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 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,
Mph2, and Ring1A/B are constituents of a multimeric protein complex similar to the polycomb repressive complex (PRC1) 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 GTAT3 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+ 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−/− 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−/− 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-γ–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 αβTCR transgenic (Tg) CD4 T cells from Bmi1+/+, Bmi1−/−, or Bmi1+/− 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+ memory Th1/Th2 cells were assessed in various organs. As shown in Fig. 1 A, a Bmi1 gene dose–dependent effect 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+/− groups showed marked effects. Although the expression levels of IL-7Rα and IL-2RB were slightly lower on Bmi1−/− memory Th2 cells as compared with wild-type cells, other surface marker antigens were expressed normally (Fig. S1 B). Next, KJ1+ effector Th1 cells were transferred into BALB/c nu/nu mice. The memory Th1 cell generation from Bmi1−/− effector Th1 cells was also impaired in all tissues tested (Fig. 1 B). The reduction in the Bmi1−/− 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−/− 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−/− 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−/− effector Th2 cells by a competition analysis using Bmi1+/− (Thy1.1) and Bmi1−/− (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+/−/Bmi1−/− (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−/− cells was similar in all tissues tested. The defect in semi-acute survival in Bmi1−/− 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−/−/ memory Th2 cells as compared with wild type (14.3 ± 0.4% vs. 9.1 ± 1.5%) (Fig. 1 E). Next, we assessed the extent of ongoing apoptotic cell death in vivo by a TUNEL assay. In Bmi1−/−/ 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+PI+ cells were detected in Bmi1−/− memory Th2 cells after restimulation in vitro, indicating that these memory cells are highly prone to cell death (Fig. S1 C).

Deletion of p16^ink4a and p19^arf 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^ink4a and p19^arf (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−/− effector Th2 cells, we confirmed the increased expression of
unchanged (Fig. 2 B). To examine whether the increased expression of 
Ink4a/Arf
in 
Bmi1+/H11002/H11002/H11002
Th2 cells plays a role in the 
Bmi1-mediated maintenance of memory Th2 cells, 
Bmi1+/H11002/H11002/H11002
mice were crossed with 
Ink4a+/H11002/H11002/H11002
Arf+/H11002/H11002/H11002
mice, and effector Th2 cells generated from 
Bmi1+/H11002/H11002/H11002
Ink4a+/H11002/H11002/H11002
Arf+/H11002/H11002/H11002
with Ly5.2 background were transferred into Ly5.1 mice. 5 wk later, the number of Ly5.2+ 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 1. Impaired generation of memory Th1/Th2 cells from 
Bmi1+/H11002/H11002/H11002
effector Th2 cells. (A and B) 
Bmi1+/+/
, 
Bmi1+/+/
, or 
Bmi1+/−
 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+ memory Th2 cells was determined by flow cytometry. Typical staining patterns of KJ1/CD4 and percentages of KJ1+ cells are shown. Four independent experiments were performed with similar results. (C) 
Bmi1+/+/
, 
Bmi1+/+/
, or 
Bmi1+/−
 effector Th2 cells with DO11.10 Tg background were intravenously transferred into BALB/c mice (n = 5). (D) Impaired semi-acute survival of 
Bmi1+/−
Th2 cells. Thy1.1 
Bmi1+/+/
[Ly5.2 background] and Thy1.2 
Bmi1+/−/[Ly5.2 background] effector Th2 cells were mixed (1:1) and transferred (3 × 107 cells/mouse) into syngeneic C57BL/6 mice with a Ly5.1 background. The ratio of Thy1.2+/Thy1.1+ cells in Ly5.2+ 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+ memory Th2 cells prepared 5 wk after cell transfer are shown. The percentages of BrdU+ cells with standard deviation are shown (n = 5). Two 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+ cells. Two independent experiments were performed with similar results.

