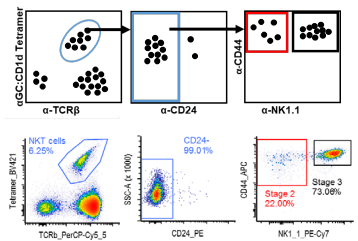
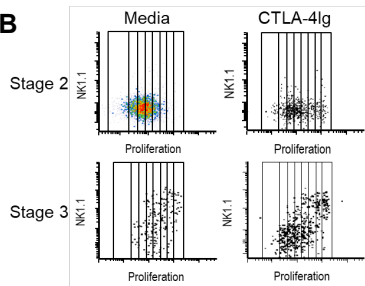
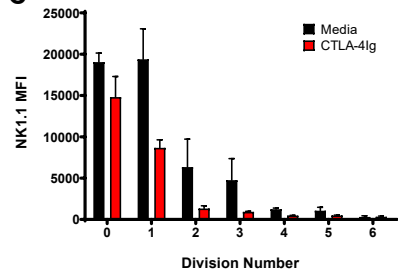


Supplementary Information

Thymic resident NKT cell subsets show differential requirements for CD28 co-stimulation during antigenic activation

Susannah C. Shissler, Nevil J. Singh, Tonya J. Webb

Department of Microbiology and Immunology and the Marlene and Stewart Greenebaum Comprehensive Cancer Center, University of Maryland School of Medicine, Baltimore, MD 21201

A**B****C**

Supplementary Figure 1. Stage 3 NKT cells downregulate NK1.1 after activation. (A) NKT cells were sorted as α -GalCer:CD1d tetramer⁺ TCR β Int⁺ CD24⁻ CD44⁺ and NK1.1⁻ (Stage 2) or NK1.1⁺ (Stage 3). Sorted NKT cells were co-cultured with splenocytes pre-loaded with vehicle or α -GalCer in the presence of media or CTLA-4Ig for 72 hours. (B) Representative plots of NKT cell division and expression of NK1.1 after co-culture with splenocytes loaded with 1ng/mL α -GalCer. (C) Mean fluorescence intensity (MFI) of NK1.1 on stage 3 NKT cells during successive rounds of division after co-culture with splenocytes loaded with 1ng/mL α -GalCer.