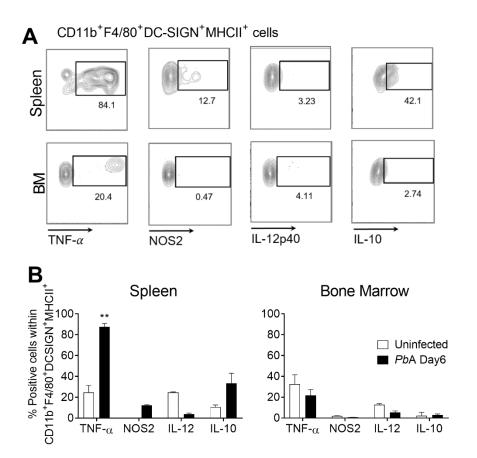
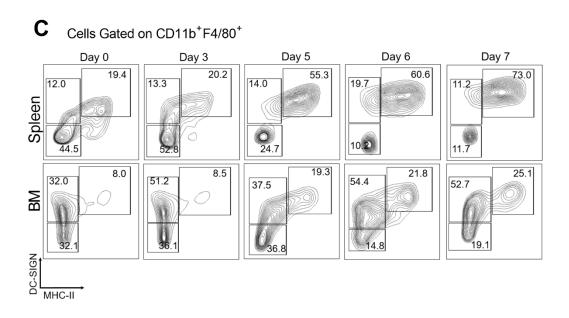
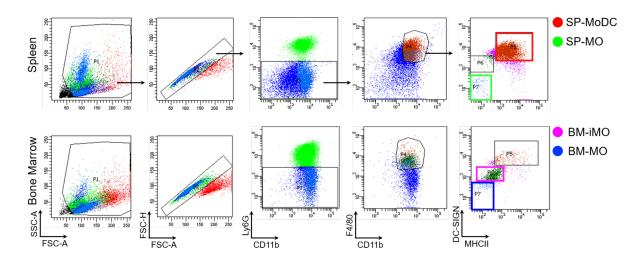
Supplementary Figure 1





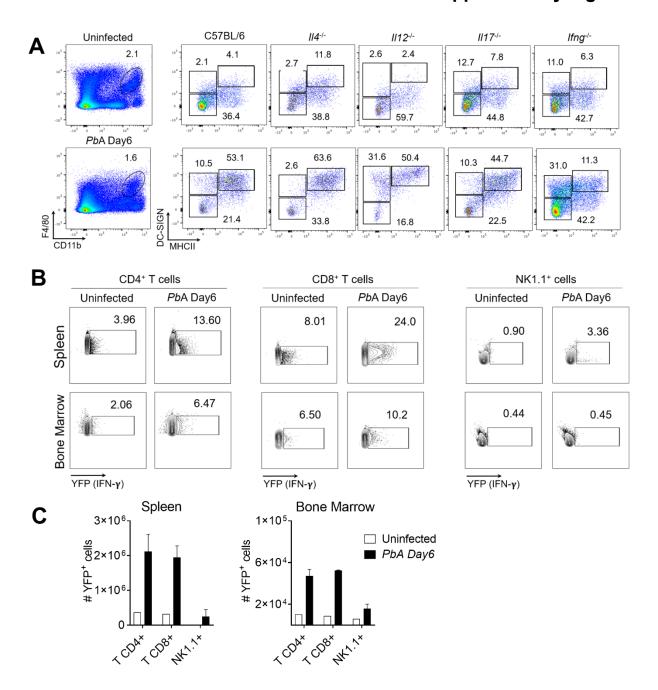
Supplementary Figure 1. Cytokine production, NOS2 expression and kinetics of MoDC differentiation during *PbA* infection. (A) Cytokine production by CD11b+F4/80+DC-SIGNhiMHCIIhi cells from spleen and BM. Total splenocytes and BM cells where cultured for 24h with 100 ng/mL of LPS and stained for flow cytometry quantification of the intracellular cytokines TNF α , IL-12p40 and IL-10 and iNOS (NOS2). (B) Frequency of CD11b+F4/80+DC-SIGN+MHCII+ cells positive for the cytokines TNF α , IL-12p40, IL-10 and NOS2. The data shown are representative of two independent experiments. Results are expressed as average ± s.e.m. **P<0.001. (C) Splenic and BM cells where collected from mice on days 0, 3, 5, 6 and 7 post-infection with PbA. Contour plot show the frequency of MODCs, iMOs and MO within total CD11b+F4/80+ cells. Data shown is representative of two independent experiments.

Supplementary Figure 2



Supplementary Figure 2. Sorting of splenic and bone marrow populations. Splenocytes and BM cells were pulled from 5 C57BL/6 mice and isolation of specific performed by cell sorting: SP-MO and populations was BM-MO (Ly6G⁻CD11b⁺F4/80⁺DC-SIGN⁻MHCII⁻) were isolated from uninfected mice; BM-iMO (Ly6G⁻CD11b⁺F4/80⁺DC-SIGN^{|0}MHCII⁻) and SP-MoDC (Ly6G-CD11b+F4/80+DC-SIGN^{hi}MHCII^{hi}) were isolated from mice 6 days post-*Pb*A infection. Three samples from each cell population were obtained and cells were later used for RNA or miRNA extraction, mRNA library preparation and sequencing.

Supplementary Figure 3



Supplementary Figure 3. Role of IFN γ on the differentiation of MoDCs during *PbA* infection. (A) Spleens of uninfected and 6 days *PbA*-infected mice were collected from C57BL/6, IL-4^{-/-}, IL-12^{-/-}, IL-17^{-/-} and IFN γ ^{-/-}. Dot plot shows the frequency of DC-SIGN+MHCII+ cells within total monocytes CD11b+F4/80+ in uninfected and infected mice.

(B) Splenocytes and BM cells from uninfected and infected GREAT mice uninfected or infected cultured for 4 h with PMA (50 ng/mL) and ionomycin (500 ng/mL) in the presence of brefeldin A and analyzed by flow cytometry. Cells were gated on CD4⁺, CD8⁺ or NK1.1⁺. YFP-positive cells were considered as IFN γ (C) Bar graphs indicate the absolute number of YFP-positive cells within each population.