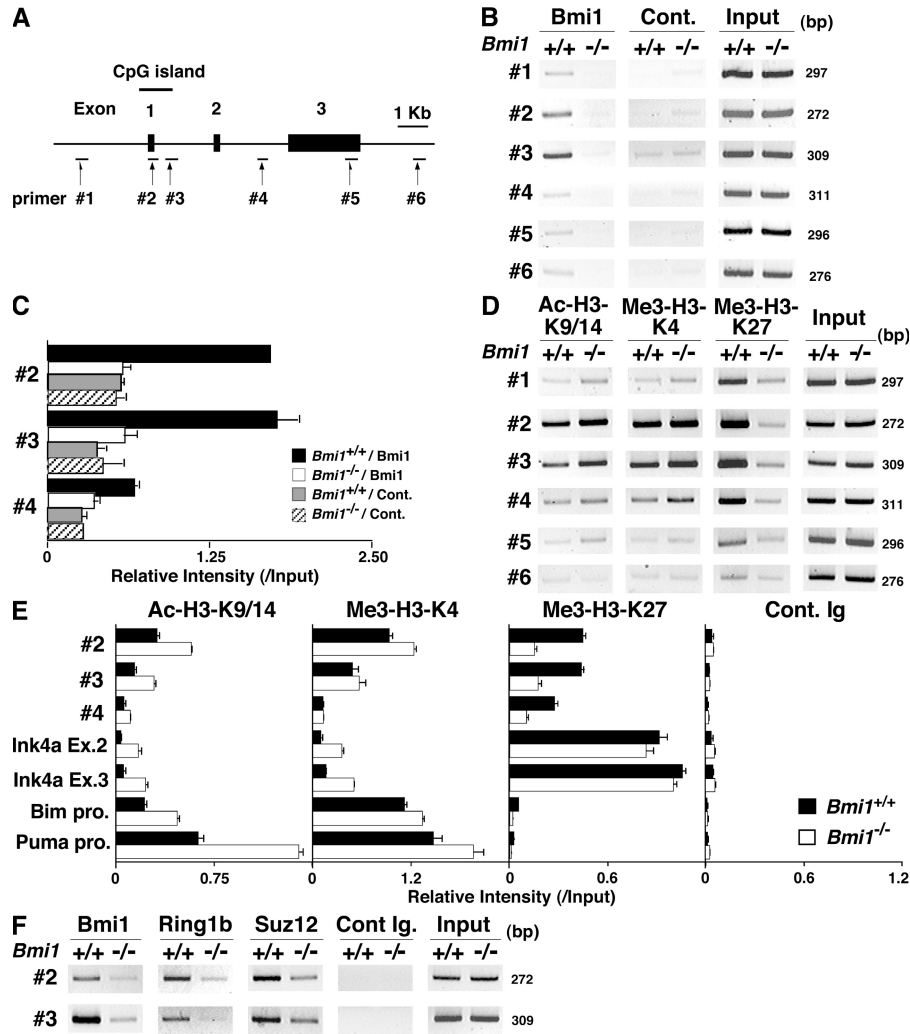
**Figure 3. Expression level of Noxa controls the generation of memory Th2 cells.** (A) Enforced expression of Noxa-induced cell death in effector Th2 cells after cytokine depletion. Effector Th2 cells infected with a *Noxa-IRES-hNGFR*-containing retrovirus were cultured in vitro for 24 h without cytokines. hNGFR profiles (left) and annexin V staining profiles of the electronically gated hNGFR<sup>+</sup> (gate #2) and hNGFR<sup>-</sup> (gate #1) populations are shown. Three independent experiments were performed with similar results. (B) KJ1<sup>+</sup> effector Th2 cells infected with *Noxa-IRES-EGFP*-containing retrovirus were transferred into BALB/c *nu/nu* mice. 5 wk later, memory Th2 cell generation was determined by KJ1/EGFP expression. Expression of EGFP in pretransferred effector Th2 cells (top left) and a typical KJ1/GFP profile of freshly prepared memory Th2 cells (top right) are shown. In the bottom panels, the percentages of KJ1<sup>+</sup> cells and GFP<sup>+</sup> Noxa-overexpressing cells and the mean fluorescence intensity of the GFP<sup>+</sup> cells are shown with standard deviations ( $n = 4$ ). The experiments were performed twice with similar results. (C) The effector Th2 cells from *Bmi1*<sup>+/+</sup>/*Noxa*<sup>+/+</sup>, *Bmi1*<sup>+/-</sup>/*Noxa*<sup>+/+</sup>, and *Bmi1*<sup>+/-</sup>/*Noxa*<sup>-/-</sup> mice (Ly5.2) were transferred into Ly5.1 host mice, and the number of Ly5.2<sup>+</sup> memory Th2 cells was determined. A typical staining pattern of CD4/Ly5.2 (top) and the percentages of Ly5.2<sup>+</sup> cells among CD4 T cells are shown with standard deviations ( $n = 5$ ; bottom). Three independent experiments were performed with similar results. (D) Deletion of the *Noxa* gene enhanced the generation of memory Th2 cells. In vitro-generated *Noxa*<sup>-/-</sup> effector Th2 cells (Ly5.2) were transferred into Ly5.1 host mice. 5 wk after cell transfer, the number of Ly5.2<sup>+</sup> memory Th2 cells was determined. A repre-

sentative CD4/Ly5.2 profile (left) and the mean values with standard deviations ( $n = 5$ ; right) are shown. The experiments were performed twice with similar results.

