

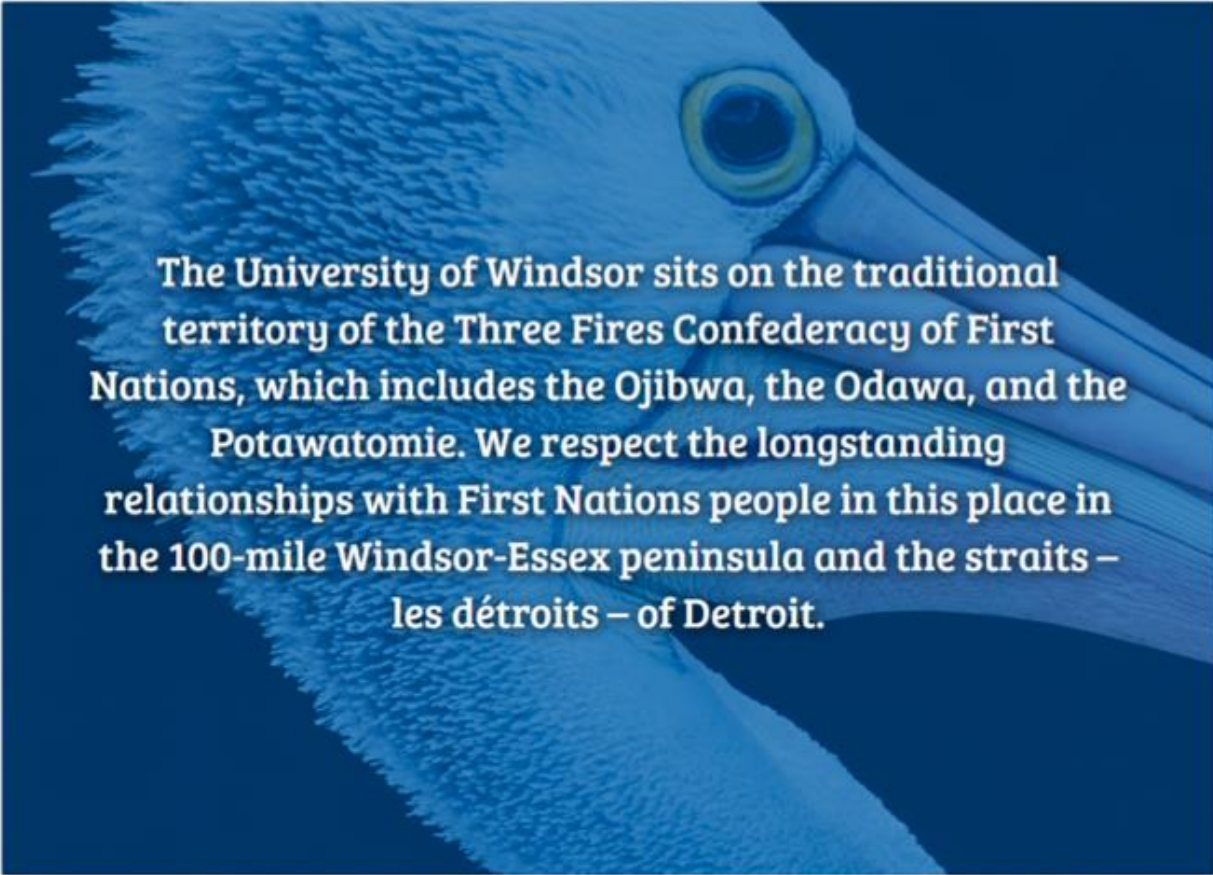
Pulsed Laser Deposition of a Silver Film For Laser-Induced Breakdown Spectroscopy Emission Enhancement of Bacteria

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The University of Windsor sits on the traditional territory of the Three Fires Confederacy of First Nations, which includes the Ojibwa, the Odawa, and the Potawatomie. We respect the longstanding relationships with First Nations people in this place in the 100-mile Windsor-Essex peninsula and the straits – les détroits – of Detroit.



