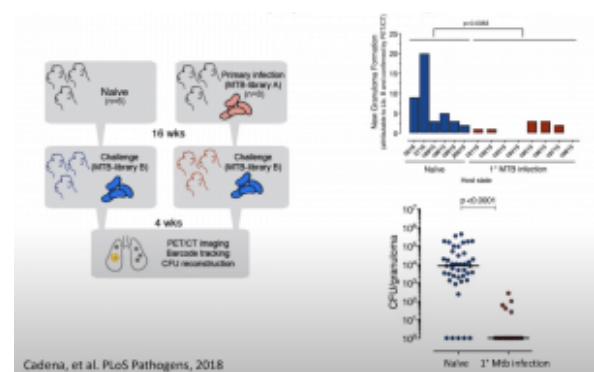
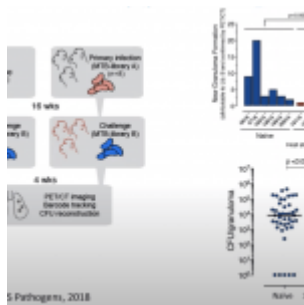


# Flynn Webinar: What immune cells play a role in protection against M.tb re-infection?



Adapted from [Cadena et al., 2018](#)

Approximately a quarter of the global population is estimated to have *Mycobacterium tuberculosis* infection (tuberculosis etiological agent). However, only 5-10% of infected individuals progress to disease. On the 8th of March, the IUIS-Immunopaedia-Frontiers webinar series featured a talk by tuberculosis immunologist Dr. JoAnne Flynn. In honour of World TB Day on the 24th of March, we shall provide three summaries highlighting findings shared during Dr Flynn's talk.

In this summary, we shall highlight research she presented that provided evidence for the role of CD4 T cells in protection against M.tb reinfection. Previous findings by Dr

Flynn and collaborators demonstrated that prior M.tb infection is associated with protection against M.tb reinfection ([Cadena et al., 2018](#)). This study showed that non-human primates (NHPs) that were re-infected had a significantly lower M.tb bacterial burden and a significantly higher proportion of sterile granulomas compared to M.tb infected naïve NHPs. Further, they showed that adaptive Th1 immune responses were not different between M.tb reinfected and naïve (primary M.tb infected) NHPs. As a follow-up to this study, Dr Flynn's research group aimed to determine if CD4 T cells are necessary for protection against reinfection. They focused on CD4 T cells because they are *important for the control of TB, and depletion results in worse acute infection and latent TB reactivation in NHPs.*

*To test this hypothesis, CD4 T cells were depleted in infected and treated (4-5 months of full TB therapy) NHPs before a second Mtb challenge. As controls, one group received IgG and a naïve group received only Lib B. The number of Lib B granulomas in the IgG group was significantly fewer than in the naïve group, while the  $\alpha$ CD4 group was in between the two. The IgG group had the lowest lung bacterial burden followed by the  $\alpha$ CD4 group (16-fold higher than IgG) and lastly the naïve group (57.5-fold higher than IgG). Both reinfected groups ( $\alpha$ CD4 and IgG) had a higher frequency of CD8 T cells producing IFN $\gamma$  and/or TNF in Lib B granulomas compared to naïve NHPs but there was no significant difference when compared to each other. (Source: [Ganchua et al., 2020](#)).*

Single-cell RNAseq analysis of tuberculosis granulomas from reinfected IgG treated NHPs had higher proportions of regulatory T cells and CD8 T cells that expressed Granzyme H and K than naïve infected NHPs. Further, depletion of CD4 T cells in reinfected NHPs negatively impacted the CD8 T cell compartment: lower proportions granzyme H and K expressing CD8 T cells. Based on these finding Dr Flynn proposes that the depletion of CD4 T cells, specifically IL-21+ CD4 T cells,

leads to the impaired induction and maintenance of cytotoxic CD8 T cell activity.

In summary, results presented in the webinar suggest that CD4 T cells play an important role in protection against re-infection. This protective effect is not restricted to CD4 T cells function but is associated with the induction and maintenance of CD8 T cells that produce granzyme H and K.

*Summary by Cheleka AM Mpande*