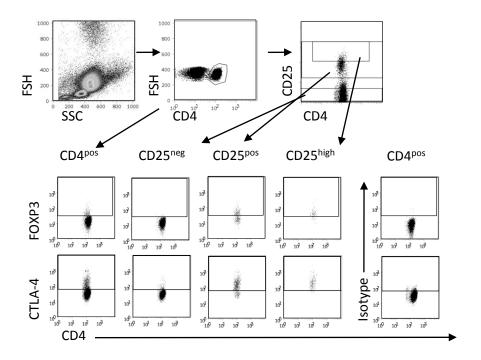
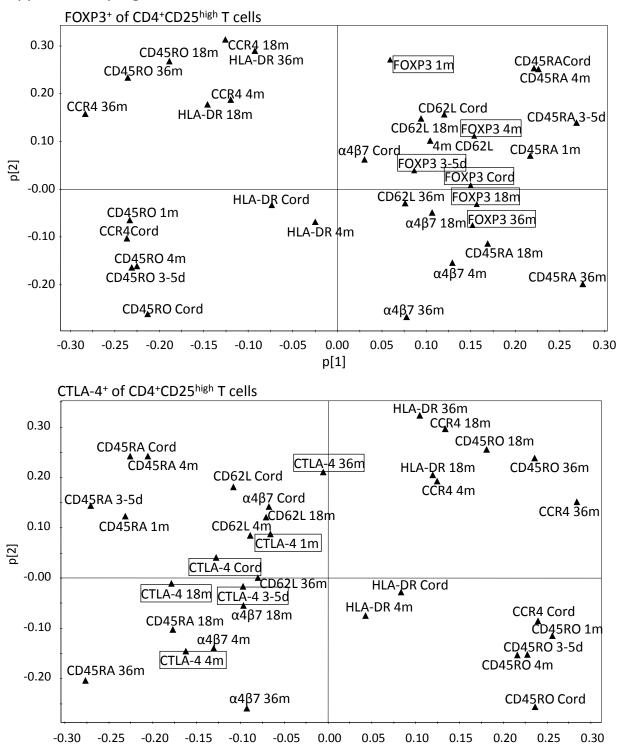
## Supplementary Figure 1



**Supplementary Figure 1.** Gating strategies for FOXP3 and CTLA-4 in CD4+, CD25<sup>neg</sup>, CD25+ and CD25<sup>high</sup> T cells. The lymphocyte population was recognized using forward (FSH) and side (SSC) scatter. Next, CD4+ lymphocytes where gated followed by the gates for CD25<sup>neg</sup>, CD25+ and CD25<sup>high</sup> (2% of CD4+ T cells that expressed high CD25). FOXP3+ and CTLA-4+ T cells were identified by comparing the expression of these markers within the CD4+, CD25<sup>neg</sup>, CD25+, CD25<sup>high</sup> T-cell populations and isotype control and setting the gates accordingly. The representative dot plots are derived from one infant at birth.

## Supplementary Figure 2



Supplementary Figure 2. Relationship between FOXP3<sup>+</sup> or CTLA-4<sup>+</sup>  $T_{regs}$  and expression of memory markers and homing receptors on CD4<sup>+</sup> T cells in blood during infancy. Principal component analysis (PCA) plot showing an overview of the data set including percentage of FOXP3<sup>+</sup> or CTLA-4<sup>+</sup>  $T_{regs}$  within the CD4<sup>+</sup>CD25<sup>high</sup> T cells and the fraction of CD4<sup>+</sup> T cells that express CD45RA, CD45RO at birth, 3-5 days, 1, 4, 18 and 36 months and HLA-DR,  $\alpha_4\beta_7$ , CD62L or CCR4 at birth, 4, 18 and 36 months of age.

p[1]