Motivation

Current methods of bacterial identification in clinical settings are contributing to the ongoing antibiotic resistance crisis:

→ They are slow (take 2-3 days)

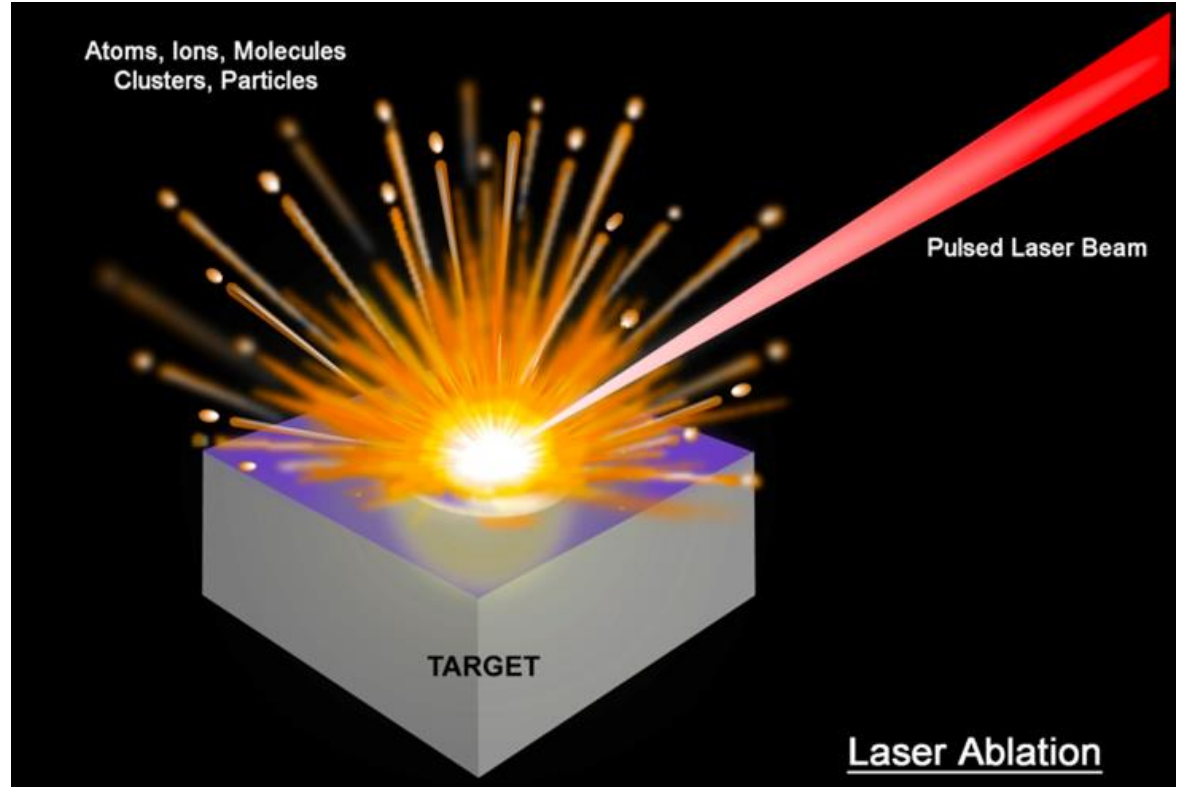
We use laser-induced breakdown spectroscopy (LIBS) on bacteria to identify and classify the bacteria species

→ It's very fast (under 1 min)

