

Laser Carbonized Electrodes for Foodborne Pathogen Detection in Large Sample Volumes via Vacuum Driven Flow

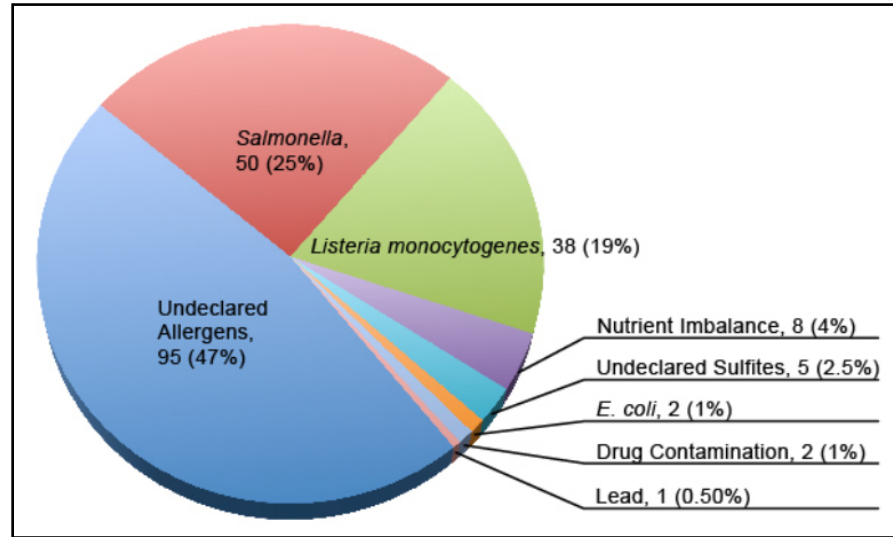
Nicholas D. Cavallaro,
C. Gomes, E. S. McLamore

Foodborne Pathogens

Everyone deserves
access to safe food.

- 1 400 Deaths
- 9.4M Illnesses
- 3.6M Illnesses from Pathogenic Bacteria
- 56K Hospitalizations

Distribution of Primary RFR Entries by Food Safety Hazard, Year 5



Years 1-4 Pie Charts can be accessed in [The Reportable Food Registry: Targeting Inspection Resources and Identifying Patterns of Adulteration Fourth Annual Report: September 8, 2010 - September 7, 2013](#).

- **Small Farmers?**
- **Big Agriculture?**
- **Organic?**
- **Non-organic?**
- **GMOs?**

Producer

- **Are the processing plants safe?**
- **What additives are in my foods?**
- **Contamination?**

Processor

??

Retailer

Distributor

- **Supermarket?**
- **Farmer's Market?**
- **Proper Sanitation?**
- **Proper refrigeration?**

