

## **SUPPLEMENTAL MATERIAL**

## St. Leger et al., https://doi.org/10.1084/jem.20170369

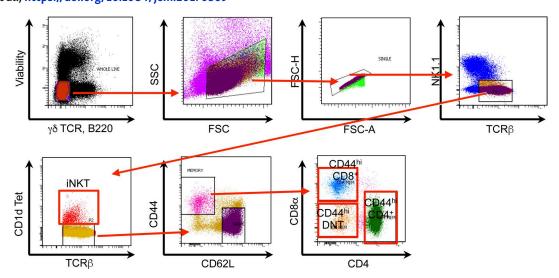


Figure S1. **Gating strategy for isolating CD44<sup>hi</sup> CD4<sup>+</sup>, CD44<sup>hi</sup> CD8<sup>+</sup>, CD44<sup>hi</sup> DNT, and iNKT cells.** Spleen and LN cells were initially sorted by negative selection for MHC II and B220 by MACS or isolated for T cells using T cell isolation columns from R&D Systems. After initial sorting, cells were stained with fluorescent antibodies against  $\gamma\delta$  TCR, B220, TCR $\beta$ , NK1.1, CD1d tetramer, CD62L, CD44, CD8, and CD4. Red boxes indicate populations of interest for the outlined studies.



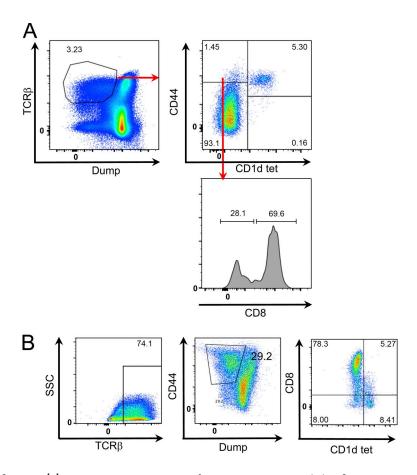


Figure S2. **Gating strategy for Fig. 5. (A)** Representative gating strategy for Fig. 5 A. Dump gate includes  $\gamma\delta$  TCR, B220, CD4, and CD62L. PLZF expression and IL-17 production was based off of the CD44<sup>hi</sup> CD8+, CD44<sup>hi</sup> DNT, or CD1d tet+ (iNKT). **(B)** Representative gating strategy to identify IL-17–producing cells in Fig. 5 B. Dump gate includes CD62L, B220,  $\gamma\delta$ TCR, and CD4.