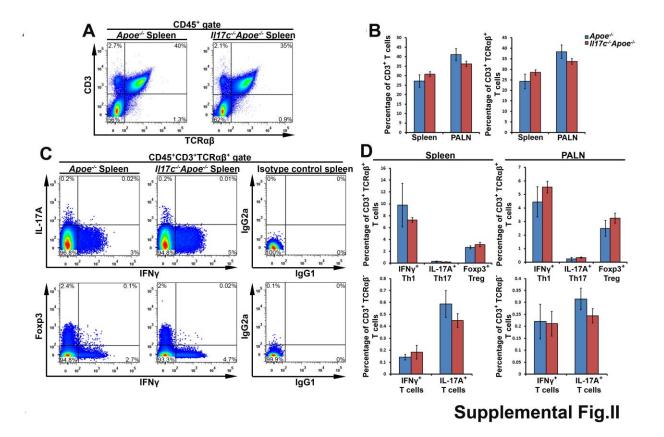


Supplemental Figure I: Aortic vascular cell flow cytometry gating scheme. Representative scheme for gating on vascular cell subsets for the flow cytometry experiments in Figure 1. Single CD45- vascular cells were gated and subdefined as follows: CD45-CD31+: endothelial cells, CD45-CD31- α SMA+: Smooth muscle cells, CD45-CD31- α SMA-CD29+: fibroblasts, CD45-CD31- α SMA-CD29-: remaining vascular cells.



Supplemental Figure II: Peripheral T cells and T cell subsets are not impacted by IL-17C deficient conditions.

(A-D) 12 week WD *Apoe*^{-/-} and *II17c*^{-/-}*Apoe*^{-/-} spleens and peri-aortic lymph nodes were stained with anti-CD45, CD3, TCR $\alpha\beta$, IL-17A, IFN γ , and Foxp3 antibodies or appropriate isotype controls, and assessed via flow cytometry. (A) Representative CD3 and TCR $\alpha\beta$ staining of *Apoe*^{-/-} and *II17c*^{-/-}*Apoe*^{-/-} splenocytes. (B) Quantification of the percentage of splenic and peri-aortic lymph node CD3⁺ T cells and CD3⁺TCR $\alpha\beta^+$ T cells. (C) Representative IL-17A, IFN γ , and Foxp3 staining within *Apoe*^{-/-} and *II17c*^{-/-}*Apoe*^{-/-} splenic CD3⁺TCR $\alpha\beta^+$ T cells. (D) Quantification of the percentage of splenic and peri-aortic lymph node Th1, Th17, Treg, CD3+ TCR $\gamma\delta^+$ IL-17A+ T cell and CD3+ TCR $\gamma\delta^+$ IFN γ + T cell contents. Red – *Apoe*^{-/-}, Blue – *II17c*^{-/-}*Apoe*^{-/-} mice. n=7-11 mice/genotype, 4 independent experiments, bars depict means±SEM.