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+/−
effector Th2 cells (Fig. 2 A). In 
Bmi1+/−
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 
Mcl1
was unchanged (Fig. 2 B). To examine whether the increased expression of 
Ink4a/Arf
in 
Bmi1+/−
Th2 cells plays a role in the 
Bmi1-mediated maintenance of memory Th2 cells, 
Bmi1+/−
 mice were crossed with 
Ink4a+/−/Arf+/−
 mice, and effector Th2 cells generated from 
Bmi1+/−/Ink4a+/−/Arf+/−
 with Ly5.2 background were transferred into Ly5.1 mice. 5 wk later, the number of Ly5.2+ 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
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+ cells was assessed.
Th2 cells (Ly5.2) were transferred into Ly5.1 host mice. 5 wk after cell generation of memory Th2 cells. In vitro – generated performed with similar results. (D) Deletion of the standard deviations (are shown. In the bottom panels, the percentages of KJ1 + cells and GFP + typical KJ1/GFP profile of freshly prepared memory Th2 cells (top right)

Expression of EGFP in pretransferred effector Th2 cells (top left) and a memory Th2 cell generation was determined by KJ1/EGFP expression. containing retrovirus were transferred into BALB/c nu/nu mice, and the numbers of Noxa-expressing (KJ1*GFP*) memory Th2 cells were assessed. Although the generation of KJ1* memory Th2 cells was not affected, the number of KJ1*GFP* 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 +/+ and Bmi1 −/− memory Th2 cells (Fig. S8, available at http://www.jem.org/cgi/content/full/jem.20072000/DC1). Bmi1 −/− Noxa −/− mice were not born despite extensive breeding attempts. Consequently, we used Bmi1 +/+ Noxa −/− mice and found that memory Th2 cell generation in the mice transfected with Bmi1 +/+ 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 −/− effector Th2 cells (Fig. 3 D). Thus, we concluded that the reduction in the number of Bmi1 −/− memory Th2 cells is at least in part due to the increased expression of Noxa in Bmi1 −/− 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 −/− 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 −/− Th2 cells (Fig. 4 E) and Bmi1 −/− 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 Noxa −/− effector Th2 cells (Ly5.2) were transferred into Ly5.1 host mice. 5 wk after cell transfer, the number of Ly5.2 + memory Th2 cells was determined. A representative 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.

**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 + (gate #2) and hNGFR − (gate #1) populations are shown. Three independent experiments were performed with similar results. (B) KJ1 + 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/EGFP profile of freshly prepared memory Th2 cells (top right) are shown. In the bottom panels, the percentages of KJ1 + cells and GFP + Noxa-overexpressing cells and the mean fluorescence intensity of the GFP + cells are shown with standard deviations (n = 4). The experiments were performed twice with similar results. (C) The effector Th2 cells from Bmi1 +/+ , Bmi1 −/+ , and Bmi1 −/− mice (Ly5.2) were transfected into Ly5.1 host mice, and the number of Ly5.2 + memory Th2 cells was determined. A typical staining pattern of CD4/Ly5.2 (top) and the percentages of Ly5.2 + 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 −/− effector Th2 cells (Ly5.2) were transferred into Ly5.1 host mice. 5 wk after cell transfer, the number of Ly5.2 + memory Th2 cells was determined. A representative 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.
Bmi1 regulates the CpG methylation at the Noxa gene locus and represses the miRNA expression of Noxa

Next, to study the levels of CpG DNA methylation around the CpG island of the Noxa gene, a methylated DNA immunoprecipitation (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

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 miRNA expression of Noxa

Next, to study the levels of CpG DNA methylation around the CpG island of the Noxa gene, a methylated DNA immuno-
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 $Bmi1^{-/-}$ 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 $Bmi1^{-/-}$ 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 $Bmi1^{-/-}$ 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 $Bmi1^{-/-}$ 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 $Bmi1^{-/-}$ memory Th2 mice (Fig. 7 A and B). Furthermore, the airway hyperresponsiveness measured using a whole-body plethysmograph was not significantly induced in the $Bmi1^{-/-}$ 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 $Bmi1^{-/-}$ memory Th2 mice (Fig. 7 D). Collectively, these results indicate that the memory Th2 cell–dependent allergic responses were thus compromised in the $Bmi1^{-/-}$ memory Th2 mice. We also assessed the eosinophilic infiltration in DO11.10 Tg $Bmi1^{+/-}$ and $Bmi1^{-/-}$ mice without cell transfer, and as expected, the level of eosinophilic infiltration was significantly milder in the $Bmi1^{-/-}$ mice (not depicted).

