

Transcription factor Bcl11b sustains iNKT1 and iNKT2 cell programs, restricts iNKT17 cell program, and governs iNKT cell survival

Mohammad Nizam Uddin^a, Dil Afroz Sultana^a, Kyle J. Lorentsen^a, Jonathan J. Cho^a, Mariana E. Kirst^a, Mark L. Brantly^a, Danielle Califano^b, Derek B. Sant'Angelo^c, and Dorina Avram^{a,1}

^aDivision of Pulmonary Medicine, Department of Medicine, College of Medicine, University of Florida, Gainesville, FL 32610; ^bCenter for Immunology and Microbial Disease, Albany Medical Center, Albany, NY 12208; and ^cDepartment of Pediatrics, Rutgers Robert Wood Johnson Medical School and The Child Health Institute of New Jersey, New Brunswick, NJ 08901

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Invariant natural killer T (iNKT) cells are innate-like T cells that recognize glycolipid antigens and play critical roles in regulation of immune responses. Based on expression of the transcription factors (TFs) Tbet, Plzf, and Ror γ t, iNKT cells have been classified in effector subsets that emerge in the thymus, namely, iNKT1, iNKT2, and iNKT17. Deficiency in the TF Bcl11b in double-positive (DP) thymocytes has been shown to cause absence of iNKT cells in the thymus and periphery due to defective self glycolipid processing and presentation by DP thymocytes and undefined intrinsic alterations in iNKT precursors. We used a model of cre-mediated postselection deletion of *Bcl11b* in iNKT cells to determine its intrinsic role in these cells. We found that Bcl11b is expressed equivalently in all three effector iNKT subsets, and its removal caused a reduction in the numbers of iNKT1 and iNKT2 cells, but not in the numbers of iNKT17 cells. Additionally, we show that Bcl11b sustains subset-specific cytokine production by iNKT1 and iNKT2 cells and restricts expression of iNKT17 genes in iNKT1 and iNKT2 subsets, overall restraining the iNKT17 program in iNKT cells. The total numbers of iNKT cells were reduced in the absence of Bcl11b both in the thymus and periphery, associated with the decrease in iNKT1 and iNKT2 cell numbers and decrease in survival, related to changes in survival/apoptosis genes. Thus, these results extend our understanding of the role of Bcl11b in iNKT cells beyond their selection and demonstrate that Bcl11b is a key regulator of iNKT effector subsets, their function, identity, and survival.

iNKT cell program | transcription factor Bcl11b | iNKT1 effector cells | iNKT2 effector cells | iNKT17 effector cells

Invariant natural killer T (iNKT) cells recognize glycolipid antigens presented by the MHC class I-like molecule CD1d and have been shown to play an important role not only in the immune response to bacterial pathogens, but also in antitumor immune responses (1, 2). iNKT cells bear a T-cell receptor (TCR) composed of V α 14–J α 18 chain paired with V β 7, β 8, and β 2 in mice, and V α 24 and V β 11 in humans (3). Following stimulation with glycolipid antigens or cytokines, iNKT cells quickly respond by producing cytokines, including IFN γ , IL-4, IL-13, IL-17, IL-10, and GM-CSF (4–9). This quick response gives them the innate-like attribute. Thymic iNKT precursors are selected on double-positive (DP) thymocytes, which present self glycolipids on CD1d molecules (10–12). Following selection, iNKT precursors go through four developmental stages: 0 (NK1.1[–]HSA^{hi}CD44^{lo}), 1 (NK1.1[–]HSA^{lo}CD44^{lo}), 2 (NK1.1[–]HSA^{lo}CD44^{hi}), and 3 (NK1.1⁺HSA^{lo}CD44^{hi}) (13). iNKT cell migration out of the thymus occurs at stages 2 and 3 (13, 14). Similar to T helper cells and innate lymphoid cells (ILCs), iNKT cells have been classified into three distinct effector subsets, based on the expression of the TFs Tbet, PLZF, and Ror γ t, namely, iNKT1 (Tbet^{hi}PLZF^{lo}), iNKT2 (Tbet^{lo}PLZF^{hi}), and iNKT17 (Tbet^{lo}PLZF^{lo}Ror γ t⁺) (15). In B6 mice, the iNKT2 and iNKT17 subsets are found predominantly within developmental stage 2, whereas the iNKT1 subset is confined

