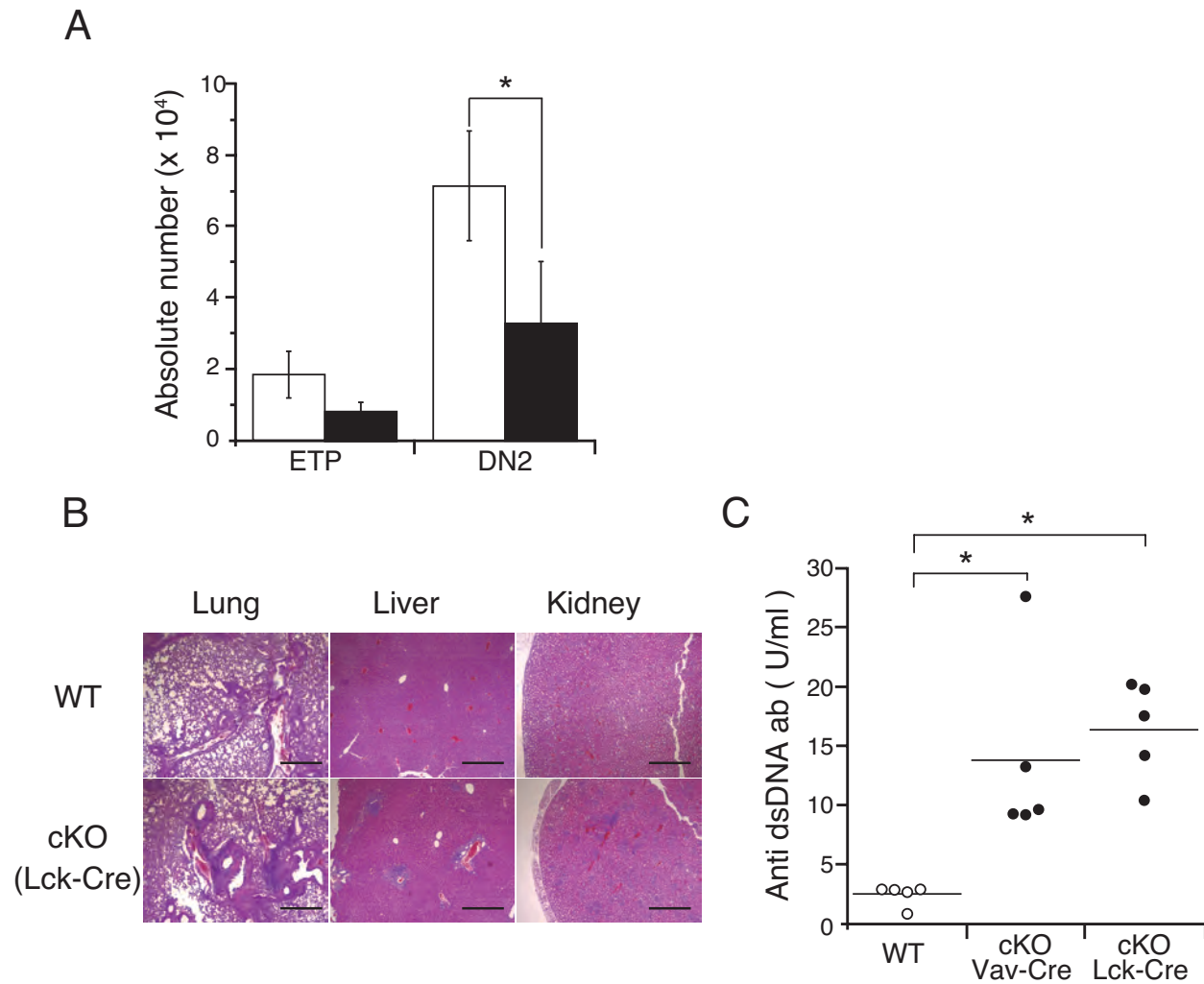
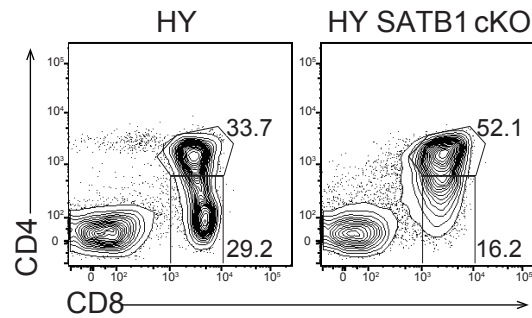