- **Testing for consumer safety?**
- **Proper storage?**
- **Safe handling?**



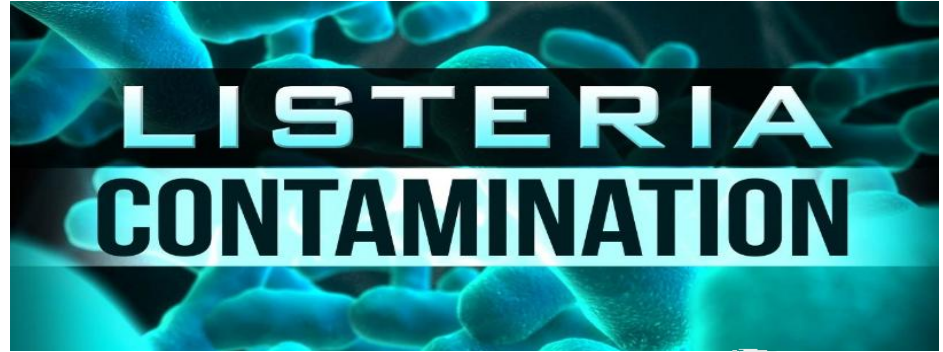
Listeria monocytogenes

Listeriosis

High Risk Populations

Recalls

Need for preventative
detection

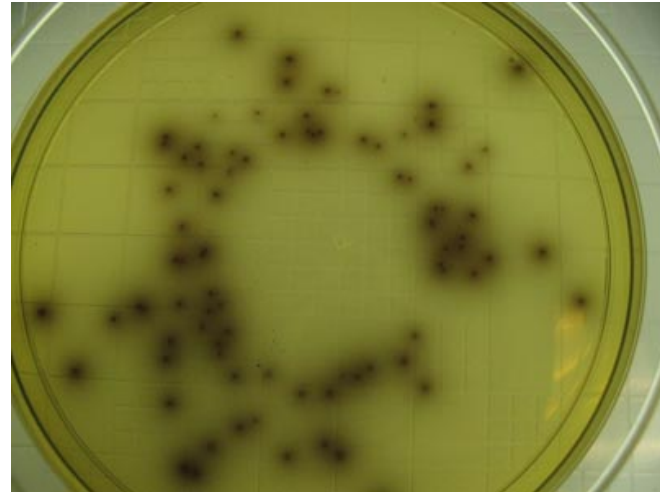


Food Safety Regulation



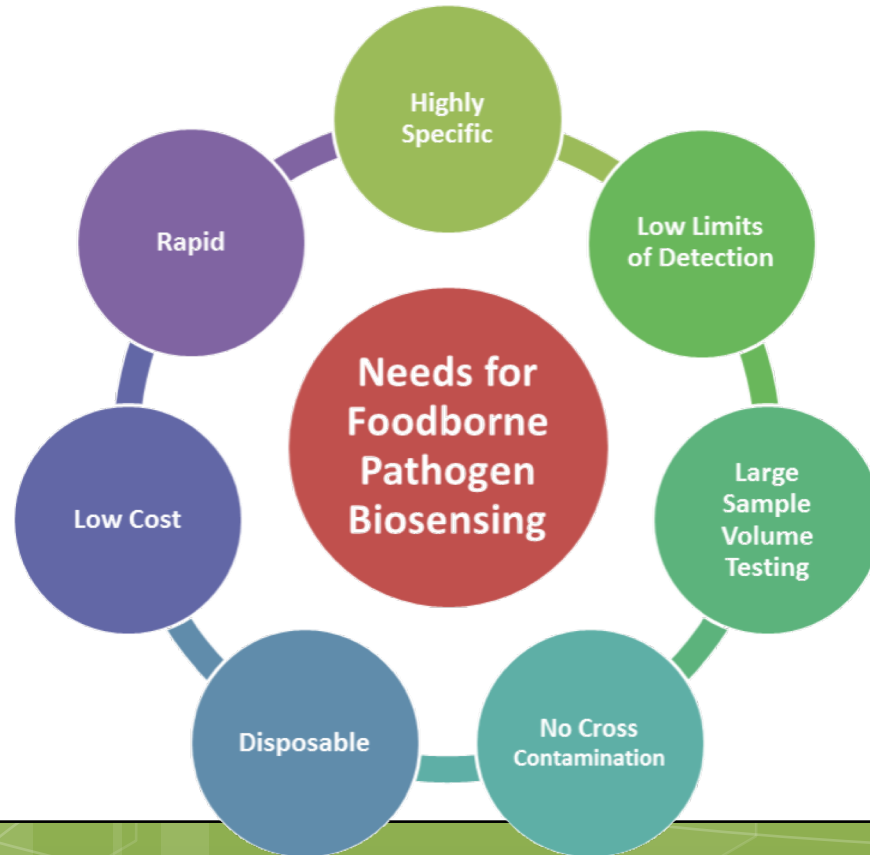
Currently Accepted Detection Techniques

- Analytical Sampling
- Incubation Period
- Colony Counting
- Polymerase Chain Reaction (PCR)



This research aims to develop the fundamental chemistry and physics required for establishing a biosensor technology for rapid and accurate microbial detection, potentially improving the efficacy of current HACCP systems in future studies.

