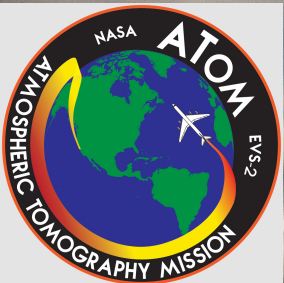



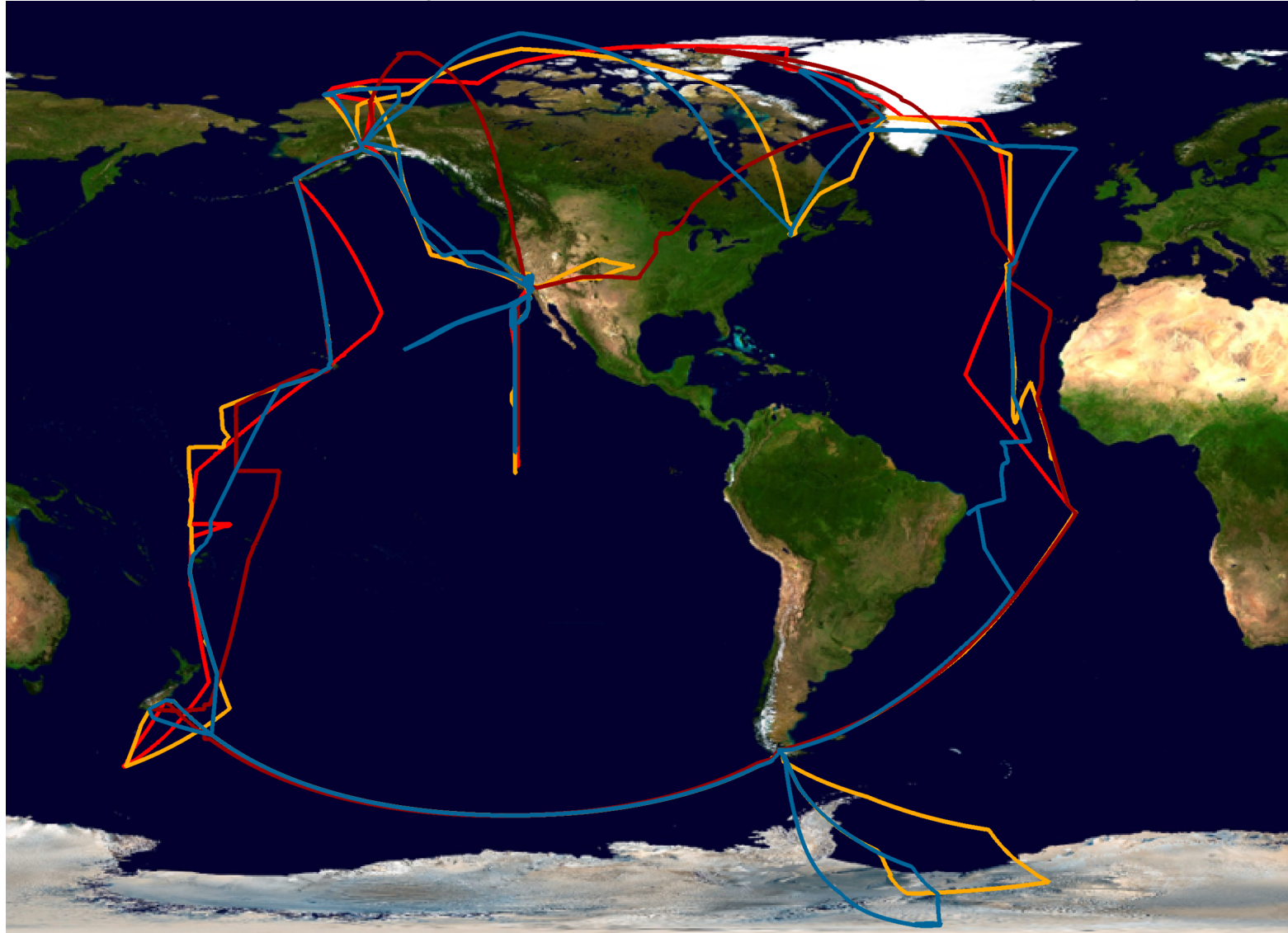
# Global-Scale in-situ measurements of Atmospheric Aerosol from the NASA Atmospheric Tomography Mission



 @chasingcloudsCW

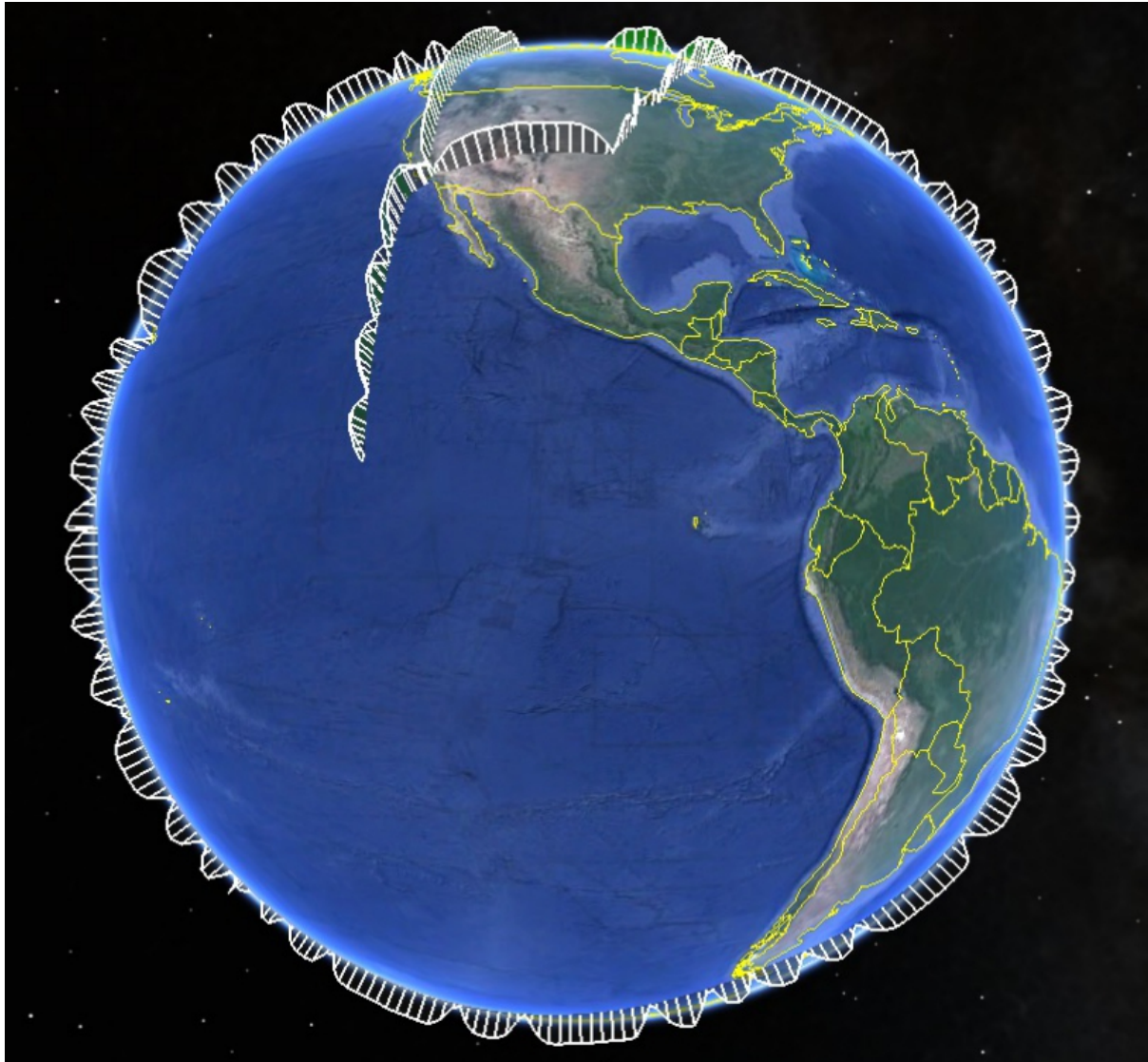
**Christina Williamson**, Hannah Allen, Huingshen Bian, Charles Brock, Taopaul Bui, Pedro Campuzano-Jost, Mian Chin, Simon Clegg, Peter Colarco, John Crouse, Doug Day, Maximillian Dollner, Karl Froyd, Jose Jimenez, Joseph Katich, Michelle Kim, Agnieszka Kupc, Eloise Marais, Daniel Murphy, Ben Nault, Jeffrey Pierce, Karen Rosenlof, Gregory Schill, Jason Schroder, Joshua Schwarz, Simone Tilmes, Kostas Tsigaridis, Pengfei Yu, Bernadett Weinzierl *Photograph: Samuel Hall, NCAR*

# The Atmospheric Tomography Mission (ATom)



■ ATom-1	Aug 2016
■ ATom-2	Feb 2017
■ ATom-3	Oct 2017
■ ATom-4	May 2018

# The Atmospheric Tomography Mission (ATom)



**Vertical Profiling**

~0.2 – 12km

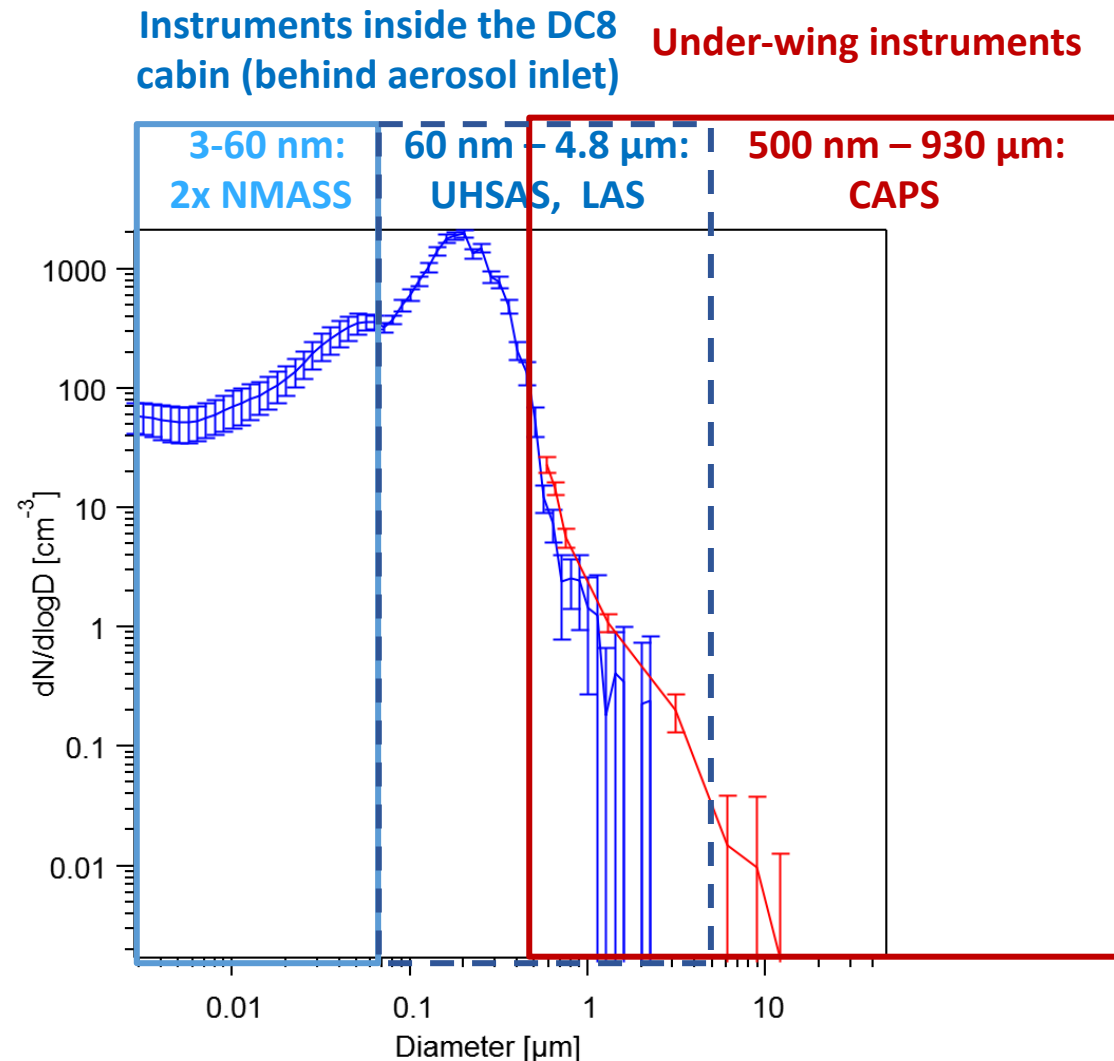
# ATom Aerosol Measurements

- Size Distributions
- Composition
  - Bulk non-refractory (AMS)
  - Bulk (filters)
  - Single Particle (PALMS)
  - Volatility
- Black Carbon (SP2)
- Brown Carbon
- Optical Properties (scattering, extinction)

## Gas Phase:

- Tracers
- VOCs
- GHGs
- NO<sub>x</sub> and NO<sub>y</sub>
- ...