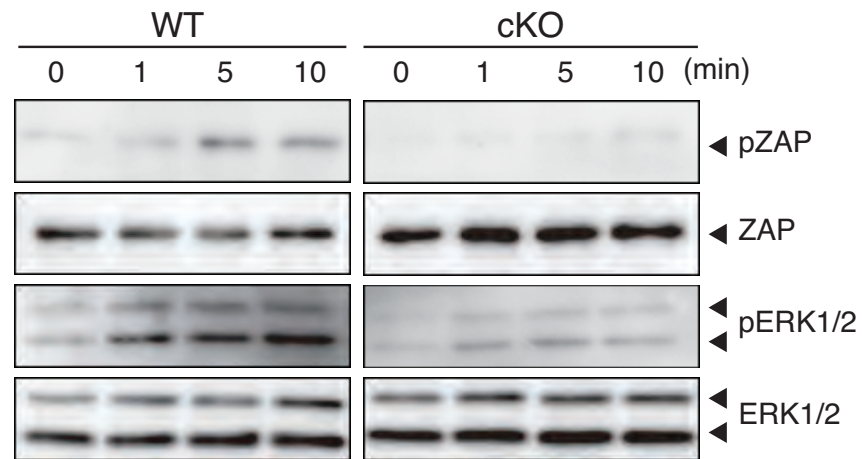
Supplemental Figure 1. (A) T cell development was arrested at the CLP stage in BM in the absence of SATB1. The number of HSC (Flt3⁺c-Kit^{hi}Lin⁻Sca-I⁺ cells), MPP (Flt3⁺c-Kit^{hi}Lin⁻Sca-I⁺ cells) and CLP (IL-7R α ⁺c-Kit^{lo}Lin⁻Sca-I^{lo} cells) cells in the bone marrow from one femur of a WT (open bars, n=3) or a SATB1cKO (closed bars, n=3) mouse is shown. Student's *t*-test was used for statistical analysis (**p*<0.05). **(B)** Treg cell numbers in the spleen of WT (open bars, n=5) and SATB1cKO mice (closed bars) at different ages (**p*<0.05).



Supplemental Figure 2. Autoimmune disorders occur in *Lck-Cre SATB1^{fl/fl}* mice. **(A)** The number of ETP (c-Kit⁺CD25⁻CD44⁺ DN) or the DN2 (c-Kit⁺CD25⁺CD44⁺ DN) fraction of thymocytes in WT mice (open bars, n=5) and in *SATB1^{fl/fl}* mice with *Lck-Cre* transgenes (closed bars, n=5) is shown (*p<0.05). **(B)** H&E staining of sections of various organs from WT and *Lck-Cre SATB1^{fl/fl}* mice. **(C)** Concentration of anti-dsDNA antibodies in the serum of WT, *Vav-Cre SATB1^{fl/fl}* and *Lck-Cre SATB1^{fl/fl}* mice.



Supplemental Figure 3. Transition from a DP to a CD8⁺ SP population is perturbed in female HY-TCR⁺ mice on a SATB1 cKO background. Expression of CD4 and CD8 on HY-TCR-expressing T3.70⁺ thymocytes from female HY-TCR TG mice on a WT (left) or a SATB1cKO background (right) was analyzed on FACS. Representative results from more than 5 analyses are shown.



Supplemental Figure 4. Signal strength via TCR is weak in T cells from SATB1cKO mice. Naïve CD4⁺ T cells were purified from the spleen of WT and SATB1cKO mice and stimulated with anti-CD3 and anti-CD28 antibodies for indicated periods. Whole cell lysates were subjected to SDS-page. Separated proteins were transferred to PVDF membrane, which was blotted with antibodies indicated in the figure.