



Wild type (WT) pr CD11a-/- (KO) mice were infected with either 1×10^3 or 5×10^4 CFU of LM-Ova. Three days later spleens were removed and bacterial burdens were measured. Data is derived from five mice per group and statistical significance was determined using the Mann-Whitney test. ns=not significantly different from the control; * = p< 0.01 as compared to control.



Figure S2. LM infection dose affects the requirement for CD11a in the CD8 T cell response. Wild type (black bars) or CD11a-/- (white bars) mice were infected with either 1×10^3 or 5×10^4 CFU of LM-Ova. Eight days later spleens were removed and the CD8 and CD4 T cell responses were measured by flow cytometry after staining with H-2K^b-OVA or I-A^b-LLO tetramers, anti-CD8, anti-CD4 and anti-CD44. Data is derived from three mice per group and statistical significance was determined using a two-tailed unpaired t test. ns=not significantly different from the control; * = p< 0.01, ** = p<0.001 as compared to control.



Figure S3. T cell responses are defective after infection with ActA-deficient LM. Wild type (black bars) or CD11a-/- (white bars) mice were infected with 1×10^6 CFU of ActA-deficient LM-Ova. Seven days later spleens were removed and the CD8 and CD4 T cell responses were measured by flow cytometry after staining with H-2K^b-OVA or I-A^b-LLO tetramers, anti-CD8, anti-CD4 and anti-CD44. Data is derived from three mice per group and statistical significance was determined using a two-tailed unpaired t test. ns=not significantly different from the control; * = p< 0.01.