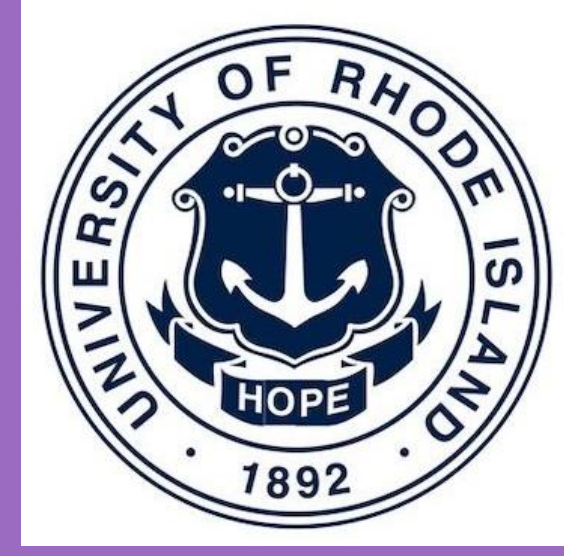


# Investigating the Antimicrobial Activity of the Sesquiterpene Lactone Laurenobiolide

Oli Horyn<sup>1</sup>, Hannah Trautmann<sup>2</sup>, Sierra Schmidt<sup>2</sup>, Kathryn M. Ramsey<sup>2</sup>, Matthew Bertin<sup>3</sup>



<sup>1</sup>Department of Pharmacy Practice, University of Rhode Island, Kingston, RI 02881

<sup>2</sup>Department of Cell and Molecular Biology, University of Rhode Island, Kingston, RI 02881

<sup>3</sup>Department of Biomedical and Pharmaceutical Sciences, University of Rhode Island, Kingston, RI 02881



## INTRODUCTION

- Antibiotic resistance is a significant threat to public health
- Novel antimicrobials may be developed based on molecules derived from plants

### Laurenobiolide

- Sesquiterpene lactone
- Isolated from North American tulip tree *Liriodendron tulipifera*
- Used by indigenous tribes as a treatment for malaria
- Bioassay-guided approach identified laurenobiolide as effective against MRSA