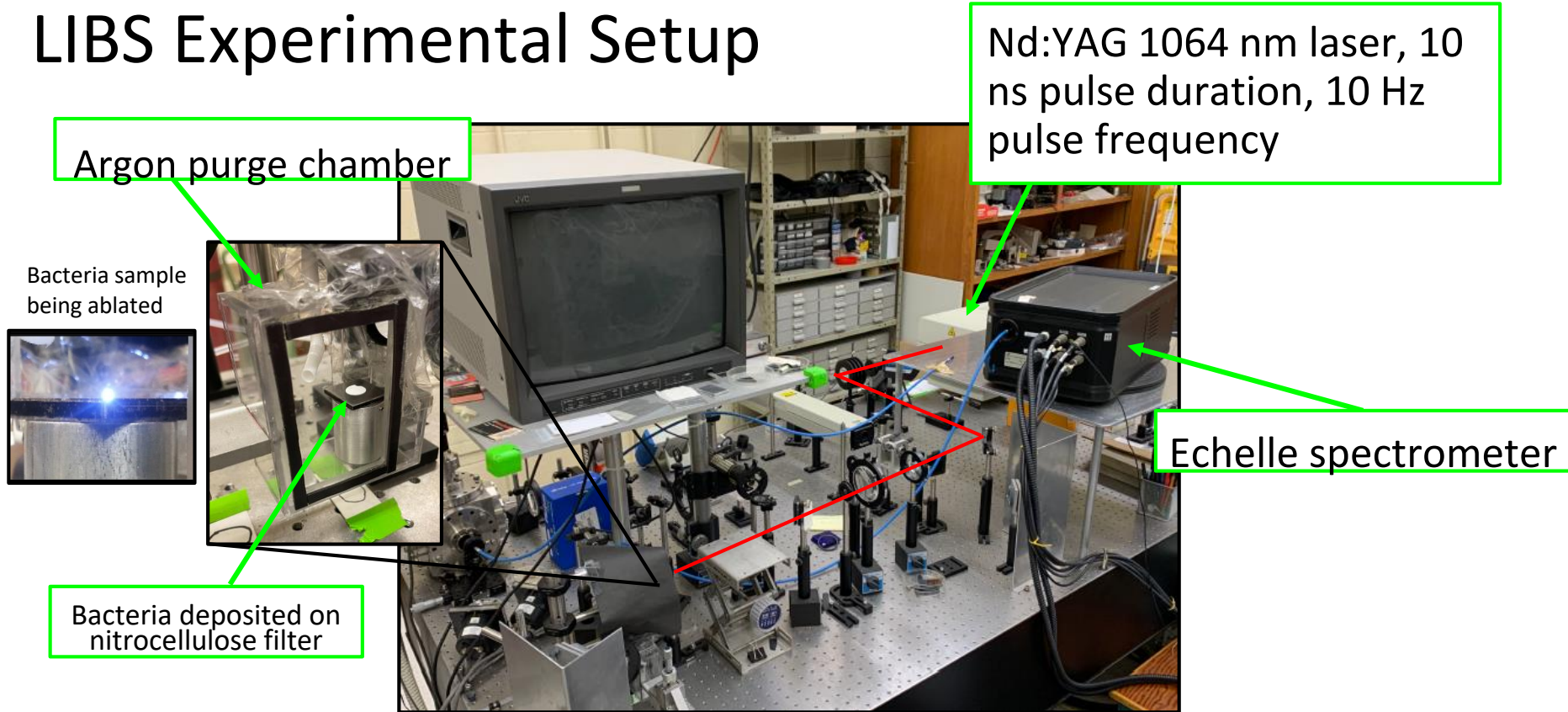


What is LIBS?

