FOUNTAIN OF YOUTH

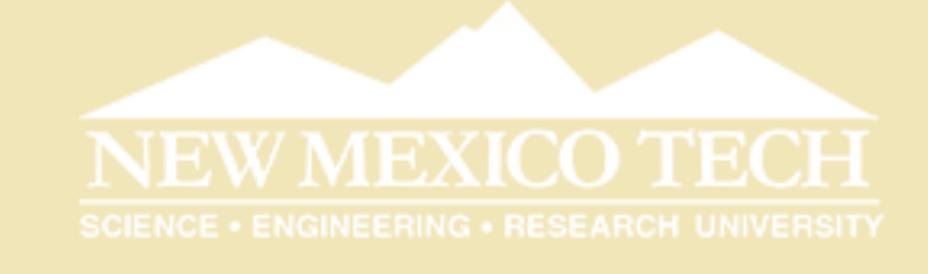
Exploring the Effects of a Telomerase Activator in the Alfalfa Leaf Cutting Bee

Gwen Sailer¹, Ella Eleven², Josh Rinehart¹, Jeff Kittilson¹, Joe Rinehart¹, Julia Bowsher¹

¹Department of Biological Sciences, North Dakota State University, Fargo, ND

²Department of Biological Sciences, New Mexico Institute of Mining and Technology, Socorro, NM