# Aerosol Size Distributions



- Minimizing inlet effects on coarse mode data
- Full size distribution between 3 nm and 930 μm
- Quantification of size distribution uncertainties

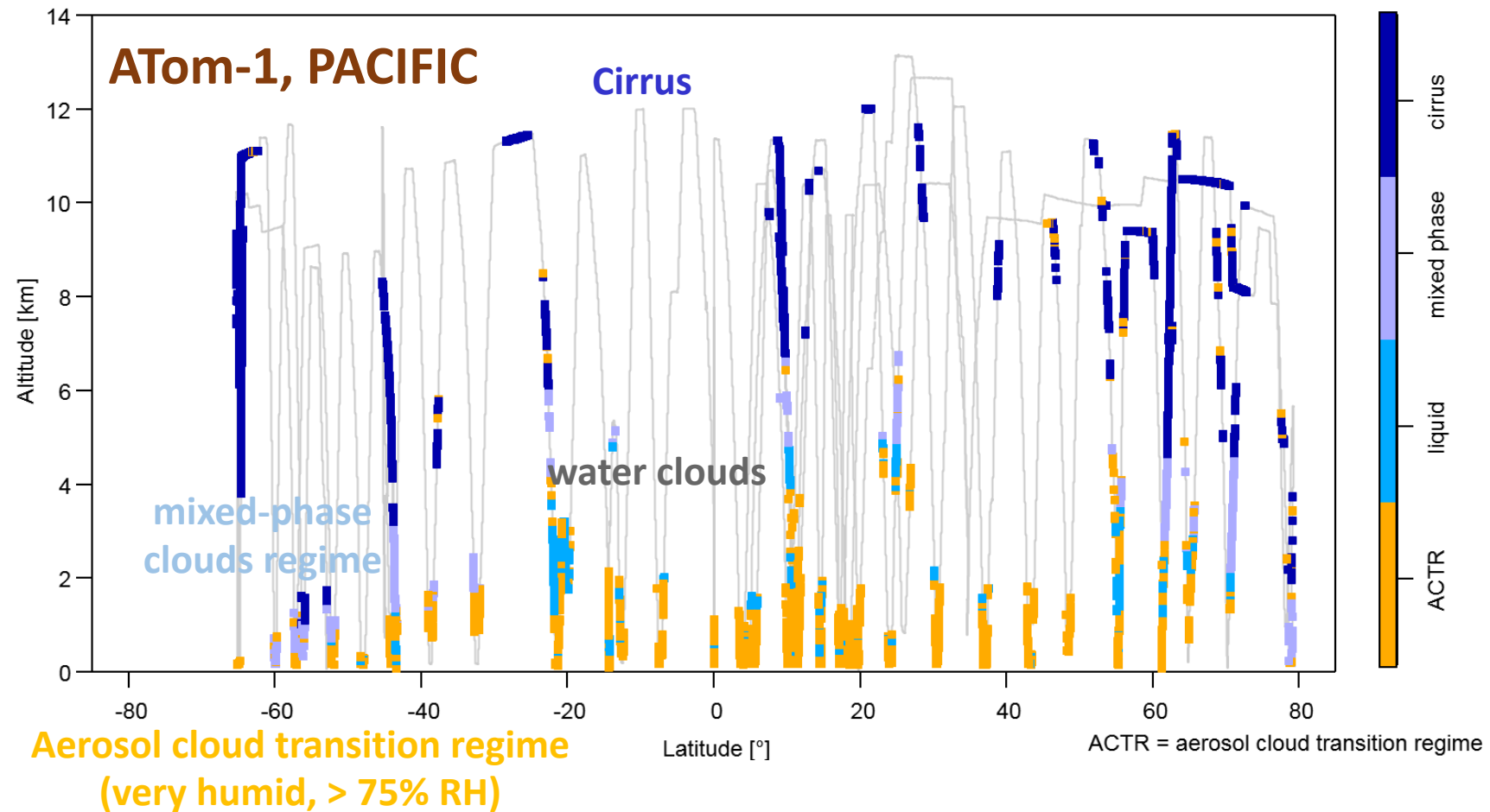
CAPS: **Bernadett Weinzierl** and Maximilian Dollner, University of Vienna

AMP: **Christina Williamson** and Charles Brock, NOAA/CIRES and Agnieszka Kupc, University of Vienna

# Using ATom Data

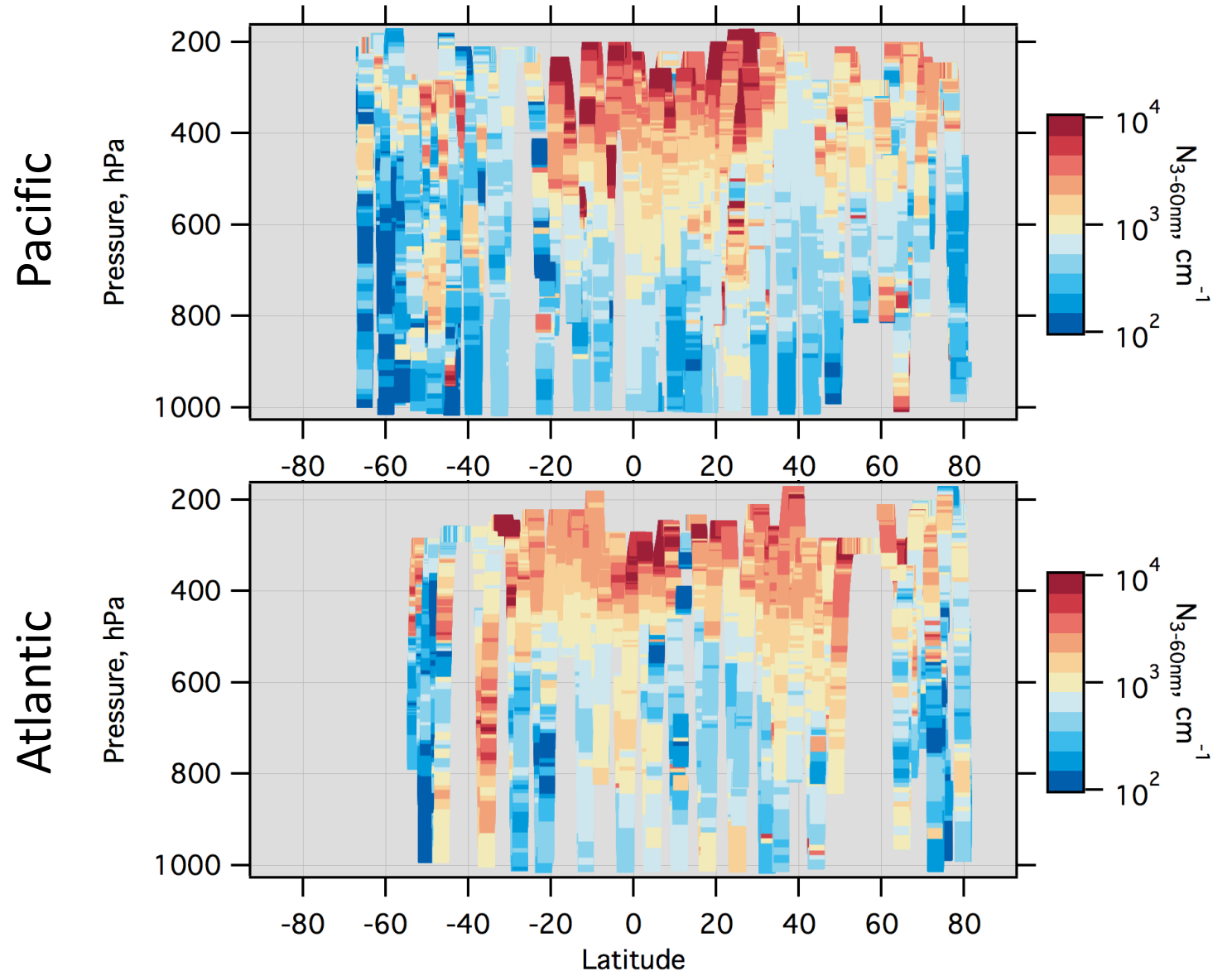
- Data publicly available on the NASA ESPO archive
- Contact and collaboration with instrumentalists (contact details on ESPO archive) strongly advised before beginning a study with any ATom data

# Derivation of cloud phase over global scales



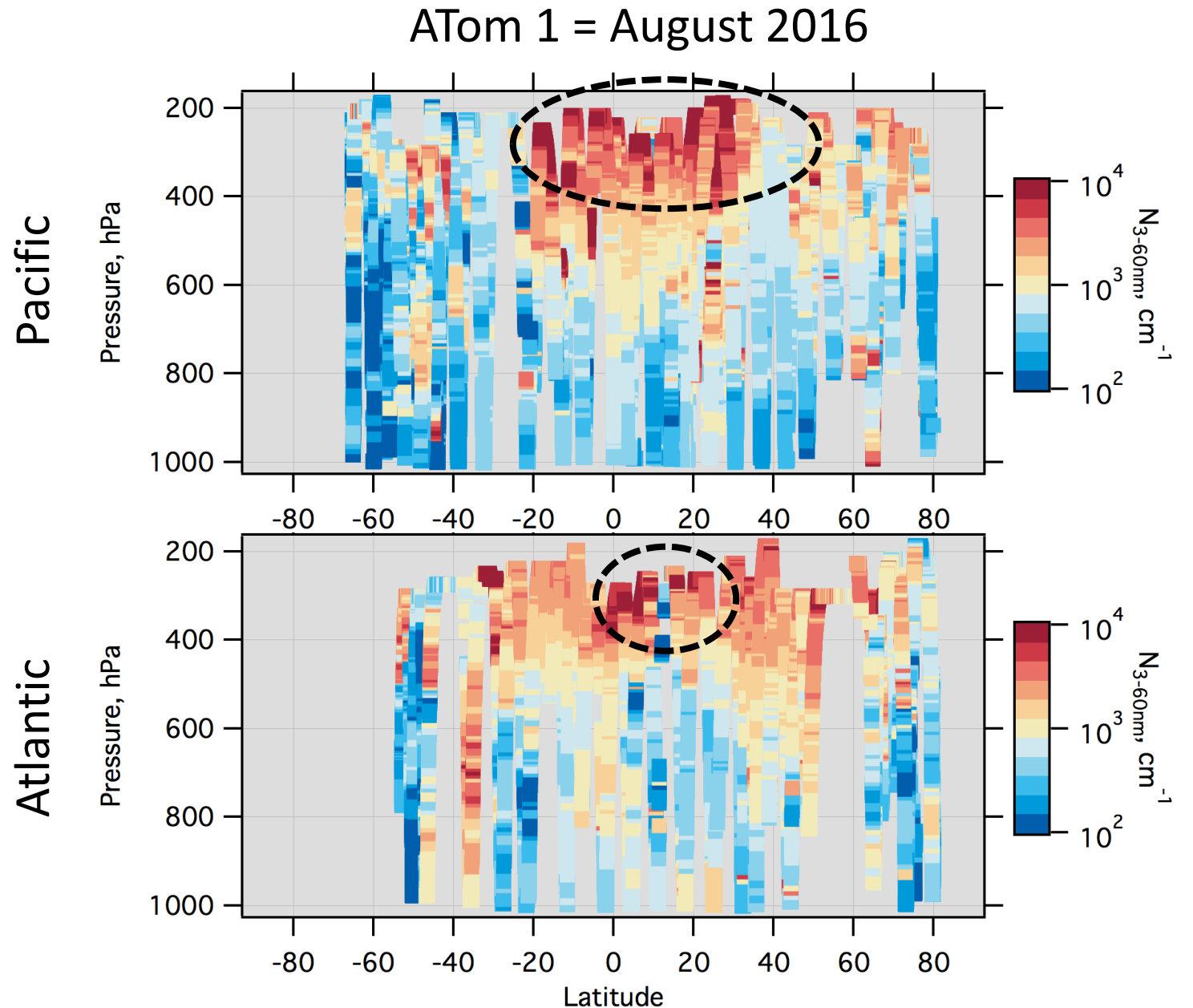
The global distribution of particles from nucleation