A spectrochemical technique used to rapidly determine elemental composition

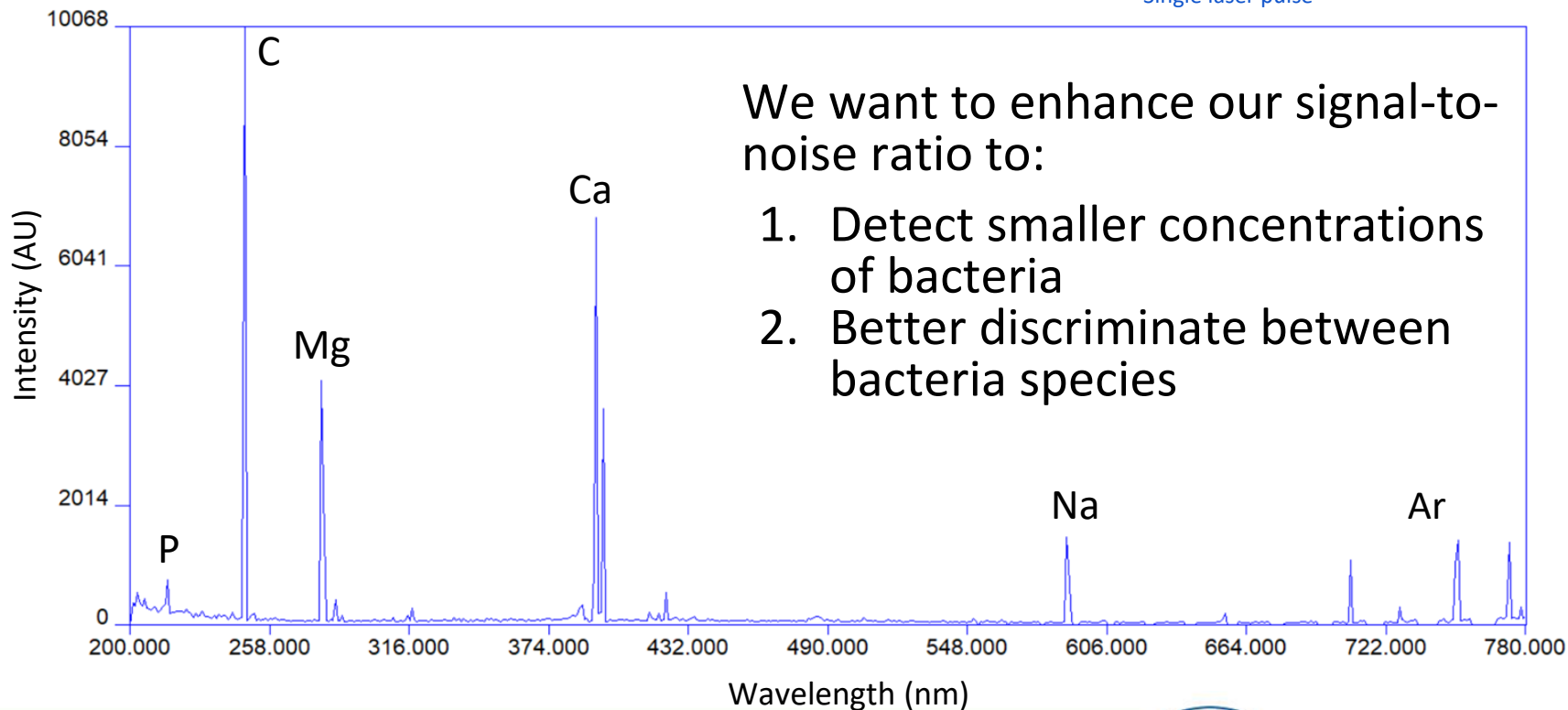


LIBS Experimental Setup

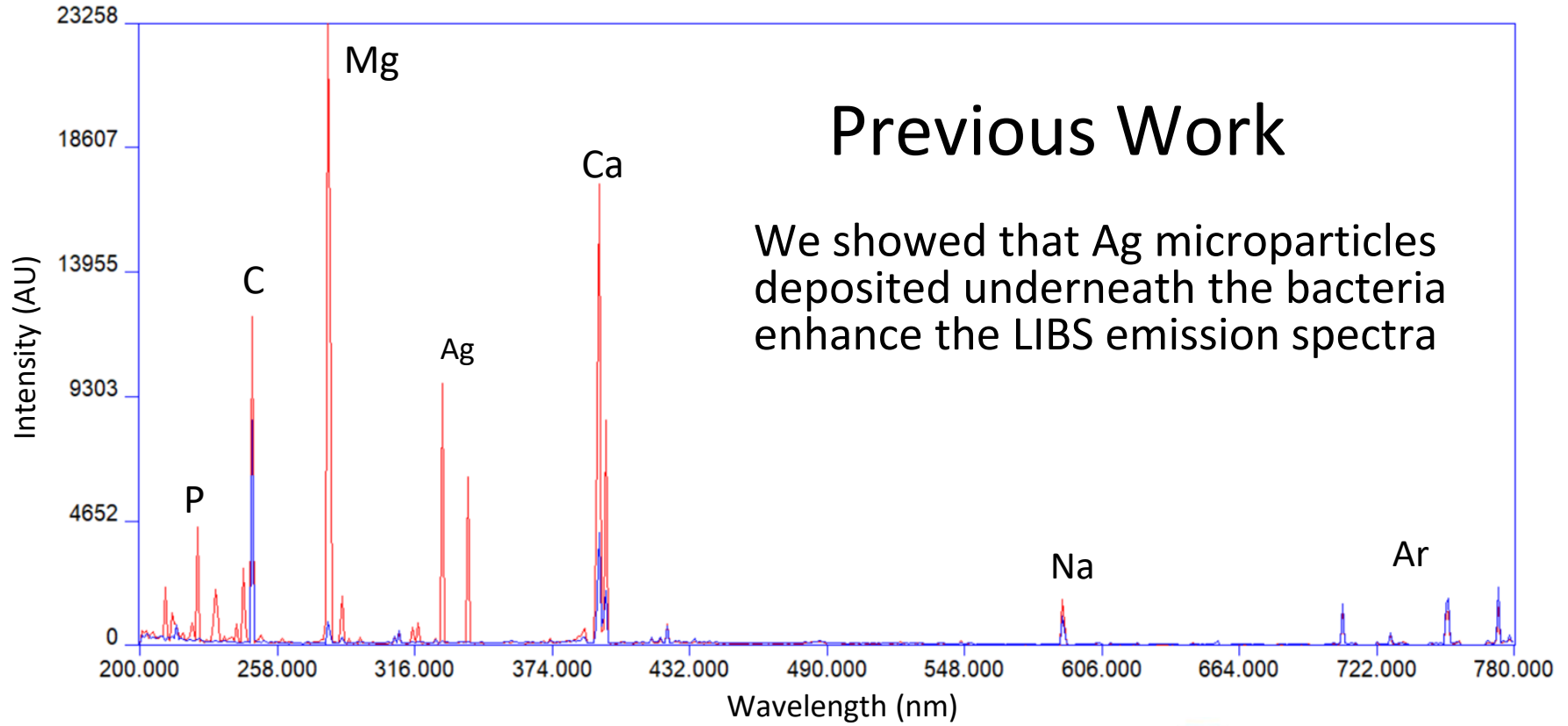


Typical LIBS Bacteria Spectrum

E. coli
2 μ s delay after plasma initiation
20 SCFH Argon environment
Single laser pulse



Red: with silver microparticles
Blue: without silver microparticles



Problems:

- Silver microparticles were displaced due to plasma shockwave resulting in decreasing intensity in subsequent pulses
- Because silver was not deposited uniformly, enhancement was very inconsistent between laser pulses

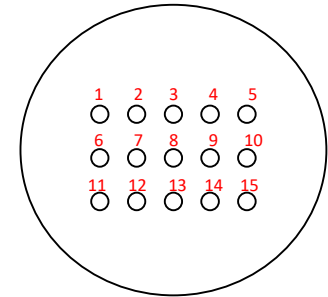
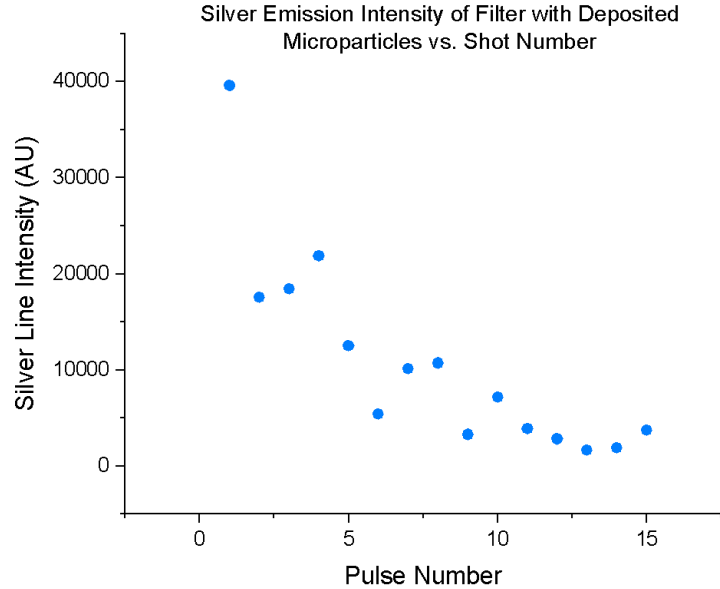


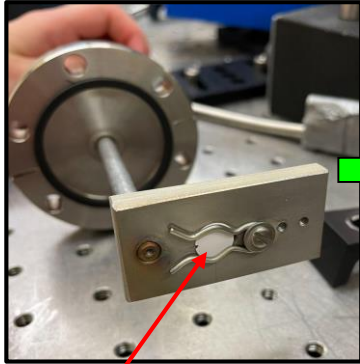
Illustration of 15 pulses on a filter

Proposed solution: pulsed laser deposition of a thin film (not microparticles)