³Insect Genetics and Biochemistry Research Unit, USDA-ARS, Fargo, ND



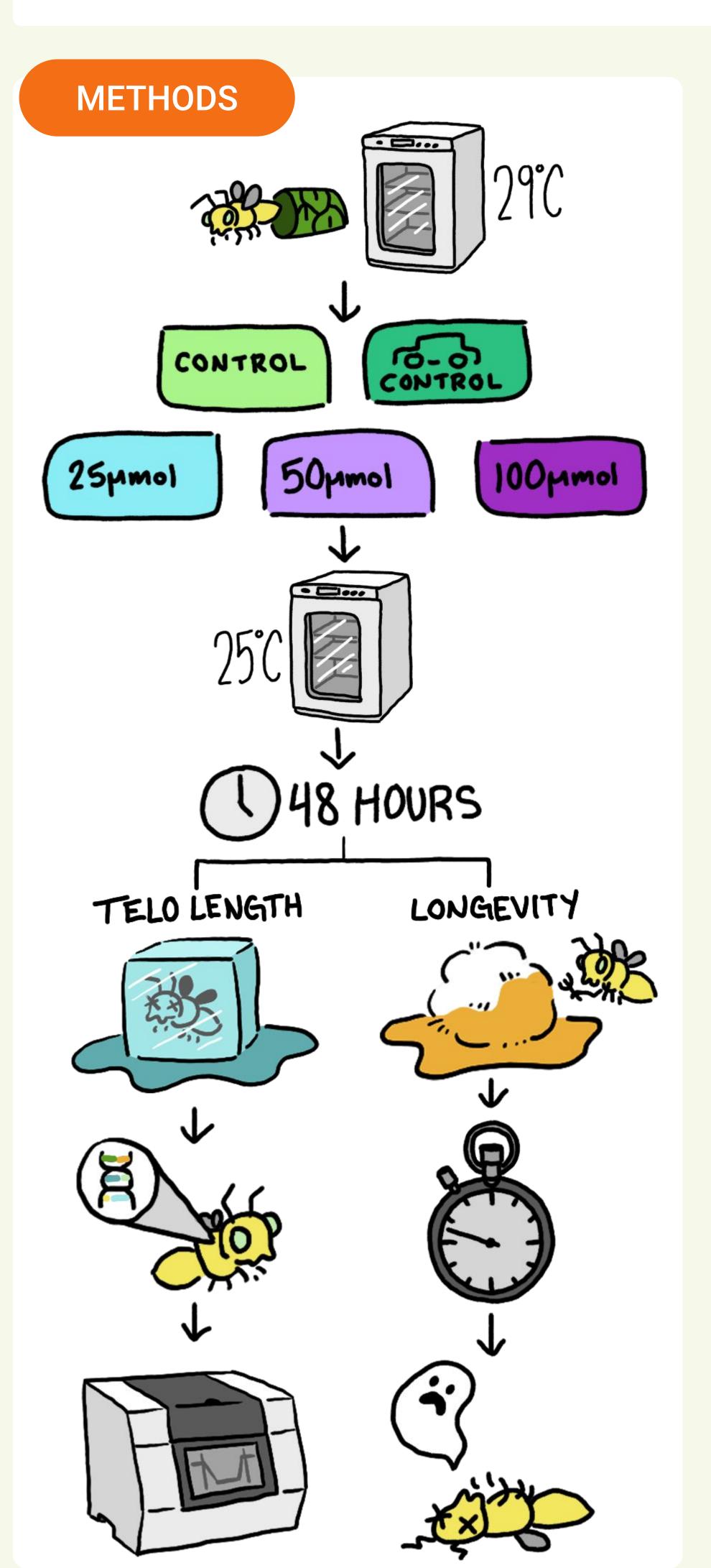


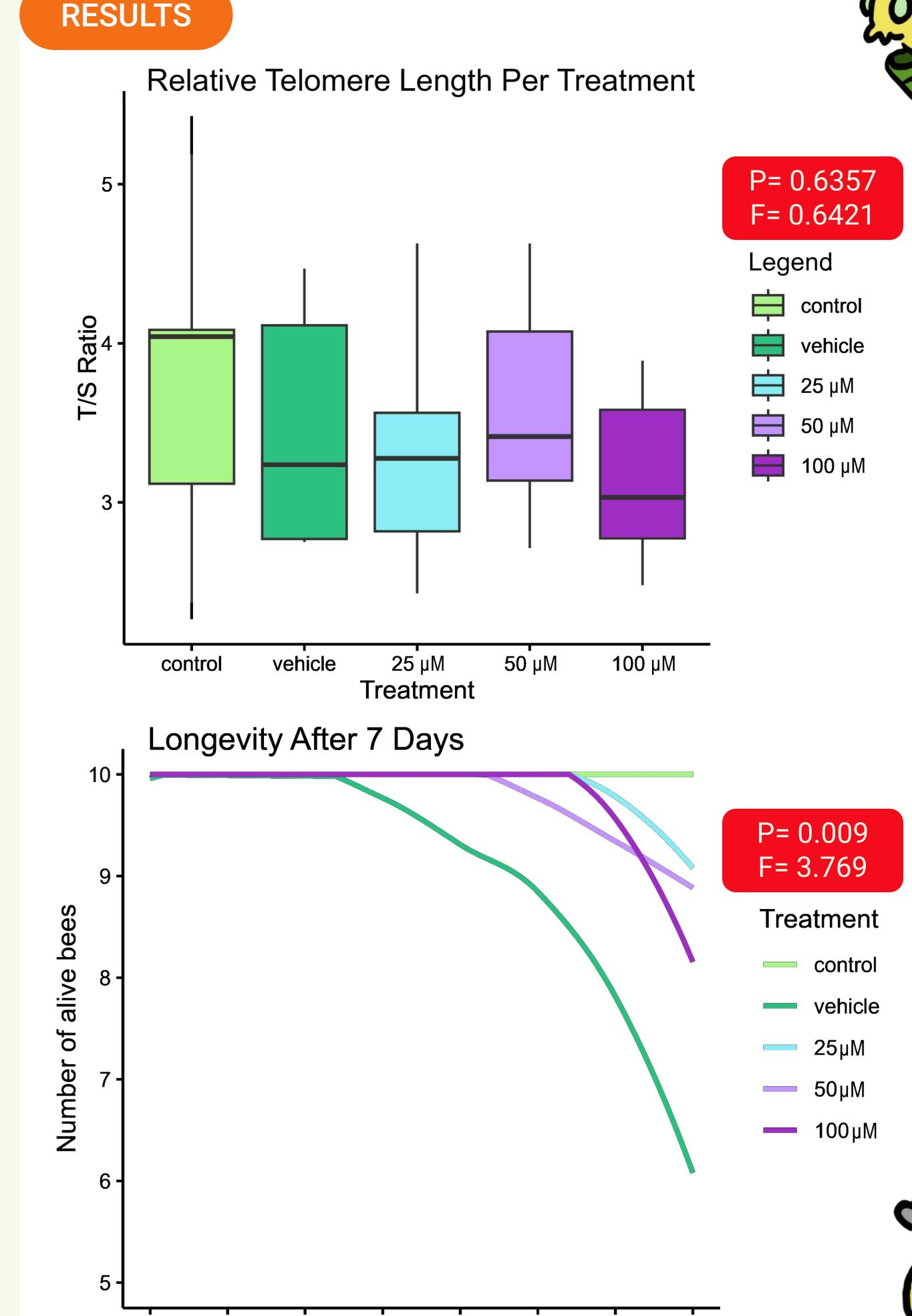
NDST NORTH DAKOTA STATE UNIVERSITY

INTRODUCTION

- Telomeres, repetitive, non-coding DNA sequences on the end of chromosomes, protect DNA during cell replication
- Telomeres shorten during cell replication, due to this, it is thought to be a biological marker of aging
- Critically short telomeres can lead to cell death
- Telomeres are repaired by the enzyme, telomerase, which declines in activity with age
- Cycloastragenol claims to promote telomerase activity which could lead to an increase in lifespan
- This project aims to examine the effects of cycloastragenol on the relative telomere length in alfalfa leaf cutting bee

HYPOTHESIS Feeding cycloastragenol to alfalfa leaf cutting bees will change telomere length and lifespan





Time in days

DISCUSSION

 T/S ratio is commonly used as a relative measurement of telomere length

Telomeric DNA primer (T)

Reference gene (S)

- Cycloastragenol did not have a significant impact on the relative telomere lengths in our treatment groups
- Given our 7 day preliminary results for the longevity study, it suggests that the drug is not increasing telomerase activity when fed to our bees
