

H-ferritin produced by myeloid cells is released to the circulation and plays a major role in liver iron distribution during infection

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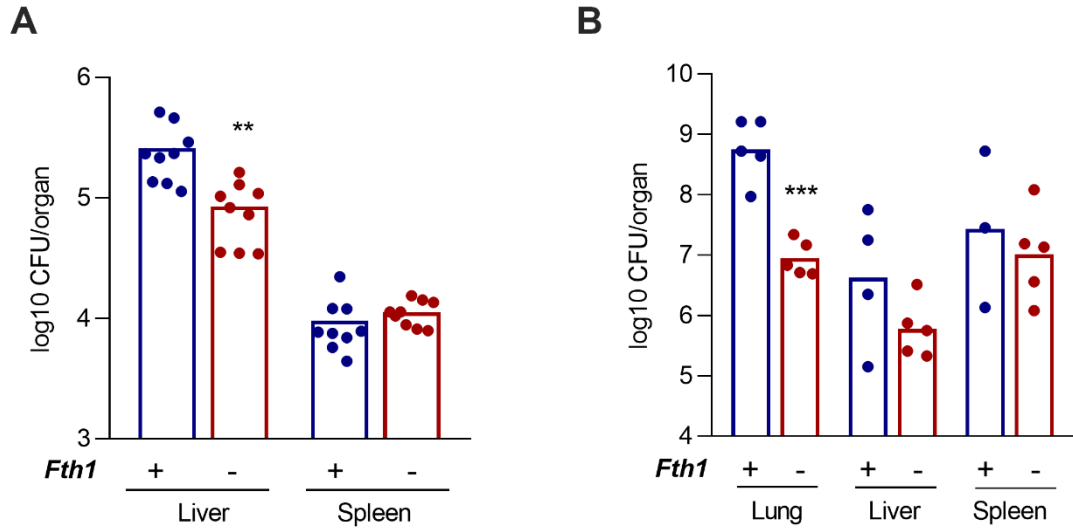


Figure S1. Mice deficient in H-ferritin in the myeloid cells are more resistant in different models of *M. avium* infection. (a) *Fth1*^{+/+} (blue) and *Fth1*^{-/-} (red) mice were intravenously infected with 10⁶ CFU of *M. avium* 2447 SmT. Bacterial burden in the liver and spleen, 60 days post infection (N=9). (b) *Fth1*^{+/+} (blue) and *Fth1*^{-/-} (red) mice were infected via aerosol with a low dose (~100 CFU/lung) of *M. avium* 25291. Bacterial burden in the lung, liver and spleen, 125 days post infection. (N=4 to 5). Bars represent the mean and the circles represent each animal. Statistics: multiple t-test. **P<0.01, ***P<0.001 when comparing between genotypes.