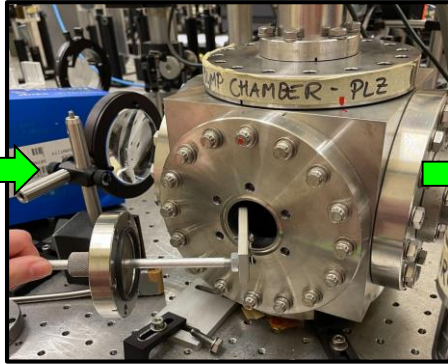
Pulsed Laser Deposition

A high-power pulsed laser beam is focused onto a target (silver) inside a vacuum chamber. The target is vaporized in a plasma plume which deposits it as a thin film on the substrate (filter).

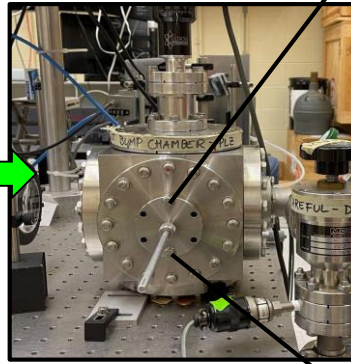
Experimental setup:



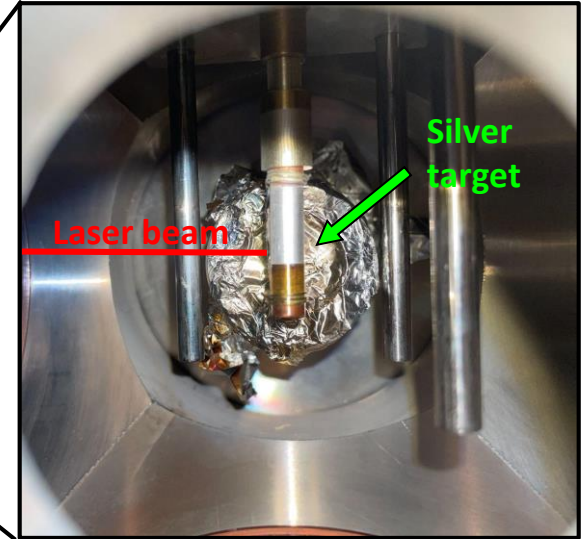
Filter on custom aluminum filter holder



Filter holder being inserted into 10 mTorr evacuated chamber



Closed vacuum chamber apparatus



Inside the chamber

Determining Filter Location and Deposition Time

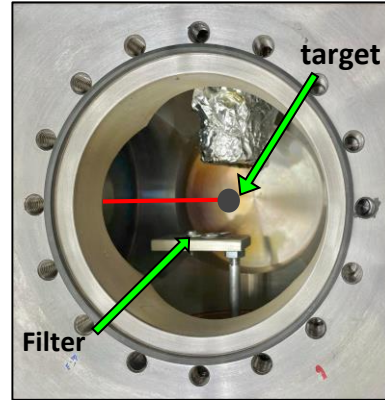
Filter location:



Brass rod



Brass coverage in various locations



Top view of chamber

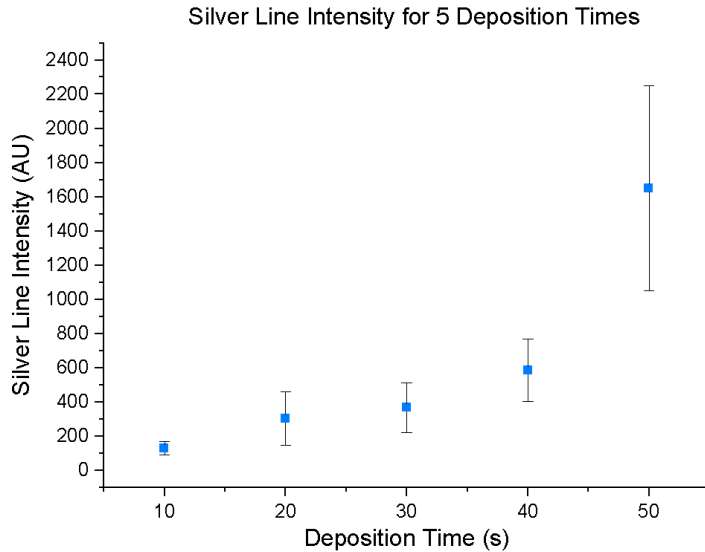


2 min deposition of silver

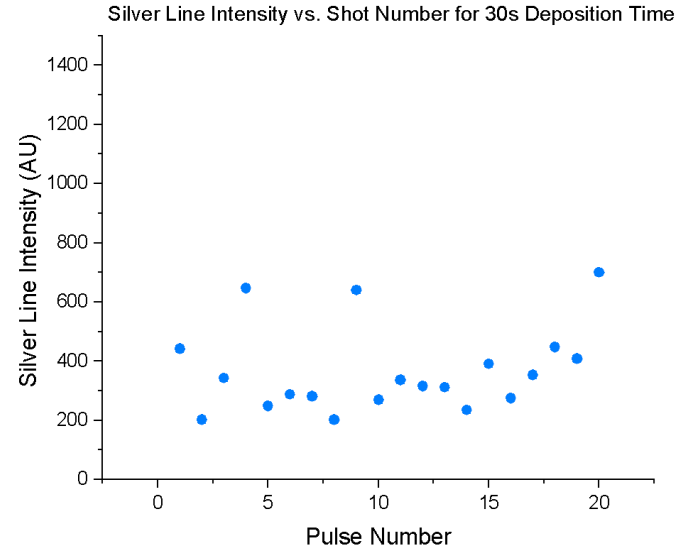
Deposition time:

- 2 min, 4 min, and 8 min sputtering times were investigated
 - ↳ Appeared to be too long, under 1 min would be investigated further with silver

Results: Uniformity of Silver Film with LIBS



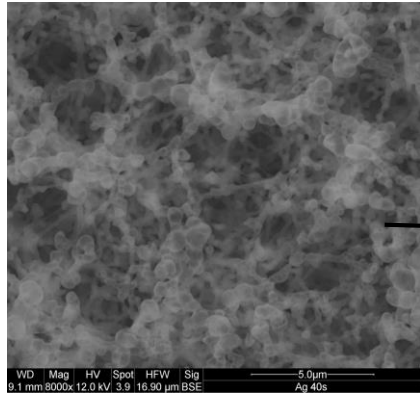
Silver line intensity increases with deposition time as expected



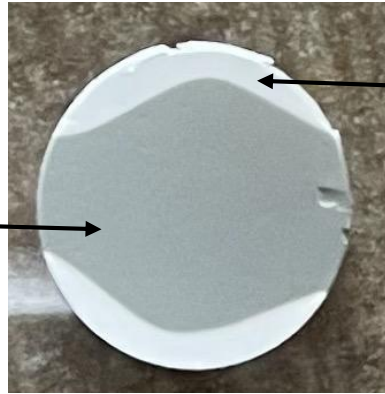
Silver line intensity shows no decrease with subsequent pulses

Results: Uniformity of Silver Film with Scanning Electron Microscope (SEM)

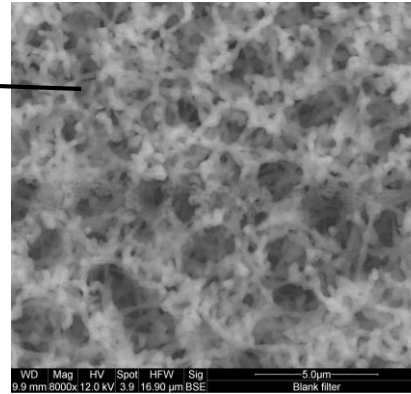
The deposition is so uniform and thin it appears the same as a blank filter



Filter with 40s Ag

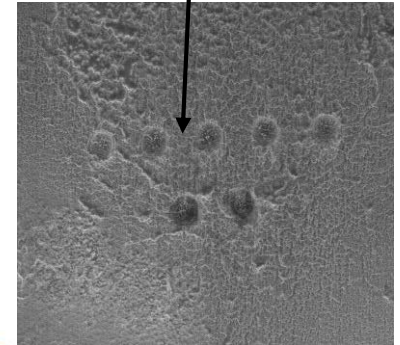
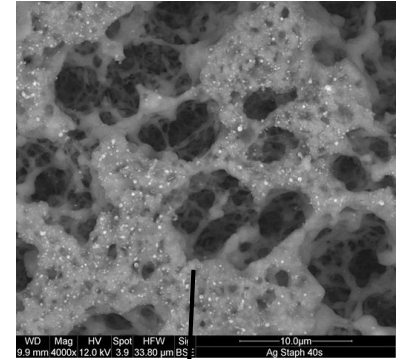


