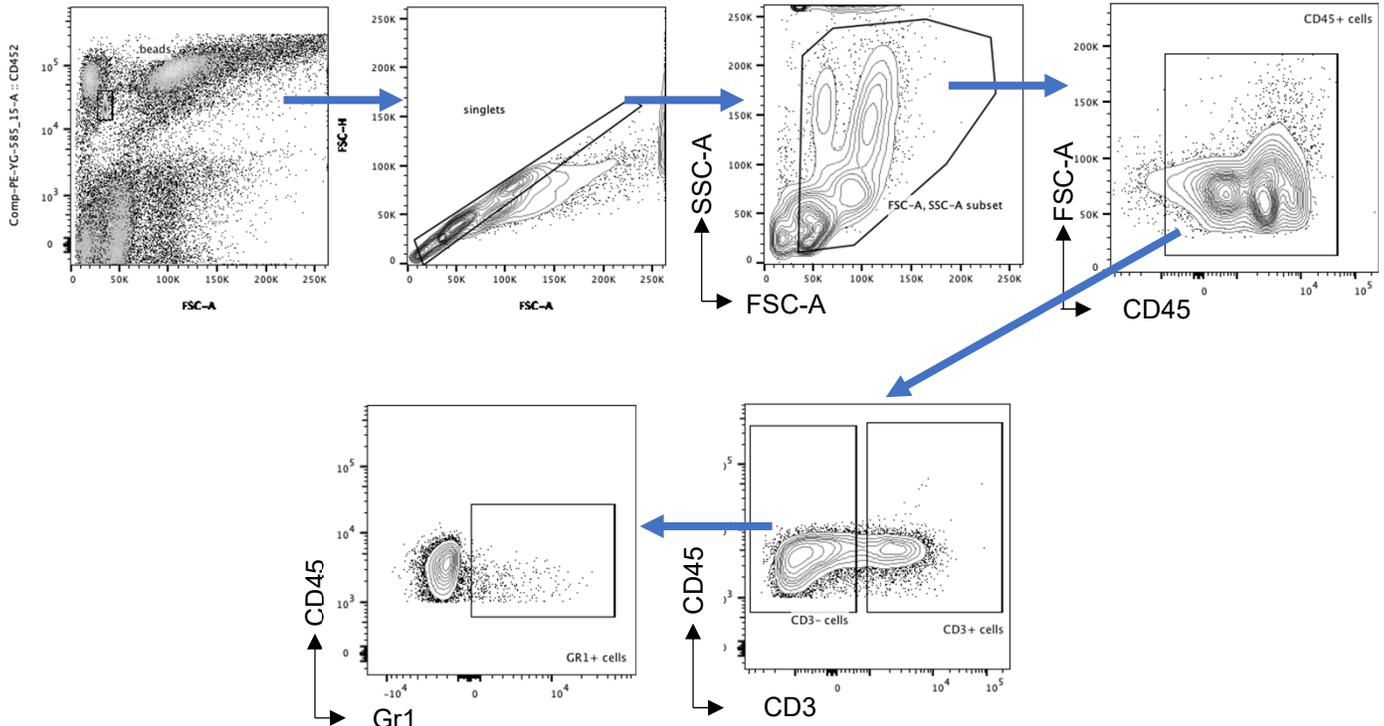
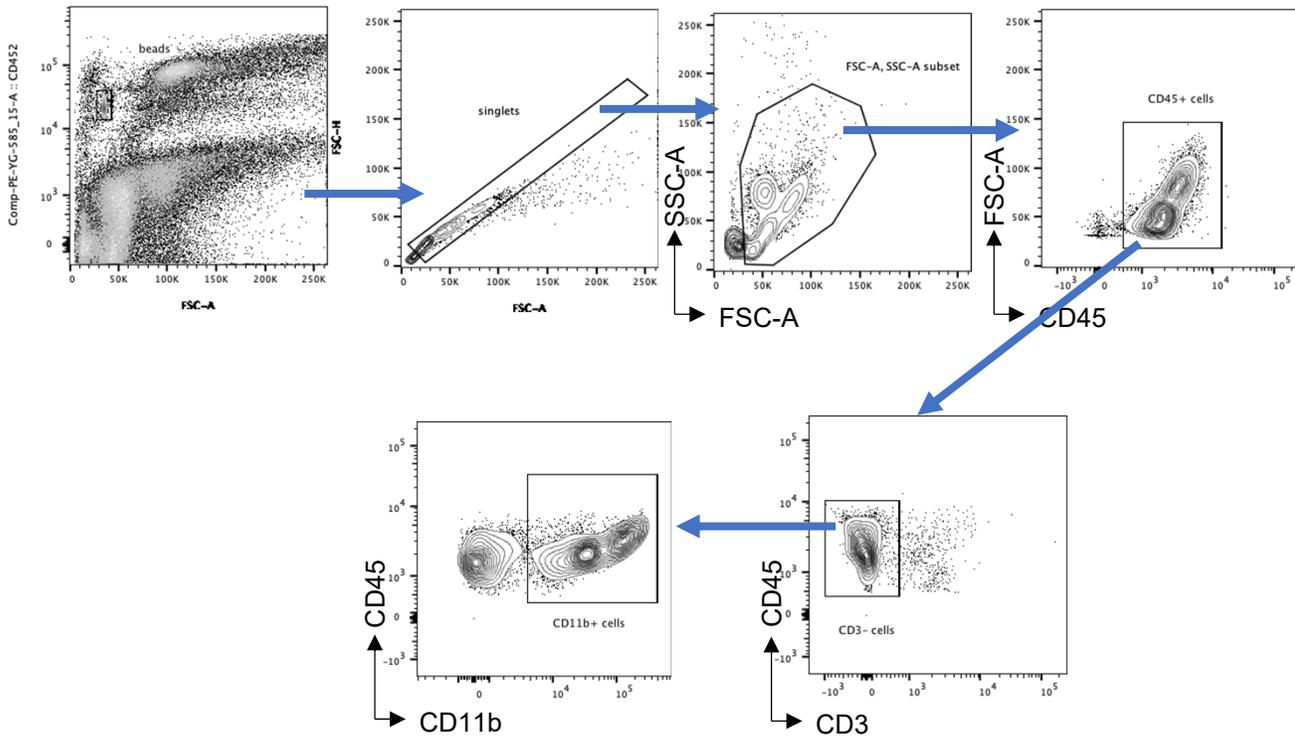