to stage 3 (15). Several transcription factors (TFs) have been found essential for iNKT cell progression through developmental stages, as well as for their effector functions. Tbet is critical for iNKT1 cell function and for terminal maturation and homeostasis (15, 16). Ror γ t not only controls the iNKT17 pathway, but together with Runx1, regulates iNKT cell development (12, 15, 17). PLZF is expressed postselection and directs the development and effector program of iNKT cells (18, 19). E and Id proteins are important for both lineage choice between iNKT and T cells during selection and differentiation into iNKT1 and iNKT2 subsets (20–22). c-myb regulates CD1d levels on DP thymocytes, as well as Slamf1, Slamf6, and SAP on iNKT cells (23). Hobit controls maintenance of mature iNKT cells and their effector functions (24). Recently Lef1 was found to be essential for iNKT2 subset formation and function, and to regulate Gata3 and Thpok (25), both known to control CD4⁺ iNKT cells (26). TF Bcl11b plays a crucial role in T-cell lineage commitment (27, 28), selection, differentiation, and survival of thymocytes (29, 30), clonal expansion and effector function of CD8⁺ T cells (31), as well as suppression function of Treg cells (32). Additionally, Bcl11b restricts expression of Th2 lineage genes in Th17 cells in experimental autoimmune encephalomyelitis (EAE) (33). Bcl11b was recently found to sustain innate lymphoid type 2 cell (ILC2) program (34, 35, 36) and to suppress ILC3 program in ILC2s

Significance

Invariant natural killer T (iNKT) cells are innate-like T cells that recognize lipid antigens and play important roles in antimicrobial and tumor immunity. Functionally, iNKT cells have been classified in three effector subsets based on expression of specific transcription factors (TFs) and cytokine genes. We previously demonstrated that the TF Bcl11b controls glycolipid processing and presentation by double-positive thymocytes to iNKT precursors and thus their formation. Using a model that allows bypassing those defects, here we provide evidence that the TF Bcl11b is critical for effector iNKT1 and iNKT2 subsets and overall survival of iNKT cells. Additionally we provide evidence that Bcl11b sustains cytokine production by iNKT1 and iNKT2 cells and restricts expression of the iNKT17 cell program in all effector subsets.

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¹To whom correspondence should be addressed. Email: Dorina.Avram@medicine.ufl.edu.

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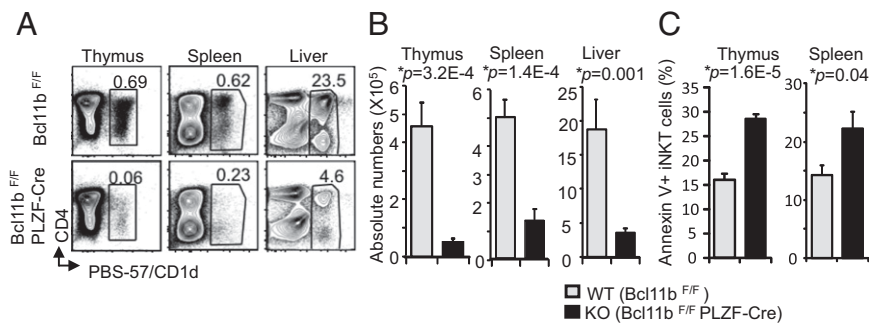


Fig. 1. *Bcl11b*'s deficiency in iNKT cells causes a major reduction in thymic and peripheral iNKT cells associated with increased apoptosis. (A and B) Frequencies (A) and absolute numbers (B) of iNKT cells from *Bcl11b*^{F/F}/PLZF-Cre (KO) and WT mice in thymus, spleen, and liver. The gated population in A shows percentages of CD1d-PBS-57⁺ cells. (C) Frequencies of Annexin V⁺ iNKT cells from *Bcl11b*^{F/F}/PLZF-Cre (KO) and *Bcl11b*^{F/F} (WT) mice in thymus and spleen. Data are representative of several independent experiments with 10 pairs of mice (A and B) and 5 pairs of mice (C). *P* values determined by unpaired two-tailed Student's *t* test are indicated. Means \pm SEM.

(36). *Bcl11b*'s deficiency in DP thymocytes resulted in lack of iNKT cells in the thymus and periphery (37, 38), despite the fact that the V α 14J α 18 TCR was normally rearranged (38). The defect was caused by the inability of *Bcl11b*^{-/-} DP thymocytes to support the selection of iNKT precursors, due to defective glycolipid self-antigen processing/presentation. Additionally, *Bcl11b*^{-/-} iNKT precursors, even when normally presented with glycolipid self-antigens, failed to generate iNKT cells, due to unidentified intrinsic defects (38). Here we set up a system to study the defects caused by the absence of *Bcl11b* in iNKT cells, using the *PLZF*-Cre mouse strain, which promotes removal of floxed alleles postselection of iNKT cells, namely starting with developmental stage 1 (18, 39, 40). Our study demonstrates that *PLZF*-Cre-mediated iNKT cell deletion of *Bcl11b* resulted in significantly reduced iNKT cells in the thymus and periphery, associated with reduction in survival in relation to changes in survival/apoptosis genes. iNKT1 and iNKT2 effector subsets were numerically reduced both in thymus and spleen, suggesting that these two subsets need *Bcl11b*. Additionally, levels of IFN γ and IL-4 within *Bcl11b*^{-/-} iNKT1 and iNKT2 subsets, respectively, were reduced, demonstrating that these cells also have functional alterations. Although numbers of *Bcl11b*^{-/-} iNKT17 cells were normal, IL-17 production was up-regulated together with increased levels of iNKT17 subset molecules, namely, IL23R and Nr1p. IL23R and Nr1p were up-regulated not only in total iNKT cells and in iNKT17 subset, but also in the other two effector subsets, suggesting an additional role for *Bcl11b* in restricting the iNKT17 program in iNKT1 and iNKT2 subsets.

