Francisella tularensis is a highly pathogenic bacterium that can be used as a biological weapon. This bacterium causes tularemia which is also known as Rabbit Fever. Previously, our laboratory identified three F. tularensis loci predicted to encode transcriptional regulators (FTL_0671, FTL_1199, and FTL_1665) that exhibited transcriptional changes in the presence of human erythrocytes. We hypothesized that because transcriptional modulations were associated with the presence of host cues, these potential transcriptional regulators may be important for pathogenesis. Therefore, mutants of these three loci and the isogenic wild-type bacterium (strain LVS) were used to infect chicken embryos. Healthy chicken embryos were injected with $10^4$, $10^5$, or $10^6$ colony forming units of each strain, and chicken embryos were cycled daily for viability. The chicken embryos were housed in an egg incubator (~37°C with occasional rocking) and were observed over a span of 10 days. Neither FTL_0671 nor FTL_1199 affected the pathogenesis of F. tularensis. However, chicken embryos infected with the FTL_1665 deletion strain exhibited mortality more rapidly compared to those infected with wild-type bacteria suggesting that this locus may be repressing pathogenesis. Future studies will evaluate the function of FTL_1665 to determine if the protein encoded by this locus truly mediates transcriptional regulation.