ATom 1 = August 2016

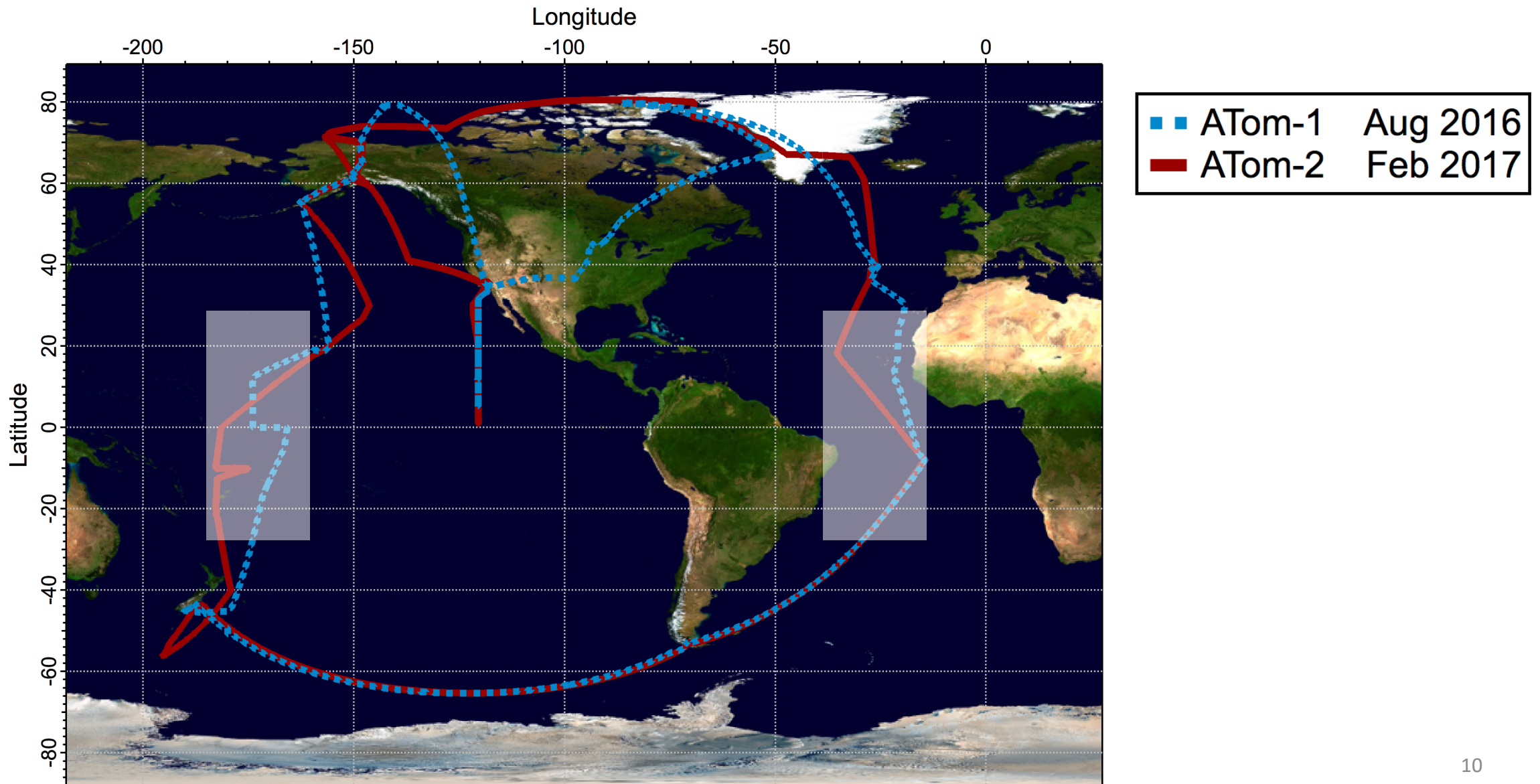




# The global distribution of particles from nucleation

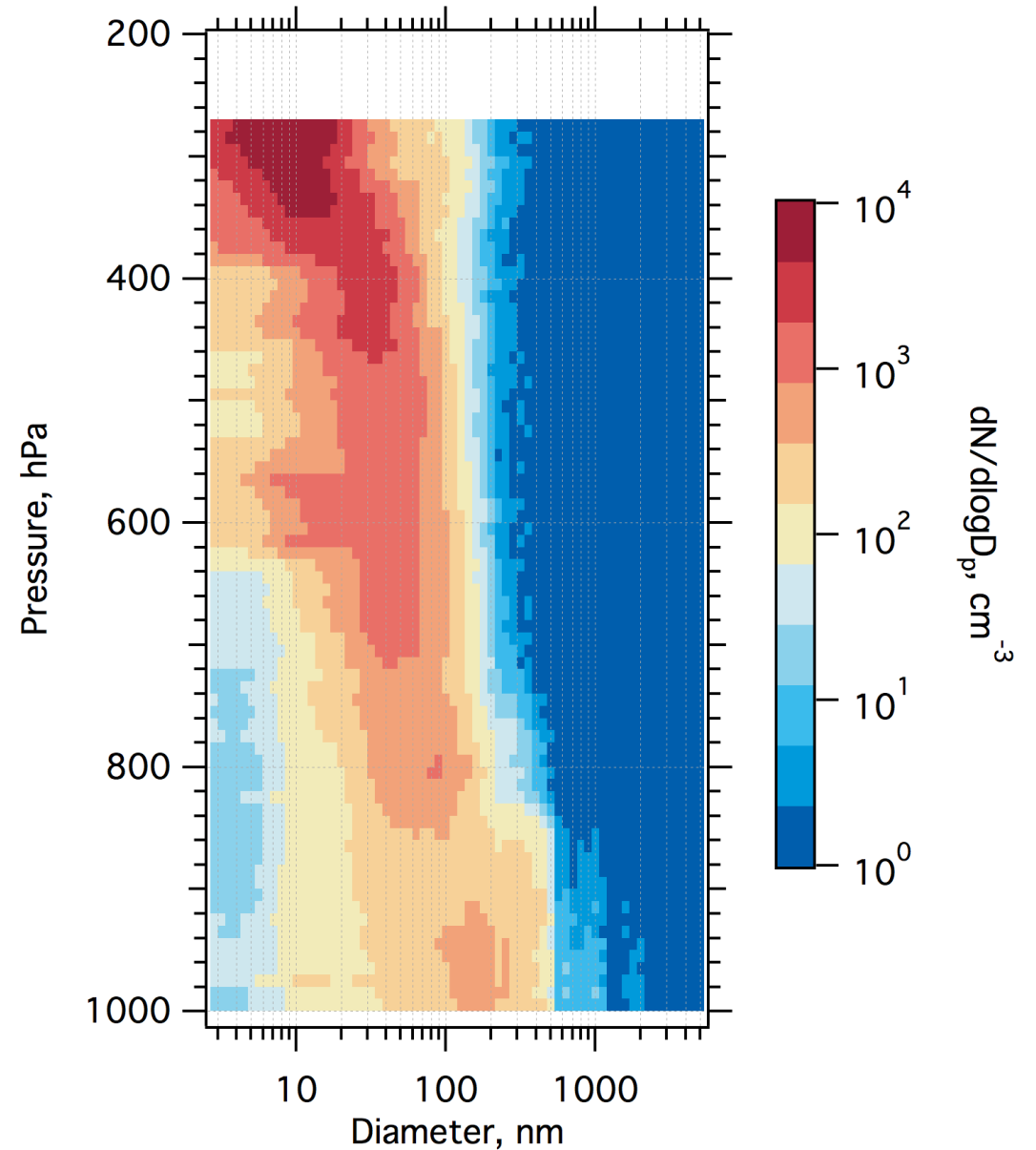


# New Particle Formation and Growth in the Tropics



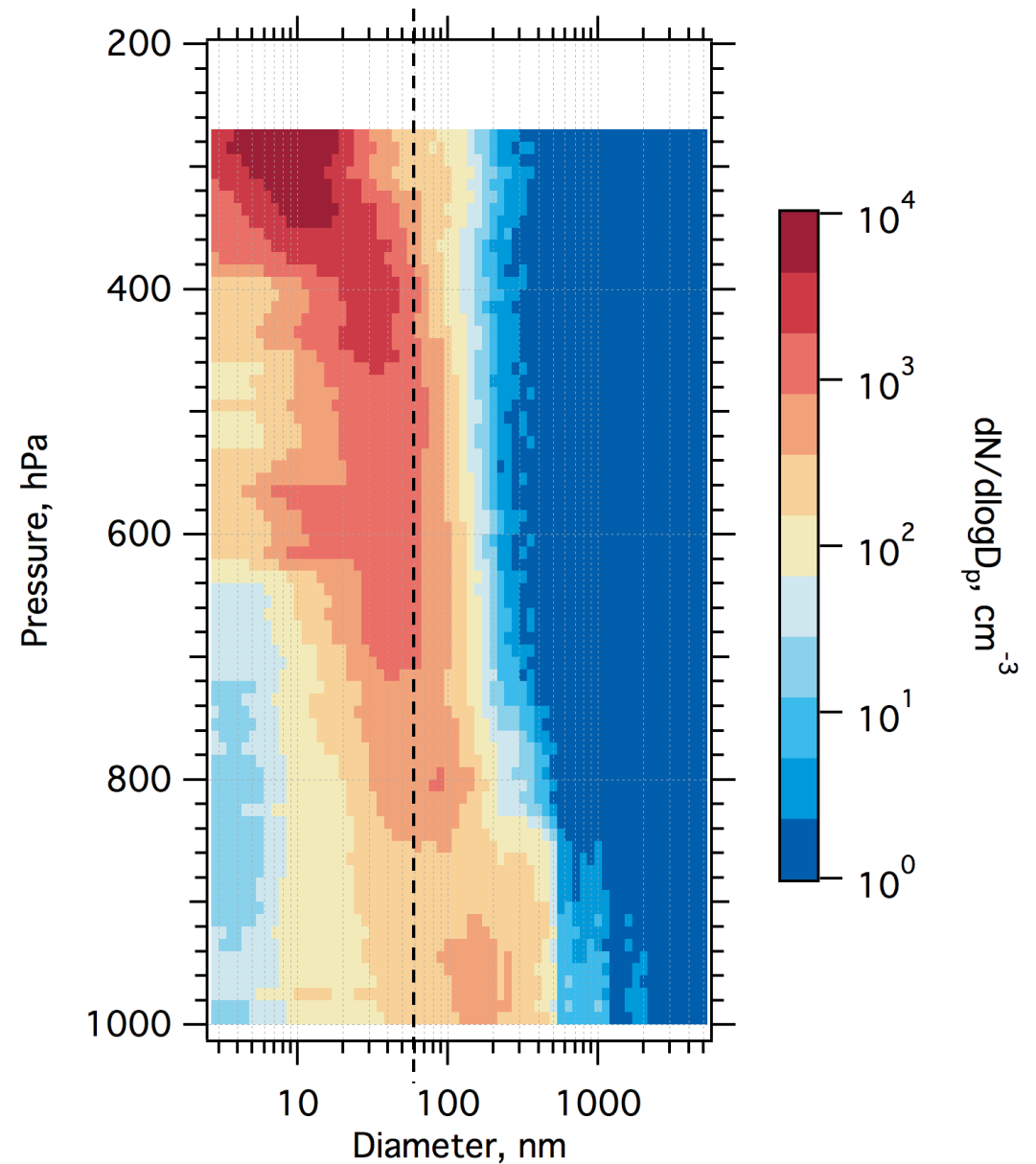
Particles formed in the tropical upper troposphere grow to Cloud Condensation Nuclei sizes on descent to the boundary layer

Tropics, August and February, Atlantic and Pacific



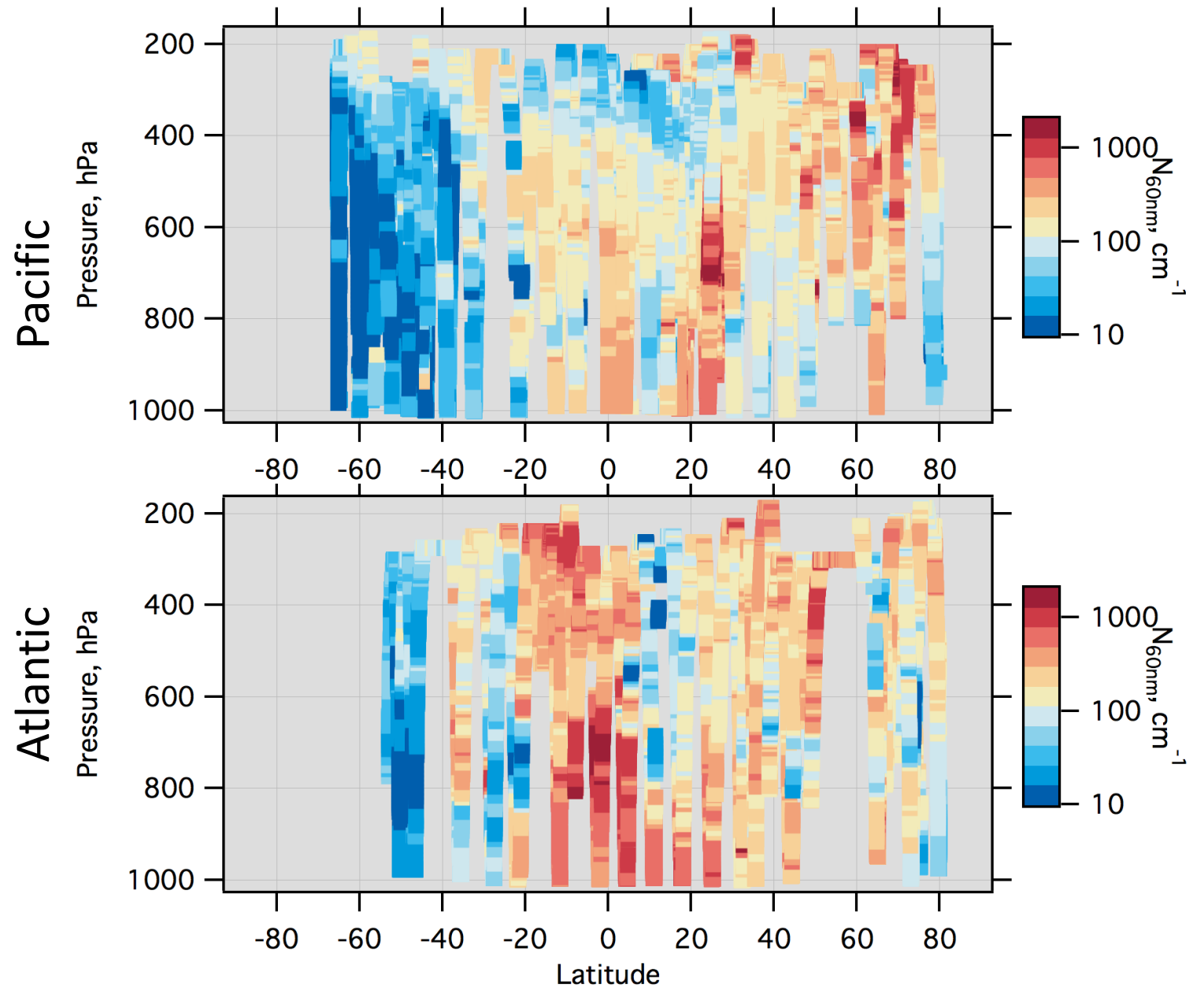
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Tropics, August and February, Atlantic and Pacific



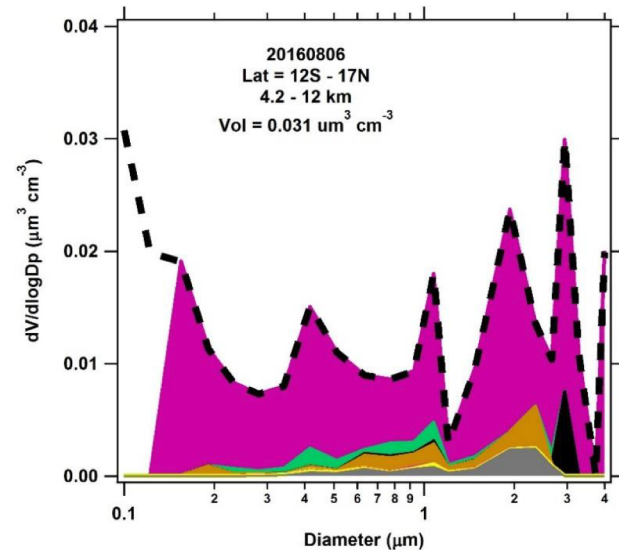
# The global distribution of Cloud Condensation Nuclei-sized particles

