

Supplementary Figure 2: Gating strategy applied for the flow cytometry analysis of mICcl2 cells. In order to quantify  $I\kappa B\zeta$  protein levels in mICcl2 cells, cell debris (FSC-A/SSC-A), cell doublets (FSC-A/FSC-H) and dead cells (fixable viability dye <sup>+</sup>) were excluded from further analysis. Mean fluorescence of  $I\kappa B\zeta$  was determined in single, viable cells. Depicted here are representative histograms for  $I\kappa B\zeta$  MFI in mICcl2 cells stimulated with PBS (mock), *B. vulgatus* (BV) and *E. coli* (EC).