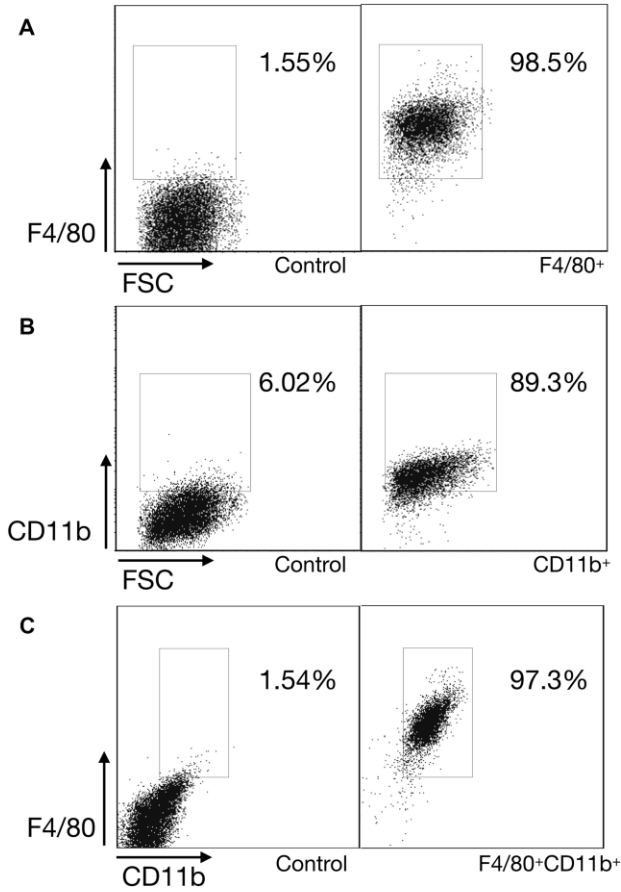
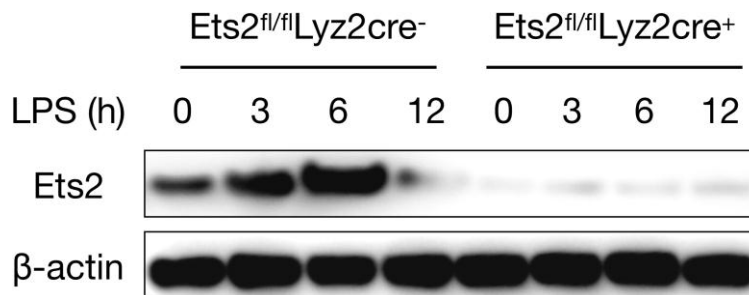