Nanobiosensors



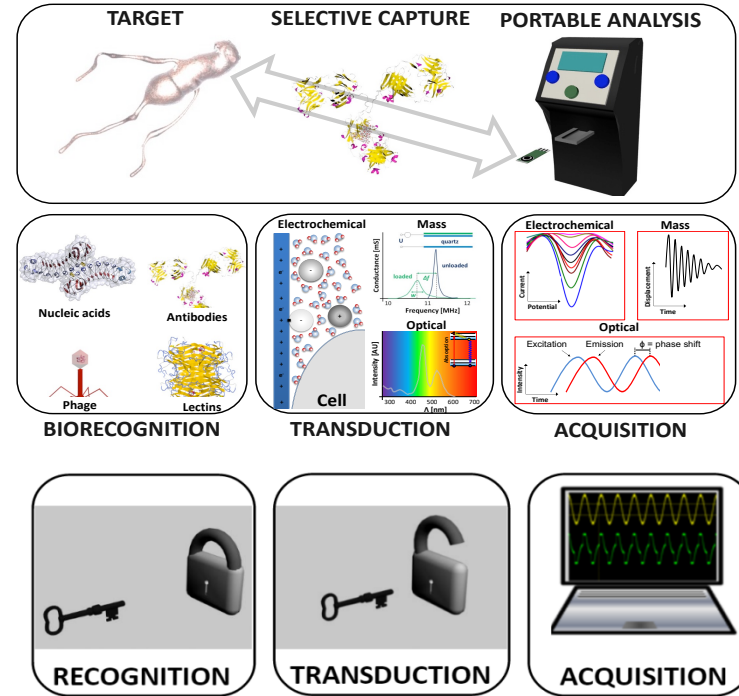
Biosensor Variety

Performance tradeoffs among biorecognition elements

Varied transduction approaches

Nanomaterials enhance biorecognition/transduction

Source: Vanegas et. al. (2017)

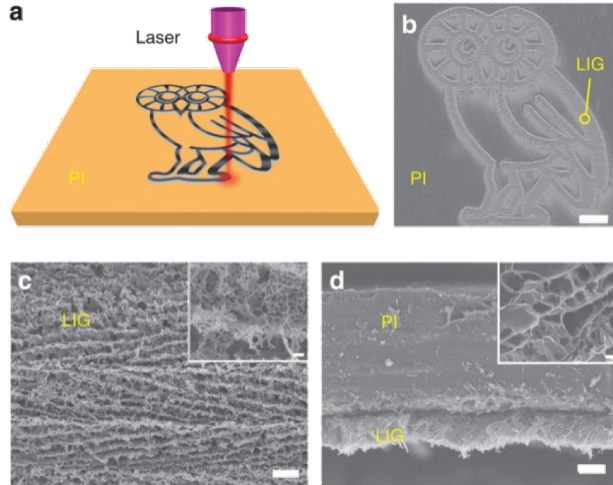


Research Goal

The overall goal of this research is to develop low cost, laser carbonized electrodes for rapid, point of use detection of *Listeria monocytogenes* in large sample volumes via vacuum driven flow directly through the electrodes.

Laser Induced/Scribed Graphene

Lin et. al. (2014)
develops LIG from
polyimide films



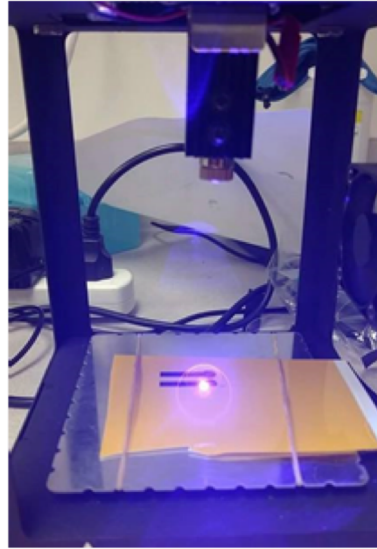
LIG/LSG for Nanosensing and Biosensing

- Tehrani et. al. (2016)
 - Glucose sensing
- Vanegas et. al (2018)
 - Biogenic amine sensing
- Garland et. al. (2018)
 - Nitrogen sensing in soil