after in vitro suspension culture for 24 h without cytokines. *Noxa*-introduced cells showed increased annexin V<sup>+</sup> staining (70.0%) as compared with mock-infected (24.0%) or noninfected cell populations (Fig. 3 A). Next, KJ1<sup>+</sup> effector Th2 cells infected with a retrovirus containing the *GFP* and *Noxa* genes were transferred into BALB/c *nu/nu* mice, and the numbers of Noxa-expressing (KJ1<sup>+</sup>GFP<sup>+</sup>) memory Th2 cells were assessed. Although the generation of KJ1<sup>+</sup> memory Th2 cells was not affected, the number of KJ1<sup>+</sup>GFP<sup>+</sup> memory Th2 cells and their GFP mean fluorescence intensity were clearly reduced in the Noxa-transduced group (Fig. 3 B). A gene dose-dependent increase in the expression of Noxa was detected in *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>-/-</sup> memory Th2 cells (Fig. S8, available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>). *Bmi1*<sup>-/-</sup>*Noxa*<sup>-/-</sup> mice were not born despite extensive breeding attempts. Consequently, we used *Bmi1*<sup>-/-</sup>*Noxa*<sup>-/-</sup> mice and found that memory Th2 cell generation in the mice transferred with *Bmi1*<sup>+/-</sup> effector Th2 cells was restored by the deletion of the *Noxa* gene (Fig. 3 C). Furthermore, enhanced generation of memory Th2 cells was observed in the mice that received *Noxa*<sup>-/-</sup> effector Th2 cells (Fig. 3 D). Thus, we concluded that the reduction in the number of *Bmi1*<sup>-/-</sup> memory Th2 cells is at least in part due to the increased expression of Noxa in *Bmi1*<sup>-/-</sup> Th2 cells.

### Bmi1 directly binds to the *Noxa* gene locus and regulates the histone modification

To assess the molecular mechanisms underlying the *Bmi1*-mediated repression of the *Noxa* gene, we performed chromatin immunoprecipitation (ChIP) assays with six primer pairs covering the *Noxa* gene (Fig. 4 A). The accumulation of *Bmi1* was observed around the CpG island (Fig. 4 B, #2 and #3) of the *Noxa* locus. The binding of *Bmi1* was confirmed by a ChIP assay with a quantitative PCR system (Fig. 4 C). Equivalent binding of *Bmi1* was observed in Th1 cells (Fig. S9, available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>). Histone H3-K9/14 acetylation and tri-methylation of histone H3-K4 were observed also around the CpG island (Fig. 4 D, #2 and #3). More interestingly, histone H3-K27 was highly tri-methylated in wild-type Th2 cells over broader regions in the *Noxa* locus, and the tri-methylation was apparently reduced in the *Bmi1*<sup>-/-</sup> Th2 cells (Fig. 4 D). The changes in histone modifications were also assessed using a quantitative PCR system. Modest increase in histone H3-K9/14 acetylation and a substantial decrease in H3-K27 tri-methylation at the *Noxa* gene locus were confirmed in the *Bmi1*<sup>-/-</sup> Th2 cells (Fig. 4 E) and *Bmi1*<sup>-/-</sup> Th1 cells (Fig. S10). Interestingly, the levels of histone H3-K27 tri-methylation at the *Ink4a* gene locus, another target gene of *Bmi1*, were not affected by the depletion of *Bmi1* (Fig. 4 E). The levels of histone H3-K27 tri-methylation at the promoter regions of the



**Figure 4. Bmi1 binds to the *Noxa* gene and regulates tri-methylation of H3-K27.** (A) Schematic representation of the *Noxa* gene locus. The location of primers (#1 to #6), a CpG island, and exons are indicated. (B) A ChIP assay of the *Noxa* gene locus was performed using anti-Bmi1 antibody (*Bmi1*) and control antibody (Cont.) in *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>-/-</sup> Th2 cells. Three independent experiments were performed with similar results. (C) A ChIP assay of the *Noxa* gene locus was performed as in B, and the levels of binding were assessed by a quantitative PCR analysis. (D) Histone modifications (Ac-H3-K9/K14; acetylation of histone H3-K9/K14 and Me3-H3-K4; tri-methylation of histone H3-K4 and Me3-H3-K27; tri-methylation of histone H3-K27) at the *Noxa* gene locus in *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>-/-</sup> effector Th2 cells. (E) A ChIP assay of the *Noxa* (#2 to #4), *Ink4a* (Ex.2, exon 2; Ex.3, exon 3), *Bim* (*Bim* pro., *Bim* promoter), and *Puma* (*Puma* pro., *Puma* promoter) locus was performed as in D, and the levels of binding were assessed by a quantitative PCR analysis. (F) The association of Ring1B and Suz12 at the *Noxa* gene locus in *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>-/-</sup> Th2 cells. The association of Bmi1, Ring1B, and Suz12 was determined by a ChIP assay using specific antibodies.

*Bim* and *Puma* were not significantly detected in the presence or absence of Bmi1. The binding of other PcG gene products, such as Ring1b and Suz12, was detected around the same regions (Fig. 4 F, #2 and #3), and the binding was substantially decreased in the absence of Bmi1. Thus, a PcG gene product complex containing Bmi1 appears to bind to the *Noxa* gene directly and regulate histone modifications, such as tri-methylation of H3-K27 in Th1 and Th2 cells.

