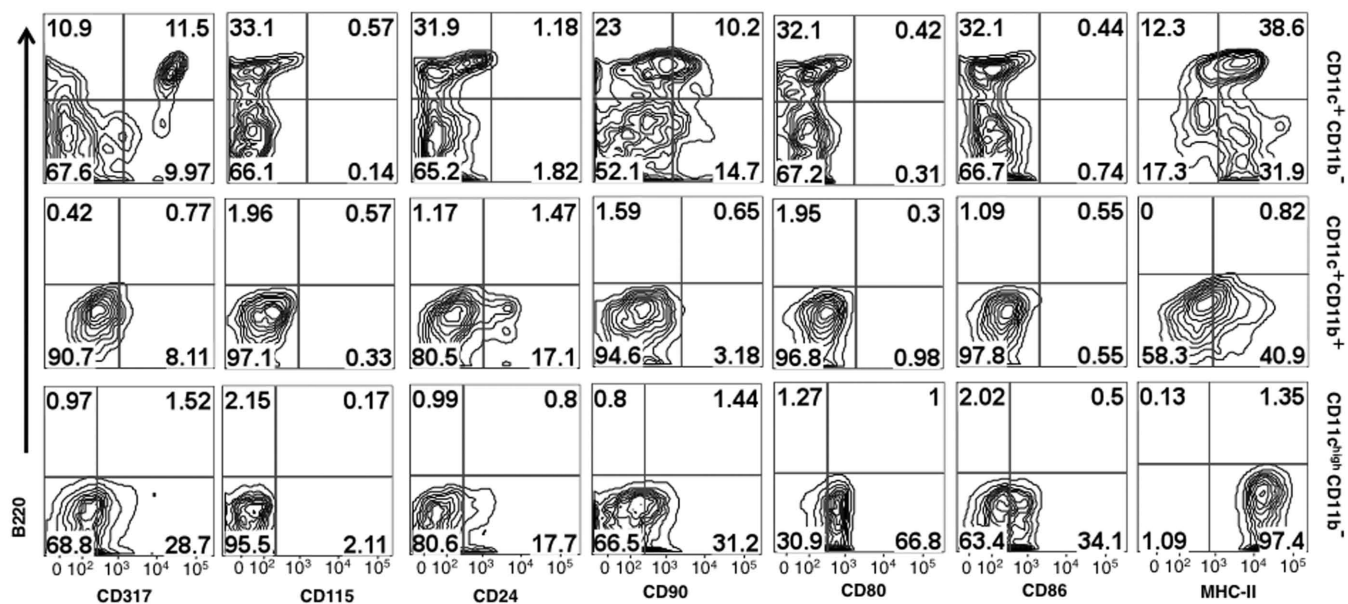
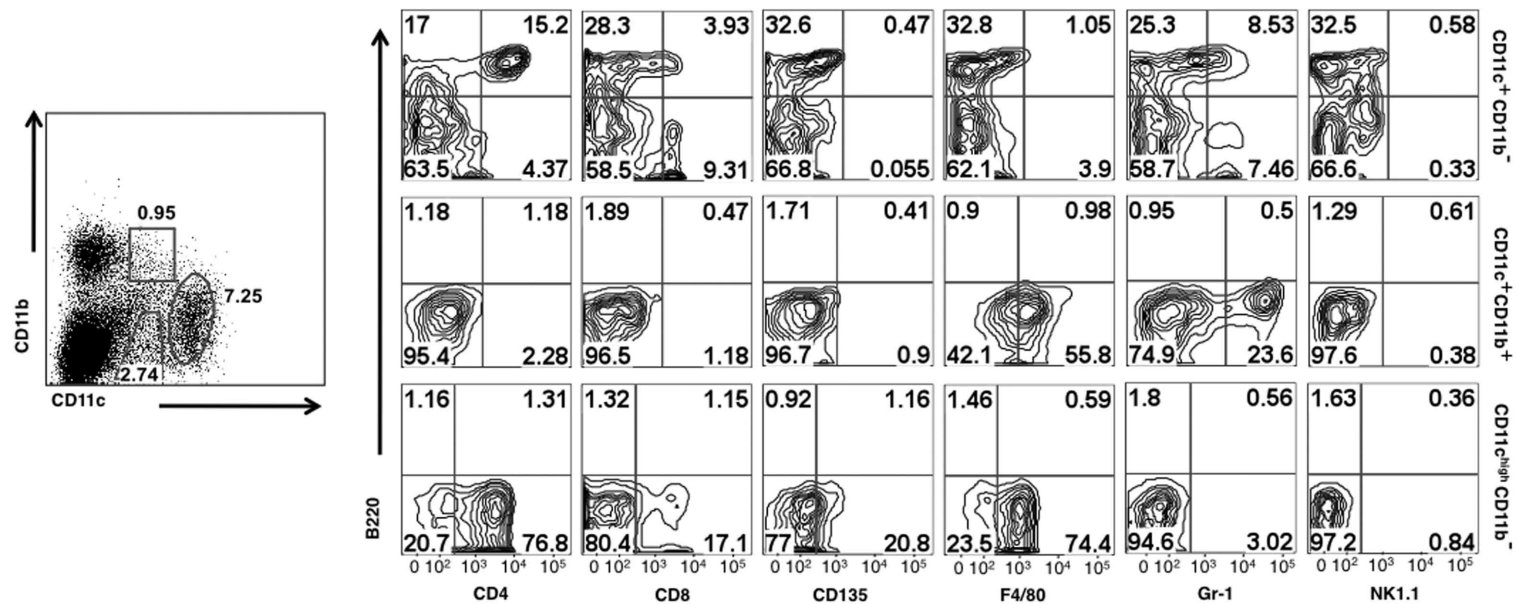