ATom 1 = August 2016

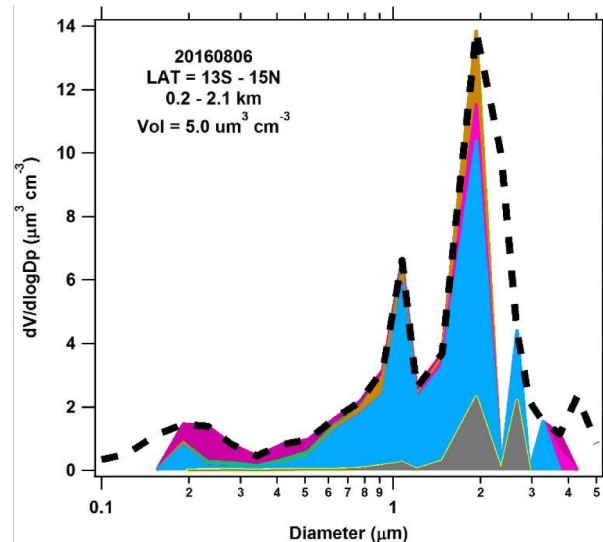


# Origins of Cloud Condensation Nuclei -sized particles

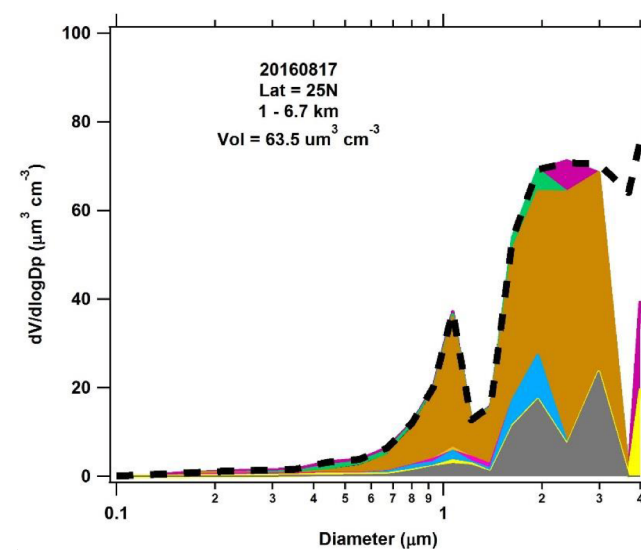
## Sulfate/Organic/Nitrate



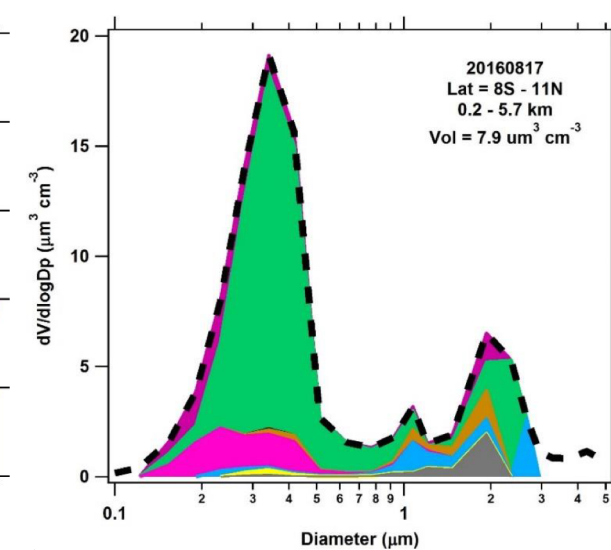
## Sea-Salt



## Mineral Dust



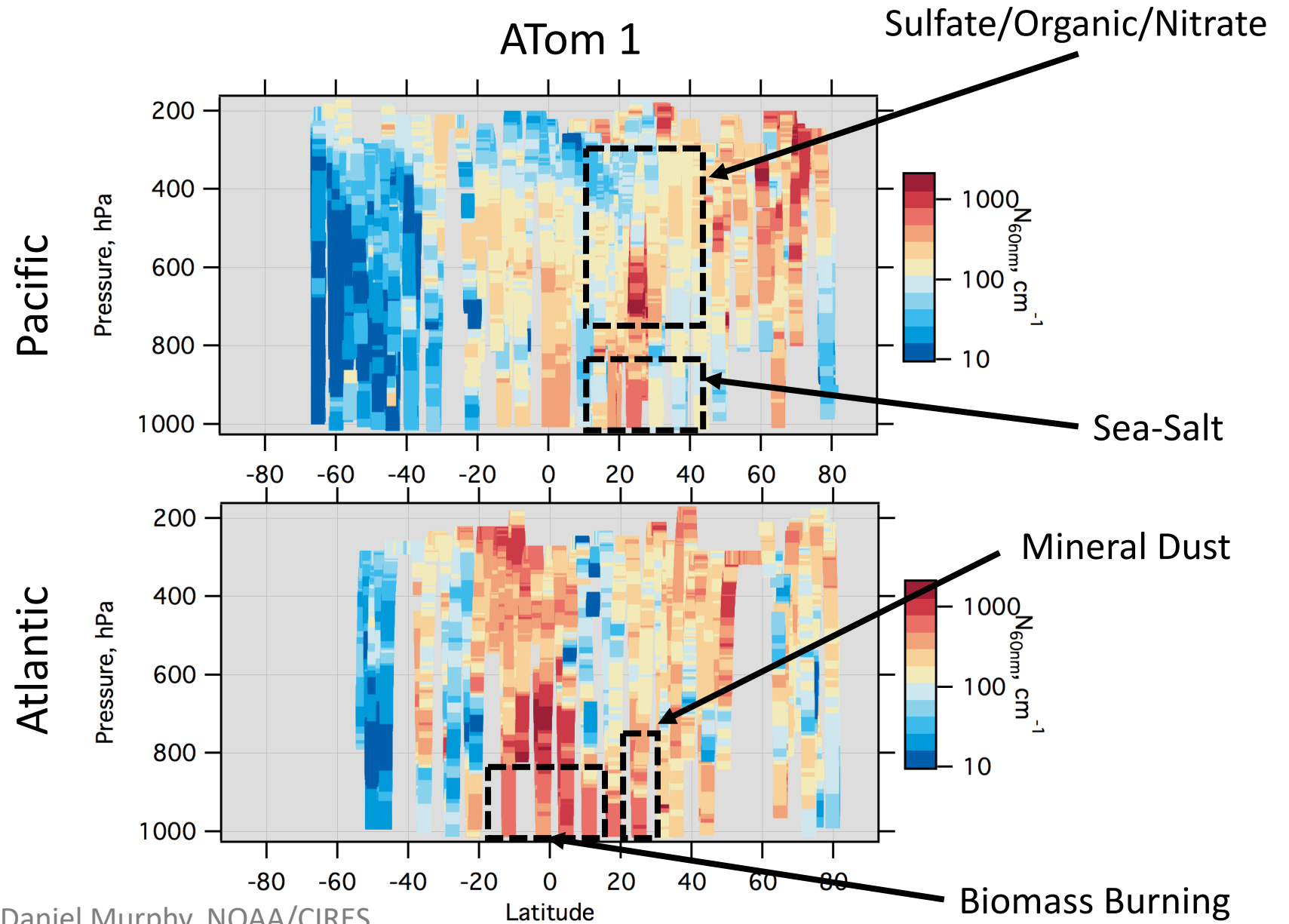
## Biomass Burning



PALMS: Karl Froyd, Gregory Schill, Daniel Murphy, NOAA/CIRES

AMP: Christina Williamson and Charles Brock, NOAA/CIRES and Agnieszka Kupc, University of Vienna 14

# Origins of Cloud Condensation Nuclei-sized particles

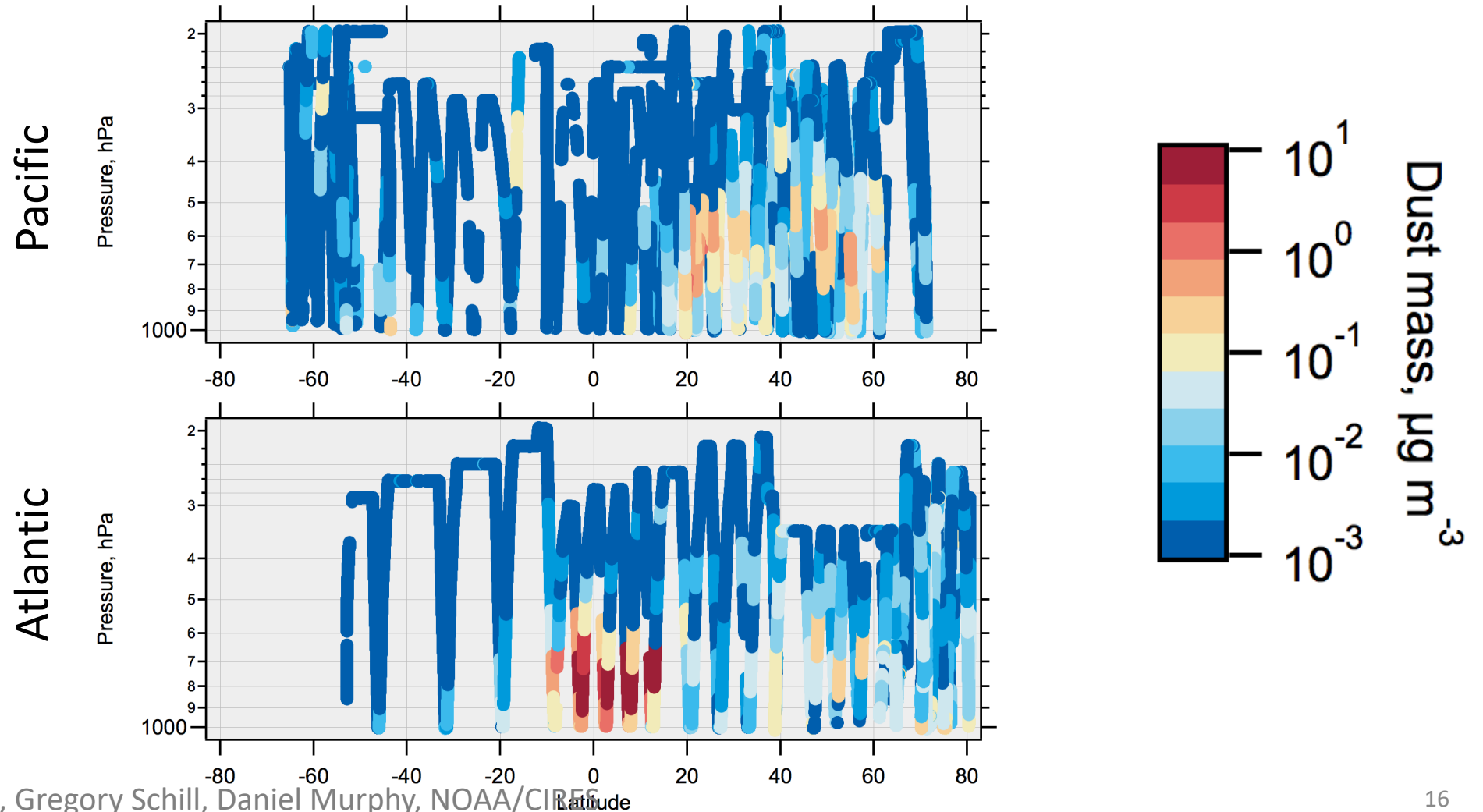


PALMS: Karl Froyd, Gregory Schill, Daniel Murphy, NOAA/CIRES

AMP: Christina Williamson and Charles Brock, NOAA/CIRES and Agnieszka Kupc, University of Vienna

