

Undergraduate Research Scholars LAUNCH: UNDERGRADUATE RESEARCH

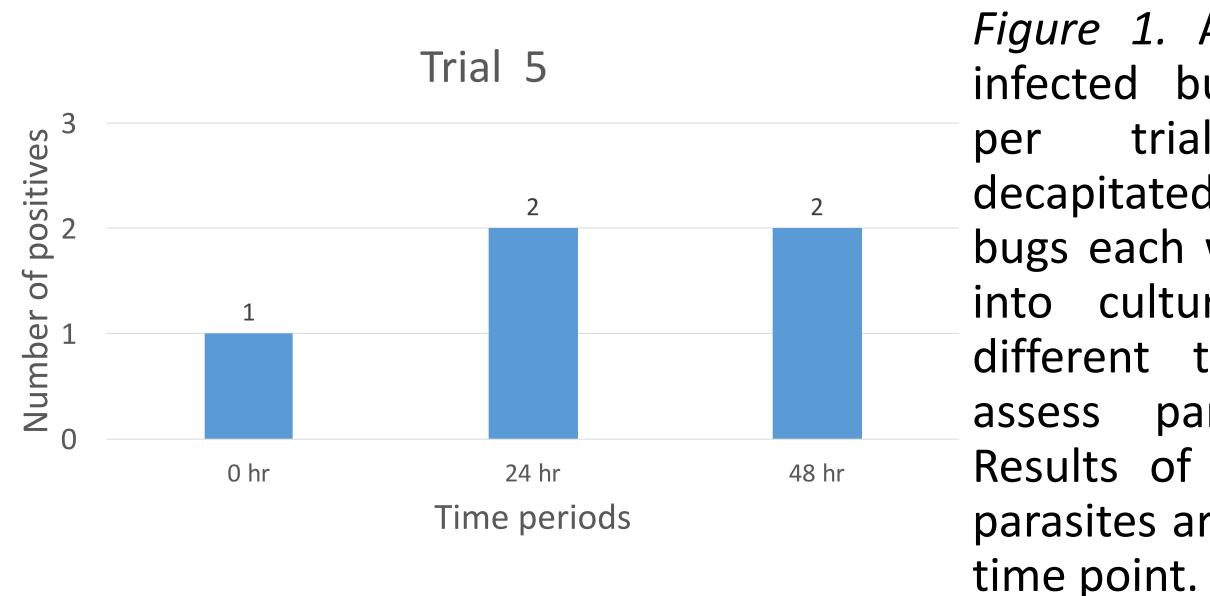
Background

- *Trypanosoma cruzi* is a protozoan parasite and the causative agent of Chagas Disease, transmitted by triatomine bugs, commonly known as kissing bugs.^{1,2,3,4}
- Affected humans and animals symptoms range from asymptomatic to severe cardiac, neurological issues, and death.
- Insecticide use can help control the vector of Chagas Disease
- However, this leaves dead bugs in the environment which poses an unknown risk of parasite transmission to dogs or other animals that may contact or consume them

Research Question

The objective of this experiment was to determine how long T. *cruzi* can survive inside of infected, dead *Triatoma gerstaeckeri* in order to determine the point at which the parasite is no longer pathogenic. The reproducibility of parasites was examined after bugs were left out for three different periods of time (0 hrs, 24 hrs, and 48 hrs).

Results



Duration of survival of the Chagas disease parasite (Trypanosoma *cruzi*) in deceased triatomine 'kissing bug' vectors



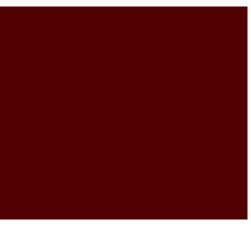




Figure 1. A total of nine infected bugs were used with all trial, decapitated at TO. Three bugs each were inoculated into culture after three different time points to assess parasite viability. Results of the trial show parasites are viable at each

Methods



In five trials, Triatoma gerstaeckeri nymphs were experimentally infected using blood spiked with the protozoan parasite, T. cruzi.



At T=0, all nine bugs were decapitated. Group one bugs were used immediately to inoculate cultures with fecal sample through abdominal compression and a portion of gut material.



The remainder of bugs in groups two and three were left out for twenty-four or forty-eight hours before being inoculated into culture.



Cultures were inoculated with fecal sample through abdominal compression and a portion of gut material from dissection.



Cultures were inspected weekly for up to 6 weeks.

Kaitlyn Perez, Keswick Killets, Sarah A. Hamer

Progress

Initially, there was difficulty growing *T. cruzi* in culture due to the competition with the gut bacteria. However, adjustments in the antibiotic and antifungal dosages in the culture media have allowed for *T. cruzi* to be grown in culture.

Conclusions

- forty-eight hours.

Acknowledgements

Thank you to the Hamer Lab including Lisa Auckland and Rachel Busselman. Also, thank you to my family and friends who have supported me on this wonderful journey.



[1] Bern C, Kjos S, Yabsley MJ, Montgomery SP. *Trypanosoma cruzi* and Chagas' Disease in the United States. Clinical Microbiology Reviews. 2011; 24(4): 655–681. [2] Montgomery SP, Starr MC, Cantey PT, Edwards MS, Meymandi SK. Neglected parasitic infections in the United States: Chagas disease. The American Journal of Tropical Medicine and Hygiene. 2014;90(5):814–818 [3] Chin-Hong PV, Schwartz BS, Bern C, et al. Screening and Treatment of Chagas Disease in Organ Transplant Recipients in the United States: Recommendations from the Chagas in Transplant Working Group. The American Journal of Transplantation. 2011;11(4):672-680 [4] Department of Pediatrics. Addressing the Challenges of Chagas Disease: An Emerging: Infectious Diseases in Clinical Practice. Lippincott Williams & Wilkins. 2017;25(3):118-125 [5] Lidani KCF, Andrade FA, Bavia L, et al. Chagas Disease: From Discovery to a Worldwide Health Problem. Frontiers in Public Health. 2019;7: 166.

Data suggests that the death of *T. cruzi* inside the gut material of a deceased bug occurs at some time point after

• Dead bugs should be removed from the environment whenever possible to reduce the chance of transmission.

• Further trials will explore time points beyond forty-eight hours to determine when *T. cruzi* is no longer viable.