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**Thesis Defense - Katherine Besse, MS Biology**

**July 22, 2021, 3:00 - 5:00 PM**

**VIEW EVENT**

**All are invited to attend the defense. For more information please contact Graduate Coordinator at [kharrism@gmu.edu](mailto:kharrism@gmu.edu).**

**Candidate:** Katherine Besse

**Program:** Biology, M.S.

**Date:** July 22, 2021

**Time:** 3:00 PM

**Place:** [frnq8--eks.xmmk.sq-h-75477114714=nub;L.Pt\\_CvKR0PURCTwOjK2cEH3SwrXXx.7](https://www.gmu.edu/registration/view-event.cfm?eventid=114714)

**Title:** IDENTIFICATION AND ANALYSIS OF THE ROLE OF CHITINASE SUBSTRATE IN FRANCISELLA NOVICIDA

**Committee Chair:** Dr. Monique van Hoek

**Committee Members:** Dr. Ancha Baranova, Dr. Brett Froelich

**ABSTRACT:**

*Francisella tularensis* is a gram negative facultative intracellular pathogen which is a class A biothreat according to the CDC. This bacteria codes for two chitinases (ChiA and ChiB) and one chitin binding protein (CbpA). Chitin is the most abundant oligosaccharide in marine environments and in the exoskeleton of many insects. In order to digest this material, microorganisms produce chitinases capable of cleaving this polymer. The chitinases in *Francisella novicida* have previously been found by our lab to negatively regulate the bacterial biofilm, most likely cleaving the extracellular polysaccharide substance (EPS) with  $\beta$ -1,4 glycosidic linkages. These enzymes may enable the organism to use the resulting cleaved polymers as carbon and nitrogen sources for growth. Although *Francisella* encodes for several polysaccharide synthases, it has not been found to produce chitin. Therefore, we hypothesized that the chitinase-enzyme substrate that is self-produced in the *F. novicida* EPS and biofilm must be some other molecule also containing  $\beta$ -1,4 glycosidic linkages. The biochemical characterization of ChiB activity on various substrates will be evaluated and the role of the substrate in the biofilm will be examined.