### *Staphylococcus aureus*

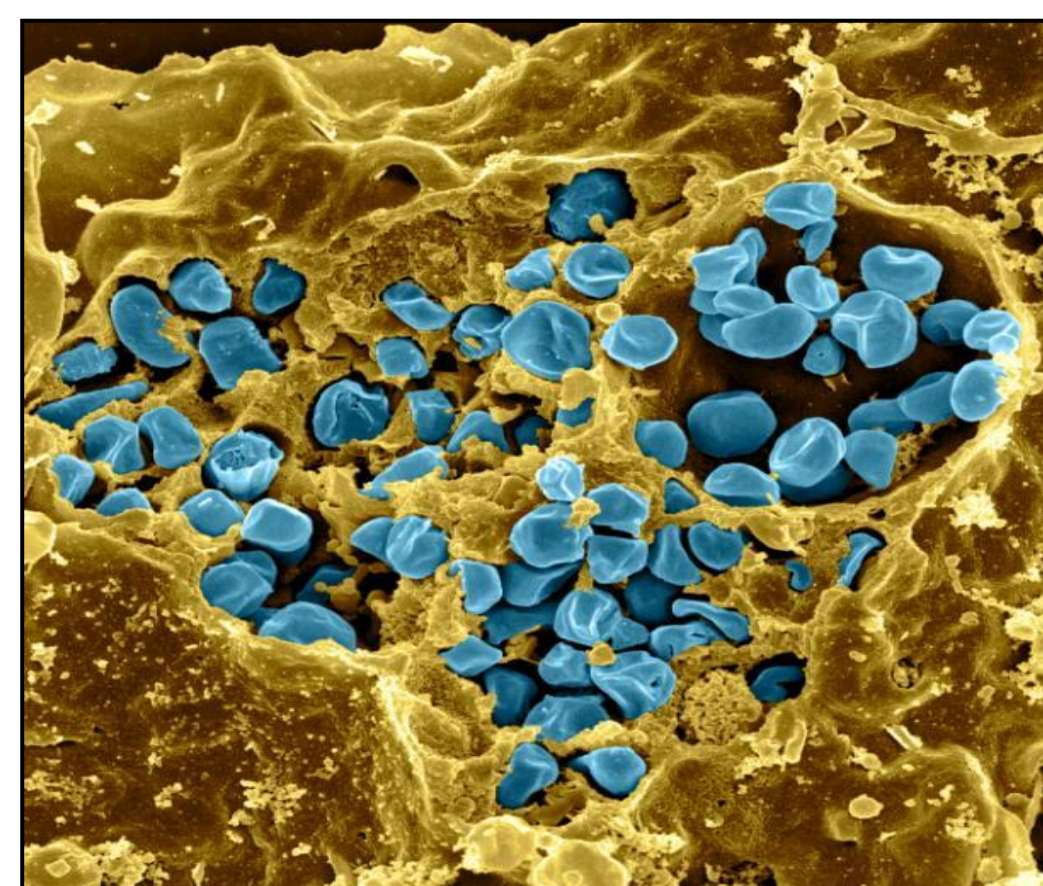
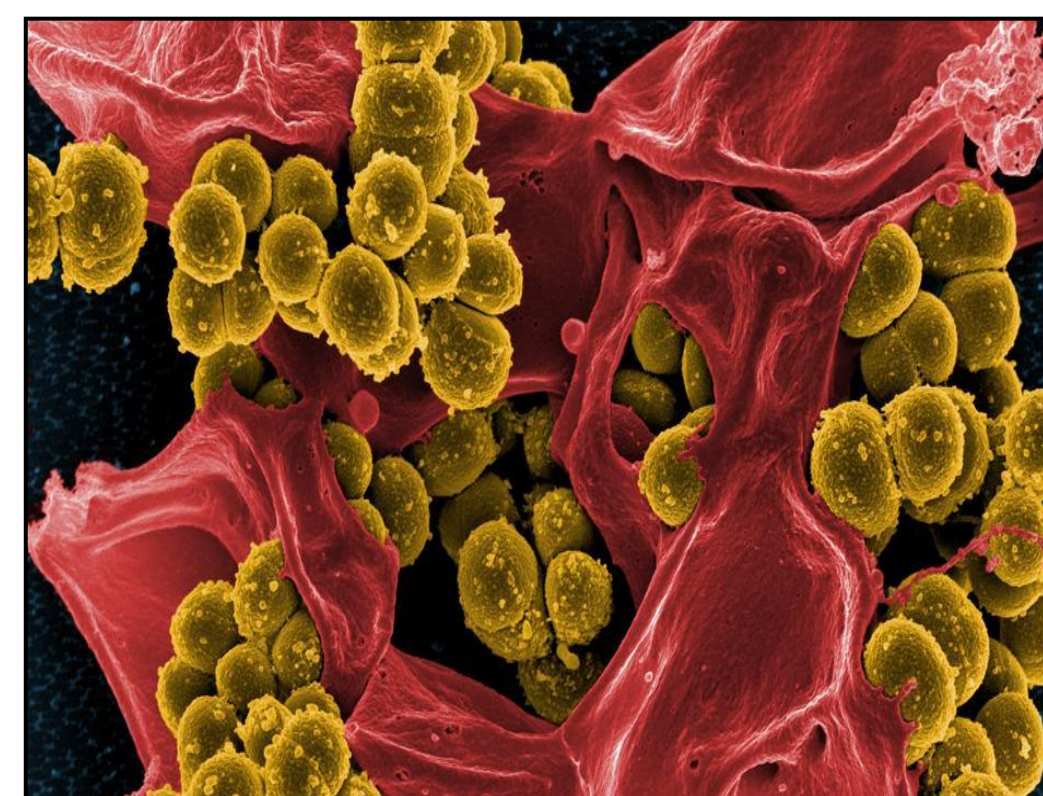