SUPPLEMENTARY FIGURE

SUPPLEMENTAL FIGURE 1. Phenotypes of CD11b⁻ and CD11b⁺ APC subsets from spleen. Splenocytes were isolated from B6 spleen and stained with biotininated lineage antibodies (CD3, CD19, Ig M, TER119, and DX5), followed by incubation with streptavidin APC, CD11c FITC, CD11b APC-Cy7, and PE conjugated antibodies for CD4, CD8, CD135 (FLT3), F4/80, Gr-1 (Ly6G/C), NK1.1, CD317 (PDCA-1), CD115, CD24, CD90, CD80, CD86 and I-A^b (MHC-II) and PE conjugated isotype matched antibody. Cells were gated on CD11c^{hi} CD11b^(lo/neg) and CD11c^{+(lo and medium)} CD11b^{neg}, and CD11c^{+(lo and medium)} CD11b⁺. FACS plots are from a single experiment representative of 5 replicate experiments.



SUPPLEMENTARY FIGURE

SUPPLEMENTAL FIGURE 2. Enhanced *ex vivo* alloantigen specific T-cell proliferation and Th1 cytokine polarization of donor T-cells from recipients of CD11b⁻ APC. B6 donor T-cells were isolated from the spleens of recipient mice on day 15 post-transplant by negative MACS selection using a cocktail of antibodies (Methods) plus anti-CD45.2. CFSE-labeled donor T-cells were cultured with irradiated (30Gy) B6 syngenic, B10 recipient-type, or B/c third-party splenocytes. **A-B**, Numbers of cell divisions in **(A)** CD4⁺ and **(B)** CD8⁺ donor T-cells following MLR co-culture with B10 splenocytes (minimal proliferation was seen for T-cells co-cultured with syngenic B6 or B/c stimulators, data not shown). **C-D**, Percentages of B6 CD4⁺ and CD8⁺ T-cells harvested day 15 post-transplant expressing IFN- γ or IL-4 after 3 days MLR culture against syngenic, recipient, or B/c third-party splenocytes, determined by intracellular cytokine staining. *p<0.05, **p<0.01, ***p<0.001, comparing results from HSC & T-cell group with those from groups that also received APC.

