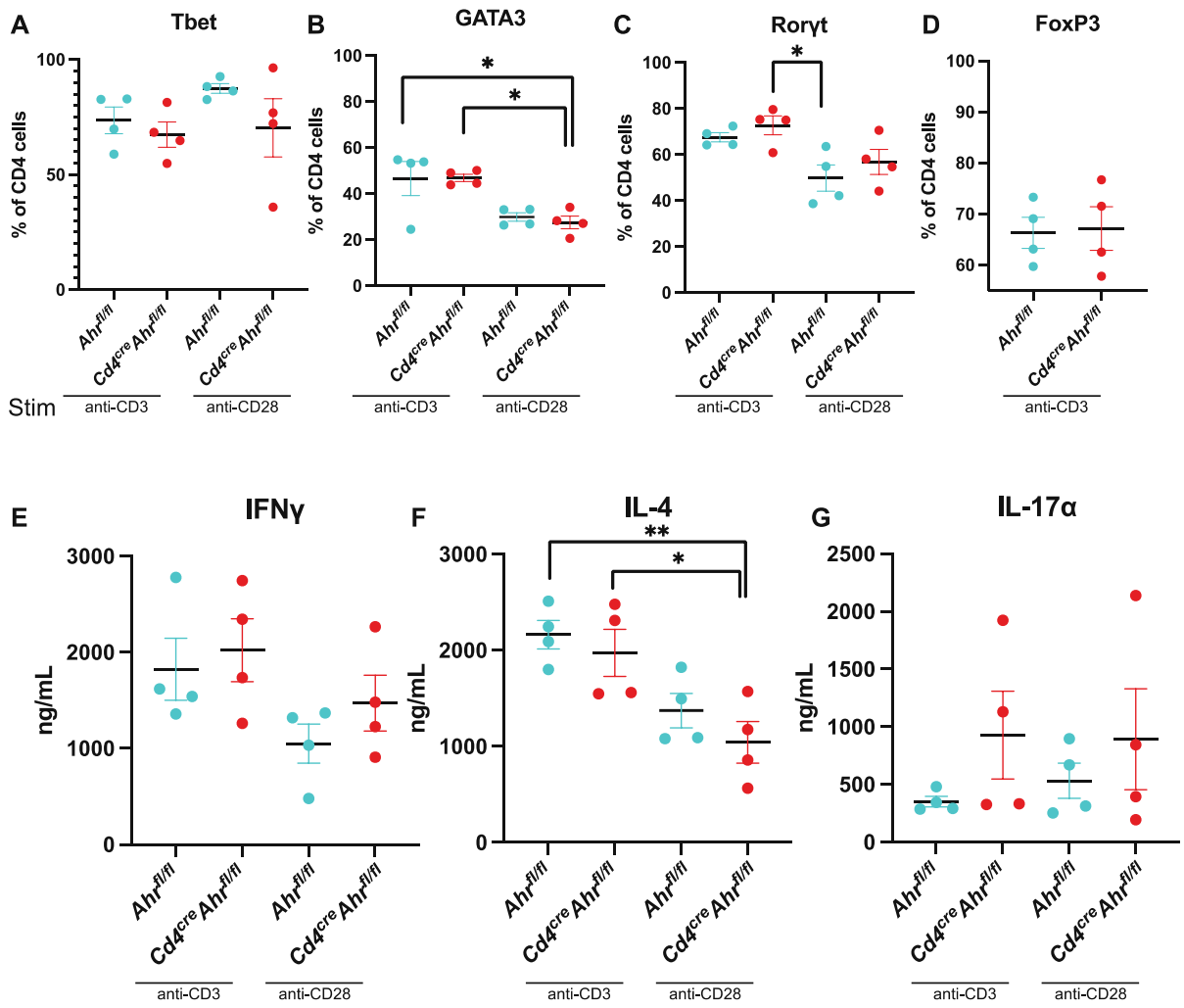

S2 Fig. CD4+ cells from separately housed *Cd4^{cre}Ahr$^{fl/fl}$* have no change in differentiation capacity or cytokine production. (**A**) Flow cytometric quantification of Tbet+ cells differentiated to T_H1 cell type and stimulated for 24 hours with either anti-CD3 or anti-CD28 (n=4 mice/group; N=1 experiment; Ordinary one-way ANOVA [p=0.2927]). (**B**) Flow cytometric quantification of GATA3+ cells differentiated to T_H2 cell type and stimulated for 24 hours with either anti-CD3 or anti-CD28 (n=4 mice/group; N=1 experiments; Ordinary one-way ANOVA [p=0.0076] followed by Tukey's Post-Hoc). (**C**) Flow cytometric quantification of RORγt+ cells differentiated to T_H17 cell type and stimulated for 24 hours with either anti-CD3 or anti-CD28 (n=4 mice/group; N=1 experiment; Ordinary one-way ANOVA [p=0.0163] followed by Tukey's Post-Hoc). (**D**) Flow cytometric quantification of FoxP3+ cells differentiated to T_{reg} cell type and stimulated for 24 hours with either anti-CD3 or anti-CD28 (n=4 mice/group; N=1 experiment; Ordinary one-way ANOVA [p=0.8801]). (**E**) IFNγ ELISA on supernatant of differentiated T_H1 cells stimulated for 24 hours with either anti-CD3 or anti-CD28 (n=4 mice/group; N=1 experiment; Ordinary one-way ANOVA [p=0.8639]). (**F**) IL-4 ELISA on supernatant of differentiated T_H2 cells stimulated for 24 hours with either anti-CD3 or anti-CD28 (n=4 mice/group; N=1 experiment; Ordinary one-way ANOVA [p=0.0064] followed by Tukey's Post-Hoc). (**G**) IL-17a ELISA on supernatant of differentiated T_H17 cells stimulated for 24 hours with either anti-CD3 or anti-CD28 (n=4 mice/group; N=1 experiment; Ordinary one-way ANOVA [p=0.4826]).