# First global maps of mineral dust by direct measurement

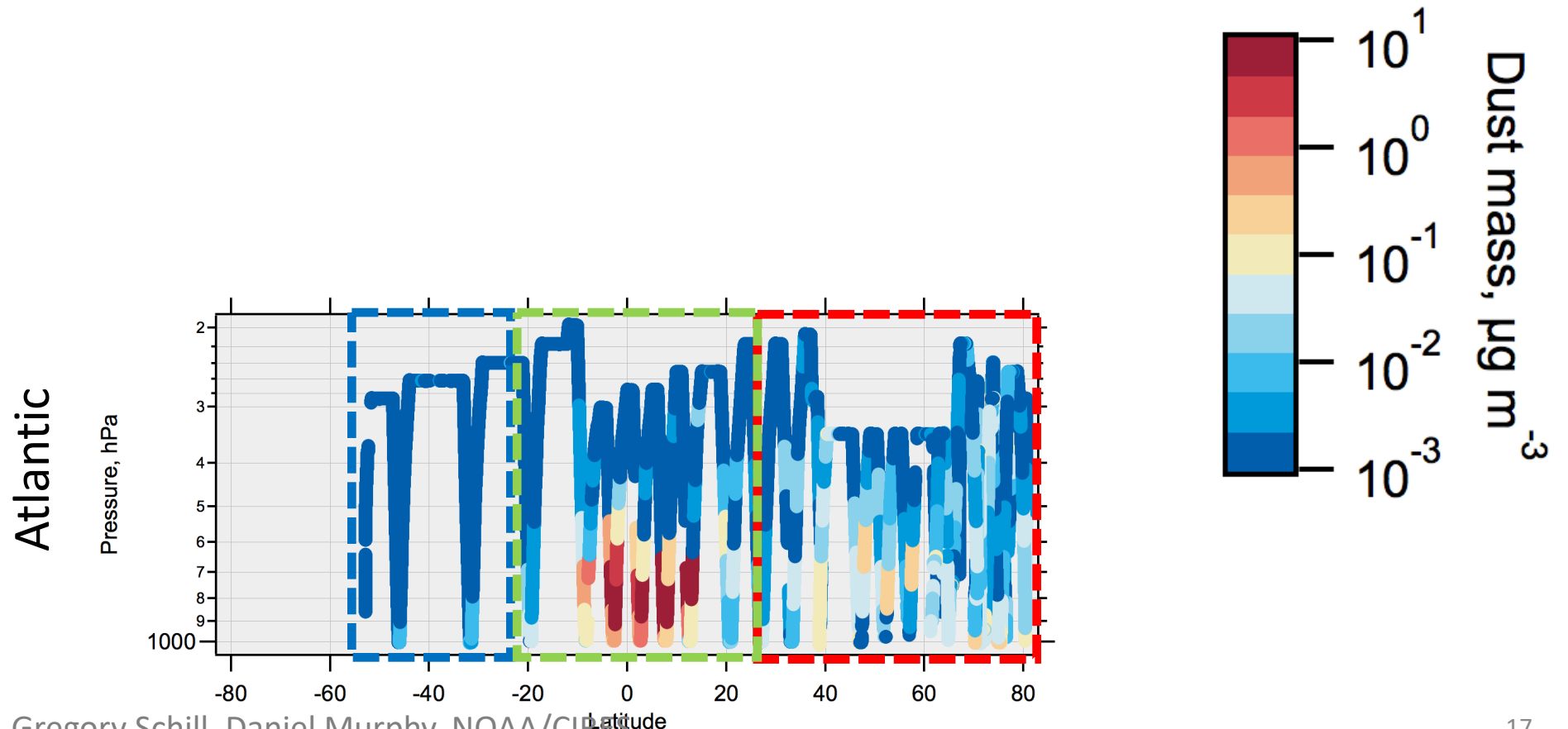
ATom2 – February 2017



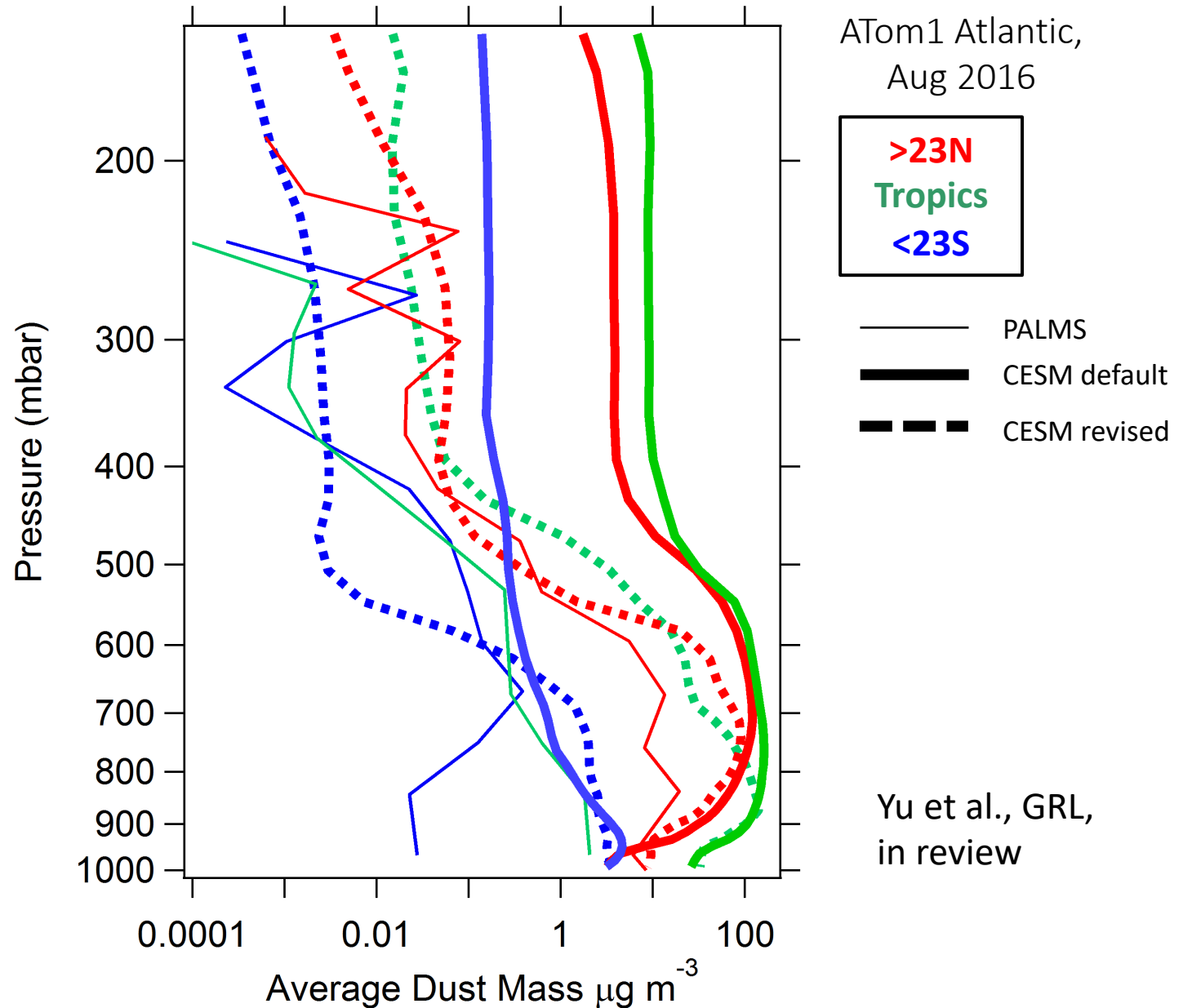


# First global maps of mineral dust by direct measurement

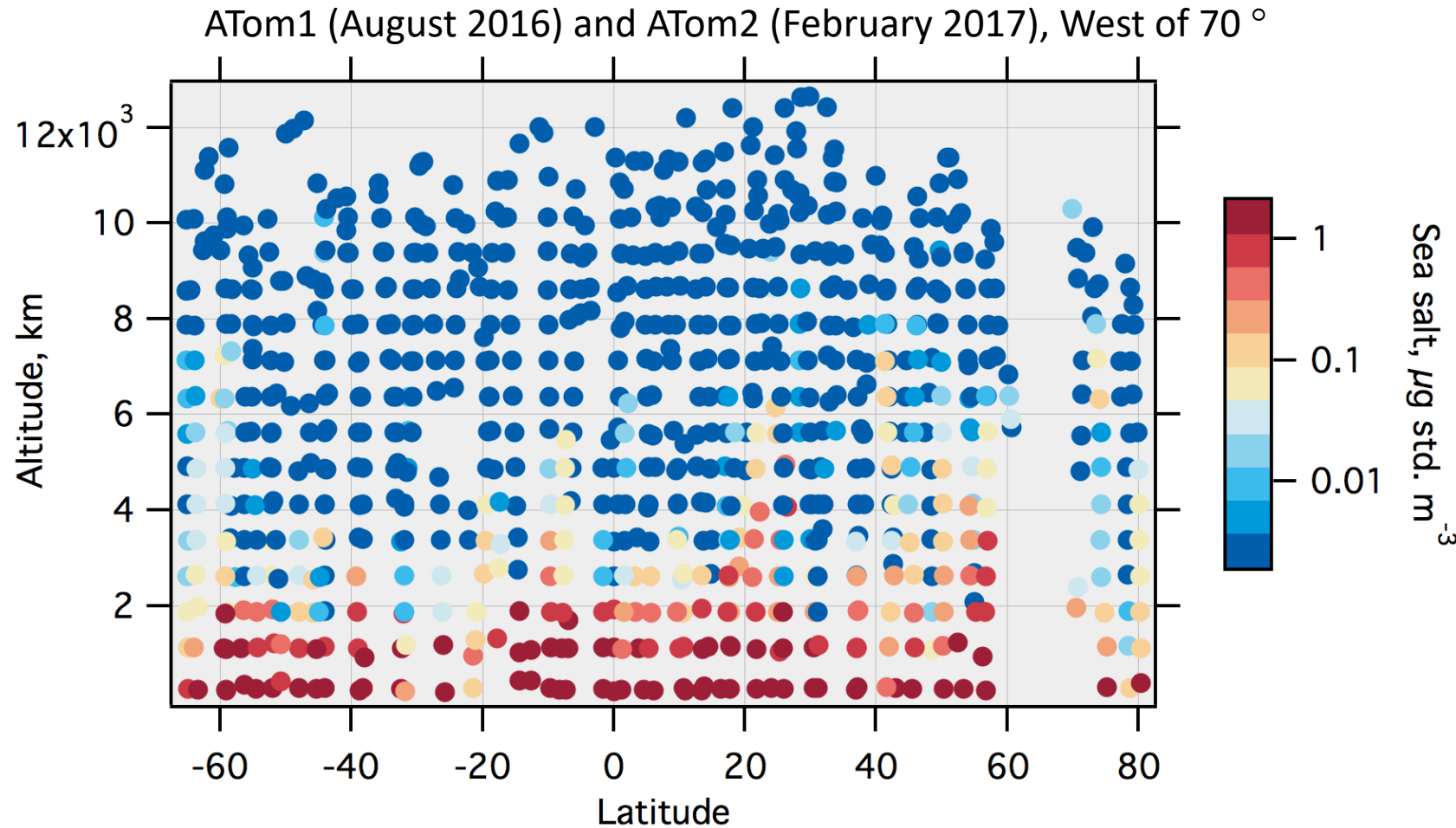
ATom2 – February 2017



Revised  
convective  
removal in  
CESM-CARMA  
better aligns  
modeled dust  
with ATom  
observations

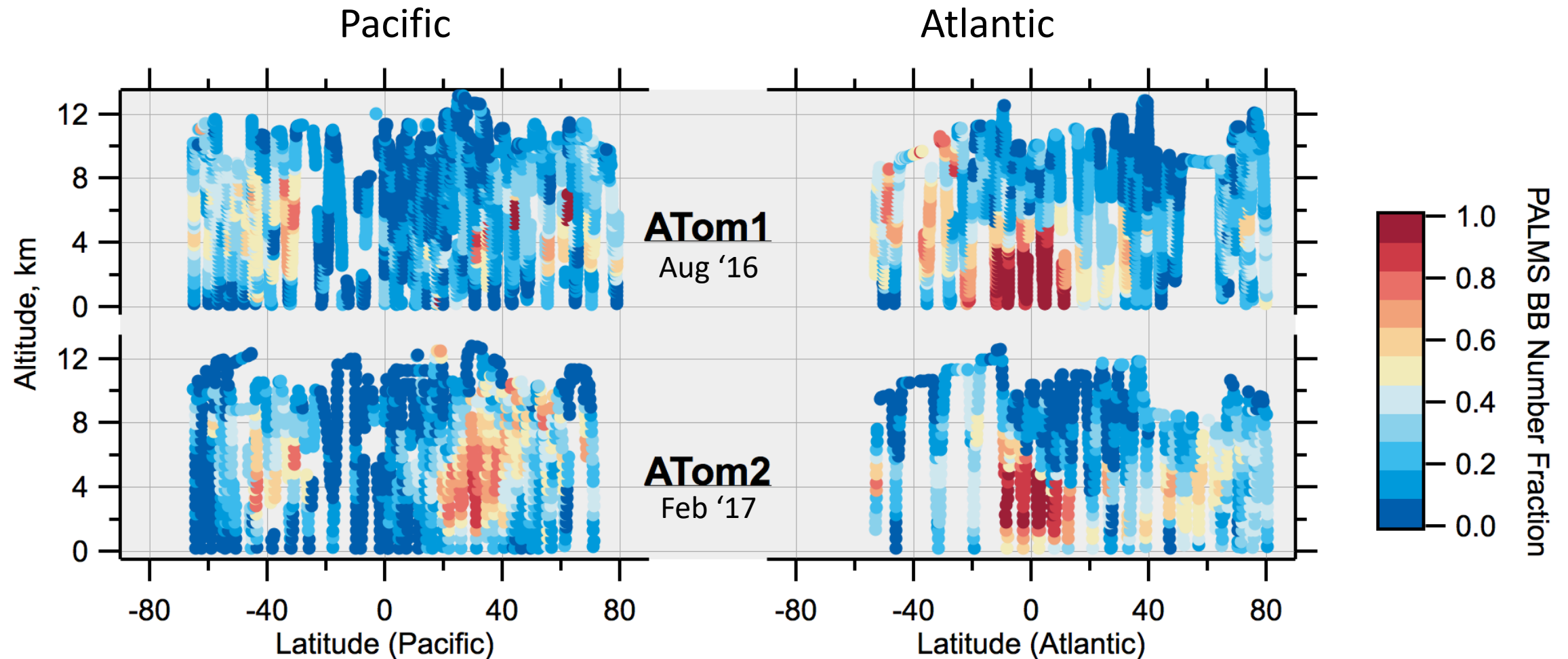


# First data on sea salt above 2 km and over a wide range of latitudes

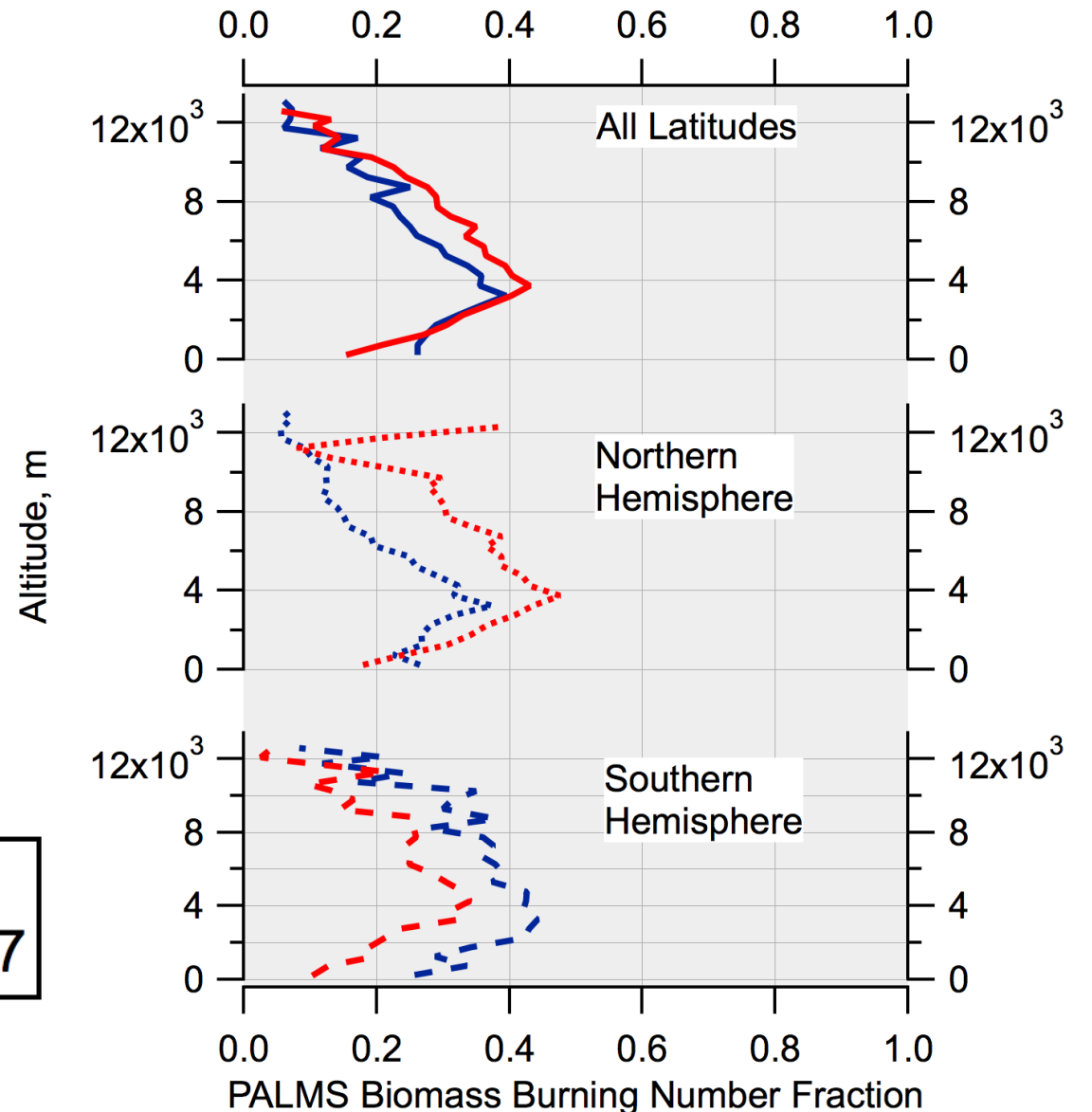
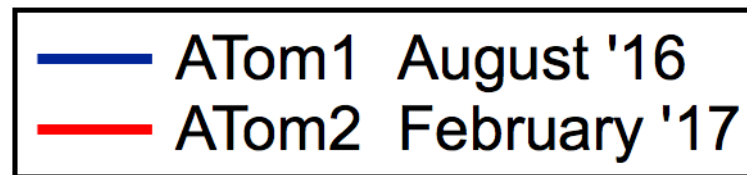


- Extremely low sea-salt ( $\sim 1\text{ppt}$ ) in upper troposphere
- New constraints on halogen chemistry