## Supplemental Figure 1

**A**



**B**

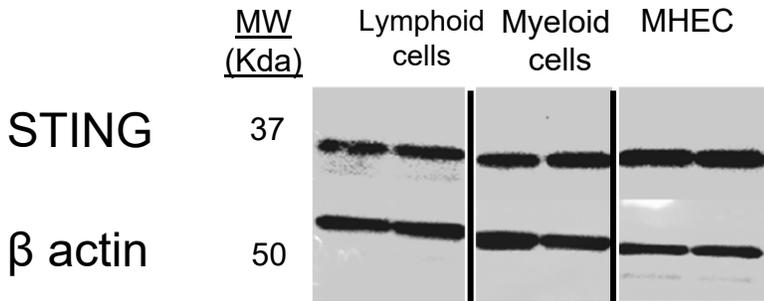


## Supplemental Figure 1

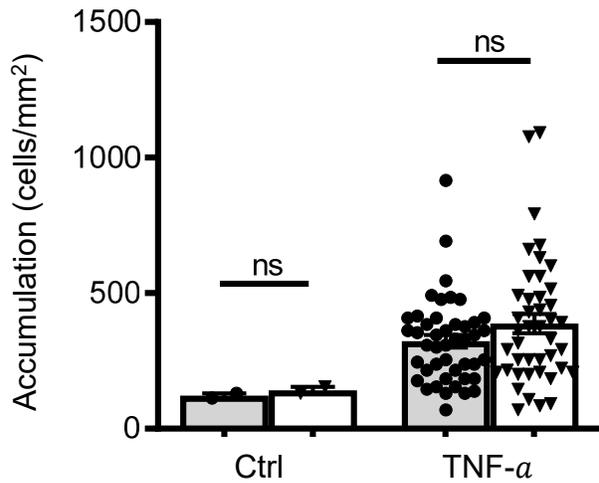
Representative gating strategy for the *in vivo* experiments of peritonitis. Plots represent the peritoneal lavage of WT mice treated with TNF- $\alpha$  24h (A) and with Thioglycollate 3 days (B).

