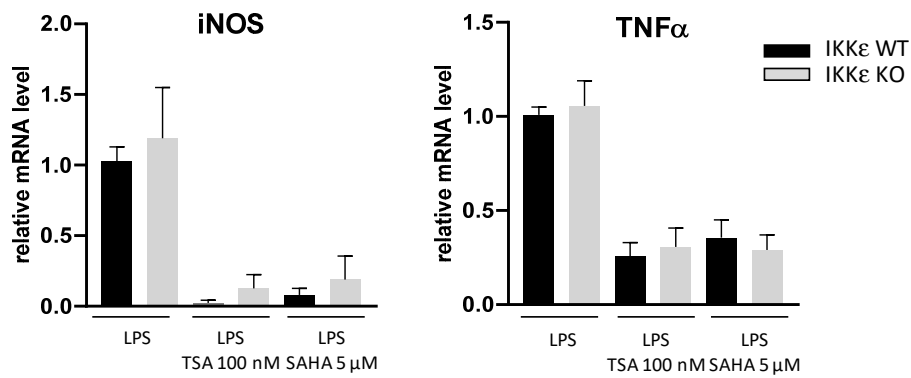
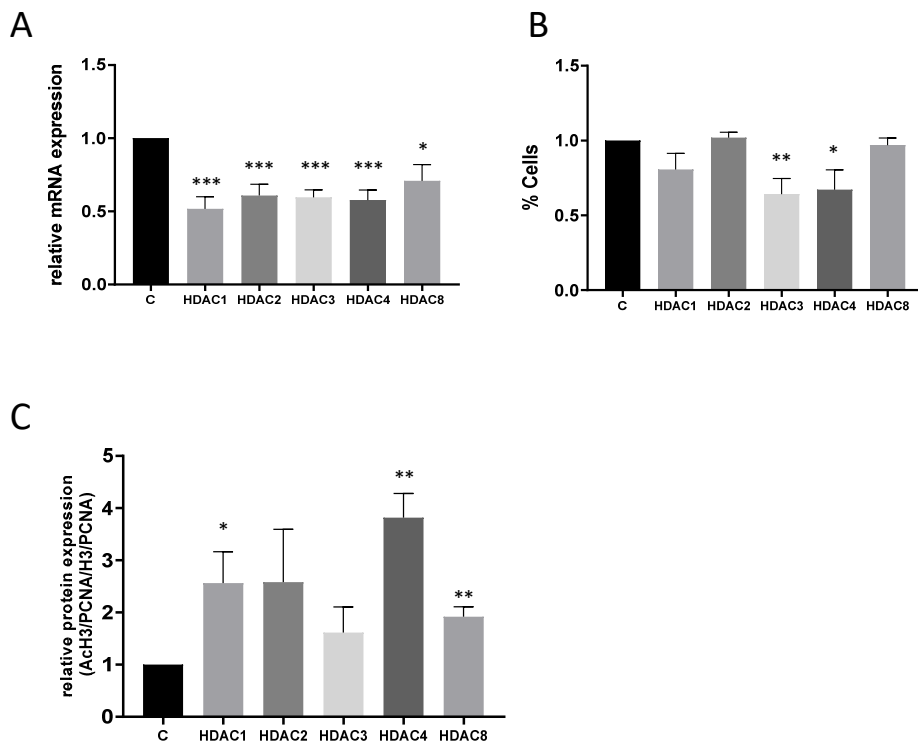


Suppl. Figure 1: Regulation of iNOS and TNF $\alpha$  mRNA in IKK $\epsilon$  wildtype and knock-out bone marrow derived macrophages



Suppl. Figure 2: Evaluation of HDAC siRNA



## Figure Legend

**Suppl. Fig. 1:** Regulation of *iNOS* and *TNF $\alpha$*  mRNA in *IKK $\epsilon$*  wildtype and knock-out bone marrow-derived macrophages

qRT-PCR analysis of *iNOS* and *TNF $\alpha$*  mRNA expression in bone marrow-derived macrophages from *IKK $\epsilon$*  wildtype and knock-out mice (n=3). For better comparison, expression of all genes was normalized to LPS-treated control cells which have been set as “1”.

**Suppl. Fig. 2:** Evaluation of HDAC siRNA

(A) qRT-PCR showing the regulation of different HDACs mRNAs after transfecting cells with the respective HDAC siRNA. Vehicle-treated cells served as control (n=9). (B) WST cell proliferation assay 48 h after cells transfection with the respective siRNAs (n=5). The proliferation rate of vehicle-treated cells was set as “1”. (C) Western Blot analysis of nuclear histone H3 acetylation 48 h after transfection of cells with HDAC siRNA. The diagram shows the densitometric analysis of all Blots (n=3). For better comparison, expression of all genes was normalized to vehicle-treated control cells which have been set as “1”. \*p<0.05, \*\*p<0.01, \*\*\*p<0.001, statistically significant difference as compared to vehicle-treated control.