Additionally, *Bcl11b* suppressed expression of some NK genes in iNKT cells. Thus, our results demonstrate a major role of *Bcl11b* in the development, function, identity and survival of iNKT cells.

Results

Bcl11b's Removal Is Specific for iNKT Cells in *Bcl11b*^{F/F}/PLZF-Cre Mice.

Given previous results in which absence of *Bcl11b* caused a developmental block in iNKT cells due to defective iNKT precursors (38), as well as defective glycolipid processing and presentation by DP thymocytes, we used the *PLZF*-Cre mouse strain to remove targeted alleles restrictively in iNKT cells, postselection (40). The *PLZF*-Cre deleter efficiently removed *Bcl11b* in splenic and thymic iNKT cells, starting with developmental stage 1 (Fig. S1A–C). Given that ~25% of T cells were previously reported to be positive for the Rosa 26-tdTomato reporter in *PLZF*-Cre mice (40), we evaluated removal of *Bcl11b* in PBS-57/CD1d⁻ thymic CD4⁺ single positive (SP) and splenic CD4⁺ T cells and found no removal of *Bcl11b* in these cells (Fig. S1A and B). Moreover, thymic SP and peripheral CD4⁺ and CD8⁺ T-cell populations remained numerically unaffected (Fig. S1D and E). Considering that *PLZF* was found to be expressed in a precursor population of helper ILCs (41), and removal of *Bcl11b* in ILC2s caused down-regulation of St2 and derepression of Ror γ t (36), we investigated whether *Bcl11b* is removed from ILC2s in *Bcl11b*^{F/F}/PLZF-Cre mice. Our results show no removal of *Bcl11b* in these cells (Fig. S1F). Additionally, ILC2s of *Bcl11b*^{F/F}/PLZF-Cre mice did not down-regulate St2 or up-regulate Ror γ t (Fig. S1G). These results

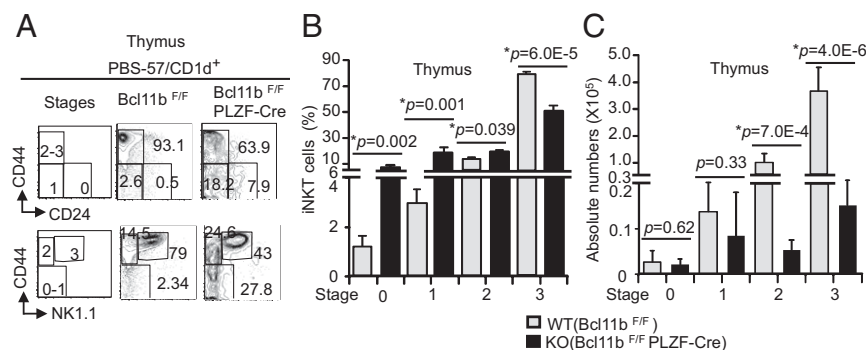


Fig. 2. *Bcl11b*'s removal in iNKT cells causes developmental alterations. (A) Frequencies of iNKT cells from *Bcl11b*^{F/F}/PLZF-Cre and WT mice in the four developmental stages (Left) based on surface CD24, CD44, and NK1.1 within the CD1d-PBS-57⁺ cells. (B and C) Quantification of percentages (B) and absolute numbers (C) of iNKT cells within the four developmental stages. Data are representative of several independent experiments with 10 pairs of mice. *P* values are indicated. Means \pm SEM.

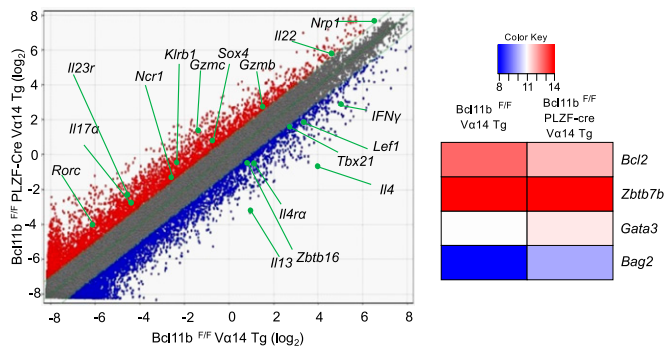


Fig. 3. *Bcl11b*-deficient iNKT cells show altered expression of genes critical for effector iNKT subsets and survival. Scatterplot of \log_2 expression values for mRNAs isolated from sorted iNKT cells of *Bcl11b*^{F/F}/PLZF-Cre Va14 Tg (KO) versus Va14 Tg control mice (WT) treated with α -GalCer. Data are representative of two independent arrays. Relevant genes are listed. The heat map shows relevant genes off the scale in scatterplot.

taken together demonstrate that *Bcl11b*'s removal is specific for iNKT cells in *Bcl11b*^{F/F}/PLZF-Cre mice.

