

Interdisciplinary **Renewable & Environmental Collaborative REU**

at

sius

Cel: eati

rees Nucl

6g

 \square

NO

Objectives

- To collect biological atmospheric Particulate Matter (PM) that is less than 2.5µm
- To culture, isolate, amplify, and sequence DNA from aerosol bacterial samples.
- To perform ice nucleation experiments to observe the potential of individual bacterial samples to freeze water at higher temperatures relative to ultrapure water.

Background

- Atmospheric PM is comprised of dust, soot, pollen, bacteria, and fungi less 10µm in diameter.
- PM that function as ice nuclei are integral to cloud formation, precipitation production, have potential for applications in and agriculture and climate manipulation.

Beach Sand Grain



Methods

- Aerosols were collected via roof-mounted inlets on quartz filters at regulated airflow in long (7 day) and short (2 day) exposures.
- Samples were cultured on LB agar, isolated, and amplified for a region of the 16S ribosomal subunit. DNA was then sequenced characterized against the NCBI and database via BLAST searches.
- Ice nucleation tests were performed on all identified bacteria cultures relative to ultrapure water.

Analysis of Bioaerosols: DNA and Ice Nucleation Potential

Julie Hibarger, Dr. Rebecca Simmons¹ & Dr. David Delene² 1. University Of North Dakota Department of Biology, 2. University of North Dakota Department of Atmospheric Science

Aerosol Collection

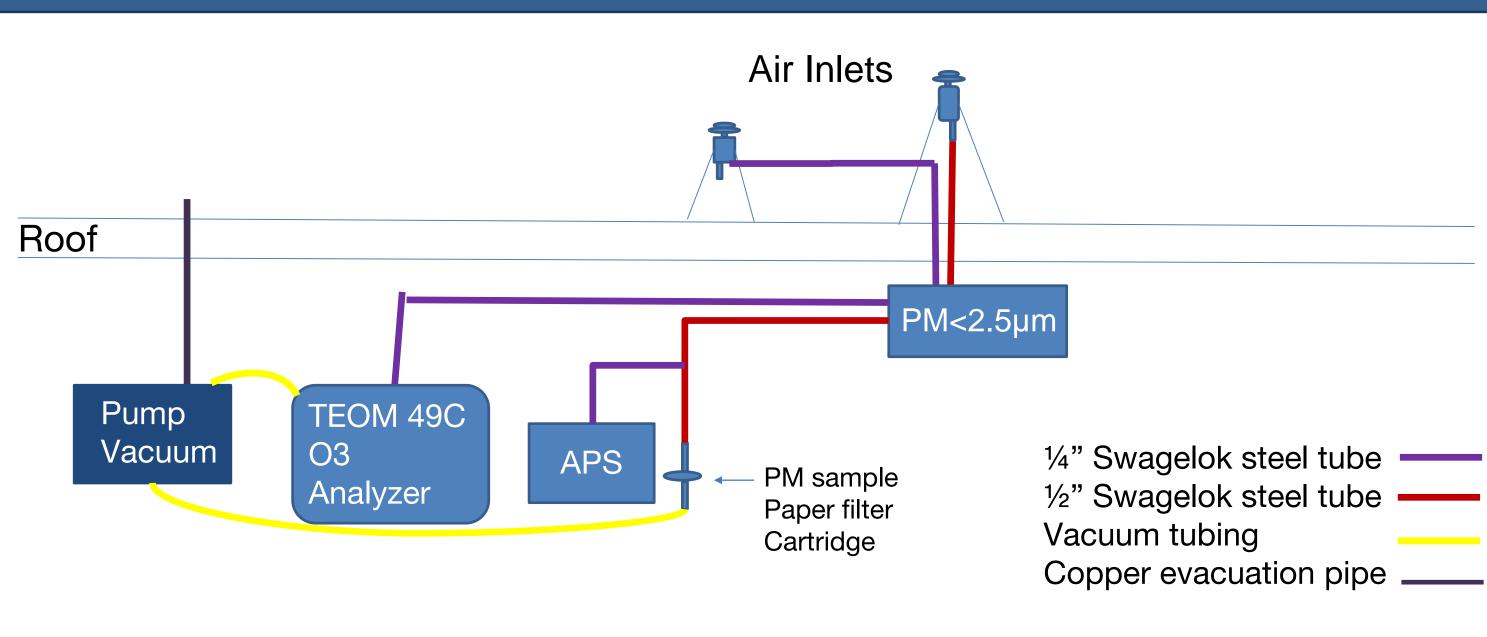


Figure 1: Bioaerosol : Collection of PM less than 2.5 µm

Results

Table 1: Bacteria Identification: BLAST results showing the closest matching species and percent sequence identity

Culture Number	Closest Match	Sequer
Culture A1	Bacillus aquimaris	1
Culture A2	Erwinia billingiae	
Culture B1	Bacillus pumilus	
Culture B2	Bacillus pumilus	
Culture B3	Bacillus pumilus	

