

Computational Measurement of *L. variegatus* Volume

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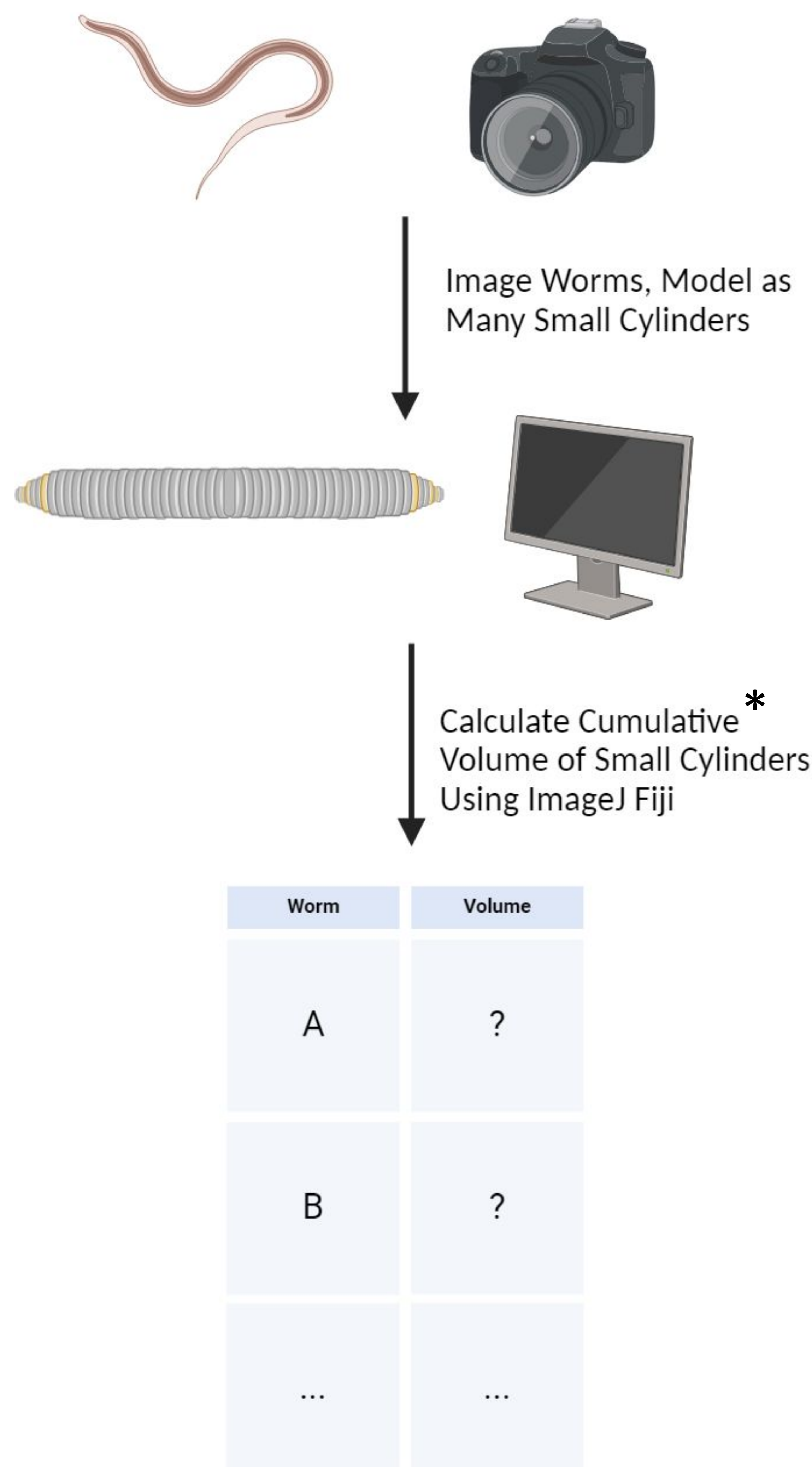


Introduction

- Lumbriculus variegatus* can fully regenerate from small segments
- Many experiments depend on measuring worm regeneration
- Current way to measure regeneration require extensive human input (time-consuming, subject to bias)