**DISCUSSION**

Here, we demonstrate that Bmi1 plays a crucial role in the generation and maintenance of memory CD4 T cells. Such 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+/−/− memory Th2 mice. (A) The serum concentrations of OVA-specific IgE and IgG1 and IgG2a were measured by ELISA. The experiments were performed twice with similar results. Bars, 100 μm. (C) On day 11, the lungs were fixed and stained with hematoxylin and eosin (HE). The number of infiltrated leukocytes in the bronchiolar region (mean cell numbers/mm2 with standard deviations; n = 5) is also shown (bottom). The experiments were performed twice with similar results. Bars, 100 μm. (D) On day 12, BAL fluid was collected and May–Grunwald–Giemsa staining was performed. The absolute cell number per milliliter and the volume of the BAL fluid recovered were determined. *, P < 0.01; **, P < 0.05. The control represents 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 IgE 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 bronchiolar region (mean cell numbers/mm2 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 number per milliliter and the volume of the BAL fluid recovered were determined. *, 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+/−/− 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+/−/− memory Th2 cells (Fig. 2B) 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. 3B), 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+/−/− 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+/−/− 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+/−/− effector and memory Th2 cells (Fig. 2, A and B). However, Bmi1+/−/−/Ink4a−/−/Arf−/− Th2 cells, in which the levels of Puma and Bim expression were not increased (Fig. 2D), failed to generate normal numbers of memory Th2 cells (Fig. 2C). Although the expression of Puma and Bim mRNA was up–regulated, the levels were considerably low compared with Noxa in Bmi1+/−/− effector Th2 cells (Fig. 2A). This might explain the predominant effect of Noxa in the cell death of Bmi1+/−/− 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
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. 1D 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. 1F). The extent of the defect in homeostatic proliferation was modest (Fig. 1E). The low number of Bmi1<sup>−/−</sup> memory CD4<sup>T</sup> 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. 4F). 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. 5A). The binding of Dnmt1 at the CpG island was Bmi1 dependent (Fig. 5B and Fig. S11). The treatment with a 5-Aza resulted in the dissociation of Bmi1, Ring1B, and Suz12 from the CpG island (Fig. 5D) and the up-regulation of Noxa mRNA (Fig. 5C and Fig. S13A). Increased expression of Noxa mRNA and reduced binding of Bmi1 were confirmed in Dnmt1 knockout 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^ink4a/p19^arf 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-αβ (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-Ly-5.2 mAb (104), and PE-conjugated KJ1 (anti-clonotypic mAb for DO11.10 TCR) were purchased from BD Biosciences. Anti-FcγRII and III mAb (2.4G2) and unconjugated anti-IL-4 mAb (11B11), anti-IL-12 mAb (C17.8), and anti–IFN-γ (R4–6A2) were used as culture supernatants. Recombiant 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^+ CD4^+ T cells from DO11.10 OVA-specific TCR Tg mice were stimulated with 1 μM of an OVA peptide (Loh15) plus APC as described previously (36). Splenic KJ1^+ CD4^+ T cells from DO11.10 OVA-specific TCR Tg mice were stimulated with 3 μg/ml of immobilized anti–TCR-β mAb plus 1 μg/ml anti-CD28 mAb. These effector Th1/Th2 cells (3 × 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-Ly-5.2).

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^+ 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 μM 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 previously 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). Reombinant lentiviruses were generated using a three-plasmid system as described previously (58). In brief, 293 T cells were transfected with self-multiplying 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 × 10^6/well) was infected and selected using puromycin.

ChIP assay. ChIP assay was performed as described previously (59). The antibodies used for ChIP assay are as follows: anti-trimethyl histone H3-K4 (ab5880; 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 antisera specific for the 5-methyl cytosine. 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 wildtype and Bmi1^−/− 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^−/− Th2 cells. Fig. S2 shows the impaired generation of memory Th1 cells from Bmi1^−/− 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^−/− and Bmi1^−/− effector Th2 cells under nonlymphopenic conditions. Fig. S5 shows a competitive analysis for memory Th2 cell generation from Bmi1^−/− and Bmi1^−/− effector Th2 cells under lymphopenic conditions. Fig. S6 shows that deletion of the p53 gene failed to restore the generation of Bmi1^−/− memory Th2 cells. In Fig. S7, mRNA expression of Noxa in Bmi1^−/− and Bmi1^−/− effector Th1, Th2, Tc1, and Tc2 cells is shown. Fig. S8 displays mRNA expression of the Ink4a/Arf and Noxa in Bmi1^−/−, Bmi1^−/−, and Bmi1^−/− 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^−/− and Bmi1^−/− effector Th1 cells. Fig. S11 shows binding of Dnmt1 at the Noxa gene locus in Bmi1^−/− and Bmi1^−/− 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 R G108, 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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