Bcl11b^{F/F}/PLZF-Cre Mice Have Reduced Numbers of Thymic and Peripheral iNKT Cells and Developmental Alterations of iNKT Cells.

Ablation of *Bcl11b* with the PLZF-Cre deleter caused a severe reduction of the percentages and absolute numbers of iNKT cells in the thymus, spleen, and liver (Fig. 1 *A* and *B* and Fig. S24), although the mean fluorescence intensity (MFI) of PBS-57-loaded CD1d tetramer bound by TCR remained similar between *Bcl11b*^{-/-} and wild-type iNKT cells (Fig. S2B). We further asked the question whether the reduced numbers of iNKT cells in the absence of *Bcl11b* is caused by their reduced survival. Annexin V staining of thymic and splenic iNKT cells showed that more *Bcl11b*^{-/-} iNKT cells stained for Annexin V compared with control, both in thymus and spleen (Fig. 1C and Fig. S2C). These results suggest that *Bcl11b*^{-/-} iNKT cells have an increased tendency to die, which is likely to contribute to their numerical decrease.

We further investigated the thymic iNKT developmental stages and found that *Bcl11b*^{F/F}/PLZF-Cre mice had higher percentages of iNKT cells in stages 0–2, whereas the percentages of stage 3 iNKT cells were reduced (Fig. 2 *A* and *B*). The absolute numbers of stages 0 and 1 iNKT cells were similar in *Bcl11b*^{F/F}/PLZF-Cre and wild-type mice, however, there was a substantial reduction of absolute numbers of iNKT cells in stages 2 and 3 (Fig. 2C). Given that iNKT cells go through a massive expansion as they transition between stages 1 and 2, we evaluated the Ki67, indicative of cells entering the cell cycle. Our results show that similar percentages of *Bcl11b*^{-/-} and wild-type iNKT cells were positive for Ki67 in stage 1, whereas during stage 2, the percentages were increased for *Bcl11b*^{-/-} iNKT cells (Fig. S3). Therefore, it is unlikely that the reduction in the numbers of iNKT cells in the absence of *Bcl11b* is caused by defective cell cycle entering, but rather by reduced survival and alterations in development.

Absence of *Bcl11b* Causes Up-Regulation of iNKT17 Genes, Down-Regulation of iNKT1 and iNKT2 Genes, and Alterations in Survival Genes.

Comparison of the mRNAs of *Bcl11b*^{-/-} and wild-type iNKT cells showed that the prosurvival gene *Bcl2* was down-regulated and the proapoptotic gene *Bag2* was up-regulated in *Bcl11b*^{-/-} iNKT cells (Fig. 3), thus supporting the observation that *Bcl11b*^{-/-} iNKT cells have a reduction in survival. The mRNAs for the *Tbx21* (Tbet), critical for the iNKT1 subset and its signature cytokine IFN γ , were reduced (Fig. 3). The mRNAs for the TFs *Zbtb16* (PLZF) and *Lef1*, important for iNKT2

subset (15, 25), were also diminished, together with the mRNAs for the iNKT2 subset cytokines IL-4 and IL-13, and the receptor IL-4R α . However, there was no change in the mRNAs for *Gata3* and *Zbtb7b* (Thpok) (Fig. 3), known to be downstream of *Lef1* (25). The mRNAs for the TFs *Roryt* (*Rorc*) and *Sox4*, known to be important for the iNKT17 subset, were up-regulated (Fig. 3), together with the iNKT17 subset mRNAs for IL-17, IL-22, IL-23R, and *Nrp1* (Fig. 3). In addition, the mRNAs for *Ncr1* (Nkp46), *Klrb1*, *Gzmb*, and *Gzmc* were elevated (Fig. 3). The results of this analysis support the conclusion that *Bcl11b* controls survival of iNKT cells and effector iNKT subset genes.

iNKT1 and iNKT2 Effector Subsets Require *Bcl11b*. Given the up-regulation of iNKT17 genes and the down-regulation of iNKT1 and iNKT2 genes, we further evaluated the effector iNKT subsets in the absence of *Bcl11b*. We first assessed the expression of *Bcl11b* in these subsets and found that *Bcl11b* was expressed in thymic and splenic effector iNKT subsets at similar levels (Fig. S4A). We further evaluated the thymic and splenic effector iNKT subsets in the absence of *Bcl11b*, and found that the percentages and absolute numbers of iNKT1 and iNKT2 cells were reduced both in the thymus and periphery (Fig. 4 *A–D* and Fig. S4 *B* and *C*). Whereas there was an increase in the percentages of iNKT17 cells, the absolute numbers of iNKT17 cells remained unchanged (Fig. 4 *A–D* and Fig. S4 *B* and *C*). These data suggest that iNKT1 and iNKT2 subsets, but not the iNKT17 subset, require *Bcl11b*.

