
SUPPLEMENTARY FIGURE LEGENDS

Supplementary Figure 1 PD-L1 mRNA level is associated with CD8+ T cells infiltration and prognosis in MSS CRC. (A) PD-L1 protein expression in CRC tissues as determined by IHC (scale bar, 100 μ m). (B) PD-L1 mRNA level in CRC tissues as determined by ISH (scale bar, 100 μ m). (C) CIBERSORT analyzes the correlation between CD274 and CD8+ T cells, activated CD4 T cells, Macrophages M1 and Tregs in CRC samples of TCGA database. (D) CIBERSORT analyzes the correlation between CD274 and CD8+ T cells, activated CD4 T cells, Macrophages M1 and Tregs in MSS CRC samples of TCGA database. *P* values and *R* values were calculated based on the analysis of Pearson's correlation. Student's *t* test. **p*<0.05, ***p*<0.01 and ****p*<0.001.

Supplementary Figure 2 miR-15b-5p is a regulator of PD-L1 at post-transcriptional level in MSS CRC. (A) PD-L1 expression in normal colon epithelial cells, inflammation-associated colon epithelial cells and cancer cells of colitis-associated cancer patients (scale bar, 200 μ m; magnification scale bar, 100 μ m). (B) miR-15b-5p, but not miR-15a-5p, inhibited PD-L1 expression in mouse intestinal cancer cell lines CT26 and MC38, but not miR-15a-5p. (C) RT-qPCR were performed to analysis the expression of PD-L1 in mRNA level after transfection of miR-15b-5p mimic in CT26 and MC38 cell lines. (D) Schematic representation of sequences of the wild-type and the mutated 3'-UTR luciferase reporters of mice. (E) Schematic representation of sequences of the wild-type and the mutated 3'-UTR luciferase

Anti-IL17A sensitizes immunotherapy in MSS CRC

reporters of human. (F) Expression of PD-L1 in protein levels were analyzed by Western Blotting analysis after transfection of miR-15b-5p mimic or inhibitor into SW620 and HT29 cell lines. (G) RT-qPCR were performed to analysis the expression of PD-L1 in mRNA level after transfection of miR-15b-5p mimic or inhibitor in human CRC cell lines. Data represent mean \pm SD. Results are representative of at least 3 separate experiments. Student's t test. * $p < 0.05$, ** $p < 0.01$ and *** $p < 0.001$.

Supplementary Figure 3 miR-15b-5p inhibits CRC tumorigenesis and sensitizes anti-PD-1 therapy by targeting PD-L1 in murine models. (A) Body weight changes of miR-15b-5p blocking and control mice were recorded throughout experiment. (B) Survival of mice miR-15b-5p blocking and control mice. (C) The weight of miR-15b-5p blocking and control group tumors were measured. (D) CT26 and MC38 cells were transfected with miR-15b-5p or miR-sc. miR-15b-5p mRNA levels were determined via RT-qPCR assay. (E) ISH analysis of miR-15b-5p expression in CT26 and MC38 tumors of subcutaneous transplantation. (F) Tumor growth of miR-sc, miR-15b-5p MC38 cells with or not PD-1 antibody treated in C57BL/6 mice (n = 7 mice per group). (G) Survival of mice bearing syngeneic MC38 tumors following treatment with PD-1 antibody. (n = 9 mice per group). (H) Intracellular cytokine staining of CD8⁺ IFN- γ ⁺ cells in the CD3⁺ T cell populations from isolated tumor-infiltrating lymphocytes. Data represent mean \pm SD. ANOVA followed by Tukey's multiple comparison test was applied. Cumulative survival time was estimated by the Kaplan-Meier method, and the log-rank test was applied to

compare the groups. * $p < 0.05$, ** $p < 0.01$ and *** $p < 0.001$.

Supplementary Figure 4 NRF1 is the major transcription factor for IL-17A to accumulate PD-L1 protein. (A) Schematic representation of predicted NRF1 binding sites within the human miR-15b-5p promoter (Left). Ch-IP analyses of NRF1 binding to the miR-15b-5p promoter using antibodies against NRF1 (Right). (B, C) Effects of NRF1/YY1 overexpression or knockdown on miR-15b-5p expression through RT-qPCR analyses of miR-15b-5p expression in MC38 cells. (D) Schematic representation of sequences of the wild-type and the mutated promoter luciferase reporters of miR-15b-5p in vector of pGL4.20. (E, F) Modulation of the miR-15b-5p promoter activity by NRF1 overexpression or knockdown in MC38 cells. Data represent mean \pm SD. Results are representative of at least 3 separate experiments. Student's t test. * $p < 0.05$, ** $p < 0.01$ and *** $p < 0.001$.

Supplementary Figure 5 Blocking IL-17A enhances the efficacy of anti-PD-1 therapy in murine model by promoting PD-L1 protein degradation. (A) Tumor growth of MC38 cells with IL-17A antibody and/or PD-1 antibody treated in C57BL/6 mice (n = 7 mice per group). (B) Survival of mice bearing syngeneic MC38 tumors following treatment with IL-17A antibody and/or PD-1 antibody (n = 9 mice per group). (C) FACS analysis of staining of CD8⁺ IFN- γ ⁺ cells in the CD3⁺ T cell populations from isolated tumor-infiltrating lymphocytes. (D) FACS analysis of staining of CD11b⁺ and Gr1⁺ cells in the tumor-infiltrating lymphocyte. (E)

Anti-IL17A sensitizes immunotherapy in MSS CRC

Immunofluorescence was used to analyze the staining of CD3+ cells and CD11b+ cells in MC38 tumors. Data represent mean \pm SD. ANOVA followed by Tukey's multiple comparison test was applied. Cumulative survival time was estimated by the Kaplan-Meier method, and the log-rank test was applied to compare the groups. * p <0.05, ** p <0.01 and *** p <0.001.