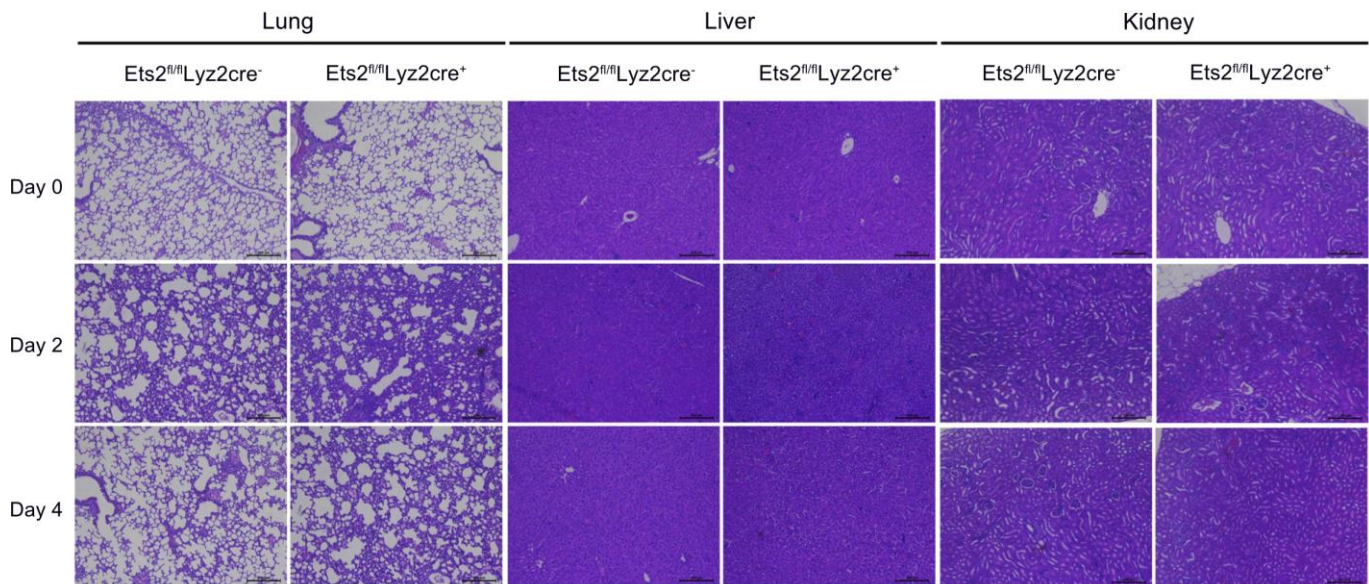
SUPPLEMENTARY FIGURES



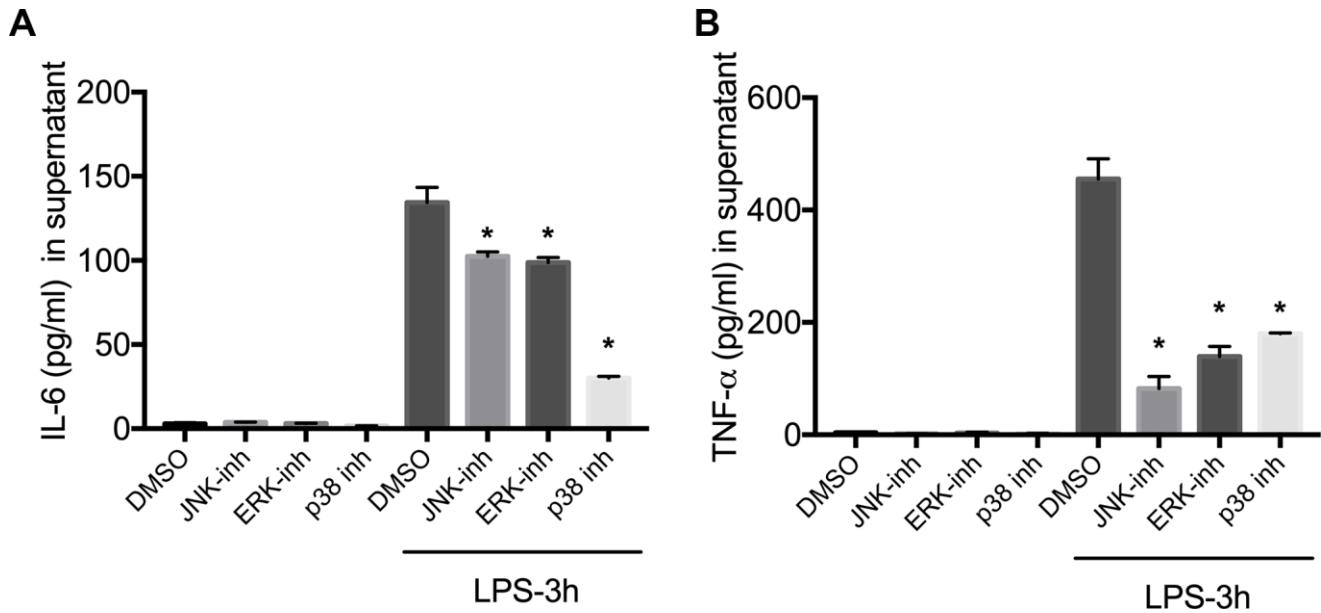
Supplementary Figure 1. Representatives of dot plots of peritoneal macrophage stained with F4/80 and CD11b in flow cytometry.