**Bmi1 controls the CpG methylation at the *Noxa* gene locus and represses the mRNA expression of *Noxa***

Next, to study the levels of CpG DNA methylation around the CpG island of the *Noxa* gene, a methylated DNA immuno-

precipitation (MeDIP) assay was performed. As shown in Fig. 5 A, the 5' region of the CpG island (promoter and #2) was methylated in wild-type effector Th2 cells, and the methylation levels were very low in the *Bmi1*<sup>-/-</sup> cells. A DNA methyltransferase Dnmt1 was found to bind at the CpG island (Fig. 5 B, #2 and #3), and the binding was also dependent on Bmi1 in Th2 cells (Fig. 5 B) and Th1 cells (Fig. S11, available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>). To assess a role of CpG methylation on *Noxa* expression, effector Th2 cells were treated with 5-Aza-2'-deoxycytidine (5-Aza), an inhibitor of CpG DNA methylation. As expected, the expression of *Noxa* mRNA was dramatically increased (Fig. 5 C) accompanied by the reduction of a tri-methylation

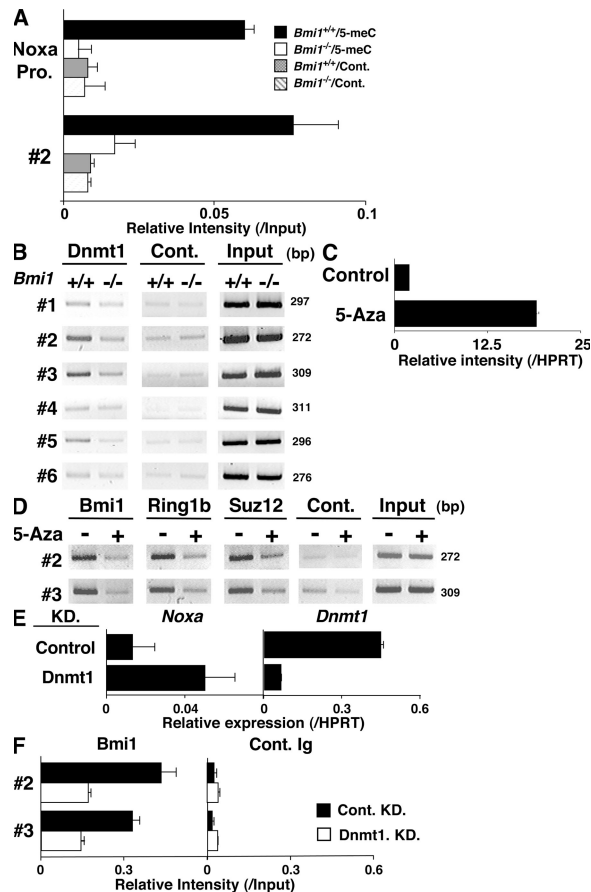
level of histone H3-K27 (Fig. S12). Furthermore, the binding of Bmi1, Ring1b, and Suz12 was substantially reduced by the treatment with 5-Aza (Fig. 5 D). A similar increase in the expression of *Noxa* and the 5-Aza-dependent reduction in the binding of Bmi1 were observed in Th1 cells (Fig. S13). Finally, we established a knockdown system for Dnmt1 using a cultured T cell line, TG40, as described in Materials and methods. Up-regulation of *Noxa* mRNA (Fig. 5 E) and the reduction of Bmi1 binding at the *Noxa* gene locus (Fig. 5 F) were induced by the introduction of shRNA for Dnmt1. Enhanced *Noxa* mRNA expression was induced in primary Th2 cells by the treatment with a DNA methyltransferase inhibitor, RG108 (Fig. S14). These results indicate that a DNA methyltransferase Dnmt1 plays an important role in the recruitment of PcG gene products and the expression of the *Noxa* gene in Th2 cells.

### The expression of Bmi1 is required for memory Th2 cell-dependent immune responses and inflammation in vivo

Finally, to examine functional defects in *Bmi1*<sup>-/-</sup> memory Th2 mice, we used a memory Th2 cell-dependent allergic airway inflammation model (36). Memory Th2 mice were generated and simply challenged by inhalation with OVA four times. The OVA-specific IgE and IgG1 (Th2-dependent isotypes) antibody production were decreased in the *Bmi1*<sup>-/-</sup> memory Th2 mice, whereas only a marginal decrease in the levels of Th1-dependent IgG2a was seen (Fig. 6 A). Next, we examined the levels of airway inflammation after OVA inhalation. The extent of inflammatory leukocyte infiltration in the peri-bronchiolar region (Fig. 6 B) and the infiltrated eosinophils, lymphocytes, and macrophages in the bronchioalveolar lavage (BAL) fluid (Fig. 6 C) was reduced significantly in the *Bmi1*<sup>-/-</sup> memory Th2 mice as compared with wild-type mice. The expression of Th2 cytokines (IL-4, IL-5, and IL-13) and Eotaxin 2 in the lung of OVA-inhaled *Bmi1*<sup>-/-</sup> memory Th2 mice was also reduced (Fig. 6 D). The periodic-acid-Schiff staining and the measurement of mRNA expression of *Gob5*, *Muc5a/c*, and *Muc5b* in the lung indicated a decrease in mucus hyperproduction in *Bmi1*<sup>-/-</sup> memory Th2 mice (Fig. 7, A and B). Furthermore, the airway hyperresponsiveness measured using a whole-body plethysmograph was not significantly induced in the *Bmi1*<sup>-/-</sup> memory Th2 mice (Fig. 7 C). In addition, by a direct invasive assay for lung resistance (RL), increase in the RL and decrease in the dynamic compliance were observed in the *Bmi1*<sup>-/-</sup> memory Th2 mice (Fig. 7 D). Collectively, these results indicate that the memory Th2 cell-dependent allergic responses were thus compromised in the *Bmi1*<sup>-/-</sup> memory Th2 mice. We also assessed the eosinophilic infiltration in DO11.10 Tg *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>+/-</sup> mice without cell transfer, and as expected, the level of eosinophilic infiltration was significantly milder in the *Bmi1*<sup>+/-</sup> mice (not depicted).