## Supplemental Figure 2

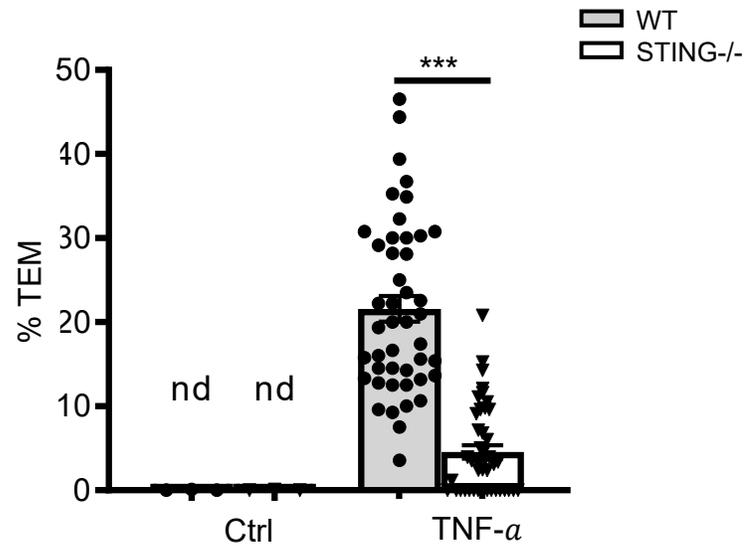
**A**



**B**



**C**

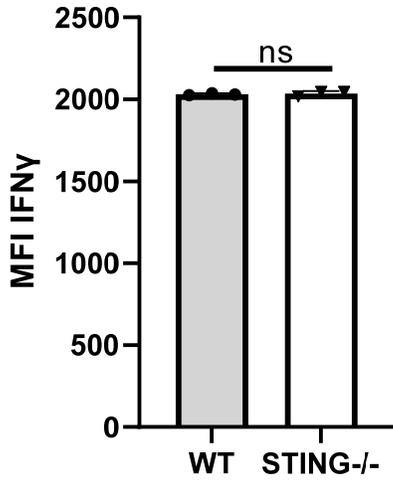


## Supplemental Figure 2

Cultured MHEC, myeloid cells and column-isolated  $\text{CD4}^+$  T cells from WT mice were lysed and analyzed by immunoblot to evaluate STING expression and  $\beta$ -actin as a loading control. (A) One representative blot is shown of 3 independent experiments; each lane is an independent cell preparation. (B-C) Quantification of accumulation and %TEM respectively of WT Th1 cells perfused across unstimulated and  $\text{TNF-}\alpha$ -stimulated WT and  $\text{STING}^{-/-}$  MHEC.  $n=6$  independent experiments ( $\text{TNF-}\alpha$ -), and  $n=2$  (non stimulated Ctrl), in duplicate or triplicate coverslips. 1-way ANOVA (B-C).