# Biomass Burning can dominate accumulation mode aerosol number, even far from sources



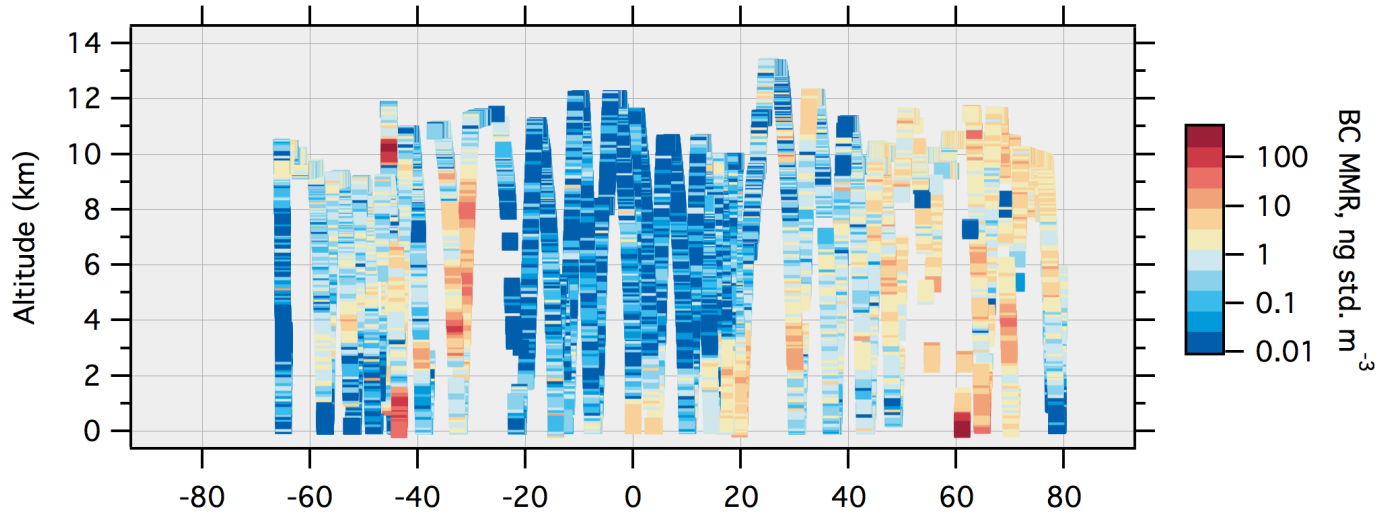
Fraction of accumulation mode aerosol from Biomass Burning is greater in each hemisphere during winter



