

Supplementary information, Figure S1. LKB1 deficiency does not affect the development of DCs, T cells or B cells. a, Real-time PCR and immunoblot analyses of Stk11 mRNA and LKB1 protein expression in splenic DCs (CD11c+MHC-II+TCRβ-CD19-CD49b-) from WT and LKB1 $^{\Delta DC}$  mice. b, Real-time PCR analysis of Stk11 mRNA expression in splenic CD11b+F4/80+ macrophages from WT and LKB1 $^{\Delta DC}$  mice. c, Flow cytometry analysis of splenic conventional DC (cDC), CD8 $\alpha$ + cDC (CD8 $\alpha$ +CD11b-), CD8 $\alpha$ - cDC (CD8 $\alpha$ -CD11b+) and pDC populations in WT and LKB1 $^{\Delta DC}$  mice. d, Statistics of frequencies and cell numbers of splenic DC populations in WT and LKB1 $^{\Delta DC}$  mice. e, Flow cytometry analysis of CD4+ and CD8+ T cell populations in the spleen, PLN, MLN and thymus, and B cells in the spleen of WT and LKB1 $^{\Delta DC}$  mice. f, Tumor growth curve in WT and LKB1 $^{\Delta DC}$  mice following inoculation of B16-OVA tumor cells (WT, n = 9; LKB1 $^{\Delta DC}$  mice, n = 7). g, Statistics of frequencies of IFN $\gamma$ -producing CD4+ and CD8+ T cells in tumor tissues of WT and LKB1 $^{\Delta DC}$  mice after inoculation of MC38 tumor cells for 13-14 days. Data in plots indicate the means ± s.e.m; each symbol represents an individual mouse. Numbers in gates indicate percentage of cells. NS, not significant; \*\*\*P < 0.001; \*\*\*\*P < 0.0001; two-tailed Mann-Whitney test (d, frequency; g), two-tailed unpaired Student's t test (d, cell number) or two-way ANOVA (f). Data are from at least three (a-e, g) independent experiments.