

FIGURE S1. Analysis of TNF-α and IL-12 mRNA stability in Pam₃CSK₄-stimulated macrophages. WT and MK2^{-/-} macrophages in 24-well microtiter plates (\sim 1 X 10⁶ cells/well) were stimulated with 10 ng/ml of Pam₃CSK₄. After 4 h, actinomycin D (10 μg/ml) was added to stop transcription, at the indicated time points cells were harvested, and total RNA isolated. The levels of TNF-α (*panel A*) and IL-12 (*panel B*) mRNA were quantified by real time RT-PCR. The mRNA levels were normalized with respect to endogenous β-actin mRNA. The experiments were performed two times each time in triplicates. Mean values \pm SEM are plotted.