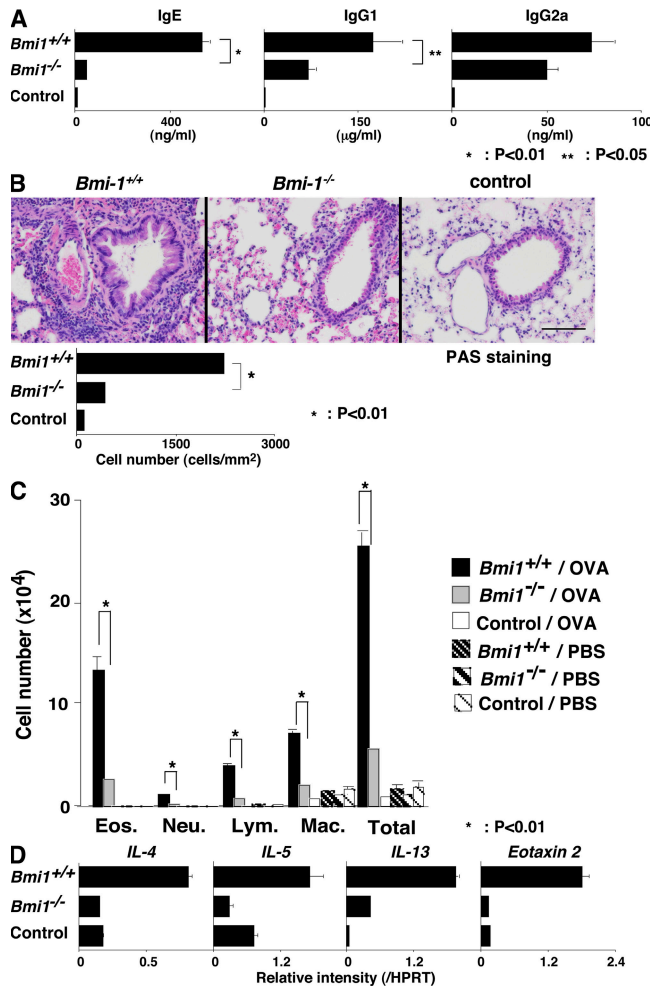
### DISCUSSION

Here, we demonstrate that Bmi1 plays a crucial role in the generation and maintenance of memory CD4 T cells. Such



**Figure 5. Bmi1 is required for the DNA CpG methylation of the *Noxa* gene.** (A) Decreased DNA CpG methylation at the CpG island of the *Noxa* gene locus in *Bmi1*<sup>-/-</sup> Th2 cells. DNA CpG methylation level was determined by MeDIP assay. MeDIP assay was performed using an anti-5-methyl cytidine antibody (5-mec) and a control antibody (Cont.). Mean values with standard deviation are shown ( $n = 3$ ). (B) The association of Dnmt1, a DNA methyltransferase, at the CpG island of the *Noxa* gene locus. The association of Dnmt1 was determined by a ChIP assay using a specific antibody. (C) Increased mRNA expression of *Noxa* after the treatment with 5-Aza, an inhibitor of DNA methyltransferase. Th2 cells were treated with 5-Aza for 2 d, and *Noxa* mRNA expression was determined by a quantitative RT-PCR system. Three independent experiments were performed with similar results. (D) Dissociation of the PcG complex from the *Noxa* gene locus after the treatment with 5-Aza. The association of Bmi1, Ring1B, and Suz12 was determined by a ChIP assay using specific antibodies. Three independent experiments were performed with similar results. (E) mRNA expression of *Noxa* and *Dnmt1* in a Dnmt1 knockdown (KD) T cell line. Dnmt1-KD and control-KD stable TG40 cell lines were generated using a lentivirus gene transduction system as described in Materials and methods. The levels of mRNA were determined by a quantitative RT-PCR. The relative intensity (/HPRT; mean of three samples) is shown with standard deviations. (F) Decreased binding of Bmi1 at the *Noxa* gene in Dnmt1-KD TG40 cells. A ChIP assay with an anti-Bmi1 antibody was performed, and the levels of Bmi1 binding were assessed by a quantitative PCR analysis.

a Bmi1-mediated regulation was seen in both memory Th1 and Th2 cells. In the absence of Bmi1, the generation of both Th1 and Th2 memory cells was impaired (Fig. 1 and Fig. S2)