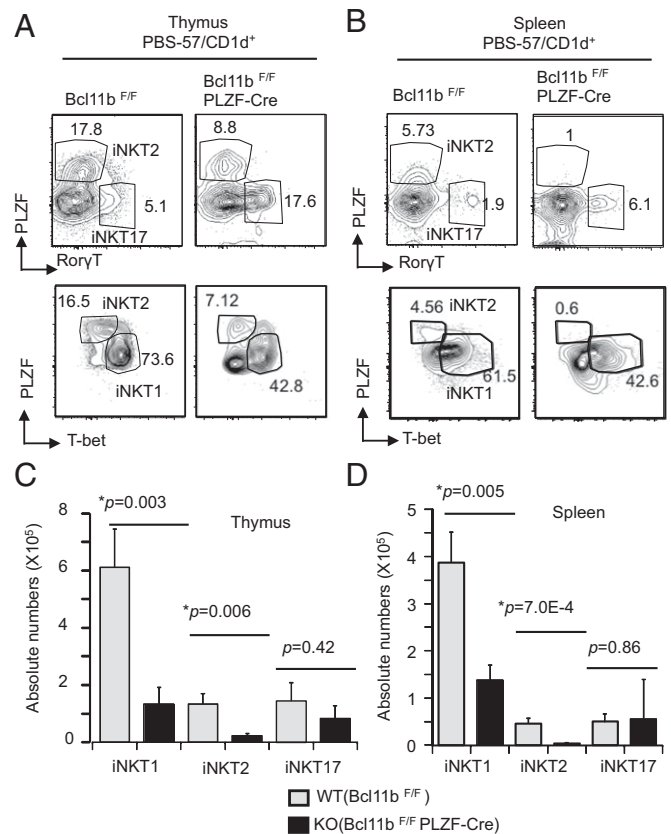


Fig. 4. iNKT1 and iNKT2 subset cell numbers are reduced in thymus and spleen of *Bcl11b*^{F/F}/PLZF-Cre mice. (*A–D*) Frequencies (*A* and *B*) and absolute numbers (*C* and *D*) of effector iNKT subsets in thymus and spleen, identified as iNKT1 (Tbet^{hi}PLZF^{lo}), iNKT2 (Tbet^{lo}PLZF^{hi}), and iNKT17 (Roryt^{hi}PLZF^{lo}) on gated CD1d-PBS-57⁺ cells from *Bcl11b*^{F/F}/PLZF-Cre (KO) and *Bcl11b*^{F/F} (WT) mice. Data are representative of 8–10 pairs of mice. *P* values are indicated. Means \pm SEM.

