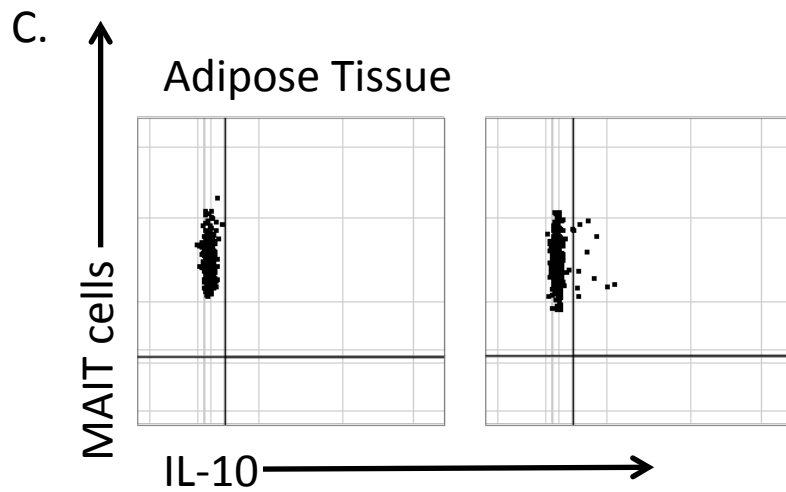
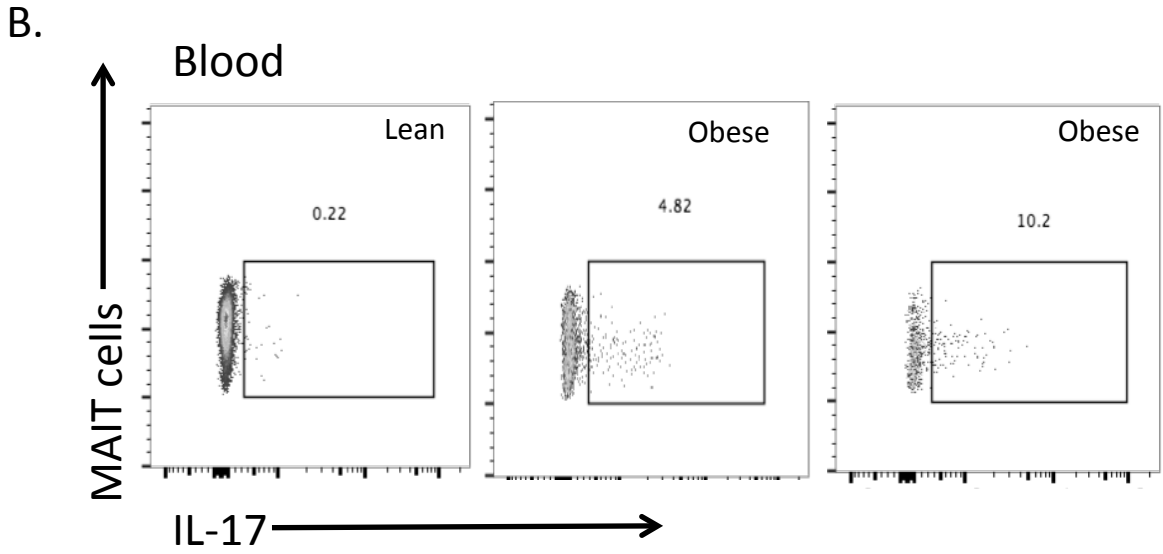
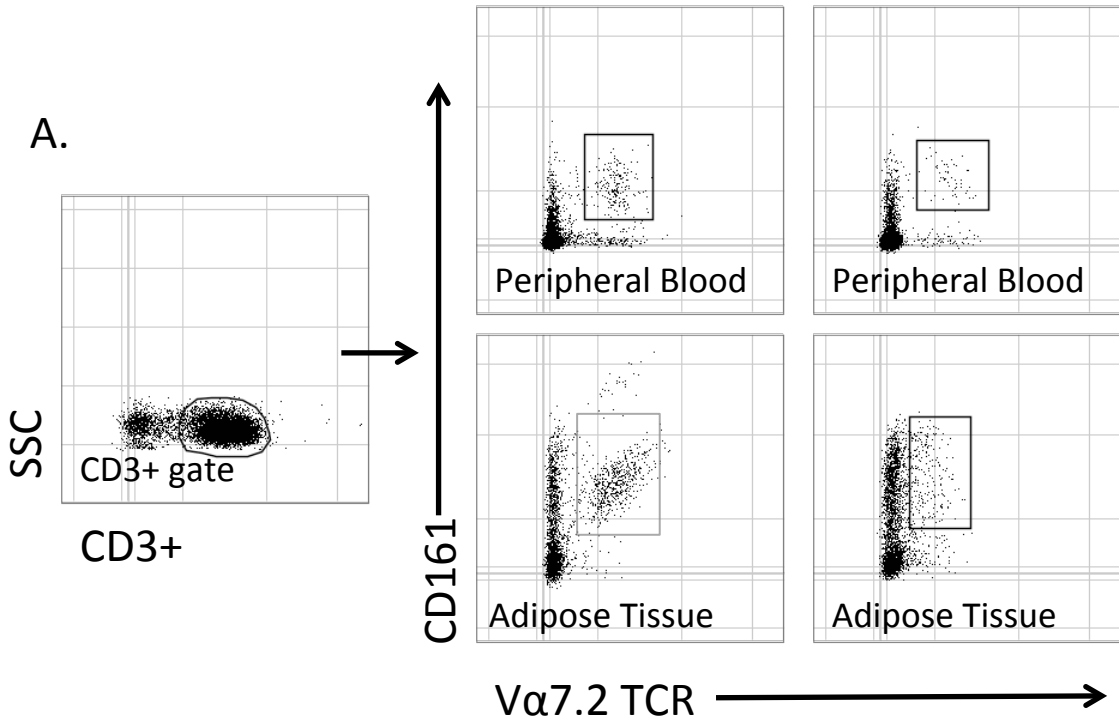


