

A comprehensive climatology of Arctic aerosol properties on the North Slope of Alaska

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Aerosol Measurements at Barrow

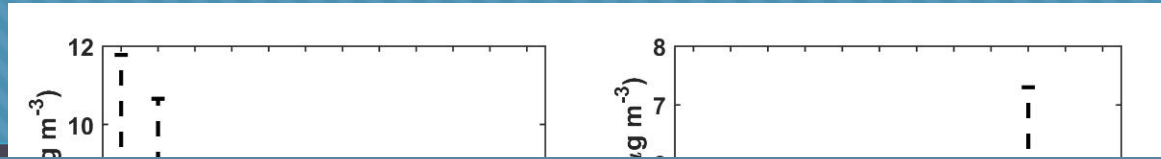
| Measurement | 1976 | 1977 | 1978 | 1979 | 1980 | 1981 | 1982 | 1983 | 1984 | 1985 | 1986 | 1987 | 1988 | 1989 | 1990 | 1991 | 1992 | 1993 | 1994 | 1995 | 1996 | 1997 | 1998 | 1999 | 2000 | 2001 | 2002 | 2003 | 2004 | 2005 | 2006 | 2007 | 2008 | 2009 | 2010 | 2011 | 2012 | 2013 | 2014 | 2015 | | | | | | | | |
|---|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|------|--|--|--|--|--|--|--|--|
| Particle number concentration (SMPS, 1 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Particle number concentration (CPC, 10 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| CCN concentration (10 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Particle size distributions (SMPS, 1 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Absorption coefficient (PSAP, 1 $\lambda_{550 \text{ nm}}$, 1 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Absorption coefficient (PSAP, 1 $\lambda_{550 \text{ nm}}$, 10 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Absorption coefficient (PSAP, 3 λ , 1 and 10 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Scattering coefficient (neph, 3 λ , 1 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Scattering coefficient (neph, 3 λ , 10 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Aerosol optical depth (AOD; CSPHOT) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Aerosol backscatter, extinction (HSRL) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| Soluble ion chemistry (1 and 10 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
| X-ray analysis (chemistry, 1 μm) | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |

Other measurements exist from field campaigns, these are continuous and long term. Data available through NOAA GMD and DOE ARM.

Background on Arctic Aerosols at Barrow

- Polissar et al.: **CN, scattering, AOD**
 - 1977-1994
- Quinn et al.: **chemical and optical properties**
 - 1997-1999
- Iziomon et al.: **size and absorption**
 - 1998-2003
- Quinn et al.: **chemistry**
 - 1976-1977; 1997-2008
- **Pollution aerosols and submicron sea salt in winter/spring**
- **Supermicron sea salt, biogenic emissions, and CN high in summer, with some influence from midlatitude fires**
- **These studies used some combination of measurements, but not all that are currently available**
- **Aerosol climate impacts vary depending on properties, thus it is important to look at chemistry, size, number, mass and optical properties**

Seasonal Trends: Mass vs. Number



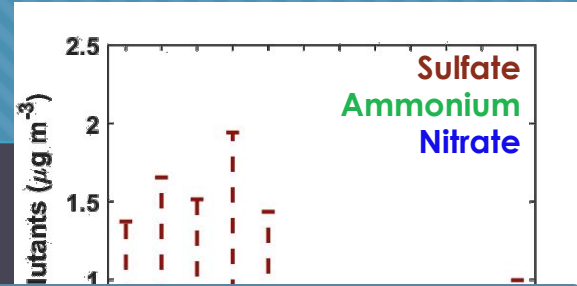
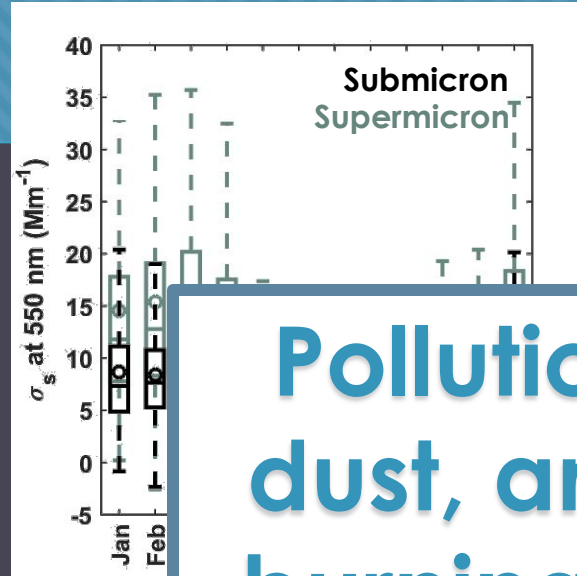
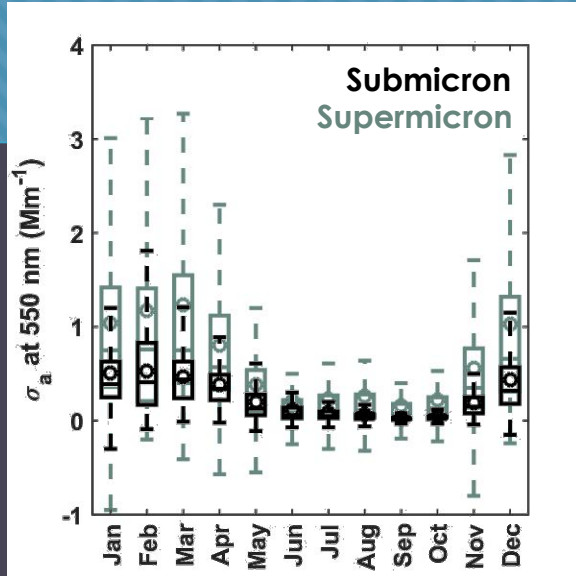
- **Number and mass follow very different seasonal trends.**
- **Mass: important for deposition on sea ice.**
- **Number: important for clouds and radiation.**

