Supplementary Material

Supplementary FIGURE LEGENDS

Figure S1 | GSK843 induces the accumulation of CD11b⁺Gr-1⁺ cells in the liver and spleen of ConA-treated mice. Mice were pre-treated i.p. with GSK843 (0.5mg/kg; MCE, USA) or vehicle 1 hour before ConA (15 mg/kg) administration; mice were sacrificed 12 hours later after ConA injection and livers and spleen were collected. Representative dot-plots (left) and the histograms (right) show the percentages of CD11b⁺Gr-1⁺ cells in HMNCs and spleen cells of mice treated with GSK843+ConA or ConA alone. All the values are shown as mean ± SD. ns., not significant; *P < 0.05, **P < 0.01.

Figure S2 | **The effect of B6-8C5 antibody on Gr-1**⁺ **MDSCs depletion.** Hepatic mononuclear cells (HMNCs) and spleen cells from IgG+GSK872+ConA treated- and aGr1+GSK872+ConA treated-mice were assayed by FACS. (A) Percentages of CD11b⁺Ly6G^{hi}Ly6C^{lo} cells (PMN-MDSCs) and CD11b⁺Ly6G^{lo}Ly6C^{hi} (Mo-MDSCs) cells in CD45⁺CD11b⁺ cells were analyzed by flow cytometry. The percentages (up) and absolute cell numbers (down) of PMN-MDSCs and Mo-MDSCs in HMNCs (B) and spleen cells (C). All the values are shown as mean ± SD. ns., not significant; **P* < 0.05, ***P* < 0.01.

Figure S3 | In vivo depletion of Gr1-positive MDSCs aggravates ConA induced liver injury. Mice were given anti-Gr1 depleting antibody ($250\mu g/mice$) or control IgG ($250\mu g$) 36 hours before ConA(15mg/kg) treatment, mice were sacrificed 12 hours after ConA-treatment and blood and liver samples were collected. (A) Percentages of CD11b⁺Gr-1⁺MDSCs and its subtypes (PMN-MDSCs and Mo-MDSCs) cells in livers were analyzed by flow cytometry. (B)The Serum levels of ALT/AST in two groups. (C) Representative photomicrographs (H&E staining; original magnification 200×; scale bars, 50 um) of livers in two groups. All the values are shown as mean ± SD. ns., not significant; **P* < 0.05, ***P* < 0.01.

Figure S1



Figure S2



Figure S3