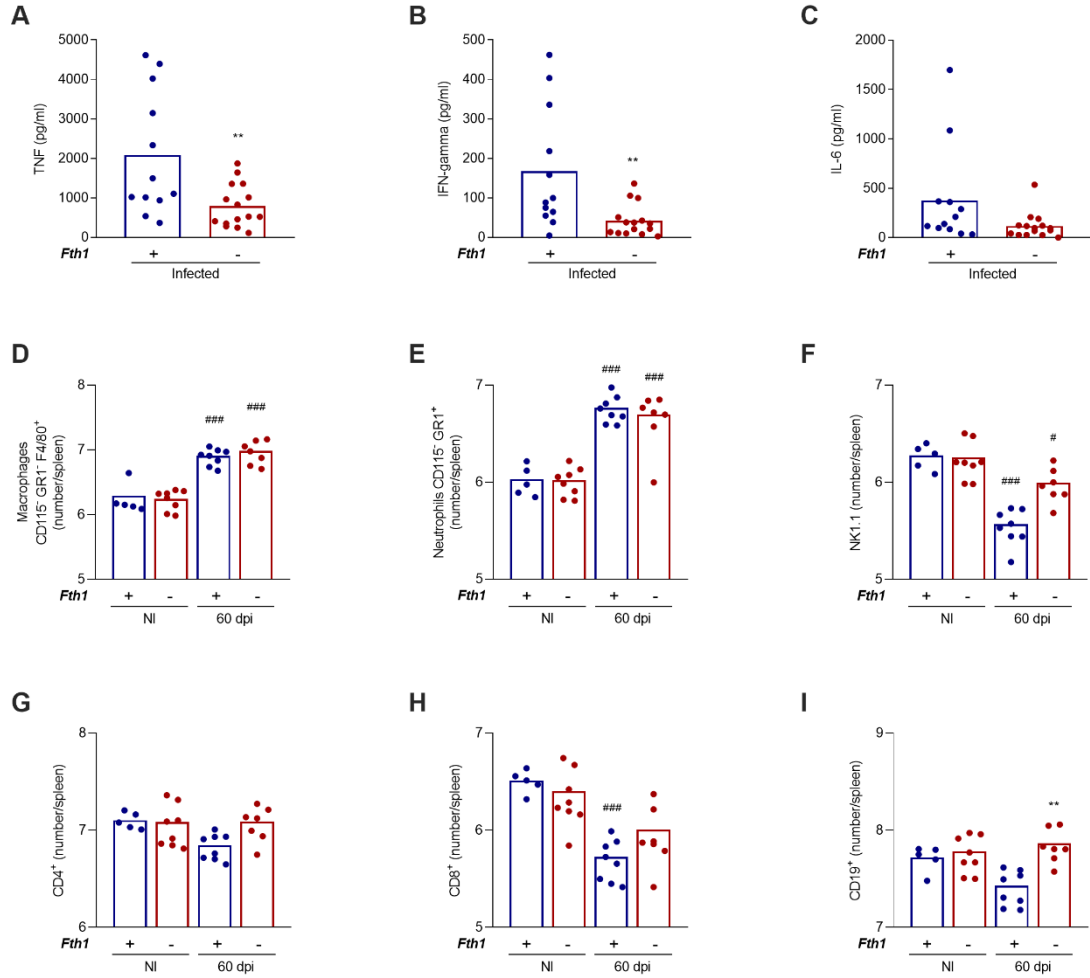


Figure S2. *Fth1*^{-/-} mice have lower levels of inflammatory cytokines in circulation and spleen cell populations are not dependent of FTH1 expression. (a) TNF-alpha, (b) IFN-gamma, (c) IL-6 were measured in the serum of the mice from the survival curve, using a cytometric bead array mouse inflammation kit (N = 12 (*Fth1*^{+/+}) or 16 (*Fth1*^{-/-})). (d-i) Number of macrophages (GR1-F4/80⁺), neutrophils (CD115-GR1⁺), NK1.1, CD4⁺, CD8⁺, CD19⁺ CD19⁺, NK1.1, CF4⁺, CD8⁺ cells in the spleen 60 days post infection (N=5 to 8). Bars represent the mean and the circles represent each animal. Statistical analysis was performed by two-way ANOVA with Sidak multiple comparison post hoc test. **P<0.01 when comparing between genotypes, ###P<0.001 when assessing the effect of infection within the same genotype.

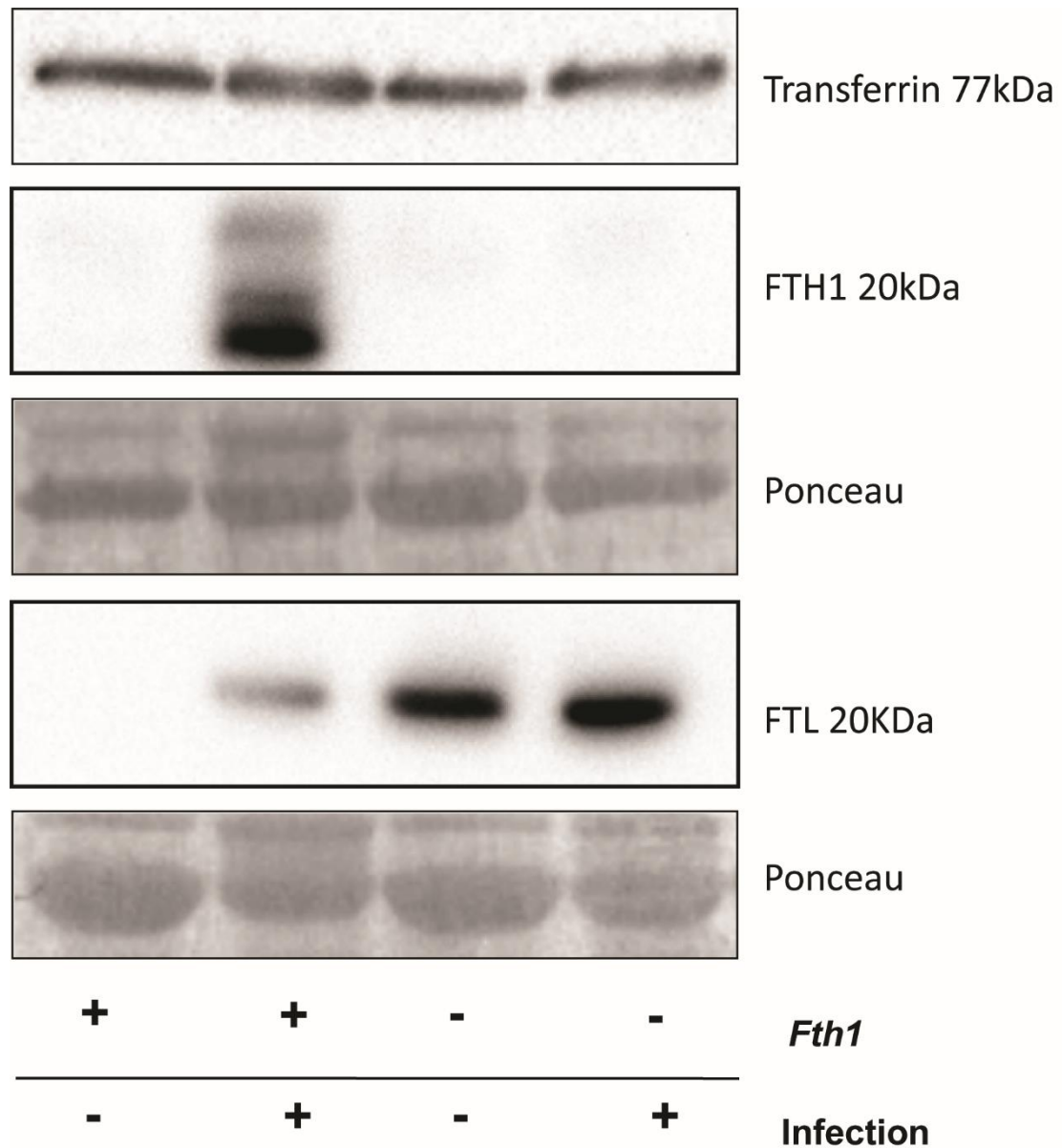


Figure S3. During *M. avium* infection, *Fth1*^{+/+} mice have high amount of FTH1 in the serum. Western Blot against transferrin, FTH1 and FTL subunits. Ponceau staining was used to stain all proteins in the gel, as a confirmation that all lanes were loaded with equivalent amounts of protein.