- **Submicron mass** high during haze
 - Pollutants, dust, biomass burning, SS
- **Submicron number** highest in spring into summer
 - Suggests the spring haze is worse than the winter (climate impact in spring)
 - Due to small transported haze particles AND biogenic

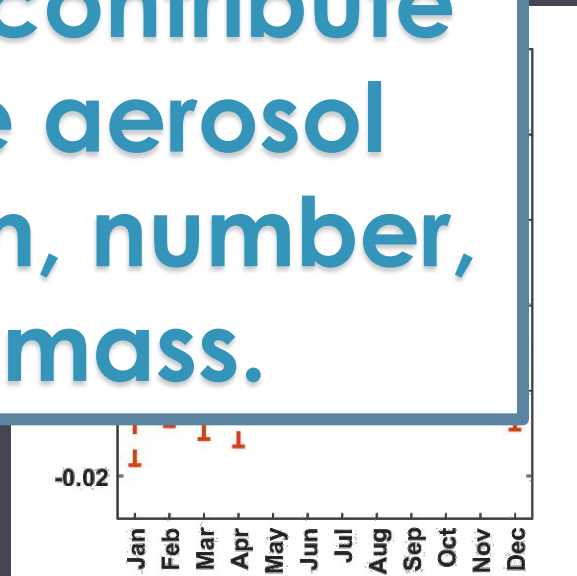
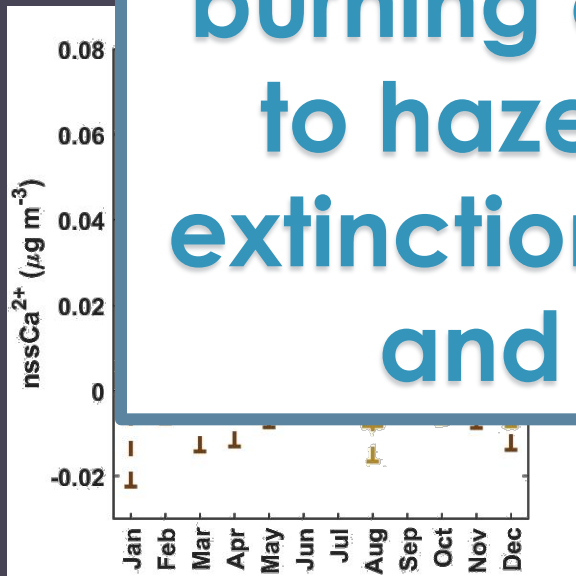
- **Supermicron mass** highest in fall, secondary peak in spring
 - SS in fall, dust in spring
- **Supermicron number** highest in late summer and during haze
 - SS starts around here... why does not continue into fall?
 - Dust in spring

Ja Fe M Apr Ma Ju J Au Se O Nc De Ja Fe M Apr Ma Ju J Au Se O Nc De

Seasonal Trends: "Haze" Aerosols



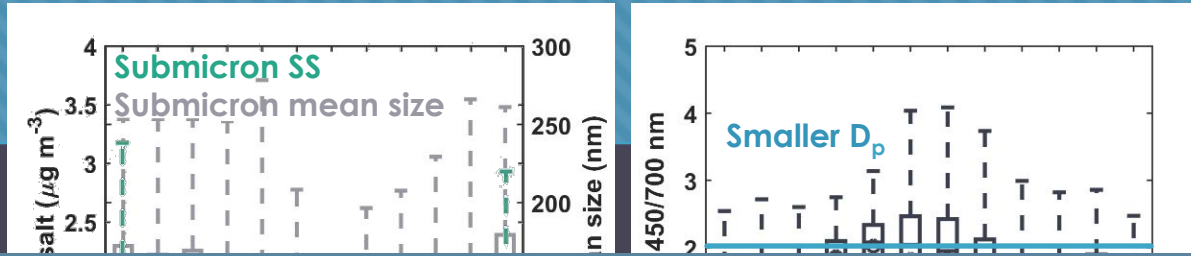
Pollution, mineral dust, and biomass burning contribute to haze aerosol extinction, number, and mass.