**Figure 6. Defects in the memory Th2 cell-dependent immune responses and inflammation in *Bmi1*<sup>-/-</sup> memory Th2 mice.** OVA-specific *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>-/-</sup> effector Th2 cells with DO11. 10 Tg background were intravenously transferred into BALB/c *nu/nu* mice. 5 wk later, the mice were challenged four times by inhalation with OVA on days 0, 2, 8, and 10. (A) The serum concentrations of OVA-specific Igs with the indicated isotype after OVA inhalation (on day 11) were determined by ELISA. The mean values with standard deviations (five animals per group) are shown. \*, P < 0.01; \*\*, P < 0.05. The control represents BALB/c *nu/nu* mice without Th2 cell transfer. The experiments were performed twice with similar results. (B) On day 11, the lungs were fixed and stained with hematoxylin and eosin (HE). The number of infiltrated leukocytes in the peribronchiolar region (mean cell numbers/mm<sup>2</sup> with standard deviations; n = 5) is also shown (bottom). The experiments were performed twice with similar results. Bars, 100 μm. (C) On day 12, BAL fluid was collected and May-Grunwald-Giemsa staining was performed. The absolute cell numbers of eosinophils (Eos.), neutrophils (Neu.), lymphocytes (Lym.), and macrophages (Mac.) in the BAL fluid are shown with standard deviations (n = 5). The results were obtained using the values of cell counting, the percentage of the cells, the total cell number per milliliter, and the volume of the BAL fluid recovered. \*, P < 0.01. The experiments were performed twice with similar results. (D) The mRNA expression of *IL-4*, *IL-5*, *IL-13*, and *Eotaxin 2* was determined by quantitative RT-PCR. The relative intensity (/HPRT; mean of three samples) is shown with standard deviations. The experiments were performed twice with similar results.

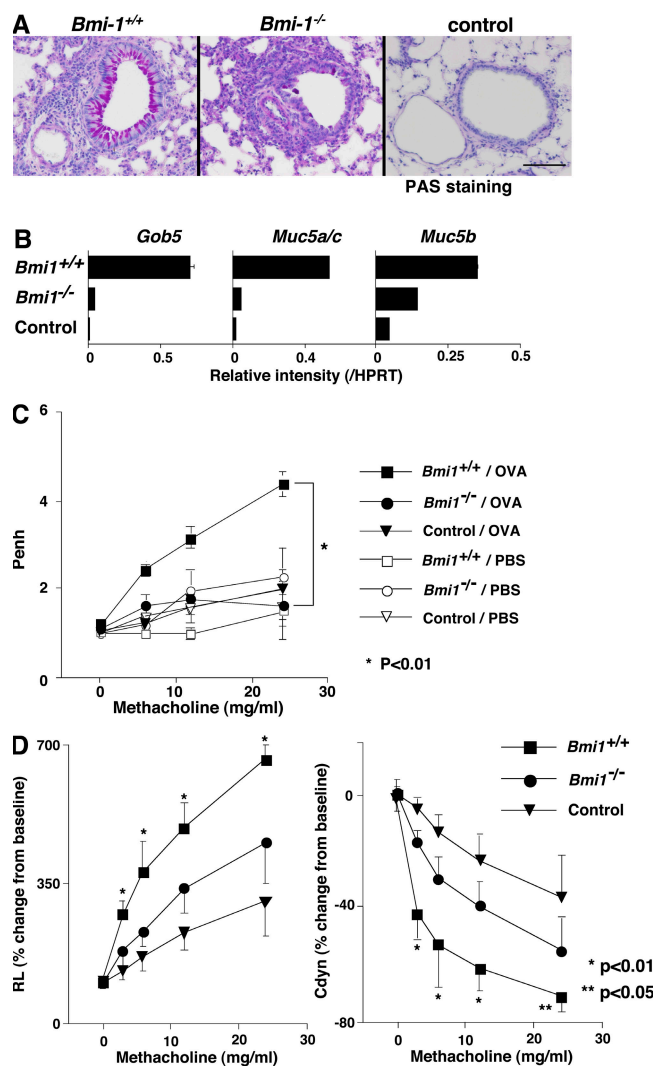
by the increased Noxa expression (Fig. 2 and Fig. S7). *Bmi1* binds to the *Noxa* gene locus in both Th1 and Th2 cells (Fig. 4 and Fig. S9), and directly represses its transcription to promote memory Th2 cell survival. The involvement of H3-K27 tri-methylation and DNA CpG methylation in the repression of *Noxa* was revealed. The Th2-dependent allergic airway inflammatory responses were compromised in *Bmi1*<sup>-/-</sup> memory Th2 mice, suggesting a physiological role of *Bmi1* in the establishment of Th2 cell-mediated memory responses.

Noxa is a member of a BH3-only protein family that initiates programmed cell death in various cells, including lymphocytes (39–41). Noxa is known to regulate selectively the pro-survival activity of Mcl1 and A1/Bfl-1 (42), and an important role of Mcl1 in the survival of lymphocytes was reported (43). The mRNA expression of antiapoptotic genes Mcl1, Bclx, and BclxL was not changed in *Bmi1*<sup>-/-</sup> memory (Fig. 2 B) and induced normally in effector Th2 cells by the IL-7 treatment (unpublished data). Enforced expression of Noxa in effector Th2 cells resulted in the decreased generation of memory Th2 cells (Fig. 3 B), suggesting that the overproduction of Noxa prevents memory Th2 cell generation even in the presence of normal abundance of antiapoptotic proteins. Thus, it is likely that *Bmi1* attenuates Noxa-mediated inhibition of Mcl1 pro-survival activity and promotes memory T cell generation.

Noxa was originally identified as a downstream target of p53 (44). Arf/Mdm2-mediated stabilization of p53 protein and the resulting mRNA expression of the p53 target genes were reported (45, 46). Therefore, it was likely that the enhancement of cell death observed in *Bmi1*<sup>-/-</sup> Th2 cells is p53 dependent. However, the deletion of the *p53* gene failed to rescue the decreased generation of memory Th2 cells in the absence of *Bmi1* (Fig. S6). Thus, an increased apoptotic cell death observed in *Bmi1*<sup>-/-</sup> memory Th2 cells appears to be p53 independent.

