# Project update

### J. Shah chimeric mouse collaboration

## Kim Dill-McFarland, kadm@uw.edu

version May 11, 2020

# Completed

- 1. Change gene-level model to contrasts of WT:TKO within uninfected or infected samples
- 2. Compare above contrasts model to previous interaction term model (~status\*cell)
  - Roughly half of genes identified as significant in either model are also significant in the other model.
  - Genes unique to the interaction model are only significant for infection status. Genes unique to the contrasts model were only significant for cell type within infected cells.
  - Thus, the contrasts model appears to be successfully removing much of the infection-only signal while amplifying cell type signal.
- 3. Modules made from 4950 genes significant (FDR < 0.3) in contrasts model
  - Resulted in 18 modules + 399 genes in module 0
- 4. Contrasts model of modules
  - All modules significant at FDR < 0.05
  - 3 modules significant in uninfected, 5 in infected (ignore module 0), and 10 for both
- 5. GSEA of significant genes and genes in each module

### To discuss

- Other methods for assigning function to modules
- Additional analyses?

#### To-do

- GSEA of fold change module groups (e.g. up in uninfected, up in infected, up in both, etc)
- Heatmap of module expression
- Modify boxplots for publication
- Point group to count files