### Supplemental Figure 3

A

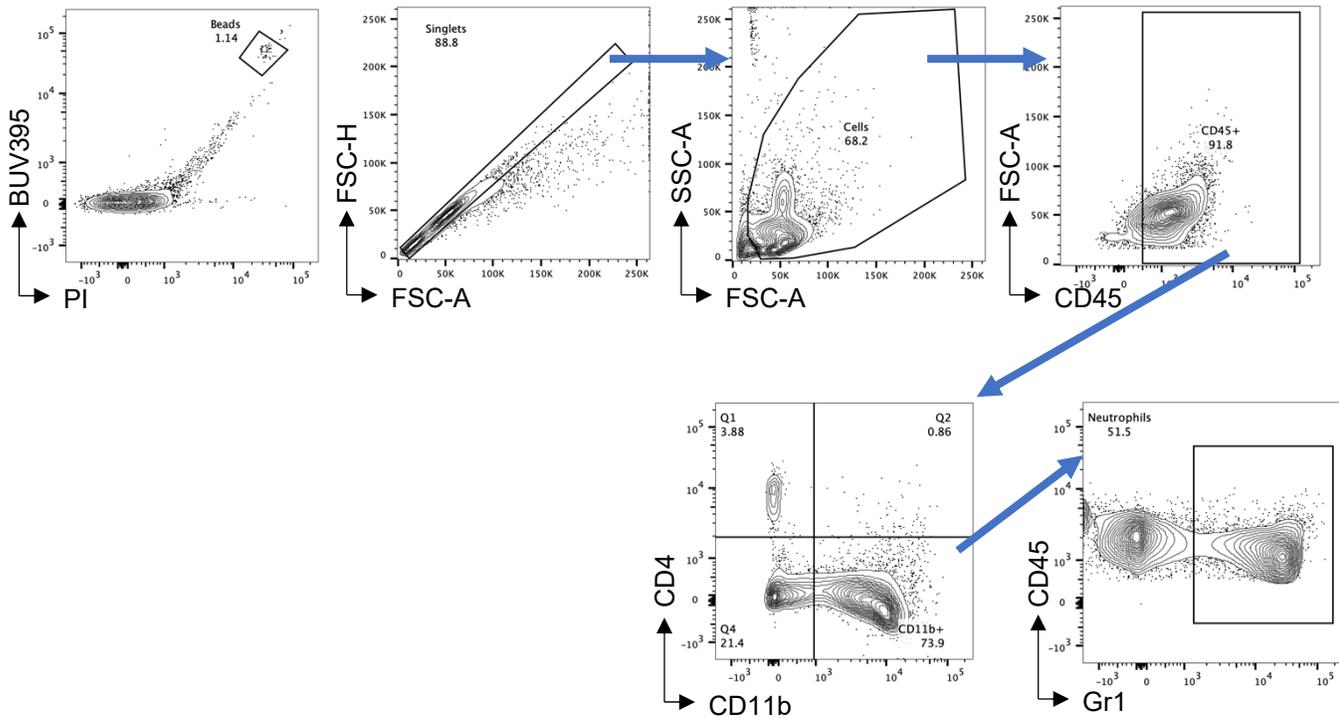


### Supplemental Figure 3

(A) WT and STING $^{-/-}$  Th1 cells were differentiated *in vitro* as described in methods, permeabilized and stained for intracellular IFN- $\gamma$ . Quantification of the median fluorescent intensity of IFN- $\gamma$  per cell is represented. n=3 independent T cell preparations/group. t-test.

## Supplemental Figure 4

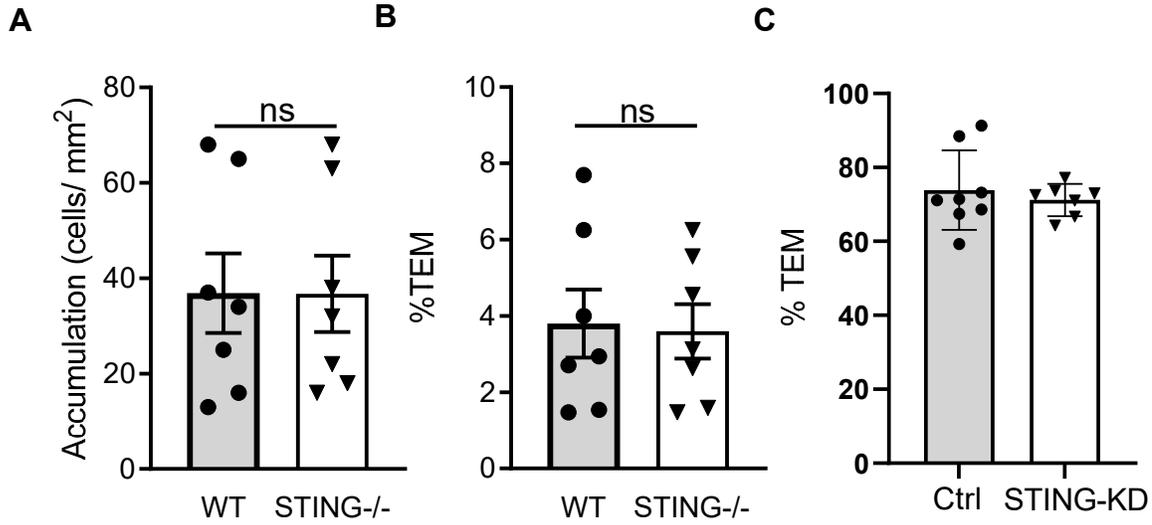
A



## Supplemental Figure 4

(A) Representative gating strategy for the *in vivo* experiments of 4h TNF- $\alpha$  induced peritonitis. Plots represent the peritoneal lavage of  $\text{Cad5}^{\text{ERTCre2+/-STING}^{\text{fl/fl}}}$  treated with tamoxifen or vehicle control 4 hours after TNF- $\alpha$  injection. The top left panel gate represents the counting beads.

## Supplemental Figure 5



**Supplemental Figure 5. STING deficiency does not result in decreased neutrophil TEM across mouse and human primary endothelial cells.** (A and B) Mouse bone marrow derived neutrophils were perfused across cultured MHEC from WT and STING<sup>-/-</sup> mice treated 4h with TNF- $\alpha$ , and quantification of accumulation (A) and %TEM (B) was determined in n= 3 independent experiments, in duplicate coverslips. (C) Neutrophils isolated from human blood were perfused under flow conditions across WT and STING KD HUVEC stimulated for 4h with TNF- $\alpha$  and %TEM was quantified. n=8 (Ctrl) and n=7 (STING-KD) coverslips from n=2 independent experiments.