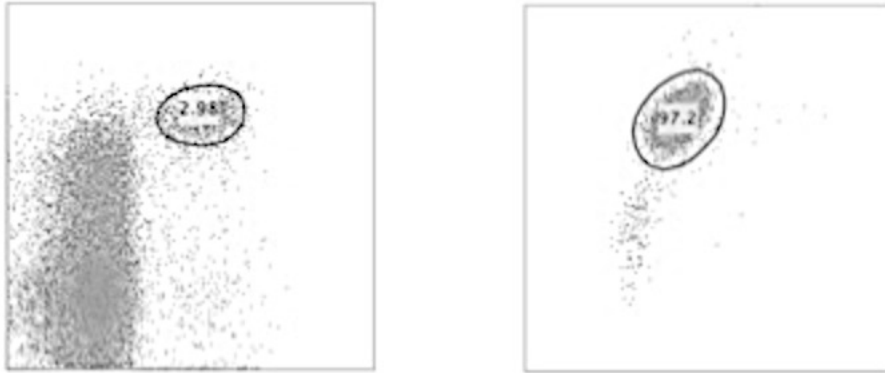
Supplementary Figure 1



Supplementary Figure 1 – MAIT cell flow cytometry gating and cytokine production. **A,** Representative flow cytometry dot plots showing the gating strategy employed to identify MAIT cells. Lymphocytes were selected according to forward and side scatter. CD3⁺ cells were then selected and the V α 7.2 TCR⁺ CD161⁺ population was determined to be MAIT cells. **B,** Sample flow plots showing IL-17 staining in 3 representative peripheral blood samples, again MAIT cells were selected as before and IL-17 positivity determined using unstained and FMO controls. **C,** Sample flow plots showing IL-10 staining in a representative adipose tissue sample, again MAIT cells were selected as before and IL-10 positivity determined using unstained and FMO controls.

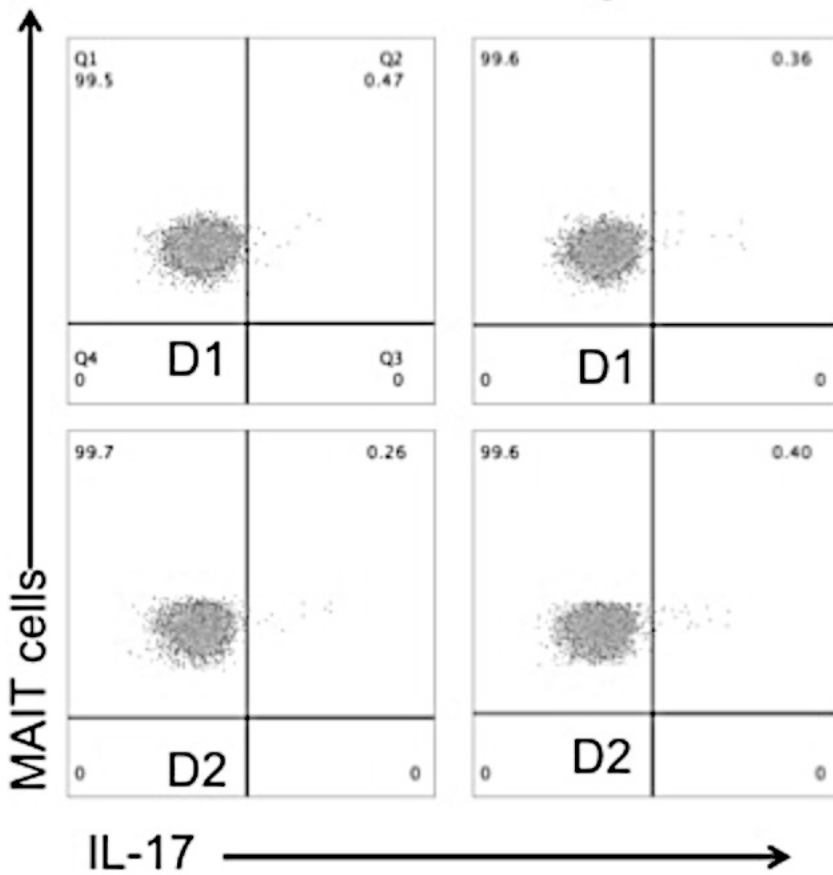
Supplementary Figure 2

A. Healthy Donor PBMC FACS sorted MAIT cells



B.

No Insulin + 38ng/ml Insulin



Supplementary Figure 2 – Insulin has no effect on MAIT cell cytokine production. **A**, Representative flow cytometry dot plots showing MAIT cells before and after cell sorting. **B**, Sample flow plots showing IL-17 intracellular staining in representative peripheral blood samples stimulated either in the absence or presence of insulin (38 ng/ml) for 24 hours. Gating was determined using unstained and FMO controls.