

Fig. S9

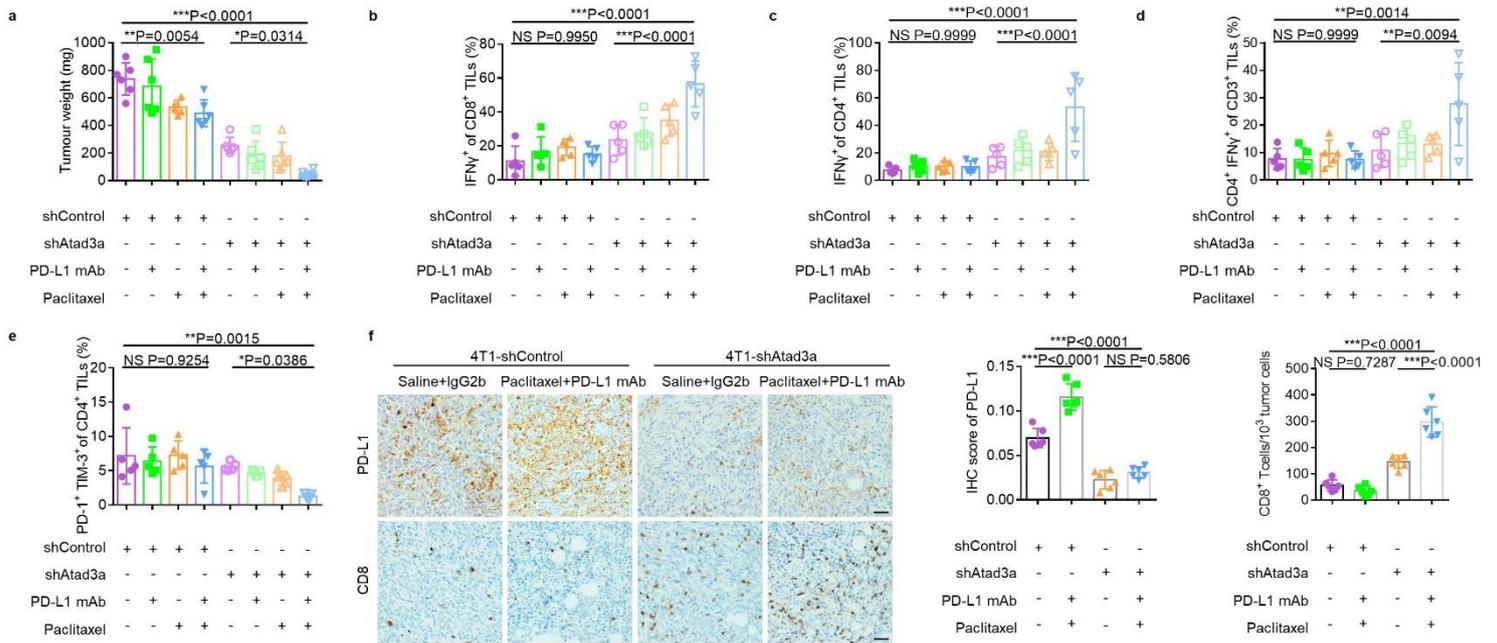


Fig. S9 Knockdown of Atad3a promotes the efficacy of combined therapy with anti-PD-L1 antibody and paclitaxel.

a Tumor volume of orthotopic 4T1 tumors formed by control and Atad3a-knockdown cells with vehicle, anti-PD-L1 monoclonal antibody (PD-L1 mAb), paclitaxel (PTX) or combined anti-PD-L1 antibody with paclitaxel treatment (PD-L1 mAb + PTX).

IgG2b and saline were used as controls ($n = 6$, one-way ANOVA). **b–e** Quantification of the percentages of tumor-infiltrating IFN γ^+ CD8 $^+$ (**b**), IFN γ^+ CD4 $^+$ (**c**, **d**) and PD-1 $^+$ TIM-3 $^+$ CD4 $^+$ T cells (**e**) in 4T1 tumors formed by control and Atad3a-knockdown cells receiving various treatments ($n = 5$, one-way ANOVA).

f Left, IHC staining of PD-L1 and CD8 on serial sections of control and Atad3a-knockdown tumors receiving vehicle or anti-PD-L1 antibody plus paclitaxel combination therapy. Scale bars, 50 μ m. Right, IHC score of PD-L1 and the quantification of the number of tumor-infiltrating CD8 $^+$ T cells in control or Atad3a-knockdown tumors with vehicle and combination therapy ($n = 6$, one-way ANOVA). Data are representative of two independent experiments and are shown as means \pm SD.