



S3 Fig. Impaired immune response promoted by pDC/Treg interactions is dependent on IDO activity. pDCs were isolated from lung infiltrating leukocytes of uninfected IDO^{-/-} and WT mice using magnetic beads anti-mPDCA, matured with *P. brasiliensis* yeasts (1:10; Pb:pDC ratio) and then co-cultured for 7 days with splenic CD3⁺lymphocytes (1:10; pDC:lymphocytes ratio) isolated by anti-CD3 magnetic beads from WT mice. (A) Frequency of CD4⁺Foxp3⁺ T cells analyzed by flow cytometry after 7 days of co-cultivation. (B) Splenic lymphocytes from uninfected WT mice were previously labeled with CFSE (5 mM) and co-cultured with *P. brasiliensis* -infected pDCs. After 7 days, the cells were adjusted to 1 × 10⁶, labeled with specific anti-CD4 and CD8 antibodies and analyzed by flow cytometry. (C) After 7 days of co-culture with infected pDCs, lymphocytes were adjusted to 1 × 10⁶, labeled with specific anti-CD4, CD8, CD25, and CD69 antibodies and analyzed by flow cytometry. The lymphocytes were gated by FSC/SSC analysis and gated cells were analyzed for the expression of CD4⁺CD25⁺ (top)

CD8⁺CD69⁺ (bottom). Bars reflect mean \pm SD of two independent experiments with eight mice per group (right) (* $p < 0.05$).