Electrode Fabrication

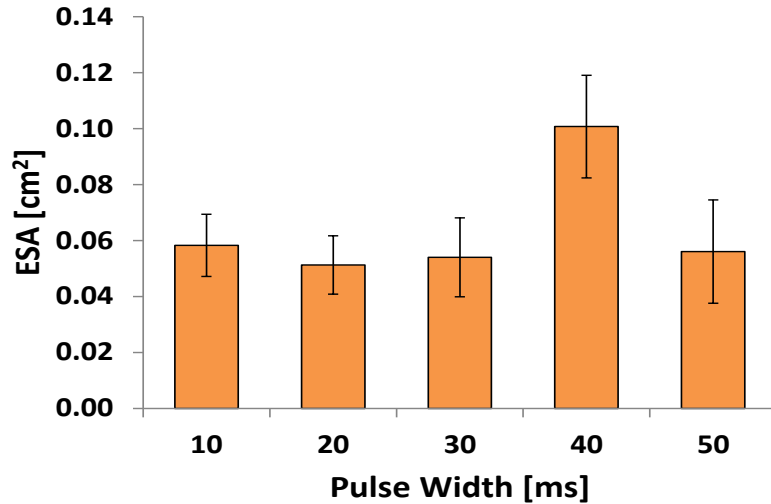
Single Working Electrode



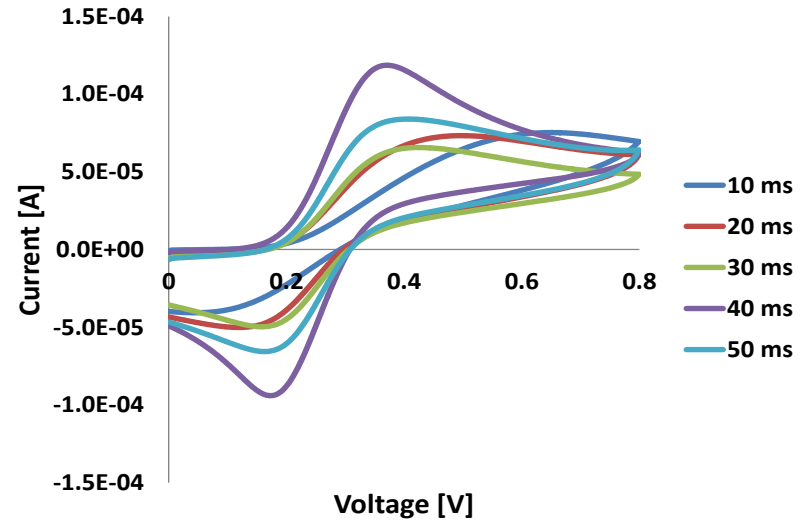
Full
Biosensor
Test Strip
w/ WE,
CE, RE

ESA Results

ESA of LSG Burned at Varying Pulse Widths in Ferrocyanide Probe

