

Figure S1. Effects of lymphocyte perturbation on acute TNF- α **-induced apoptosis.** (A) An example of a western blot for caspase 3 cleavage and β -Tubulin in the duodena of Rag1 null mice (left) and wild-type mice (right) as a time course following TNF- α (5 µg) stimulation. (B) Caspase 3 cleavage induced 2 hours post-TNF- α (5 µg) administration within the duodena of wild-type control mice (cyan), Rag1 null mice (red), TCR β/δ (Tcrb/Tcrd) null mice (dark red), and immuno-globulin heavy chain mu (Ighm) null mice (brown). Data are normalized to wild-type control mice and error bars represent SEM for 3 mice. (C) Quantification of CD8+ T cells in the duodena of wild-type mice by FACS in wild-type controls (clear) or mice pretreated with anti-MadCAM1 (2 mg/kg) for 2 hours (shaded). Treatment with anti-MadCAM1 prevents TNF- α -induced recruitment of T cells to the intestine. Error bars represent the SEM for 3 mice. (D) Caspase 3 cleavage within the duodena of wild-type mice with anti-MadCAM1 (shaded). Data are normalized to wild-type control mice pretreated with anti-MadCAM1 (shaded). Data are normalized to SEM for 3 mice (green), and wild-type mice pretreated with anti-MadCAM1 (shaded). Data are normalized to wild-type control mice and error bars represent the SEM for 3 mice. (D) Caspase 3 cleavage within the duodena of wild-type control mice (white), Rag1 null mice (green), and wild-type mice pretreated with anti-MadCAM1 (shaded). Data are normalized to wild-type control mice and error bars represent SEM for 3 mice.