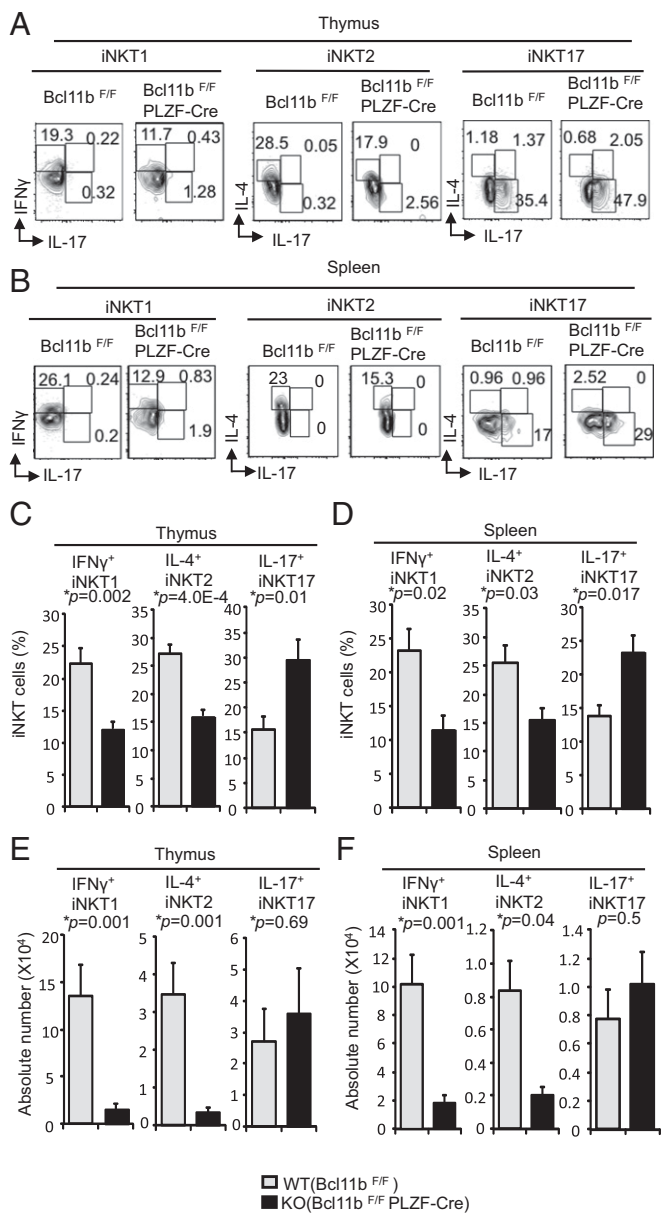


Fig. 5. *Bcl11b*-deficient iNKT cells are functionally altered. (A–F) Frequencies (A–D) and absolute numbers (E and F) of IL-4⁺, IFN γ ⁺, and IL-17⁺ iNKT cells within the indicated effector subsets, iNKT1, iNKT2, and iNKT17, defined as in Fig. 4 A and B. Data are representative of several experiments with eight pairs of mice. P values are indicated. Means \pm SEM.

***Bcl11b*^{-/-} iNKT Effector Subsets Have Functional Alterations.** We further evaluated the cytokine production by total *Bcl11b*^{-/-} iNKT cells, as well as by the effector subsets. Our results indicate that overall frequencies of IL-17-producing *Bcl11b*^{-/-} iNKT cells were increased, whereas frequencies of IL-4- and IFN γ -producing *Bcl11b*^{-/-} iNKT cells were reduced compared to wild-type iNKT cells both in thymus and spleen (Fig. S5 A and B), in agreement with the changes in the proportion of effector subsets. Within the iNKT1 and iNKT2 subsets, the percentages of *Bcl11b*^{-/-} IFN γ - and IL-4-producing iNKT cells, respectively, were reduced both in thymus and spleen (Fig. 5 A–D). Additionally, the absolute numbers of *Bcl11b*^{-/-} IFN γ - and IL-4-producing iNKT1 and iNKT2 cells, respectively, were reduced as well (Fig. 5 E and F). Within the iNKT17 subset, the percentages of *Bcl11b*^{-/-} IL-17-producing iNKT cells were increased in the thymus and spleen

and the absolute numbers were slightly increased, but not significantly (Fig. 5 A–F). These results suggest that within the *Bcl11b*^{-/-} iNKT1 and iNKT2 subsets, fewer cells produced IFN γ and IL-4, respectively, thus presenting functional