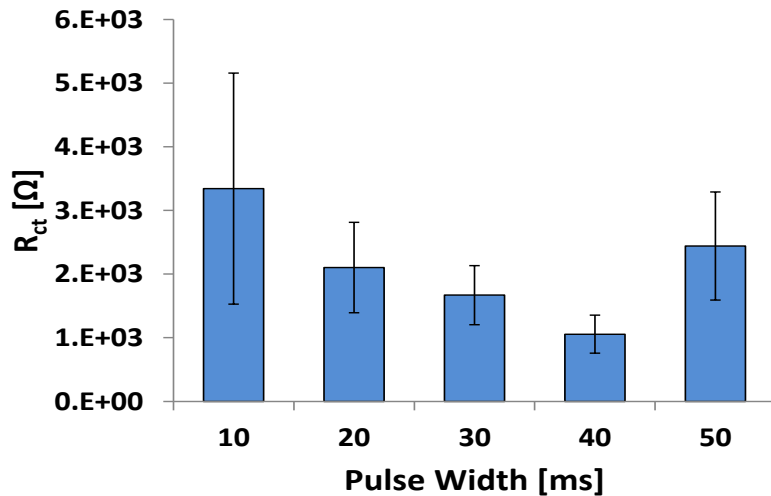


100 mV Sweep Cyclic Voltammograms Pulse Width Comparison in Ferrocyanide Probe

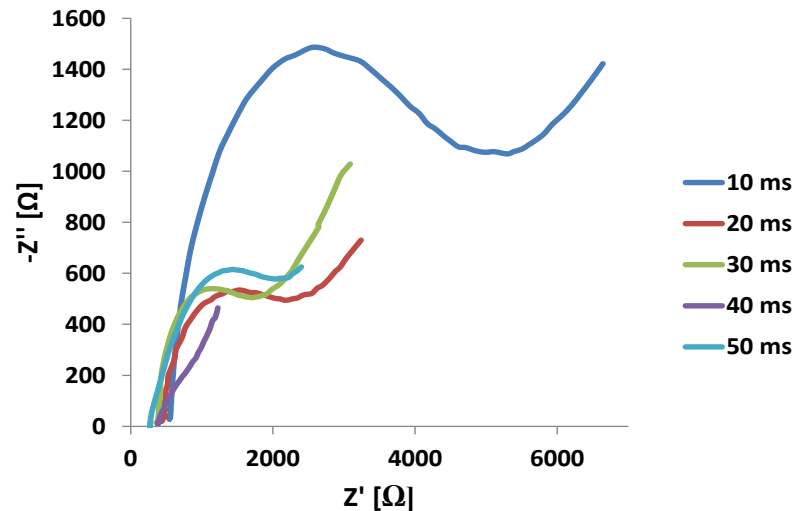


EIS Results

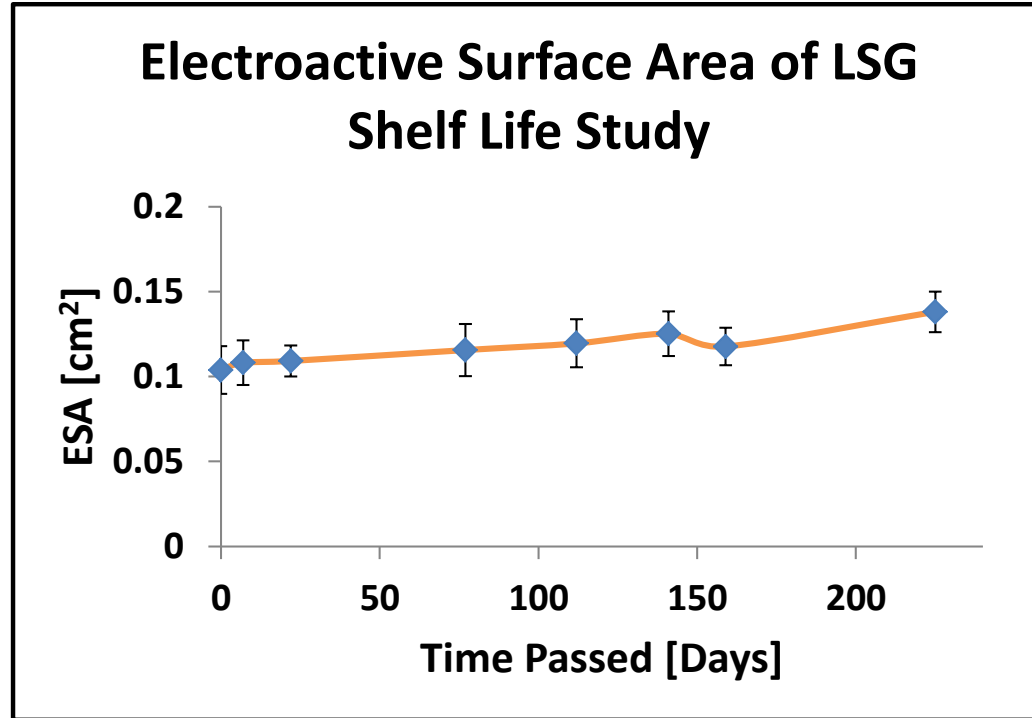
**Charge Transfer Resistance of LSG
Burned at Varying Pulse Widths**