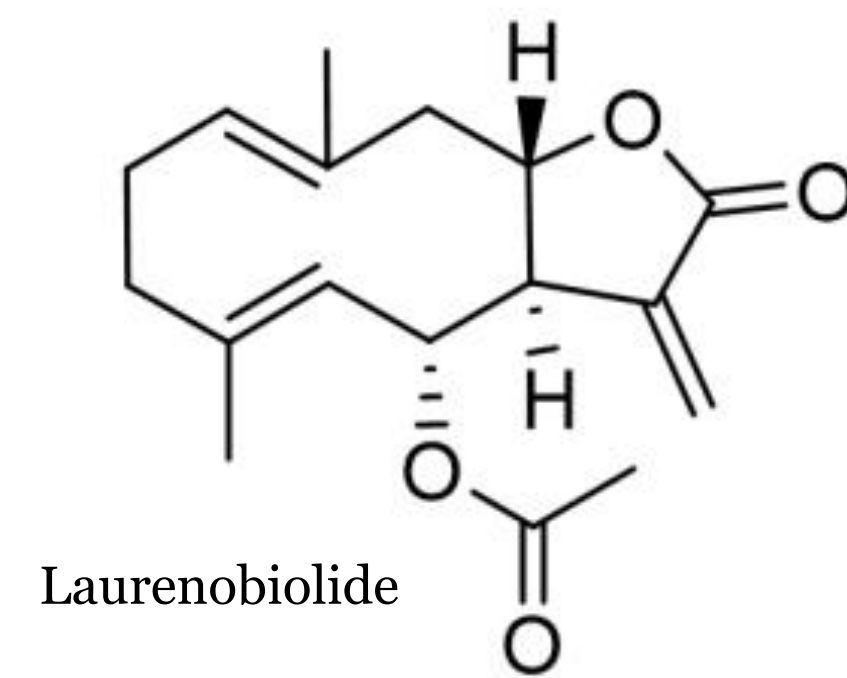
- Gram-positive
- Causes skin infections
- Can be multi-drug resistant