The *Bmi1*-mediated repression of the *Ink4a/Arf* gene observed in the hematopoietic and neural stem cells (45, 46) appears to operate in Th2 cells (Fig. 2, A and B). It is known that Arf induces p53 activation (38), resulting in the induction of p53-dependent genes, including Puma and Bim. An important role of Puma and Bim in the cell death of activated T cells was reported (47, 48), and the expression of these genes was up-regulated in *Bmi1*<sup>-/-</sup> effector and memory Th2 cells (Fig. 2, A and B). However, *Bmi1*<sup>-/-</sup>/*Ink4a*<sup>-/-</sup>/*Arf*<sup>-/-</sup> Th2 cells, in which the levels of Puma and Bim expression were not increased (Fig. 2 D), failed to generate normal numbers of memory Th2 cells (Fig. 2 C). Although the expression of *Puma* and *Bim* mRNA was up-regulated, the levels were considerably low compared with Noxa in *Bmi1*<sup>-/-</sup> effector Th2 cells (Fig. 2 A). This might explain the predominant effect of Noxa in the cell death of *Bmi1*<sup>-/-</sup> Th2 cells. However, the current experimental data do not allow us to make any conclusions to be drawn in regards to the relative contribution of each of the factors in the survival of CD4 T cells. Thus, although Puma and Bim may play a role in the survival of memory Th2 cells, Noxa appears to be a





**Figure 7. Decreased mucus hyperproduction and airway hyperresponsiveness in *Bmi1*<sup>-/-</sup> memory Th2 mice.** *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>-/-</sup> memory Th2 mice were challenged by inhalation with OVA as in Fig. 5. (A) 1 d after the last OVA inhalation (day 11), the lungs were fixed and stained with periodic-acid-Schiff (PAS). A representative staining pattern is shown. The control represents BALB/c *nu/nu* mice without Th2 cell transfer. Bars, 100  $\mu$ m. (B) On day 12, total RNA was prepared from the lung, and the expression of *Gob5*, *Muc5a/c*, and *Muc5b* (molecular makers for Goblet cell hyperplasia and mucus production) was determined by a quantitative PCR analysis. The relative intensity (HPRT; mean of three samples) is shown with standard deviations. (C) OVA-induced airway hyperresponsiveness in *Bmi1*<sup>-/-</sup> memory Th2 mice. On day 11, the airway hyperresponsiveness in response to increasing doses of methacholine was measured in a whole-body plethysmograph. The mean values ( $n = 5$ ) are shown with standard deviations. PBS, PBS-inhaled control; OVA, OVA-inhaled. \*,  $P < 0.01$ . The experiments were performed twice with similar results. (D) On day 11, changes in the RL (left) and the dynamic compliance (Cdyn; right) were assessed. Mean values (six mice per group) are shown with standard deviations. \*,  $P < 0.01$ ; \*\*,  $P < 0.05$ .

critical target for the Bmi1-mediated regulation of memory Th2 cell survival.

Semi-acute survival of transferred *Bmi1*<sup>-/-</sup> effector Th2 cells was substantially impaired (Fig. 1 D and Figs. S3–S5).

On the other hand, high numbers of apoptotic cells were detected in the spleen even 5 wk after effector *Bmi1*<sup>-/-</sup> Th2 cell transfer (Fig. 1 F). The extent of the defect in homeostatic proliferation was modest (Fig. 1 E). The low number of *Bmi1*<sup>-/-</sup> memory CD4 T cells persisted for at least 2 mo (unpublished data). These results suggest that the long-term survival of memory Th2 cells is also dependent on the expression of Bmi1. Collectively, we conclude that Bmi1 controls both semi-acute survival of effector Th2 cells and long-term survival of memory Th2 cells.

The PRC2 was reported to possess an intrinsic histone H3-K27 methyltransferase activity (26–29). The PRC1, including Bmi1 and Ring1, was shown to possess an activity of histone H2A ubiquitination (49–51), whereas there has been no report indicating that the PRC1 possess an intrinsic histone H3-K27 methyltransferase activity. Accumulating evidence supports a sequential binding model (52), in which PRC2-mediated H3-K27 methylation serves as a binding site for the recruitment of PRC1 complex through the specific recognition of the H3-K27 methyl mark by the chromo domain of the polycomb protein, such as M33 (53, 54). In our study, however, the levels of tri-methylation of histone H3-K27 were substantially decreased in *Bmi1*<sup>-/-</sup> Th1 and Th2 cells (Fig. S10 and Fig. 4, D and E). In addition, the levels of binding of Suz12, a component of PRC2, at the *Noxa* gene locus were decreased in *Bmi1*<sup>-/-</sup> Th2 cells (Fig. 4 F). Thus, in Th2 cells, Bmi1 may play an important role for the recruitment of PRC2 and subsequent histone H3-K27 methylation at the *Noxa* gene locus. Alternatively, the Bmi1-containing PcG complex in Th2 cells may associate with a unique molecule possessing an intrinsic histone H3-K27 methyltransferase activity. In any event, tri-methylation of histone H3-K27 at the *Noxa* gene locus is strictly regulated by the expression of Bmi1.