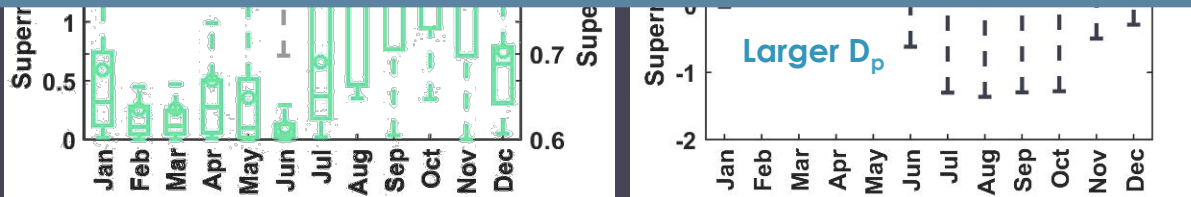
- **Pollutants** highest during haze, concurrent with σ_a and σ_s
- **Biomass burning** markers highest depending on size
 - Submicron peak in winter
 - Supermicron peak in late fall/winter

- **Mineral dust** markers highest in spring, earlier than midlatitudes

Seasonal Trends: Sea Salt Aerosols



- Small sea salt indicate transport with pollutants.
- Larger sea salt generated mechanically during open ocean, contribute to high number and mass.



- Submicron sea salt (SS) highest in winter, concurrent with large submicron mean size

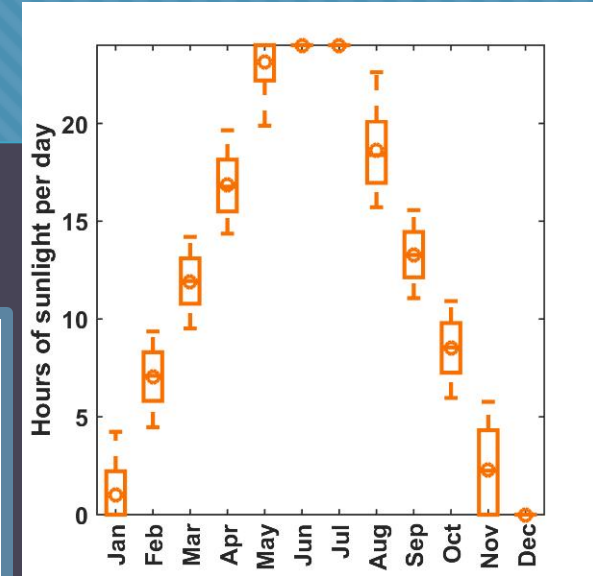
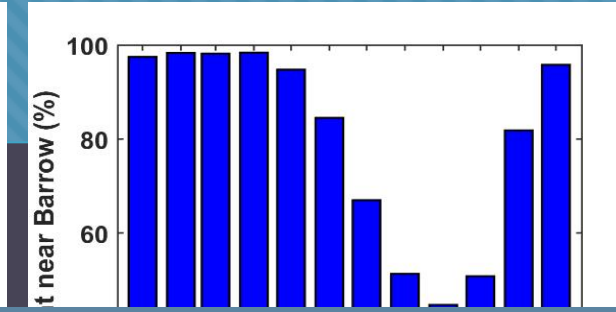
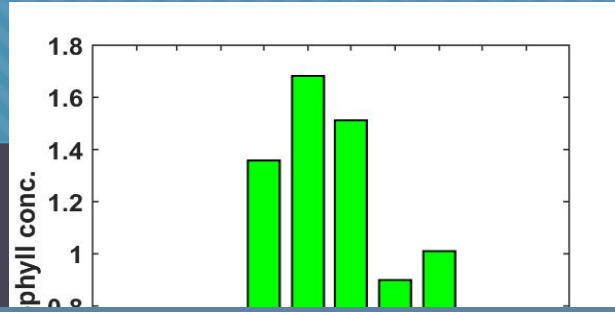
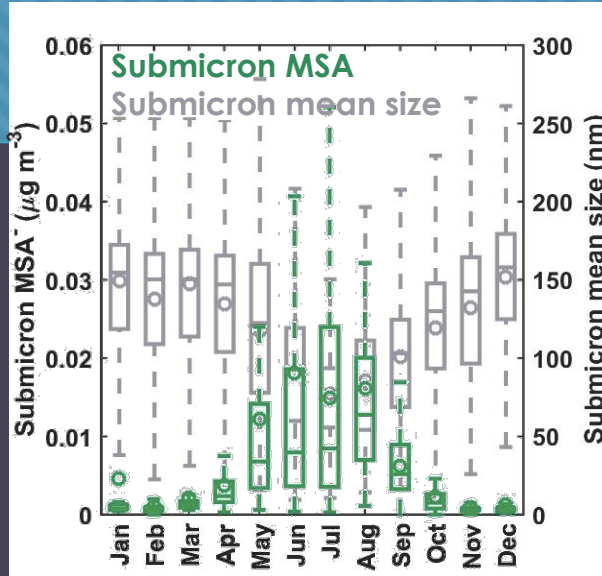
- Suggests LRT