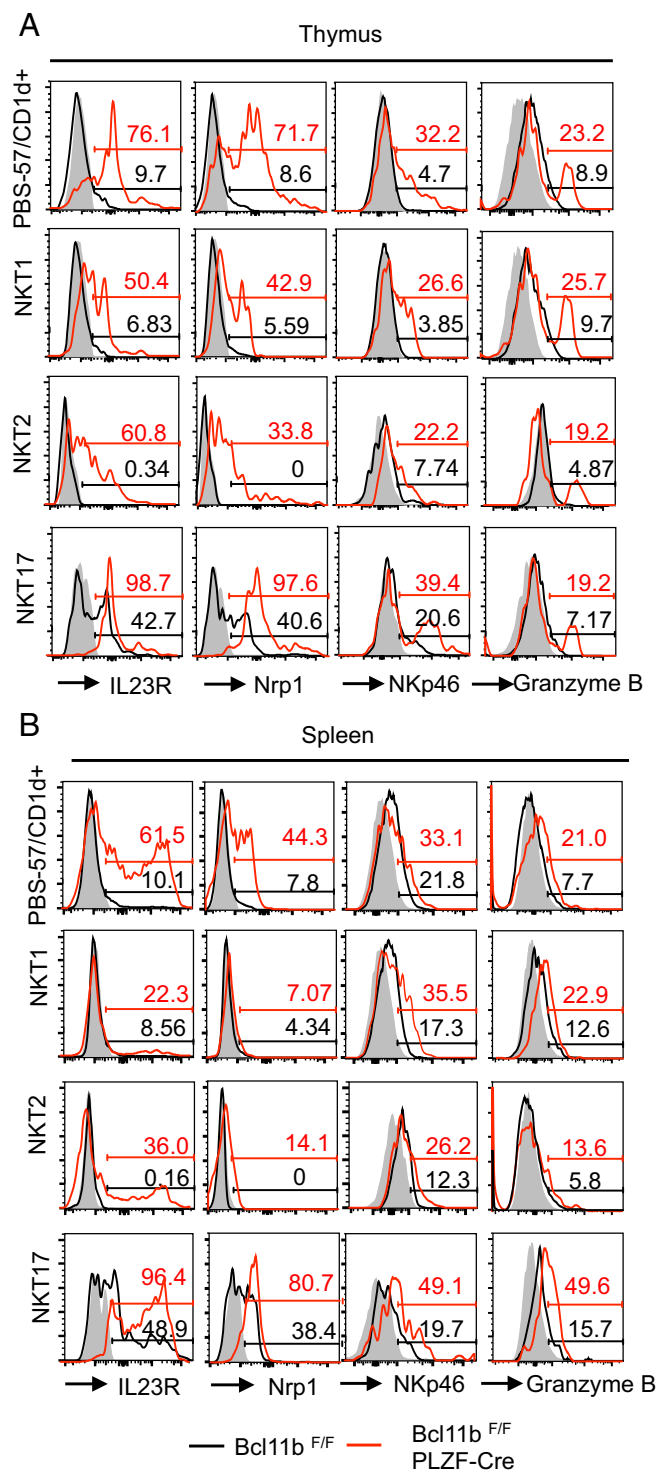


Fig. 6. *Bcl11b*-deficient iNKT cells, including iNKT1 and iNKT2 effector subsets up-regulate iNKT17 cell markers. Frequencies of IL-23R⁺, Nrp1⁺, NKp46⁺, and Granzyme B⁺ in total iNKT cells and in effector iNKT subsets of *Bcl11b*^{F/F} PLZF-Cre and *Bcl11b*^{F/F} mice in thymus (A) and spleen (B). iNKT1, iNKT2, and iNKT17 are defined as in Fig. 4 A and B. Data are representative of three independent experiments.

alterations, whereas within the *Bcl11b*^{-/-} iNKT17 subset, a larger percentage produced IL-17.

Bcl11b Restricts Expression of iNKT17 Program in iNKT1 and iNKT2 Effector Subsets. We further investigated whether other genes, dysregulated in the absence of Bcl11b from iNKT cells, were altered within the effector subsets or their difference in expression was a consequence of ratio change between the effector subsets.

Several mRNAs for iNKT17 genes were up-regulated, including for IL-23R and Nrp1 (Fig. 3). IL-23R and Nrp1 were up-regulated not only in the total iNKT cells and within the iNKT17 subset, but even in the iNKT1 and iNKT2 effector subsets (Fig. 6). These results suggest that Bcl11b may be overall implicated in restraining the iNKT17 program, including in iNKT1 and iNKT2 effector subsets.

We further evaluated Nkp46 and Gzmb, two natural killer (NK) genes up-regulated at mRNA level in total *Bcl11b*^{-/-} iNKT cells and known to be derepressed following Bcl11b's removal in other T-cell populations and progenitors (reviewed in ref. 42). Both Nkp46 and Gzmb were very modestly expressed in the wild-type thymic and splenic iNKT cells at steady state (Fig. 6), similar to what has been recently reported (43). Removal of *Bcl11b* resulted in the up-regulation of Nkp46 and Gzmb in all of the three effector subsets both in thymus and spleen (Fig. 6). As shown above, the mRNA encoding Lef1, recently found to be essential for iNKT2 cells (25), was reduced (Fig. 3). There was also a modest reduction of Lef1 in the total *Bcl11b*^{-/-} iNKT cells, however, no change in iNKT2 subset (Fig. S6A), suggesting that the reduced Lef1 mRNA level in total *Bcl11b*^{-/-} iNKT cells was simply due to the iNKT2 subset numerical reduction. In agreement with these results, no up-regulation of CD8 was observed in the *Bcl11b*^{-/-} iNKT cells above wild-type background (Fig. S6B) and Gata3 remained unchanged (Fig. S6A).

Thus these results taken together suggest that Bcl11b is required to restrict iNKT17 genes overall in iNKT cells, including in effector iNKT1 and iNKT2 subsets. Additionally, Bcl11b represses expression of some NK genes in iNKT cells.

