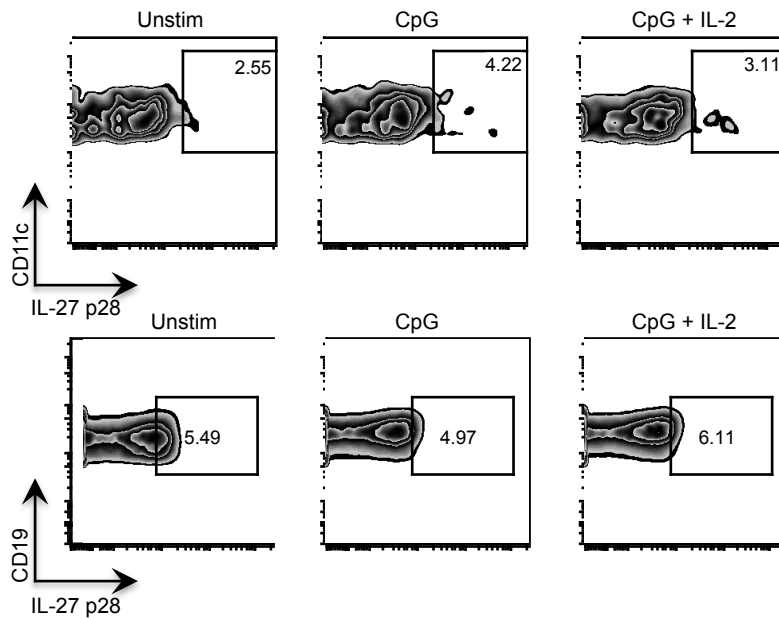


Supplemental Figure 1. Expression of IL-2 receptor chains and IL-2 promotion of IFN- γ production in CD11b⁺ CD11c⁻ cells.

A) CD11b⁺CD11c⁻ cells were isolated from the spleens of NOD and NOD.*Idd3* mice by cell sorting and stimulated with CpG for 4 h as indicated. Cells were then lysed for RNA isolation and use in real-time PCR. IL2R β (n= 3), IL2R γ c (n=3). ns=not significant. B) Expression of IL2R β and IL2R γ c on NOD CD11b⁺CD11c⁻ cells before (black line) and after stimulation with CpG (gray line). Filled histogram, fluorescence minus one control. C) CD11b⁺CD11c⁻ cells were prepared as in (A) and stimulated with either CpG or PGN. Left panel, CD25 (IL2R α) (n=4), *p=0.0003, **p=0.006, t-test. ns=not significant. Middle and right panels, data from two independent experiments. C) CD11b⁺CD11c⁻ cells were isolated from the spleens of NOD mice

and stimulated with PGN in the presence or absence of 25ng/mL IL-2 for 24h, (n=5). *p=0.0447, t-test. Cytokine measured by CBA.



Supplemental Figure 2. IL-27 p28 is mainly produced by CD11b⁺ CD11c⁻ cells.

NOD APC were T-depleted and stimulated for 18h with media, CpG, or CpG+IL-2. Cells were washed, incubated with GolgiStop for 4h and intracellular cytokine staining preformed. Data are representative of five experiments.