- The increased mortality rates may be due to the adverse effects of DMSO, the carrier of the drug

FUTURE RESEARCH

- Longevity is ongoing to examine lifespan of bees when given cycloastragenol
- Alternative methods of dosing bees:
 - Injection
 - Longer feeding time
 - Higher dosages
- Trap assay
 - Measures telomerase activity



ACKNOWLEDGEMENTS

Thank you to all the faculty at the USDA-ARS and the REU students for their contributions and support. This project could not have been completed without their help.

CONTACT INFO



Gwenyth Sailer - gwenyth.sailer@ndus.edu



Ella Eleven - rottingpiethesquid@gmail.com

REFERENCES

Blackburn, E. H. (2005). Telomeres and telomerase: Their mechanisms of action and the effects of altering their functions. *FEBS Letters*, *579*(4 SPEC. ISS.), 859–862. https://doi.org/10.1016/j.febslet.2004.11.036

Chakravarti, D., LaBella, K. A., & DePinho, R. A. (2021). Telomeres: history, health, and hallmarks of aging. In *Cell* (Vol. 184, Issue 2, pp. 306–322). Cell Press.

https://doi.org/10.1016/j.cell.2020.12.028

Grula, C. C., Rinehart, J. D., Anacleto, A., Kittilson, J. D., Heidinger, B. J., Greenlee, K. J., Rinehart, J. P., & Bowsher, J. H. (2024). Telomere length is longer following diapause in two solitary bee species. *Scientific Reports*, *14*(1). https://doi.org/10.1038/s41598-024-61613-2