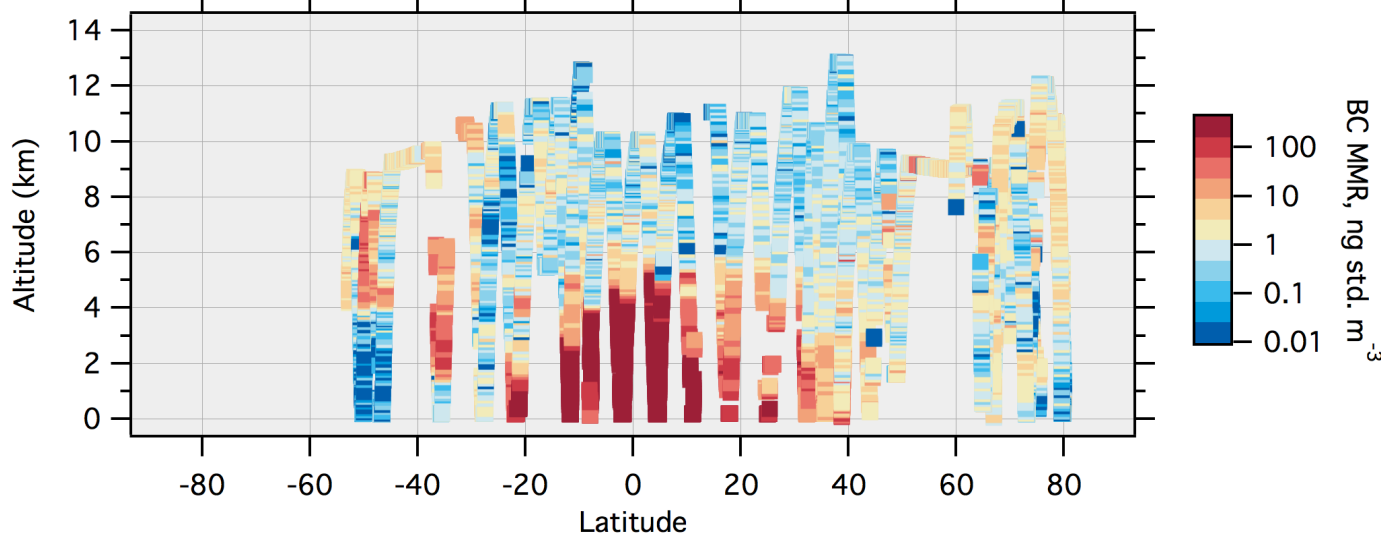
# Global Distribution of Black Carbon

ATom 1  
Aug '16

Pacific



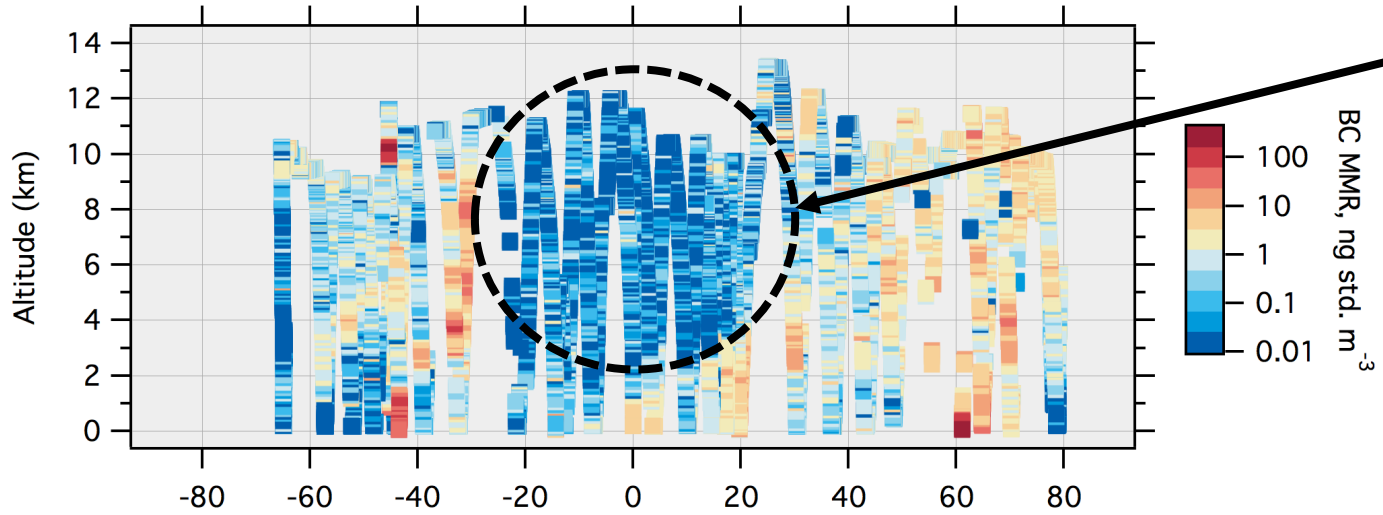
Atlantic



# Global Distribution of Black Carbon

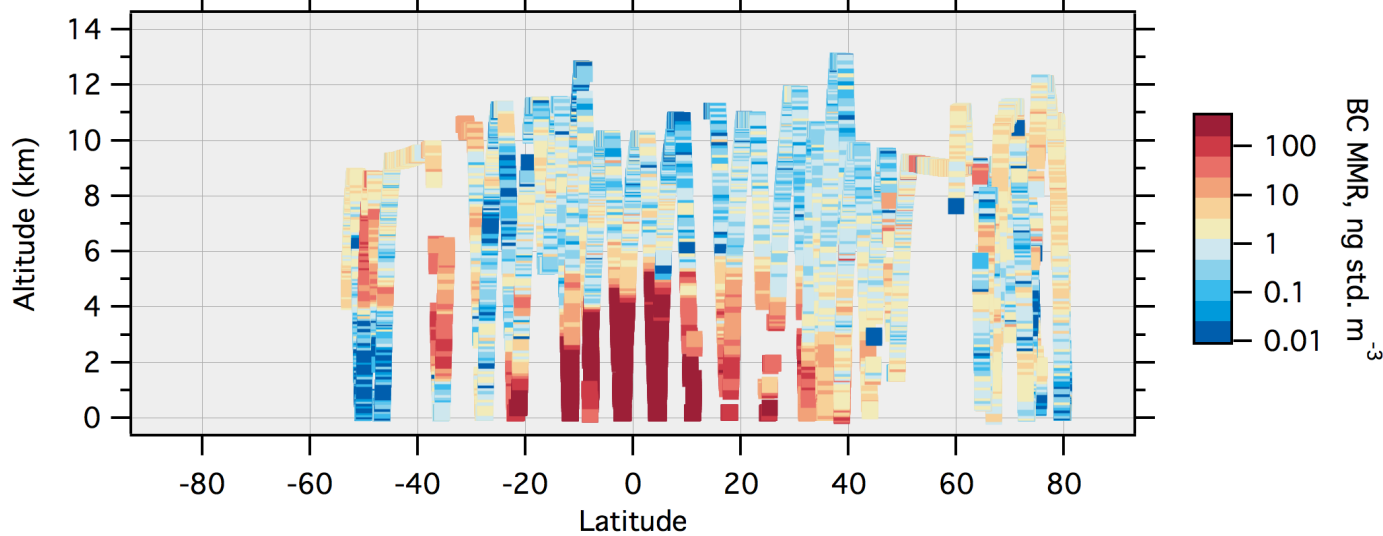
ATom 1  
Aug '16

Pacific



Convective Cleaning

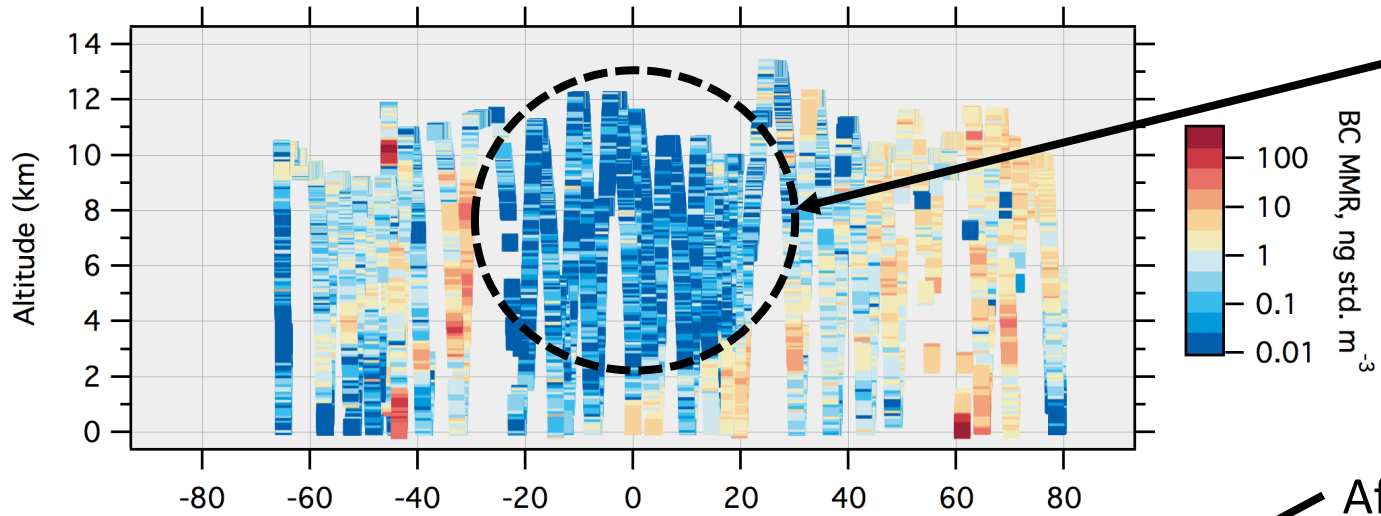
Atlantic



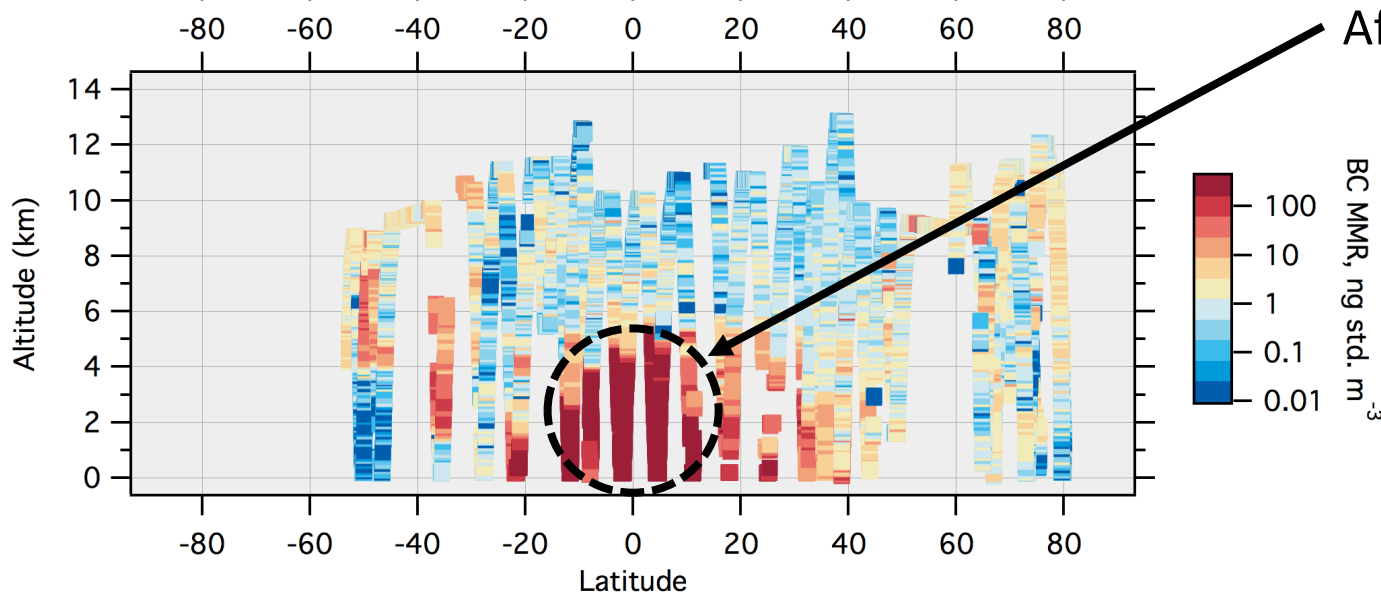
# Global Distribution of Black Carbon

ATom 1  
Aug '16

Pacific



Atlantic



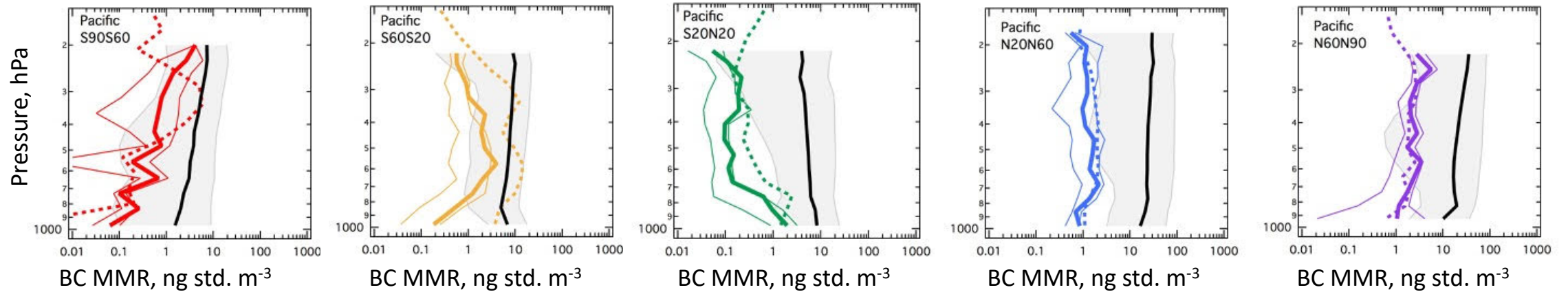


# Models tend to over-estimate black carbon at high altitudes

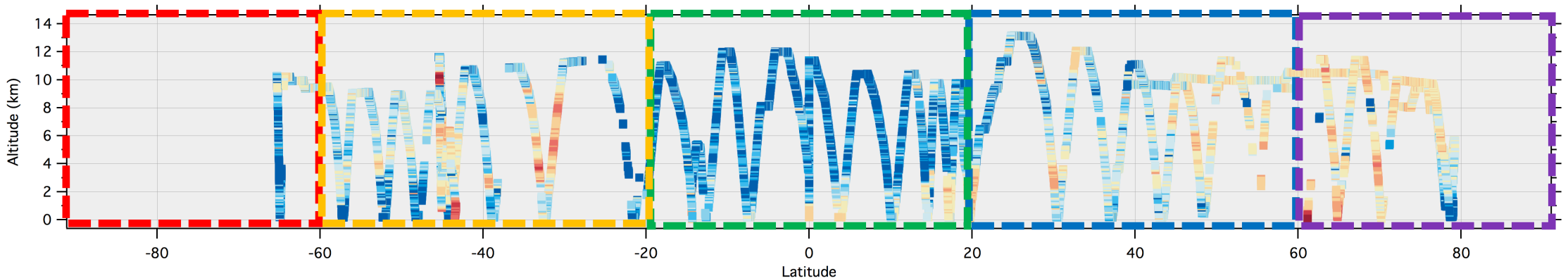
ATom 1 (Aug '16)

- ATom mean
- ATom 25<sup>th</sup> and 75<sup>th</sup> Percentiles
- - - HIPPO mean
- AeroCom Phase II ensemble average (13 models included)
- AeroCom output full range

Pacific



Pacific

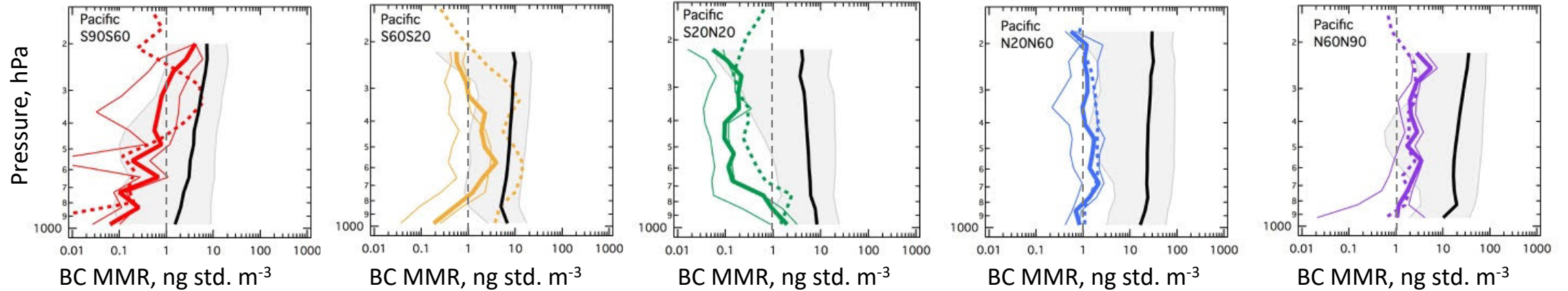


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ATom 1

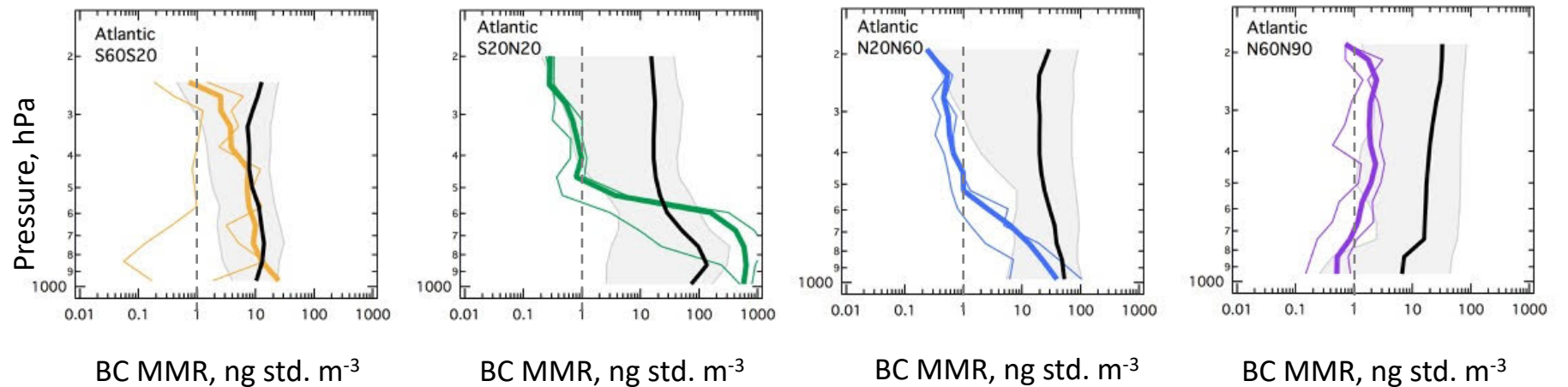
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Pacific



Katich et al. (2018), Strong contrast in remote black carbon aerosol loadings between the Atlantic and Pacific basins, *JGR*, in review

Atlantic

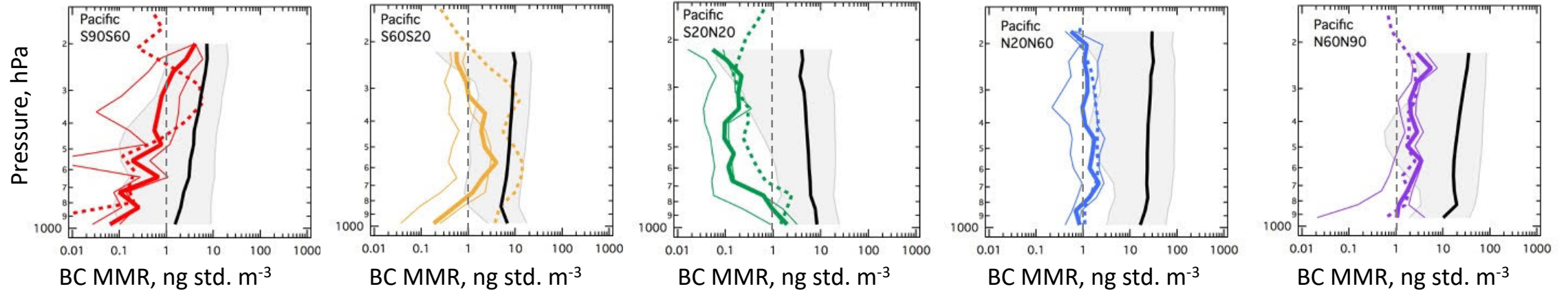


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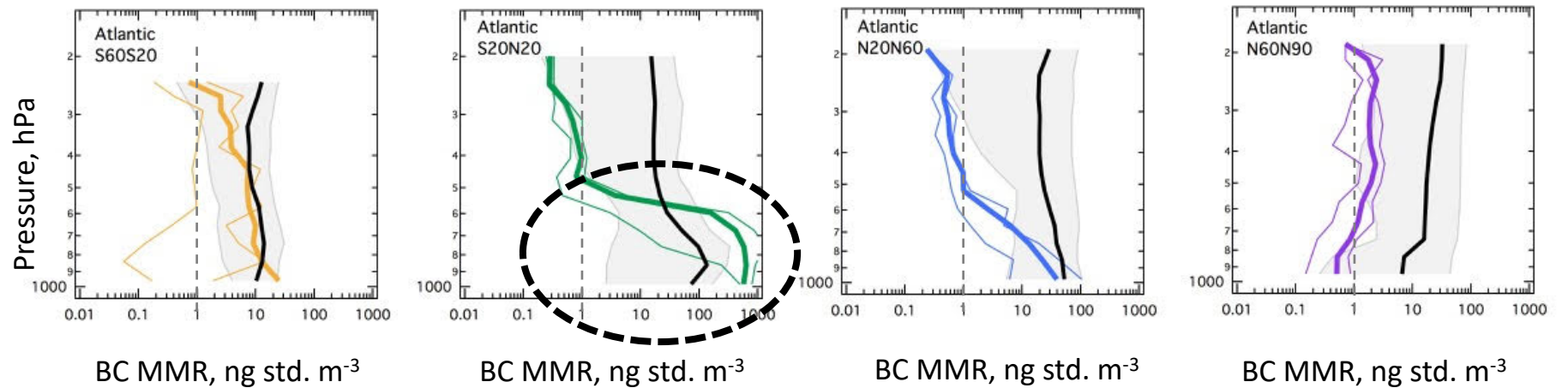
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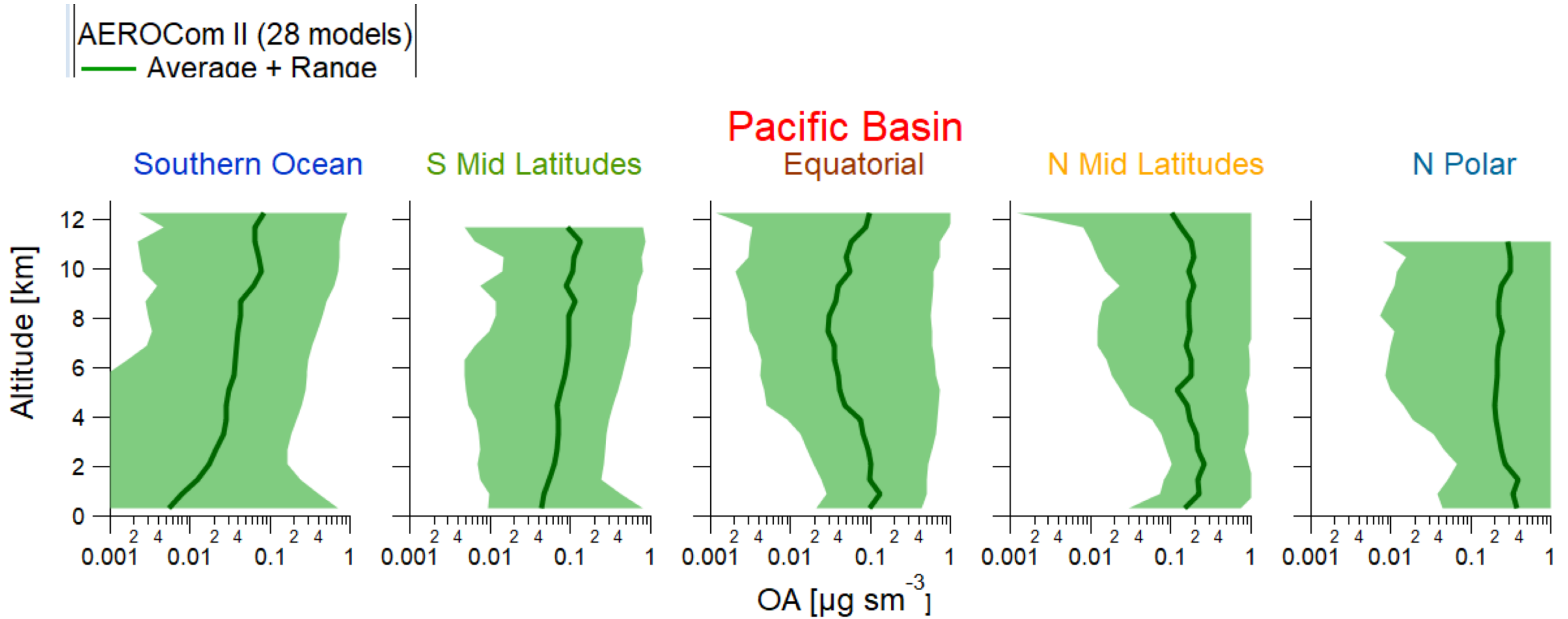