The expression of Bmi1 is required for the DNA CpG methylation of the *Noxa* gene (Fig. 5 A). The binding of Dnmt1 at the CpG island was Bmi1 dependent (Fig. 5 B and Fig. S11). The treatment with a 5-Aza resulted in the dissociation of Bmi1, Ring1B, and Suz12 from the CpG island (Fig. 5 D) and the up-regulation of *Noxa* mRNA (Fig. 5 C and Fig. S13 A). Increased expression of *Noxa* mRNA and reduced binding of Bmi1 were confirmed in Dnmt1 knock-down T cells (Fig. 5, E and F). Recently, the PcG complex was reported to be associated with DNA methyltransferases, including Dnmt1 (55, 56). Thus, a DNA methyltransferase, such as Dnmt1, may play a critical role in the recruitment and the repressive function of the PcG complex at the *Noxa* gene.

In this study, we demonstrate that the generation and maintenance of memory Th1/Th2 cells (Fig. 1) and the memory Th2 cell-dependent airway inflammation are controlled by the expression of Bmi1 (Figs. 6 and 7). Our preliminary experiments demonstrate that at least Mel-18, Mph1/Rae28, and M33 appear to be involved in the regulation of memory Th2 cell generation (unpublished data), suggesting that the survival of memory Th2 cells is regulated by an epigenetic mechanism involving the Bmi1-containing PcG complex. We have recently reported that Bmi1 stabilizes GATA3 protein

through the direct interaction with GATA3 (33). Mel-18 is involved in the induction of GATA3 expression in developing Th2 cells (32). Collectively, PcG gene products appear to control the development of effector and memory Th2 cells at multiple steps in a distinct manner and govern the Th2-type immune responses and inflammation.

## MATERIALS AND METHODS

**Mice.** *Bmi1*-deficient mice were provided by M. van Lohuizen (The Netherlands Cancer Institute, Amsterdam, Netherlands) (30). *p16<sup>Ink4a</sup>/p19<sup>Arf</sup>* double-deficient mice were provided by R.A. DePinho (Harvard Medical School, Boston, MA) (37). *Noxa*-deficient mice were provided by T. Taniguchi (The University of Tokyo, Tokyo, Japan) (44). The animals used in this study were backcrossed to BALB/c or C57BL/6 mice 10 times. Anti-OVA-specific TCR- $\alpha\beta$  (DO11.10) Tg mice were provided by D. Loh (Washington University School of Medicine, St. Louis, MO) (57). All mice were used at 4–8 wk old. BALB/c and BALB/c *nu/nu* mice were purchased from Clea Inc. Ly5.1 mice were purchased from Sankyo Laboratory. All mice used in this study were maintained under specific pathogen-free conditions. All experiments using mice received approval from the Chiba University Administrative Panel for Animal Care. Animal care was conducted in accordance with the guidelines of Chiba University.

**Reagents.** The reagents used in this study are as follows: FITC- or APC-conjugated anti-CD4 mAb (GK1.5), FITC-conjugated anti-Ly5.2 mAb (104), and PE-conjugated KJ1 (anti-clonotypic mAb for DO11.10 TCR) were purchased from BD Biosciences. Anti-FcR $\gamma$ II and III mAb (2.4G2) and unconjugated anti-IL-4 mAb (11B11), anti-IL-12 mAb (C17.8), and anti-IFN- $\gamma$  (R4-6A2) were used as culture supernatants. Recombinant mouse IL-4 was from TOYOBO. The OVA peptide (residues 323–339; ISQA-VHAAHAEINEAGR) was synthesized by BEX Corporation.

**The generation of effector and memory Th1/Th2 cells.** Effector and memory Th1/Th2 cells were generated as described previously (35, 36). In brief, splenic KJ1<sup>+</sup>CD4<sup>+</sup> T cells from DO11.10 OVA-specific TCR Tg mice were stimulated with 1  $\mu$ M of an OVA peptide (Loh15) plus APC (irradiated splenocytes) under the Th1 or Th2 culture conditions for 6 d in vitro. Th1 condition: 25 U/ml IL-2, 10 U/ml IL-12, and anti-IL-4 mAb. Th2 condition: 25 U/ml IL-2, 10 U/ml IL-4, anti-IL-12 mAb, and anti-IFN- $\gamma$  mAb. In some experiments, splenic CD4 T cells were stimulated with 3  $\mu$ g/ml of immobilized anti-TCR- $\beta$  mAb plus 1  $\mu$ g/ml anti-CD28 mAb. These effector Th1/Th2 cells ( $3 \times 10^7$ ) were transferred intravenously into BALB/c *nu/nu*, BALB/c, or Ly5.1 C57BL/6 recipient mice. 5 wk after cell transfer, the generation of memory Th2 cells was assessed using donor cell-specific mAbs (KJ1 and anti-Ly5.2).

**TUNEL assay.** TUNEL assay was performed with In Situ Cell Detection kit (Roche).

**Measurement of BrdU incorporation in vivo.** The memory Th2 mice were treated twice with 1 mg BrdU on days 0 and 2. BrdU incorporation in splenic KJ1<sup>+</sup> memory Th2 cells was assessed on day 4 using the BrdU Flow kit (BD Biosciences).

**5-Aza treatment.** Developing Th2 cells were treated with 10  $\mu$ M 5-Aza (Sigma-Aldrich) for 3 d, and then total RNA was prepared.