### *Francisella tularensis*

- Gram-negative
- Causes tularemia
- Potential bioweapon due to its highly infectious nature

### *Escherichia coli*

- Gram-negative
- Causes foodborne illness



## METHODS AND MATERIALS

### Disc Diffusion Assay Workflow

- Plated bacteria at optimized concentration in triplicate
- Two discs per plate: control (methanol/CH<sub>3</sub>OH) and disc impregnated with compound
- Incubated at 37°C
  - 24 hours for *E. coli* and *S. aureus*
  - 48 hours for *F. tularensis*

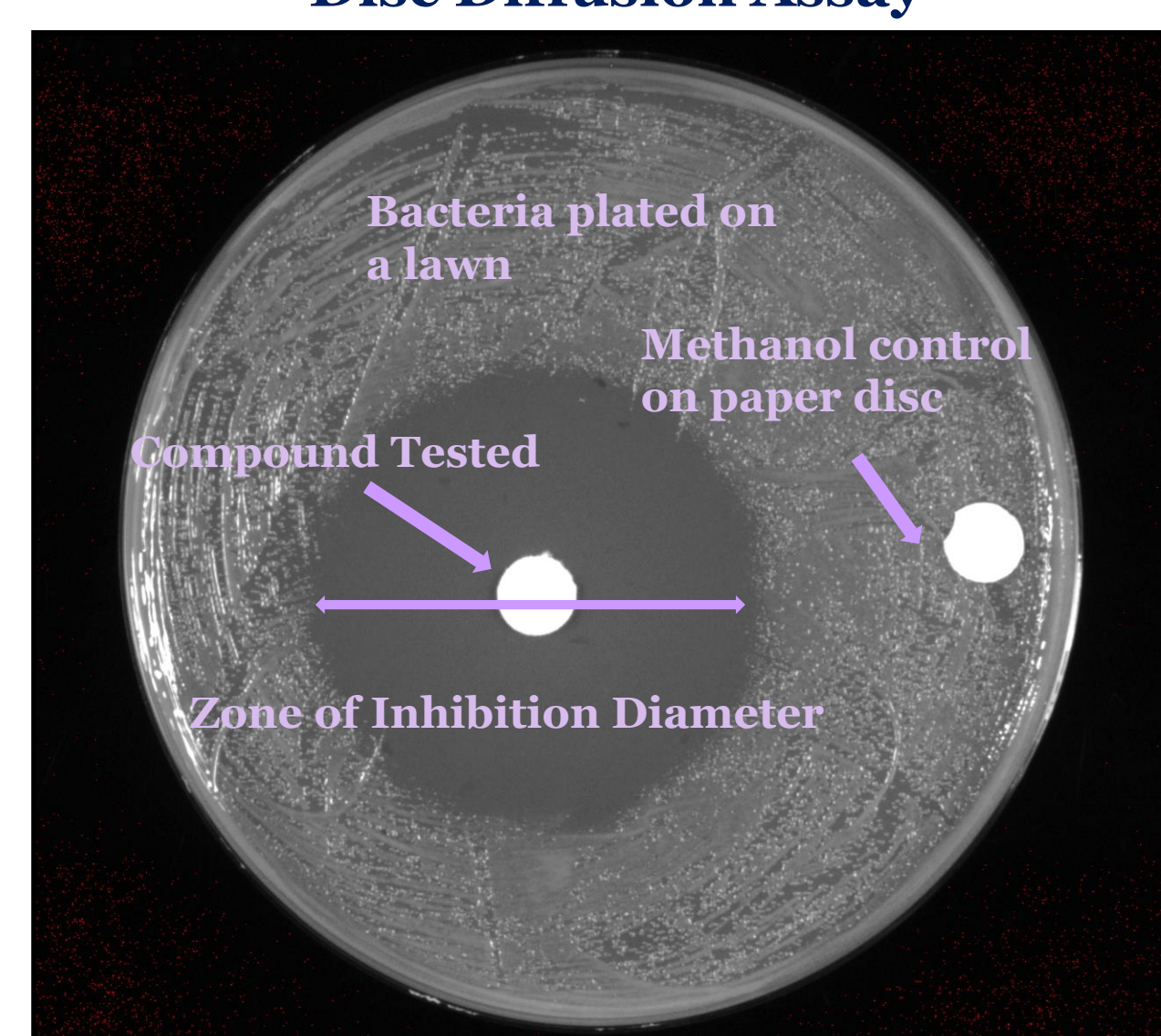
### Compounds Tested

- Laurenobiolide
- Extracts from
  - *L. tulipifera*
  - *L. chinense*
  - *L. tulipifera* – *L. chinense* hybrid

### Organisms Tested

*Staphylococcus aureus* - SA113  
*Escherichia coli* - ATCC25922  
*Francisella tularensis* – Live Vaccine Strain (LVS)

### Disc Diffusion Assay



## RESULTS

### Antimicrobial Activity of Laurenobiolide

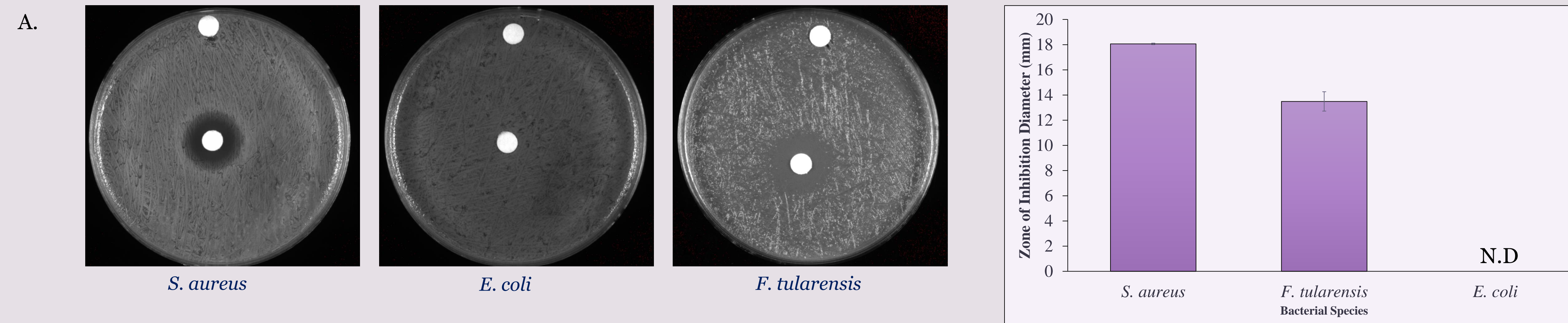


Figure 1. Multiple bacterial species are sensitive to laurenobiolide.

