

Figure S1. Expression of gut-trophic molecules on effector CD4 and CD8 T cells in the mLNs and PPs in  $mTOR^{f/f}$ -CD4Cre and mTOR<sup>f/f</sup> mice. Overlaid histograms show CD103, CCR9, and integrin  $\alpha 4\beta 7$  staining in gated CD44<sup>+</sup>CD62L<sup>+</sup> (CM) and CD44<sup>+</sup>CD62L<sup>-</sup> (EM) CD4 and CD8 T cells. Data shown are representative of three experiments.

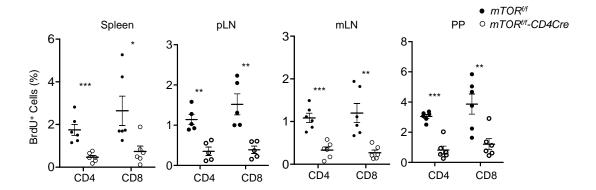
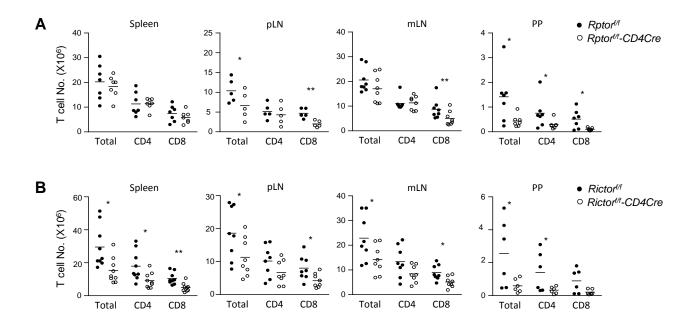


Figure S2. BrdU incorporation in CD4 and CD8 T cells in the peripheral lymphoid organds from  $mTOR^{f/f}$ -CD4Cre and  $mTOR^{f/f}$  mice. Mice were intraperitoneally (i.p) injected with BrdU and BrdU incorporation in T-cells were examined by flow-cytometry 16 hours later. Scatter plots showing percentages of BrdU<sup>+</sup> T-cells in the indicated populations. Horizontal bars represent the mean  $\pm$  SEM. Each circle represents one mouse of the indicated genotypes. Data shown represent or calculated from at least three experiments. \*, P<0.05; \*\*, P<0.01; \*\*\*, P<0.001 determined by unpaired two-tailed Student t-test.



**Figure S3. Effects of mTORC1 and mTORC2 defciency on T-cell numbers in peripheral lymphoid organs.** Scatter graphs showing the mean  $\pm$  SEM of total T, CD4, and CD8 T-cell numbers in the spleen, pLNs, mLNs, and PPs from *Raptor* and *Raptor* and *Raptor* and *Rictor* and *Rictor* mice (**B**) mice. Data shown are calculated from at least five experiments. \*, P < 0.05; \*\*, P < 0.01 determined by unpaired two-tailed Student *t*-test.

## Supplementary Fig. S4

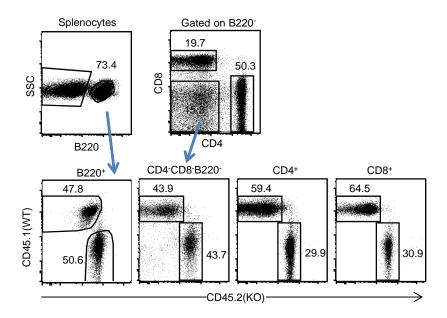


Figure S4. Role of mTORC1 in splenic T cell homeostasis in mixed bone marrow irradiation chimeric mice. WT (CD45.1<sup>+</sup>) and *Raptor*<sup>f/f</sup>-*CD4Cre* (CD45.2<sup>+</sup>) BM cells were mixed at a 1:1 ratio and adoptively transferred into sublethally irradiated TCR- $\beta^{-/-}\delta^{-/-}$  mice. Eight weeks later, splenocytes from the chimeric mice were analyzed for B220<sup>+</sup> B cells, CD4 and CD8 T-cell and B220<sup>-</sup>CD4<sup>-</sup>CD8<sup>-</sup> non-T/B cells. Data shown are representative of three experiments.