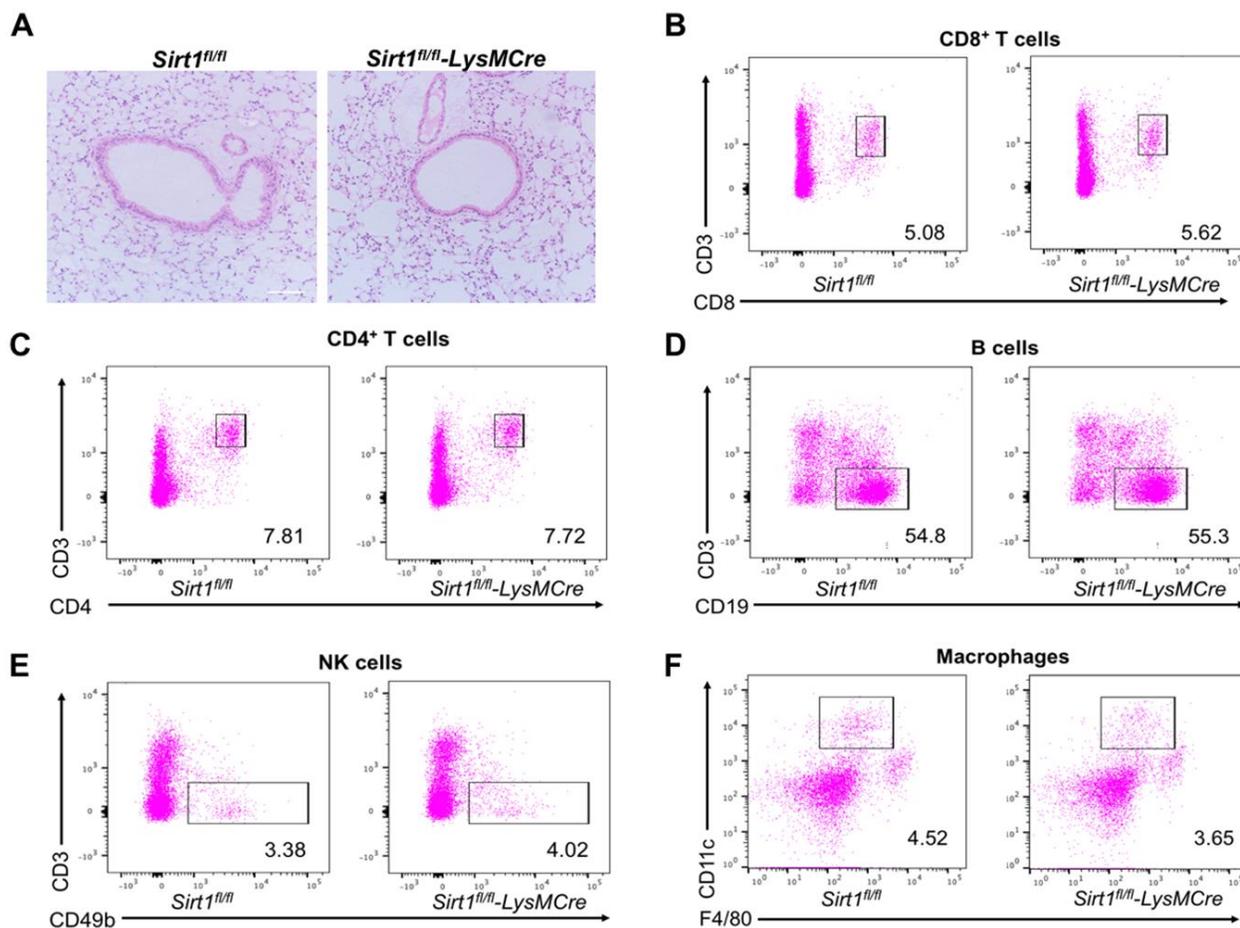
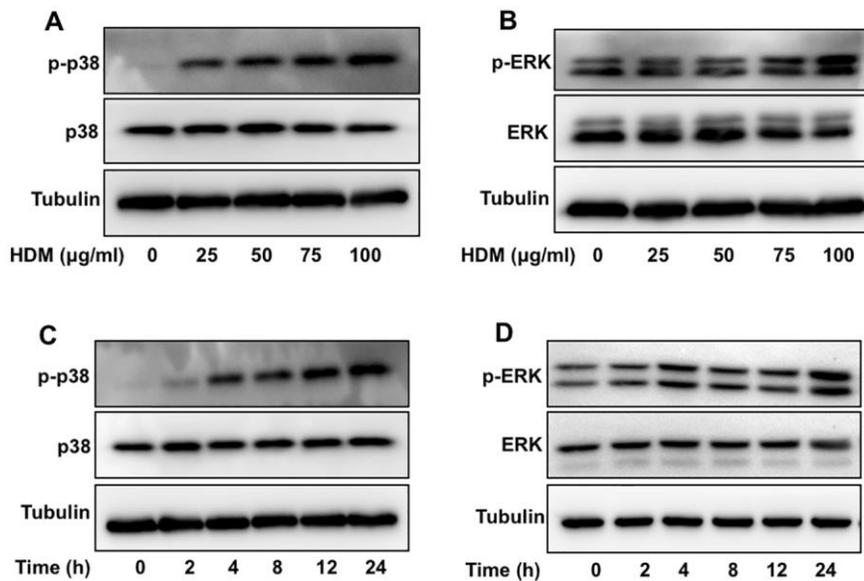


SUPPLEMENTARY FIGURES



Supplementary Figure 1. *Sirt1* deficiency has no influence on cell development and differentiation *in vivo*. (A) The lung sections were histologic analyzed using HE staining to visualize inflammatory cell recruitment (Scale bar, 100 μ m). (B–F) The CD45⁺CD3⁺ cells in the splenocytes of *Sirt1^{fl/fl}* and *Sirt1^{fl/fl}-LysMCre* mice were gated for further analysis of CD8⁺ T cells, CD4⁺ T cells, B cells, and NK cells. The CD45⁺CD11c⁺F4/80⁺ cells in the splenocytes of *Sirt1^{fl/fl}* and *Sirt1^{fl/fl}-LysMCre* mice were gated for macrophages. Representative flow cytometer of CD8⁺ T cells, CD4⁺ T cells, B cells, and NK cells, and macrophages expression are shown.



Supplementary Figure 2. HDM-induced inflammation response in BMDMs via MAPK pathways. BMDMs were treated with HDM, and the protein levels of MAPK pathways were measured by Western blot. (A, B) HDM activated p38 and ERK phosphorylation in a dose-dependent manner (0, 25, 50, 75 and 100 µg/ml for 24 h). (C, D) Stimulation of BMDMs with 100 µg/ml of HDM induced phosphorylation of p38 and ERK in a time-dependent manner (0, 2, 4, 8, 12, 24 h).