

Supplementary Figure 1. T cell development in the *Zfp91*^{fl/fl}*Cd4*-Cre mice. (A) Immunoblot analysis of ZFP91 in whole-cell lysates of splenic CD4⁺ T cells from *Zfp91*^{+/+}*Cd4*-Cre (WT) mice and *Zfp91*^{fl/fl}*Cd4*-Cre (KO) mice. (B) Cell numbers of CD4⁻ CD8⁻ (DN), CD4⁺CD8⁺ (DP), CD4⁺CD8⁻ (CD4), CD4⁻CD8⁺ (CD8) in the thymus, and percentage of CD4⁺ and CD8⁺ T cells in the spleen and peripheral lymph nodes (pLN) of 6week-old *Zfp91*^{+/+}*Cd4*-Cre and *Zfp91*^{fl/fl}*Cd4*-Cre mice (n=6). (C) Flow cytometric analysis of the percentage of memory-like (CD44^{hi}CD62L^{lo} for CD4⁺ and CD44^{hi} for CD8⁺ T cells) CD4⁺ and CD8⁺ T cells in total splenocytes from 6-week-old *Zfp91*^{+/+}*Cd4*-Cre and *Zfp91*^{fl/fl}*Cd4*-Cre mice (n=6). (D) Flow cytometric analysis of the percentage of Foxp3⁺ Treg cells in the thymus (Thy), spleen (Spl) and peripheral lymph nodes (pLN) from 6-week-old *Zfp91*^{+/+}*Cd4*-Cre and *Zfp91*^{fl/fl}*Cd4*-Cre mice (n=3). Data are representative of three independent experiments. ns, not statistically significant; two-tailed Student's t test.



Supplementary Figure 2. Antitumor immune responses in the *Zfp91*^{fl/fl}*Cd4*-Cre mice. (A) Survival curves of *Zfp91*^{+/+}*Cd4*-Cre and *Zfp91*^{fl/fl}*Cd4*-Cre mice given subcutaneous injection of 5×10^5 B16-F10 melanoma cells (n=10). (B and C) Flow cytometric analysis of T cells in tumors of *Zfp91*^{+/+}*Cd4*-Cre and *Zfp91*^{fl/fl}*Cd4*-Cre mice injected s.c. with B16-F10 melanoma cells (day 14). The data are presented as representative plots (B) and as summary graphs (C, n=5). Data are representative of two independent experiments. Data are presented as the mean \pm SEM. **, P < 0.01. (A) was analyzed by log-rank (Mantel-Cox) test and (C) was analyzed by two-tailed Student's t test.



Supplementary Figure 3. Splenic T cell activity of tumor-bearing mice with transferred OT-I cells. (A) Number of OT-I cells in spleen from MC38-OVA tumor bearing mice with transferred CFSE-labeled WT OT-I and CTV-labeled KO OT-I cells (at day 7 after OT-I cells adoptive transfer) (n=4). (B-D) Percentage and number of IFN- γ^+ OT-I cells (B, C) or Ki-67⁺ OT-I cells (D) in spleen from MC38-OVA tumor bearing mice with transferred OT-I cells (at day 7 after OT-I cells (at day 7 after OT-I cells (D) in spleen from MC38-OVA tumor bearing mice with transferred OT-I cells (at day 7 after OT-I cells adoptive transfer) (n=4). Data are representative of three independent experiments. ns, not statistically significant; two-tailed Student's t test.



Supplementary Figure 4. Th cell differentiations in *Zfp91*^{+/+}*Cd4*-**Cre and** *Zfp91*^{fl/fl}*Cd4*-**Cre T cells.** (**A**) Naïve CD4⁺ T cells from *Zfp91*^{+/+}*Cd4*-Cre mice and *Zfp91*^{fl/fl}*Cd4*-Cre mice were stimulated under standard Th1, Th17 or Treg conditions and harvested on day 5 for flow cytometric analysis. (**B**) Naïve CD4⁺ T cells from *Zfp91*^{+/+}*Cd4*-Cre mice and *Zfp91*^{fl/fl}*Cd4*-Cre mice were stimulated under standard Th0 or Th2 conditions and harvested on day 5 for qRT-PCR analysis (n=3). Data are representative of three independent experiments. ns, not statistically significant; two-tailed Student's t test.



Supplementary Figure 5. ZFP91 interacts with PP2Ac. (**A**) Lysates from HEK293T cells transfected with the indicated expression vectors were subjected to IP using an antibody against Flag. (**B**) Lysates from T cells from $Zfp91^{+/+}Cd4$ -Cre (WT) mice and $Zfp91^{fl/fl}Cd4$ -Cre (KO) mice stimulated with anti-CD3 and anti-CD28 antibodies for 4 hours were subjected to IP using an antibody against ZFP91. Data are representative of three independent experiments.