Discussion

Previous reports demonstrate that absence of the TF Bcl11b at DP stage of T-cell development caused a defect in self glycolipid processing and presentation, which impacted the selection of iNKT precursors (38). Additionally, *Bcl11b*^{-/-} iNKT precursors had unknown intrinsic defects that blocked their development even when selected on DP thymocytes able to present glycolipids (38). Here we set up a system to study the specific role of Bcl11b in iNKT cells postselection, at developmental stage 1. Importantly, our results demonstrate specific and restrictive removal of *Bcl11b* with the *PLZF*-Cre system in iNKT cells and no removal of *Bcl11b* in other T cells or ILC2s. Using this system we found that deficiency of *Bcl11b* caused a significant decrease in iNKT cells in the thymus and periphery, related to reduced iNKT1 and iNKT2 cell numbers and to an overall reduced survival of iNKT cells attributed to changes in survival genes, similar to what was previously reported for *Bcl11b*^{-/-} thymocytes (29, 30). Although iNKT1 and iNKT2 cell numbers were reduced in the absence of *Bcl11b*, there was no change in the numbers of iNKT17 cells, suggesting an essential role of Bcl11b in the control of effector iNKT1 and iNKT2 subsets. The reduction in the iNKT1 and iNKT2 subsets in the absence of *Bcl11b* is also in agreement with the reduction in developmental stage 3, predominantly represented by the iNKT1 subset, and the reduction in developmental stage 2, represented by iNKT2 and iNKT17 subsets. One possibility is that development of iNKT1 and iNKT2 subsets is dependent on Bcl11b, whereas the iNKT17 subset is not. Another possibility is that *Bcl11b*^{-/-} iNKT1 and iNKT2 cells have altered survival, however, *Bcl11b*^{-/-} iNKT17 cells survive normally. Not

only were the iNKT1 and iNKT2 subsets numerically reduced, but the percentages of IFN γ - and IL-4-producing iNKT cells were also reduced within the iNKT1 and iNKT2 subsets, respectively, suggesting that iNKT1 and iNKT2 cells have functional alterations in the absence of *Bcl11b*. The TF Lef1, recently demonstrated to control Gata3 and IL-4 expression in iNKT2 cells (25), was not reduced in *Bcl11b*^{-/-} iNKT2 cells. There was no reduction in Gata3 expression, suggesting that Bcl11b might be involved in the control of IL-4, independent of these transcriptional regulators. Also, Bcl11b might be implicated in the control of IFN γ either directly or through Tbet. Interestingly, levels of the IL-17, IL-23R, and Nrp1, all part of the iNKT17 program, were elevated in total and iNKT17 cells, which suggests a role of Bcl11b in restraining the iNKT17 program. IL-23R and Nrp1 were also up-regulated in iNKT1 and iNKT2 subsets, further suggesting that Bcl11b restricts expression of the iNKT17 program in iNKT1 and iNKT2 subsets. This is similar to the role of Bcl11b in ILC2s, in which Bcl11b represses the ILC3 program (34). As mentioned above *Bcl11b*^{-/-} iNKT17 cells were not affected numerically in the absence of Bcl11b, similar to what happened in the absence of *Bcl11b* in Th17 cells during EAE (33). However, different from *Bcl11b*^{-/-} Th17 cells in EAE (33), *Bcl11b*^{-/-} iNKT17 cells did not express IL-4. Thus, Bcl11b regulates some common themes in some immune populations, such as ILCs and iNKT cells, however, it acts also in a context-dependent manner in other populations (reviewed in ref. 42). It remains to be established whether Bcl11b plays other roles in iNKT17 cells except restricting expression of its signature genes. In addition, as in other T-cell populations, Bcl11b repressed NKp46 and Gzmb, but not NK1.1, CD244, or myeloid genes, that are derepressed in the absence of Bcl11b in early thymocytes (27, 28). This again suggests that Bcl11b governs not only common, but also context-specific programs. The suppression of NKp46 and Gzmb can be related to overall repression of NK program. NKp46 was expressed in wild-type iNKT subsets at very low level, with the highest level in the iNKT17 subset, at least in the thymus. Therefore, it is possible that NKp46 is part of the iNKT17 program, similar to a subset of ILC3s, and absence of Bcl11b causes its increase in relation to derepression of iNKT17 genes or as mentioned above as part of restricting expression of NK genes. This remains to be established in the future.

In conclusion, our results demonstrate a critical role of Bcl11b in the control of iNKT1 and iNKT2 subsets and in sustaining their function. Bcl11b also restricts expression of the iNKT17 program, including in iNKT1 and iNKT2 subsets. In addition, Bcl11b is required for overall survival of iNKT cells and repression of some NK genes.

Materials and Methods

Detailed materials and methods are given in *SI Materials and Methods*.

Mice. *Bcl11b*^{fl/fl}/*PLZF*-Cre mice were generated by breeding *Bcl11b*^{fl/fl} and *PLZF*-Cre mice, previously described (29, 30, 40). α 14 mice were previously described (44). Mice were kept under specific pathogen-free conditions. All of the experiments were conducted in accordance with animal protocols approved by the Institutional Animal Care and Use Committee of the University of Florida and Albany Medical College.

Statistical Analysis. The statistical difference between experimental groups was determined by unpaired two-tailed Student *t* test. The *P* values ≤ 0.05 were considered significant.

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