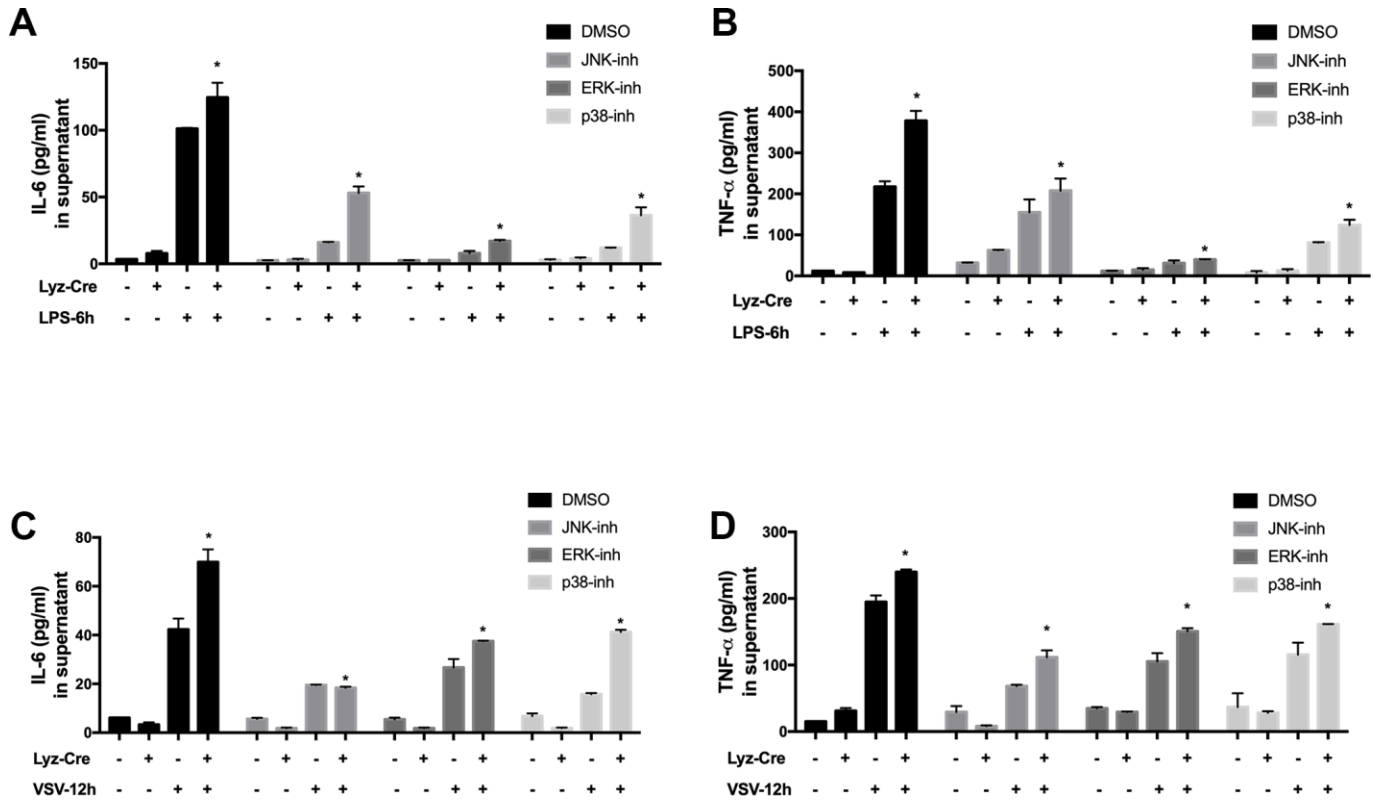
Supplementary Figure 2. Western blot of Ets2 and β-actin of Ets2-deficient (Ets2^{fl/fl}Lyz2cre⁺) or wild-type (Ets2^{fl/fl}Lyz2cre⁻) PMs stimulated with 100 ng/ml LPS in indicated times (0, 3, 6, 12 hours).



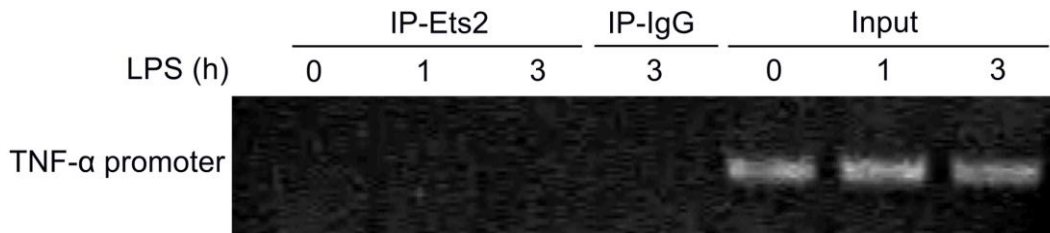
Supplementary Figure 3. Representatives of H&E staining of lung, liver and kidney in postoperative Day 2, and 4 of Ets2^{fl/fl}Lyz2cre⁻ or Ets2^{fl/fl}Lyz2cre⁺ mice underwent CLP surgery. Day 0 present the Sham surgery procedure.



Supplementary Figure 4. (A, B) ELISA of IL-6 (A) and TNF-α in the supernatant of peritoneal macrophage pre-treated with DMSO or inhibitor of ERK, JNK, and p38 respectively and stimulated with LPS for 3h. Data are shown as the mean ± s.d. of three samples. Student's t-test compared with the DMSO. *, P<0.05.



Supplementary Figure 5. (A–C) ELISA of IL-6 (A, C) and TNF- α (B, D) in the supernatant of peritoneal macrophage of *Ets2^{fl/fl}Lyz2cre⁻* or *Ets2^{fl/fl}Lyz2cre⁺* mice, pre-treated with DMSO or inhibitor of ERK, JNK, and p38 respectively and stimulated with 100 ng/ml LPS 6h (A, B) or 10 MOI VSV (C, D) for 12h. Data are shown as the mean \pm s.d. of three samples. Student's t-test compared with the DMSO. *, $P < 0.05$.



Supplementary Figure 6. ChIP analysis of the TNF- α promoter using an Ets2 antibody in mouse primary peritoneal macrophages treated with LPS for the indicated times.