**Representative Nyquist Plots
Pulse Width Comparison**

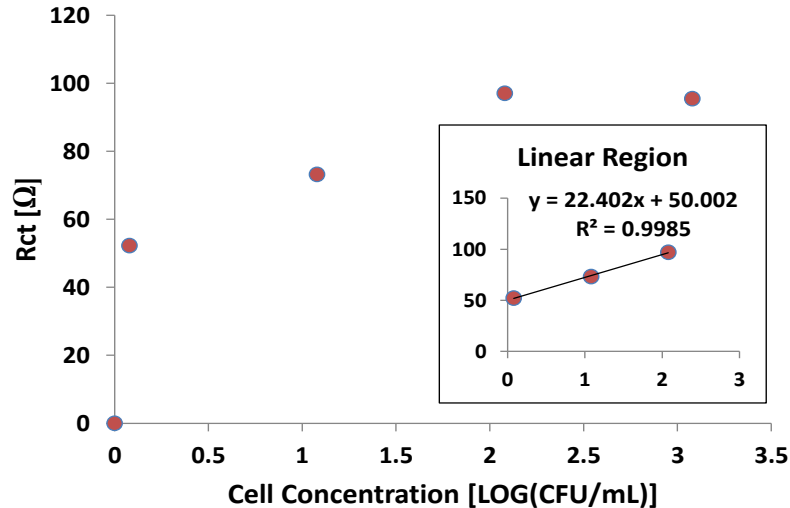


Shelf Life Study

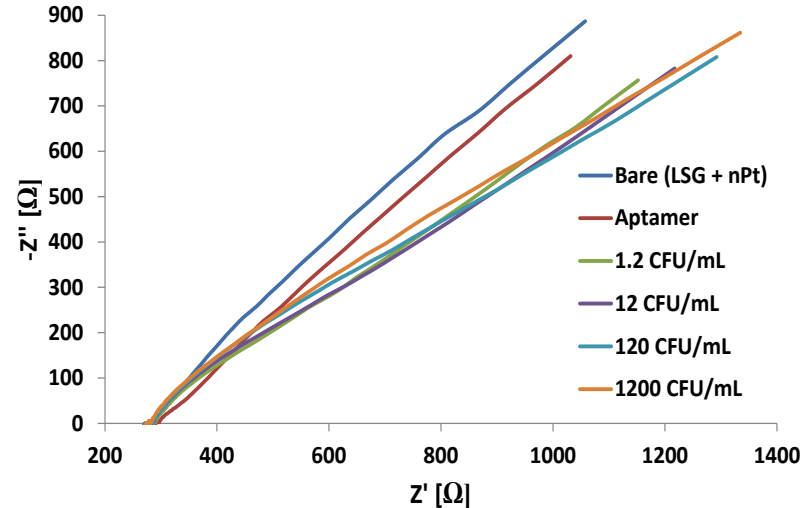


Preliminary Pathogen Detection

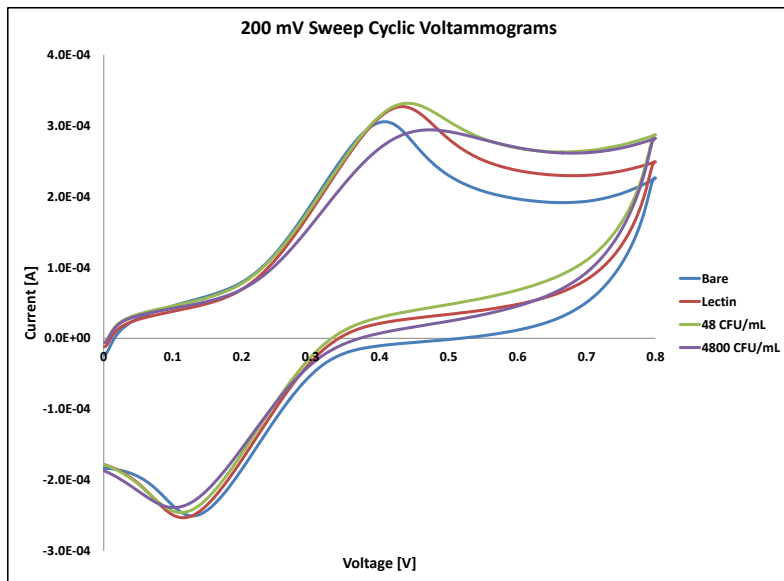
L. monocytogenes detection with LSG
Calibration Curve