Goal: Create an ImageJ¹ Fiji² Macro to Measure Volume

Methods



Created in BioRender.com

*2D Measurements Collected with Fiji Ridge Detection Package³

“Similarly Wrong” Measurements

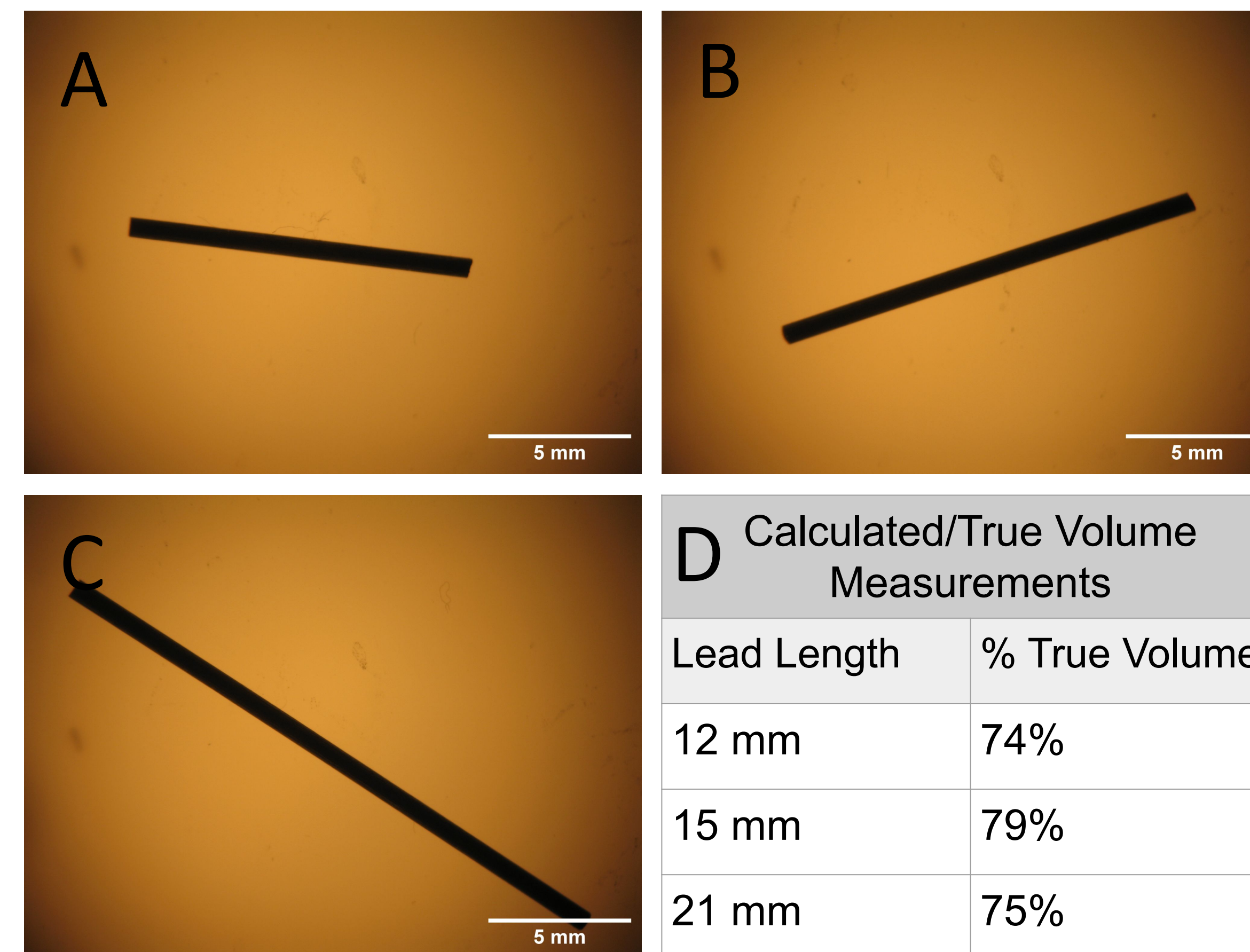


Figure 1. Calculation Consistently Inaccurate. Mechanical pencil lead 0.7 mm in diameter was sectioned, its length measured, and its true volume calculated. These lead pieces (Lengths: 12 mm [A], 15 mm [B], 21 mm [C]) were imaged and their volume was calculated using my macro. These measurements were divided by the respective true volume to calculate “% True Volume” (D). (n = 1).