A. Representative image of disc diffusion assay results, using indicated bacteria

B. Quantification of disc diffusion results

Disc diffusion assays were performed with the indicated strains in biological triplicate using discs impregnated with 8 mg/mL laurenobiolide. Plates were incubated for 24 or 48 hours (strain dependent) and zones of inhibition were measured in mm. Error bars represent standard deviation. ND indicates no zone of inhibition detected

### Antimicrobial Activity of Tree Extracts

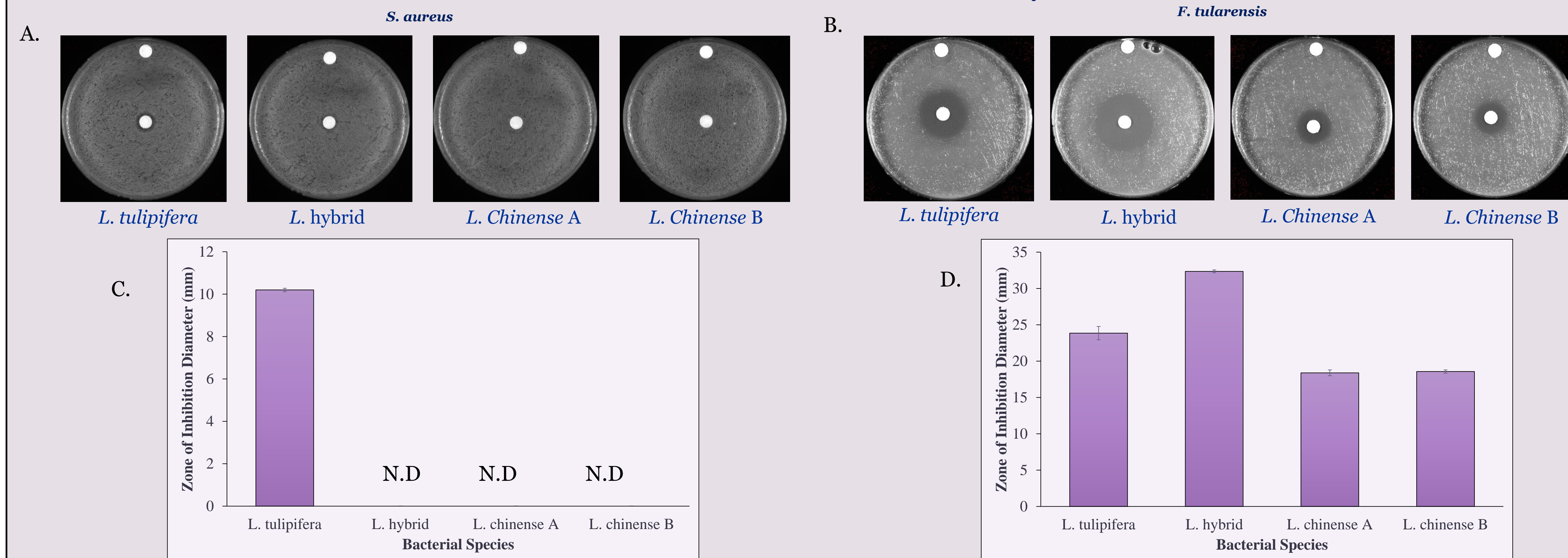


Figure 2.

