

**FIGURE 1.** T and B cell development is normal in Klf12-deficient mice. (A) Percentages and total cell numbers of CD4<sup>+</sup> and CD8<sup>+</sup> T cells in the thymus (*left panels*) and in the spleen (*right panels*). (B) Percentages and total cell numbers of developmental B cell subsets in the BM (*left panels*), spleen (*second left panels*), lymph node (*second right panels*), and peritoneal cavity (*right panels*). Subsets in the BM are defined as FrC' pro-B cells: CD19<sup>+</sup>IgM<sup>-</sup>B220<sup>+</sup>CD43<sup>+</sup>CD24<sup>+</sup>; FrD pre-B cells: CD19<sup>+</sup>IgM<sup>-</sup>B220<sup>+</sup>CD43<sup>-</sup>, FrE immature B cells: CD19<sup>+</sup>IgM<sup>+</sup>B220<sup>+</sup>AA4.1<sup>+</sup>CD24<sup>hi</sup>; and FrF mature B cells: CD19<sup>+</sup>IgM<sup>+</sup>B220<sup>+</sup>AA4.1<sup>+</sup>CD24<sup>hi</sup>; and FrF mature B cells: IgM<sup>+</sup>IgD<sup>+</sup>CD19<sup>+</sup>B220<sup>+</sup>AA4.1<sup>-</sup>CD23<sup>+</sup>; FO (follicular) B cells: IgM<sup>+</sup>IgD<sup>+</sup>CD19<sup>+</sup>B220<sup>+</sup>AA4.1<sup>-</sup>CD23<sup>+</sup>; and MZ (marginal zone) B cells: IgM<sup>+</sup>IgD<sup>+</sup>CD19<sup>+</sup>B220<sup>+</sup>AA4.1<sup>-</sup>CD23<sup>-</sup>. Subsets in the lymph node are defined as FO (follicular) B cells: IgM<sup>+</sup>IgD<sup>+</sup>CD19<sup>+</sup>CD43<sup>-</sup>. Subsets in the peritoneal cavity are defined as B2 follicular B cells: IgM<sup>+</sup>IgD<sup>+</sup>CD19<sup>+</sup>CD43<sup>-</sup>CD23<sup>+</sup>; B1a B cells: IgM<sup>+</sup>IgD<sup>+</sup>CD19<sup>+</sup>CD19<sup>+</sup>CD43<sup>-</sup>B220<sup>+</sup>CD5<sup>-</sup>; B1a B cells: IgM<sup>+</sup>IgD<sup>+</sup>CD19<sup>+</sup>CD19<sup>+</sup>CD43<sup>+</sup>B220<sup>+</sup>CD5<sup>-</sup>; D11b<sup>+</sup>. Data are representative of 4 experiments (*n* = 3 mice/genotype/experiment).



**FIGURE 2.** Upregulation of pERK is normal in Klf12-deficient T and B cells. (A) Representative histograms of pERK upregulation in lymph node CD4<sup>+</sup>, CD8<sup>+</sup> T cells, and CD19<sup>+</sup> B cells from *Klf12<sup>+/+</sup>*, *Klf12<sup>F/+</sup>*, and *Klf12<sup>F/+</sup>* mice stimulated with anti-CD3 or anti-IgM. (B) Percentage and MFI of pERK in CD4<sup>+</sup> T cells (*left panels*), CD8<sup>+</sup> T cells (*middle panels*), and CD19<sup>+</sup> T cells (*right panels*). Data are representative of 4 experiments (n = 2-3 mice/genotype/experiment).



**FIGURE 3.** T and B cell proliferation is normal in Klf12-deficient mice. Proliferation of  $Klf12^{+/+}$ ,  $Klf12^{F/+}$ , and  $Klf12^{F/+}$  lymph node CD4<sup>+</sup> T cells (*left panels*), CD8<sup>+</sup> T cells (*middle panels*), and CD19<sup>+</sup> T cells (*right panels*) after *in vitro* stimulation for 2-5 days with anti-CD3 and anti-IgM. Data are representative of 4 experiments (n = 2-3 mice/genotype/experiment).



**FIGURE 4.** Normal numbers of Klf12-deficient T cells from MCMV-infected BM chimeric mice. Relative numbers of naïve (CD44<sup>lo</sup>CD62L<sup>+</sup>) CD4<sup>+</sup> T cells, effector-memory (CD44<sup>hi</sup>CD62L<sup>-</sup>) CD4<sup>+</sup> T cells, naïve (CD44<sup>-</sup>CD62L<sup>+</sup>) CD8<sup>+</sup> T cells, effector-memory (CD44<sup>+</sup>CD62L<sup>lo/+</sup>) CD8<sup>+</sup> cells, and MCMV-specific NKG2D<sup>+</sup>CD8<sup>+</sup> T cells in the blood following MCMV infection. Data are representative of 4 experiments (n = 3-6 mice/experiment).