Testing Calculation Precision

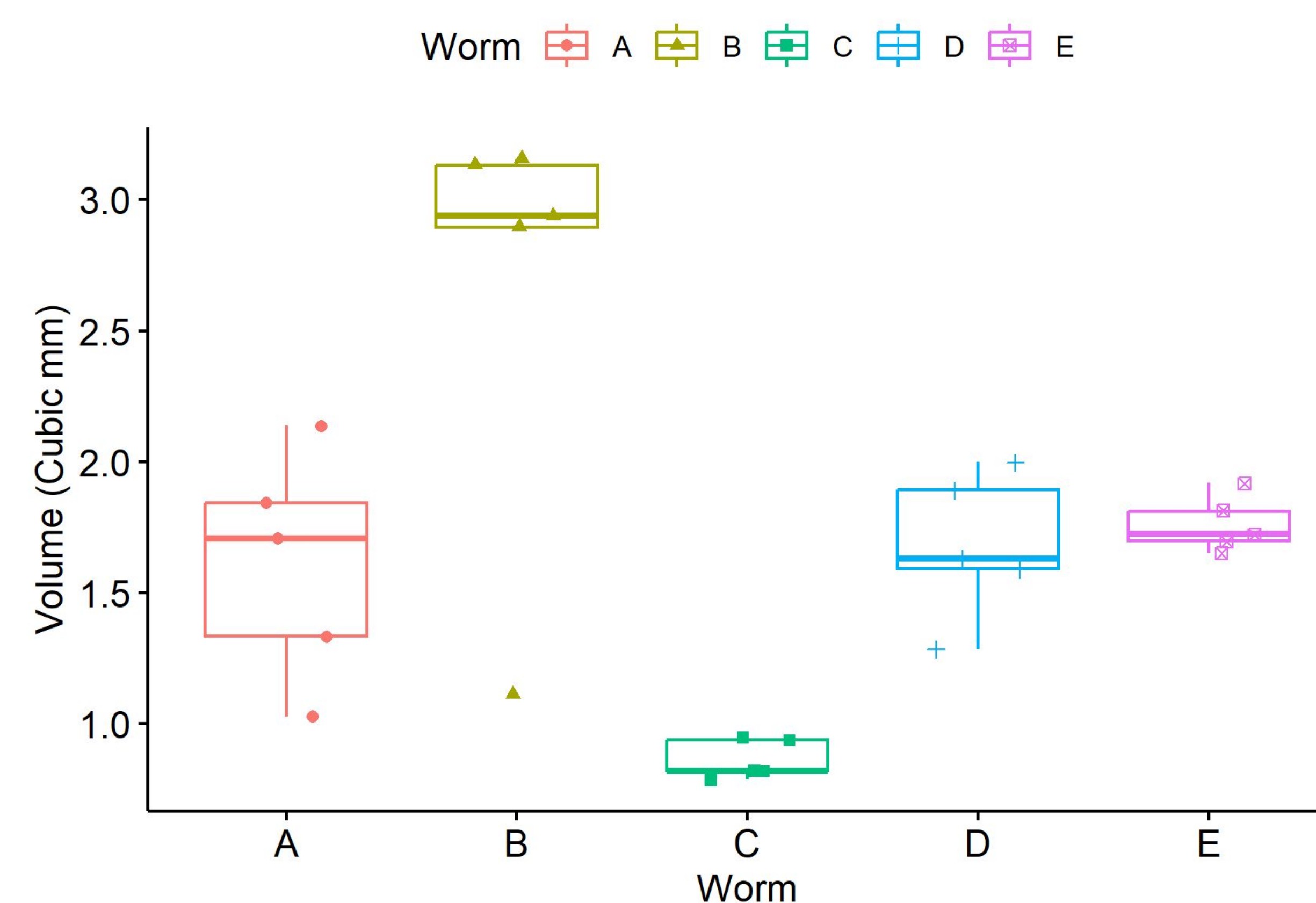


Figure 2. Worm Volume Shows Inconsistent Variance. Five *L. variegatus* were each placed under a microscope, imaged, and their volume measure using my macro. The large differences in variance between worms may suggest issues with image quality. (n = 5).

Conclusions

- Program Can Inaccurately but Consistently Measure Volume of Standard Object
 - Experiments/later code must account for inaccuracy
- Worm Movement Does Not Seem to Greatly Affect Volume Measurements
 - High variance is likely due to the use of low contrast images and shadows, possible bugs

Future Directions

- Test Program Using Higher Contrast Images
- Test Sensitivity of Program to Determine if it Can be Used to Measure Regeneration
- Further Develop Program to Measure Worm Regeneration
 - Divide volume of regenerated tissue by volume of worm right after cut

References

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