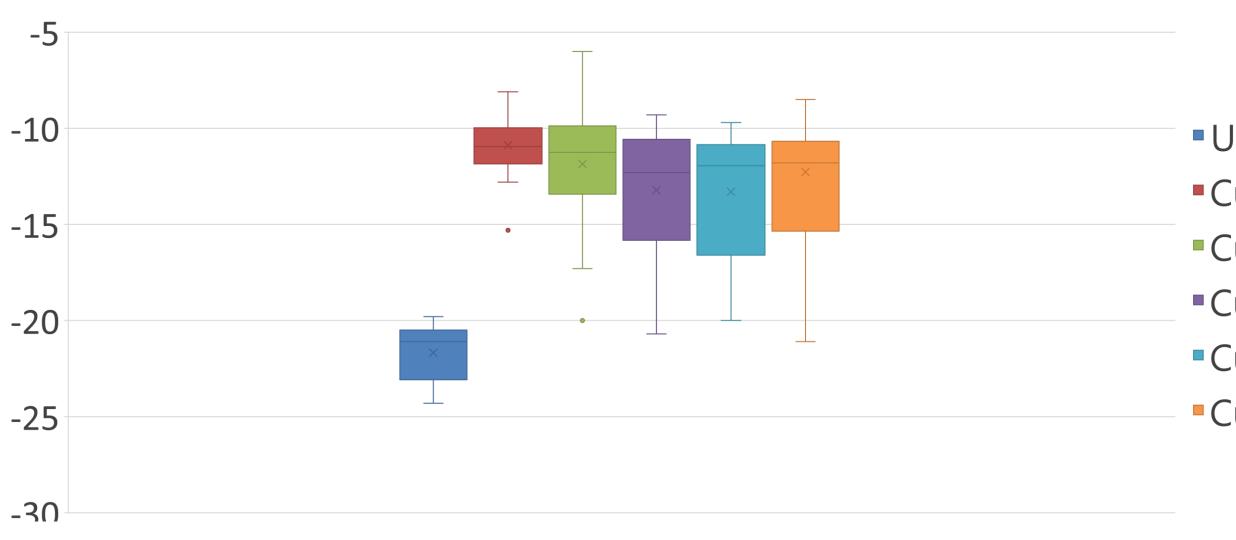
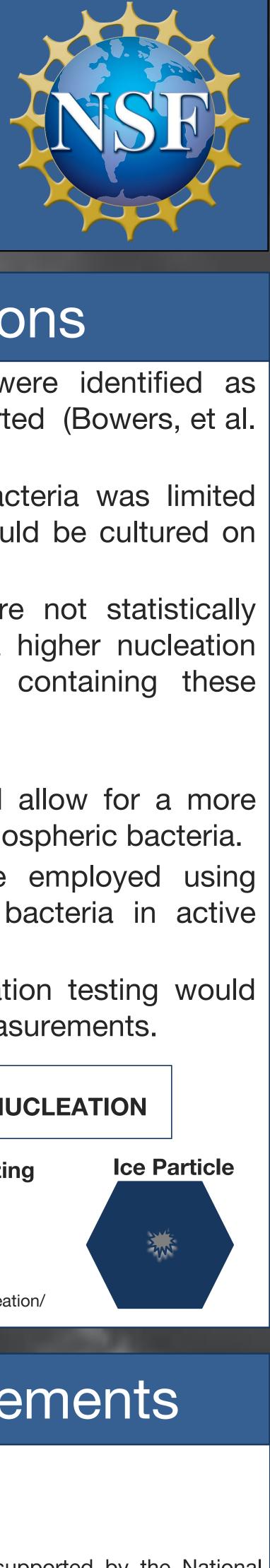


Figure 2: Temperature at ice nucleation (C°). Samples with cultured bacteria formed ice nuclei at slightly higher temperatures.



\frown	•
$($: $\cap n \cap l$	lusions
	USICIIS

- Most bacterial samples were identified as Bacillus as previously reported (Bowers, et al. 2013).
- Unfortunately, sampled bacteria was limited to aerobic species that could be cultured on LB agar.
- Ice nucleation results were not statistically significant but do imply a higher nucleation temperature for samples containing these bacteria.

FUTURE WORK:

- Illumina sequencing would allow for a more comprehensive view of atmospheric bacteria.
- These methods could be employed using aerial sampling of cloud bacteria in active storm conditions
- More sophisticated nucleation testing would improve consistency in measurements.

MECHANICS OF ICE NUCLEATION

Solid Particle

Deposition Freezing





Redrawn from http://bertram.chem.ubc.ca/ice-nucleation/

Acknowledgements

- Dr. Frank Bowman
- Dr. Brian Darby
- Nickolas S. Goodman, M.S.
- Matthew Flom, M.S.
- This material is based upon work supported by the National Science Foundation under Grant No. CHE-1757922. Any opinions, findings, and conclusions or recommendations expressed in this material are those of the authors and do not necessarily reflect the views of the National Science Foundation.

References

Robert M. Bowers, Nicholas Clements, Joanna B. Emerson, Christine Wiedinmyer, Michale P. Hannifan, and Noah Fierer. Seasonal Variability in Bacterial and Fungal Diversity of the Near Surface Atmosphere. Environmental Science and Technology. 2013. 12097-12105.

nce Identity

- 100%
- 99%
- 99%
- 99%
- 99%

Ultrapure Water (n=30)

- Culture A1 (n=30)
- Culture A2 (n=30)
- Culture B1 (n=30)
- Culture B2 (n=30)
- Culture B3 (n=30)