**ELISA.** Serum OVA-specific Ig concentrations were determined by ELISA as described previously (36).

**Quantitative RT-PCR.** Total RNA was isolated using the TRIzol reagent (Invitrogen). cDNA was synthesized using oligo (dT) primer and Superscript II RT (Invitrogen). Quantitative RT-PCR was performed as described pre-

viously using ABI PRISM 7000 Sequence Detection System (36). The primers for TaqMan probes for the detection were purchased from Applied Biosystems. The expression was normalized using the HPRT signal.

**Retrovirus infection.** Retrovirus vector, pMXs-IRES-GFP, was provided by T. Kitamura (The University of Tokyo, Tokyo, Japan). The method for the generation of virus supernatant and the infection was described previously (32). Infected cells were collected 4 d after infection and transferred into recipient mice.

**Lentivirus infection.** Lentivirus vectors, pLKO.1 (SHC002) and pLKO.1 mouse *Dnmt1* (TRCN39024), were purchased from Sigma-Aldrich. pCAG-HIVgp and pCMV-VSV-G-RSV-Rev vectors were provided by H. Miyoshi (RIKEN Bioresource Center, Ibaraki Japan). Recombinant lentiviruses were generated using a three-plasmid system as described previously (58). In brief, 293 T cells were transfected with self-inactivating lentiviral vector, pCAG-HIVgp vector, and pCMV-VSV-G-RSV-Rev vector. Virus containing culture supernatant was collected 48 h after transfection and used for infection. T cell line TG40 ( $2.5 \times 10^5$ /well) was infected and selected using puromycin.

**ChIP assay.** ChIP assay was performed as described previously (59). The antibodies used ChIP assay are as follows; anti-trimethyl histone H3-K4 (ab8580; Abcam), anti-trimethyl histone H3-K9 (Abcam), anti-trimethyl histone H3-K27 (Millipore), anti-*Bmi1* (Santa Cruz Biotechnology, Inc.), anti-*Dnmt1* (sc-20701; Santa Cruz Biotechnology, Inc.), anti-Suz12 (Abcam), and normal rabbit IgG (Santa Cruz Biotechnology, Inc.). An mAb specific for mouse Ring1b was provided by H. Koseki (Riken Research for Allergy and Immunology, Yokohama, Japan). The specific primers used in ChIP assay are shown in Table S1, which is available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>.

**MeDIP.** MeDIP was performed using a METHYL kit (Diagenode). In brief, genomic DNA was purified from effector Th2 cells and sheared by sonication to reduce DNA lengths to between 200 and 1,000 bp. The sheared DNA was diluted and incubated with antiserum specific for the 5-methyl cytidine. Next, immune complexes were precipitated with protein A agarose. The precipitated DNA was purified using QIAquick PCR Purification kit (QIAGEN).

**Assessment of memory Th2 cell function in vivo.** OVA-specific wild-type and *Bmi1*<sup>-/-</sup> effector Th2 cells ( $10^7$  cells) were intravenously transferred into BALB/c *nu/nu* mice. 5 wk after cell transfer, the mice were challenged four times by inhalation with OVA on days 0, 2, 8, and 10. The serum Ig levels, lung histology, mRNA expression in the lung, and airway hyperresponsiveness were then assessed on day 11 as described previously (36, 60). BAL fluid was collected on day 12.

**Statistical analysis.** Student's *t* test was used.

**Online supplemental material.** Fig. S1 shows the phenotypic and functional characterization of *Bmi1*<sup>-/-</sup> Th2 cells. Fig. S2 shows the impaired generation of memory Th1 cells from *Bmi1*<sup>-/-</sup> effector Th1 cells. Fig. S3 depicts the kinetic analysis of the number of Th2 cells after adoptive transfer of effector Th2 cells, and Fig. S4 displays the competitive analysis for memory Th2 cell generation from *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>+/-</sup> effector Th2 cells under nonlymphopenic conditions. Fig. S5 shows a competitive analysis for memory Th2 cell generation from *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>-/-</sup> effector Th2 cells under lymphopenic conditions. Fig. S6 shows that deletion of the *p53* gene failed to restore the generation of *Bmi1*<sup>+/-</sup> memory Th2 cells. In Fig. S7, mRNA expression of *Noxa* in *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>-/-</sup> effector Th1, Th2, Tc1, and Tc2 cells is shown. Fig. S8 displays mRNA expression of the *Ink4a/Arf* and *Noxa* in *Bmi1*<sup>+/+</sup>, *Bmi1*<sup>+/-</sup>, and *Bmi1*<sup>-/-</sup> effector Th2 cells. In Fig. S9, binding of *Bmi1* at the *Noxa* gene locus in effector Th1 and Th2 cells is shown. Fig. S10 depicts histone modifications at the *Noxa* gene locus in *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>+/-</sup> effector Th1 cells. Fig. S11 shows binding of *Dnmt1* at the *Noxa* gene locus in *Bmi1*<sup>+/+</sup> and *Bmi1*<sup>+/-</sup> effector Th1 cells, and Fig. S12

shows histone modifications at the *Noxa* gene locus after 5-Aza treatment. Fig. S12 displays *Noxa* mRNA expression and the binding of Bmi1 at the *Noxa* gene locus in effector Th1 cells after 5-Aza treatment. In Fig. S14, mRNA expression of *Noxa* in Th2 cells treated with RG108, a DNA methyltransferase inhibitor is shown. Table S1 lists the primer pairs used for the ChIP assay. The online supplemental material is available at <http://www.jem.org/cgi/content/full/jem.20072000/DC1>.

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