

Figure S1. Representative FACS plots of data presented in Fig. 2. ISTF and anti-4-1BB mAb co-treatment induces marked expansion of CD11c⁺CD8⁺ T cells in splenocytes of tumor-bearing mice. Renca tumors were established and four groups of mice were treated with ISTF and/or anti-4-1BB mAb as described in Fig. 1. On day 14, splenocytes were excised; double stained with PE-, Cy- and FITC-conjugated antibodies and analyzed by FACS. Representative flow plots showing the percentages of (A) CD3⁺CD8⁺ T cells, (B) CD11c⁺CD8⁺ T cells, (C) CD4⁺Foxp3⁺ Treg cells, (D) CD11b⁺F480⁺ macrophages, (E) CD3⁺CD4⁺ T cells, (F) CD3⁺DX5⁺ natural killer cells, (G) CD11b⁺Gr1⁺ myeloid cells, CD11b⁺Gr1⁺int (myeloid-derived suppressor cells), CD11b⁺Gr1⁺hi (neutrophils) and (H) B220⁺ B cells. ISTF, immunostimulatory factor; mAb, monoclonal antibody; FACS, fluorescence activated cell sorting.

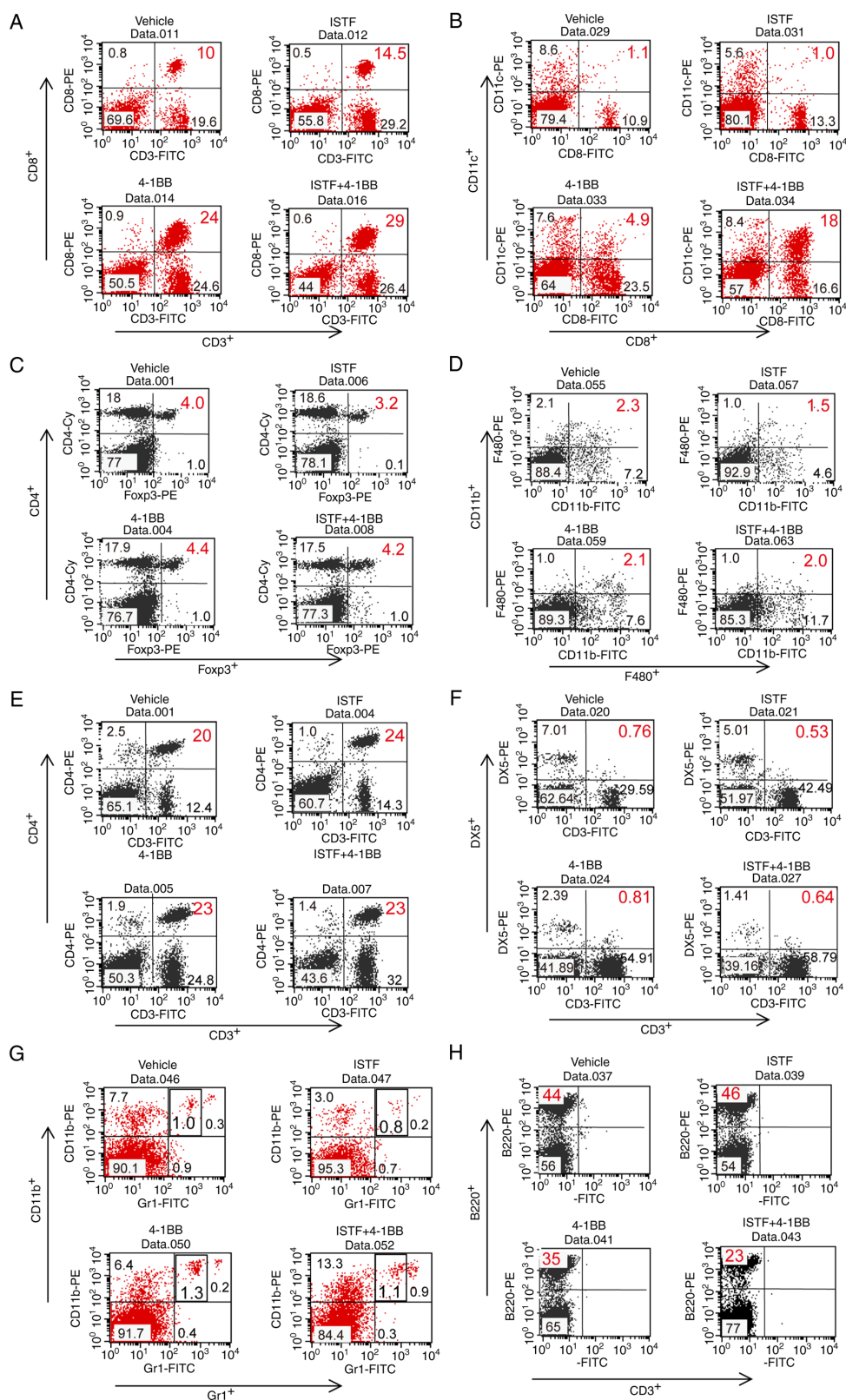


Figure S2. Representative FACS plots of data presented in Fig. 4. Combined therapy with ISTF and anti-4-1BB mAb induces marked expansion of CD11c⁺CD8⁺ T cells in TILs. Renca tumors were established and four groups of mice were treated with ISTF and/or anti-4-1BB mAb as described in Fig. 1. On day 14, the mice were sacrificed and tumors were harvested. Tumor-infiltrating lymphocytes were stained with PE-, Cy- and FITC-conjugated antibodies and analyzed by FACS. Representative flow plots showing the percentages of CD3⁺CD4⁺ T cells among TILs. ISTF, immunostimulatory factor; mAb, monoclonal antibody; FACS, fluorescence activated cell sorting; TILs, tumor-infiltrating lymphocytes.