40 s deposition time
Ag filter



Blank Filter

White specks = Silver nanoparticles

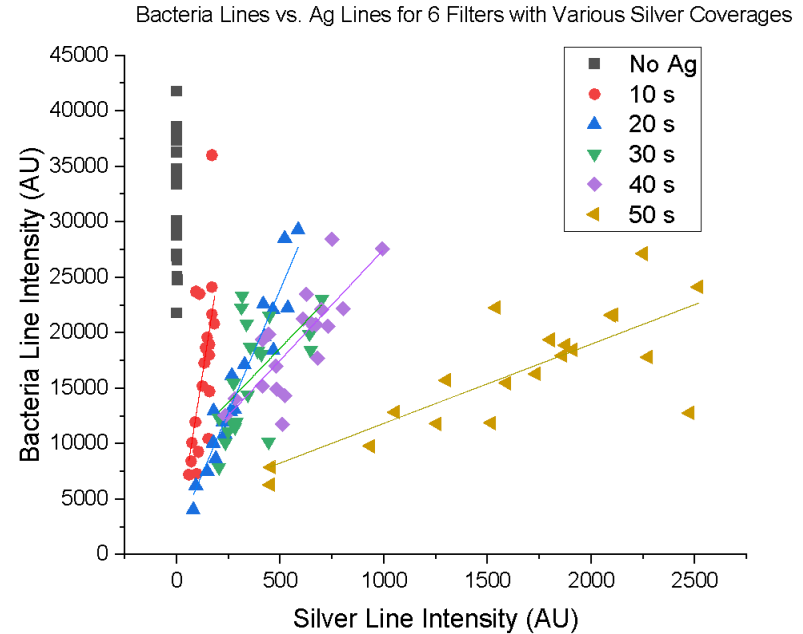


Results: Enhancement of Bacteria Spectra

Filters with more silver were not observed to have higher bacteria lines

Possible causes:

- there is not enough silver present
- the silver attenuates the laser beam before it reaches the filter



Conclusions

- A uniform silver film was deposited as shown by SEM and LIBS
- The silver film is not disrupted by subsequent laser pulses
- No bacterial enhancement was observed regardless of deposition time

Future work

- Increase amount of silver on the filter (limited by filter performance)
- Perform pulsed laser deposition in a non-vacuum environment to produce nanoparticles
- Use a gold target instead of silver

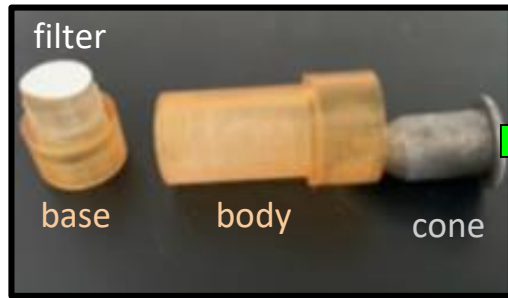


Acknowledgements

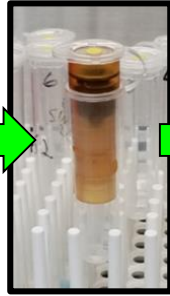
- Advisor: Dr. Steven Rehse
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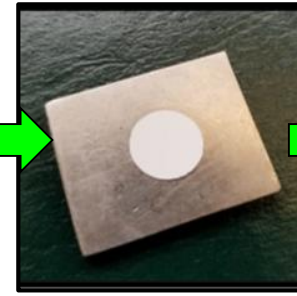
Bacterial Deposition



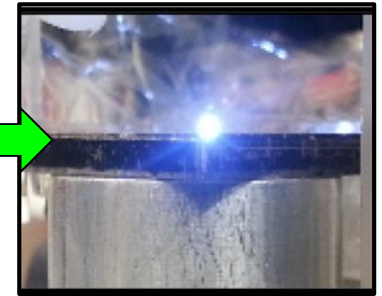
Centrifuge insert piece



Centrifugation



Sample mounted on steel piece



Sample being irradiated by laser pulse