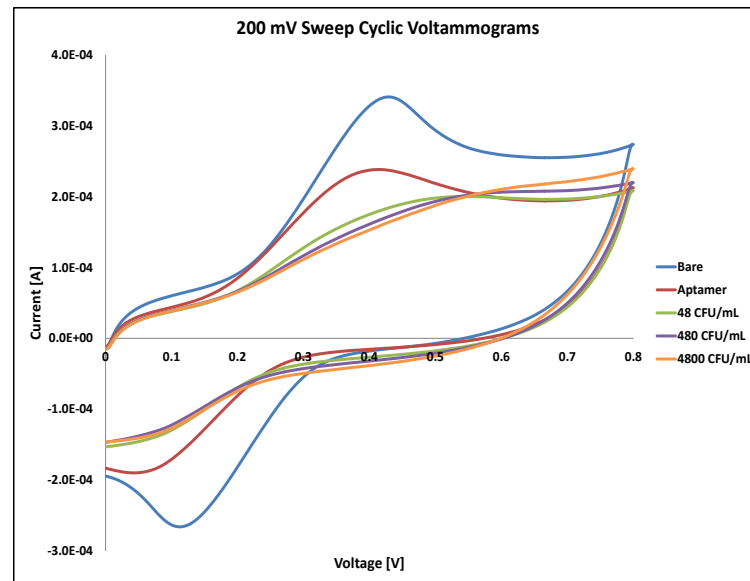
Detection of *L. monocytogenes* with LSG
Representative Nyquist Plot



Easily Change the Target: *E. coli* Detection



Electrostatically Bound Lectin



Covalently Bound Aptamer

Vacuum Driven Flow Sensors

Develop and test a technique for *Listeria monocytogenes* detection in 225 mL samples, according to FDA standards, using vacuum driven flow through laser scribed graphene.

Objective Goal

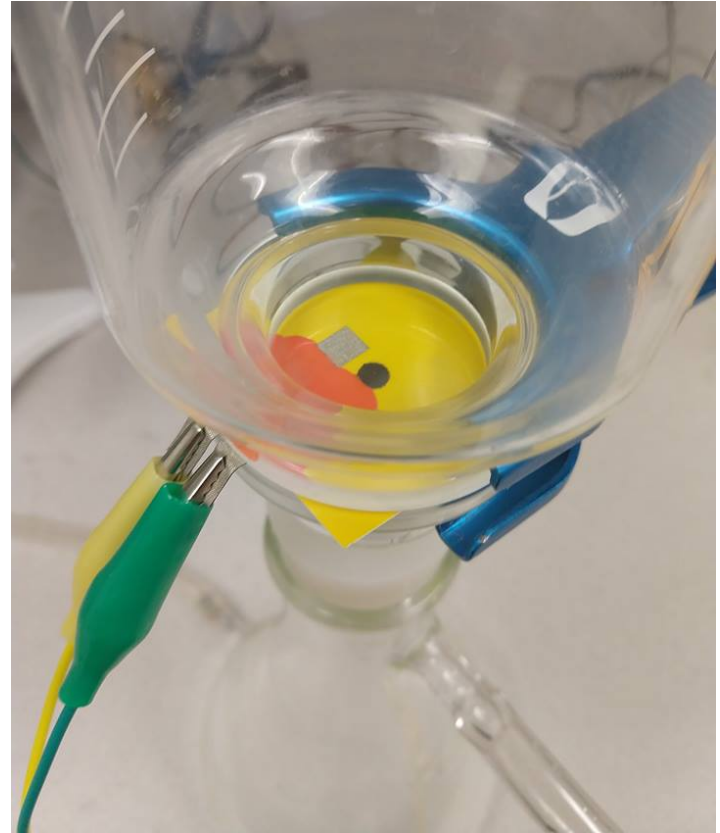
- Vacuum processing of large sample volumes directly through porous graphene electrodes will allow for greater contact of *Listeria monocytogenes* cells with the biorecognition elements on the sensor surface and lead to increased detection capability.

Significance


- This study intends to develop a new technique for processing large sample volumes in biosensing.

Vacuum Driven Flow Throw LSG

- 225 mL Testing Volume
- Durability Testing with CV
- Flow Rate Determination
- Live Potential Measurements
- Sequential Testing: Bare LSG, nPt, Aptamer, Chitosan
- *L. monocytogenes* Detection



Broader Impacts

- 
- Low cost methods for pathogen detection
 - Increased sample volume processing
 - Reduced pre-incubation time for microbial testing
 - Reduced holding times and reduction of food waste
 - Impacts beyond the food industry

Acknowledgements

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Undergraduate Assistants:

Ariel Terrand, Alyssa Bement, Zoë Davis,
Enoch Kuo, Samantha Winther





Thank You For Your Time

Questions?