- Supermicron sea salt (SS) highest in fall, when open water is exposed concurrent with large and most certain single-scattering albedo (ω_o)

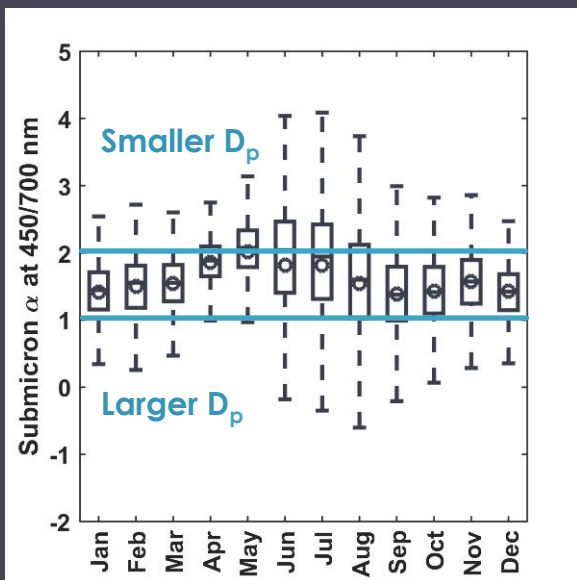
- Also concurrent with lowest submicron Angström Exponent (α) and larger D_p

- Also concurrent with lowest supermicron Angström Exponent = larger D_p

Seasonal Trends: Biogenic Aerosols



Biogenic activity in summer with most sunlight, creating large number concentrations of small particles.



- sea ice is diminishing, exposing open water
- Sunlight is available for biological production
- Smallest mean sizes are observed

Conclusions

○ Haze (winter + spring) = ↑ extinction, ↑ pollutants, dust, biomass burning, submicron sea salt, ↑ submicron number, ↑ submicron mass, ↑ submicron size

= LRT influence

○ Summer = ↑ bio emissions (chlorophyll and MSA), ↓ size, ↑ submicron number, ↑ Ångström Exponent

= local secondary emissions once sea surface and sunlight are available

○ Fall = ↑ supermicron SS, ↑ supermicron mass, ↑ supermicron number (late summer/early fall), ↑ single-scattering albedo, ↓ Ångström Exponent

= local mechanical emissions from open ocean surface

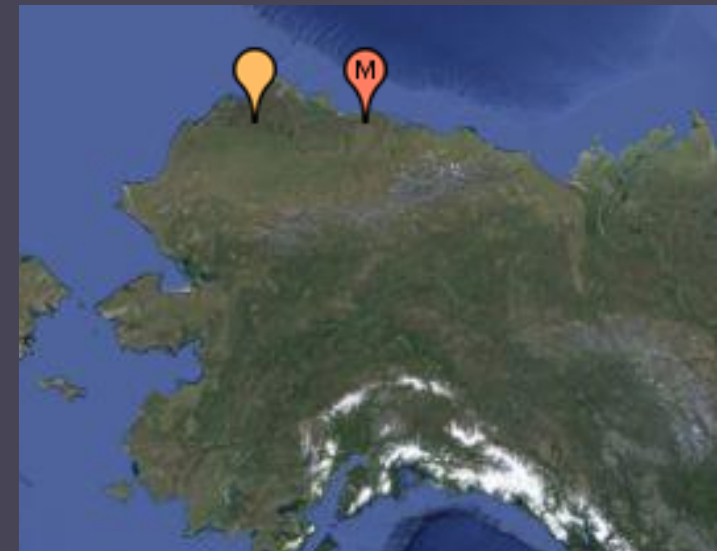
○ The different aerosol sources contribute to number and mass, which have disparate seasonality.

○ **It is important to look at multiple types of measurements to help constrain models.**

Future Directions

- **Next steps:**

- Relate seasonal trends to those in meteorology, dynamics, transport sources
 - Look at increases or decreases over time, do we see any relationships with regulations?
 - Quantitative comparison with previous work
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- Oliktok Point, AK will soon have aerosol measurements, very interesting comparison for Barrow!



Acknowledgements

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- **References:**

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