A/B. Representative image of disc diffusion assay results, using indicated bacteria

C/D. Quantification of disc diffusion results

Disc diffusion assays were performed with the indicated strains in biological triplicate using discs impregnated with 10 mg/mL sample extract. Plates were incubated for 24 or 48 hours depending on strain, and zones of inhibition were measured in mm. Error bars represent standard deviation. ND indicates no zone of inhibition detected

## CONCLUSIONS AND FURTHER STEPS

### Laurenobiolide Antimicrobial Activity

- *S. aureus* confirmed sensitive
- Laurenobiolide active against *F. tularensis*, a Gram-negative bacterium
- Suggests possibility for broad spectrum antibiotic
- *E. coli* did not exhibit sensitivity at concentration of 8mg/mL

### Antimicrobial Activity of Tree Extracts

- *S. aureus* determined only sensitive to *L. tulipifera* extract, consistent with presence of laurenobiolide.
- *F. tularensis* sensitive to all tested extracts, suggesting the presence of other inhibitory compounds

### Isolated Laurenobiolide Resistant Mutants

- Next step into determining mode of action

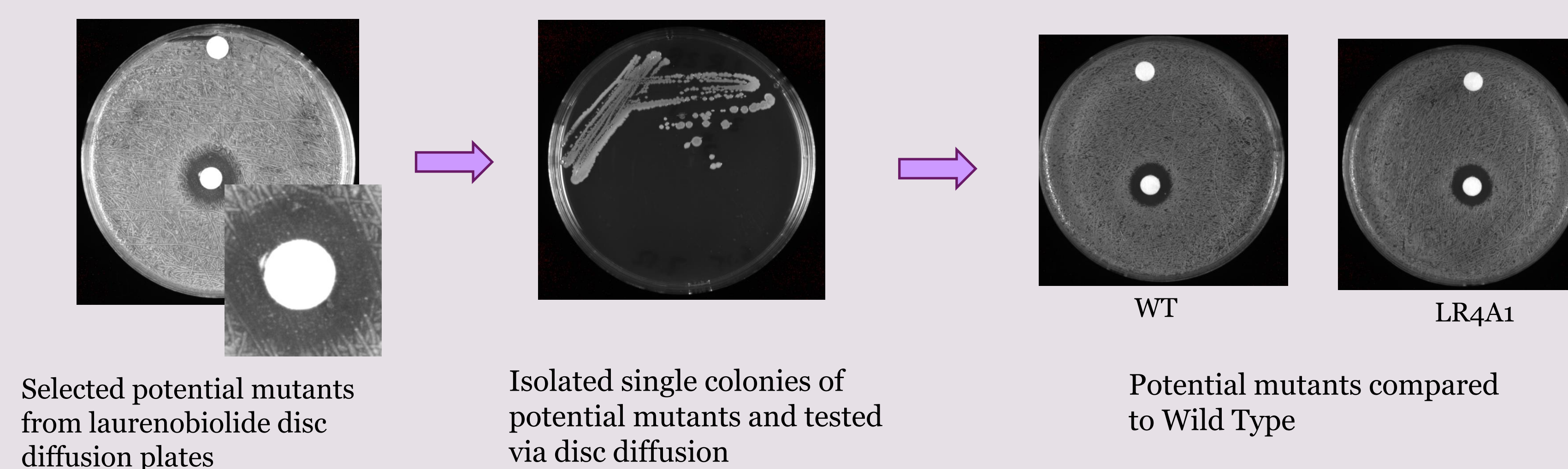
### References

Kirk, Riley D., "EVALUATING NATURAL PRODUCT LIBRARIES WITH EMPHASIS ON IN VITRO PERMEABILITY WORKFLOWS" (2021). *Open Access Dissertations*. Paper 1294. [https://digitalcommons.uri.edu/oa\\_diss/1294](https://digitalcommons.uri.edu/oa_diss/1294)

### Acknowledgments

Research reported in this presentation was supported by the Rhode Island Institutional Development Award (IDEA) Network for Biomedical Research Excellence from the National Institute of General Medical Sciences of the National Institutes of Health under grant number P20GM103430. We would like to thank Dr. Gregory and lab for productive joint lab meetings.

### Identification of Laurenobiolide Resistant Mutants



(waiting on more data, place holder(?) data)

Preliminary screening was performed with *S. aureus* with one biological replicate using discs impregnated with 8 mg/ml sample extract. Plates were incubated for 24 hours, and zones of inhibition were measured in mm. Replicates were compared against Wild Type to determine mutants of note to send for whole genome sequencing. Error bars represent standard deviation.