Atlantic

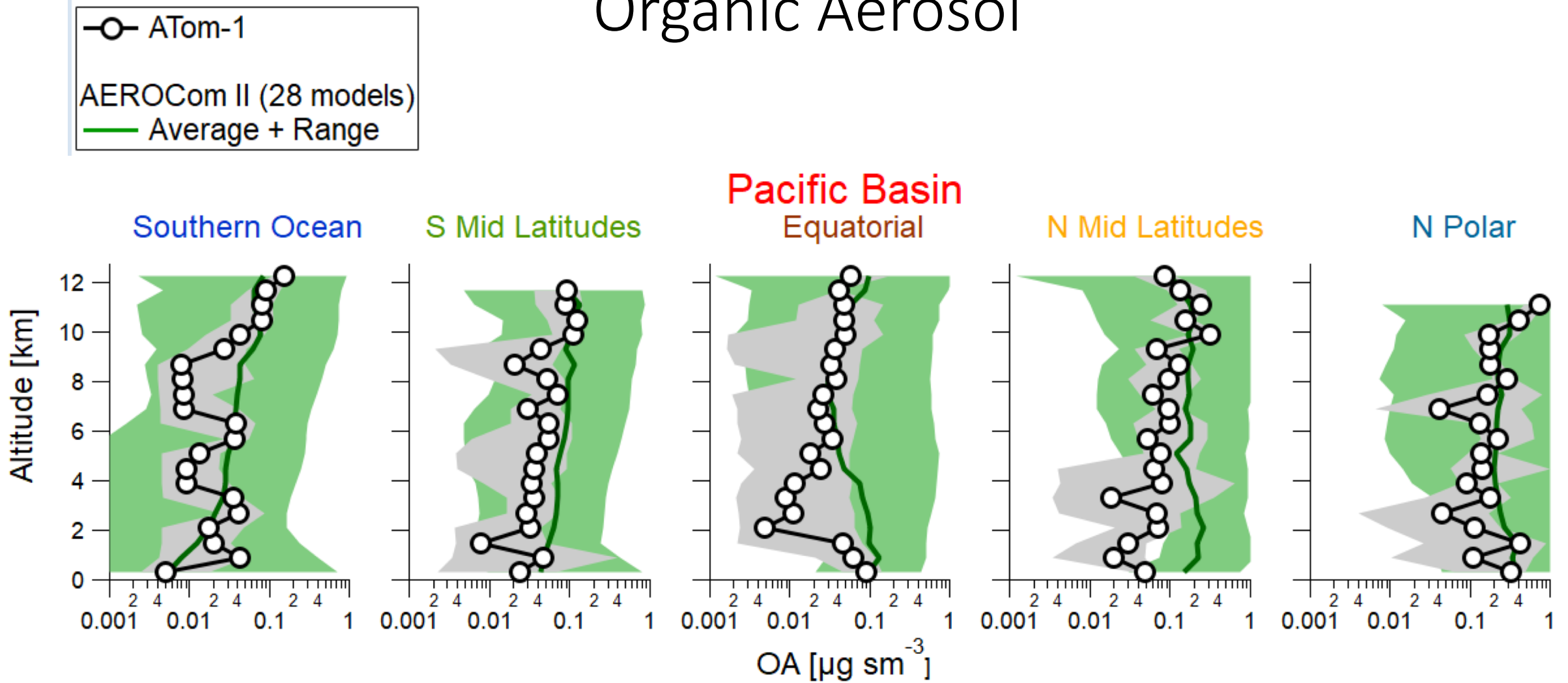


Katich et al. (2018), Strong contrast in remote black carbon aerosol loadings between the Atlantic and Pacific basins, *JGR*, in review

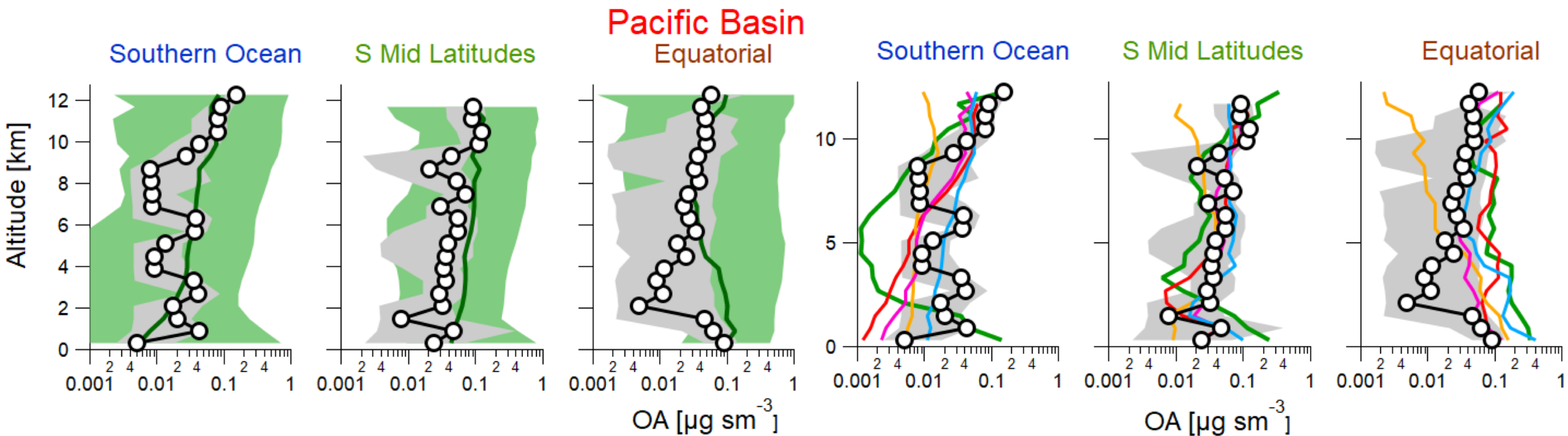
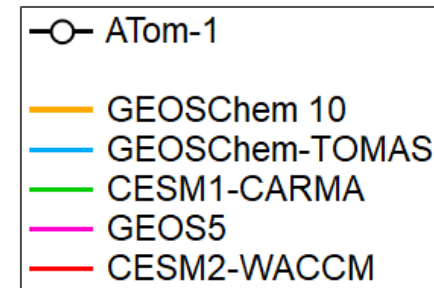
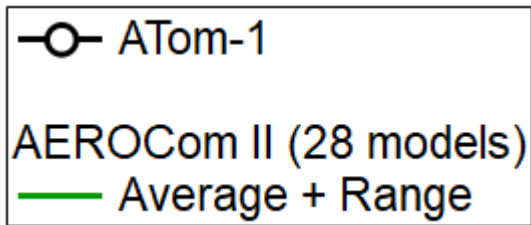
# AEROCOM II Models tend to overestimate remote Organic Aerosol



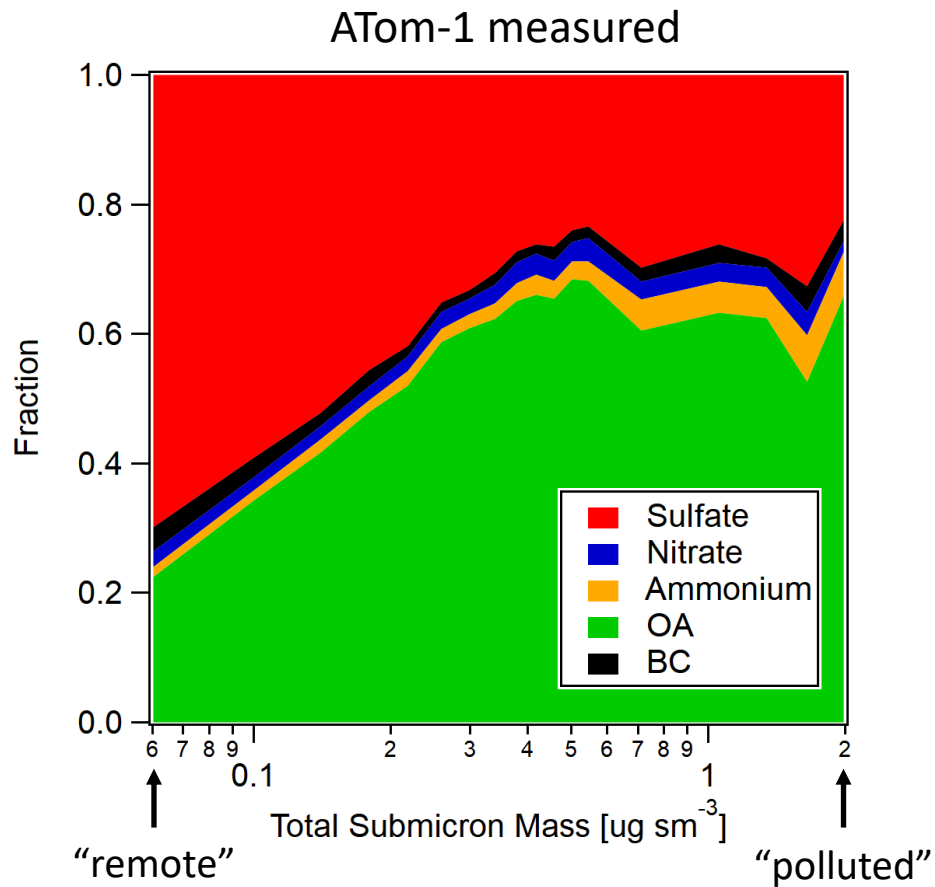
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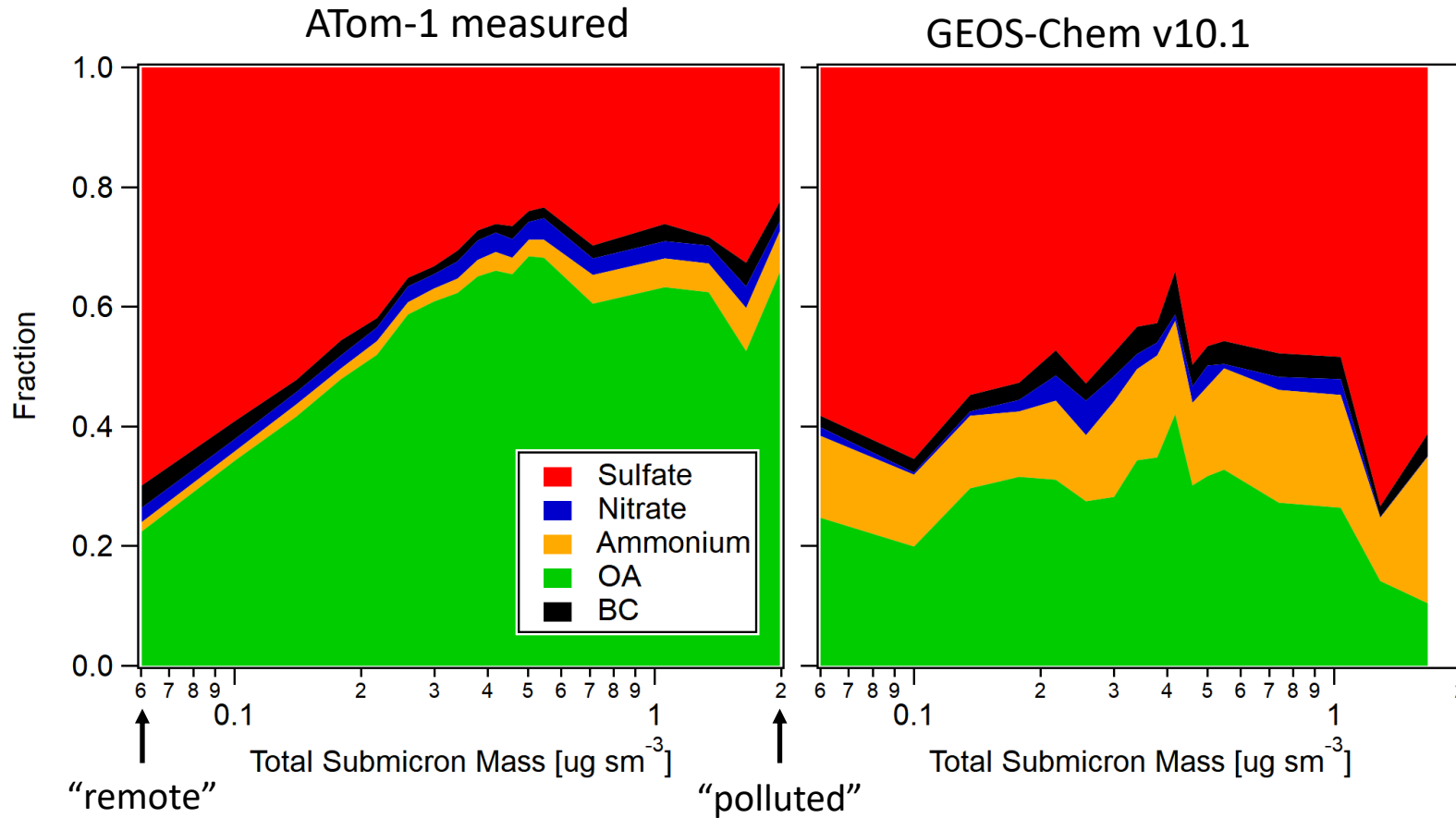
# A missing chemical loss process for organic aerosol?



Cleanest areas have lower  $f(\text{OA})$ , suggesting:

- Strong local sulfate production or
- Removal of OA at faster rates than other species

# A missing chemical loss process for organic aerosol?



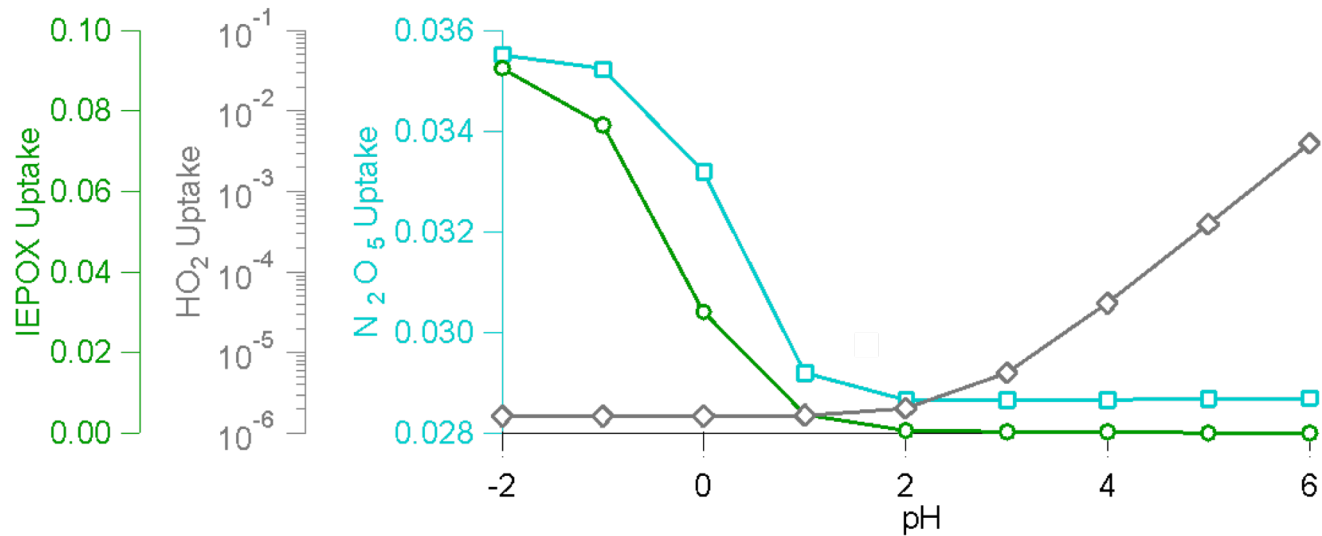
Cleanest areas have lower  $f(\text{OA})$ , suggesting:

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**Models without OA-specific removal fail to reproduce this trend**

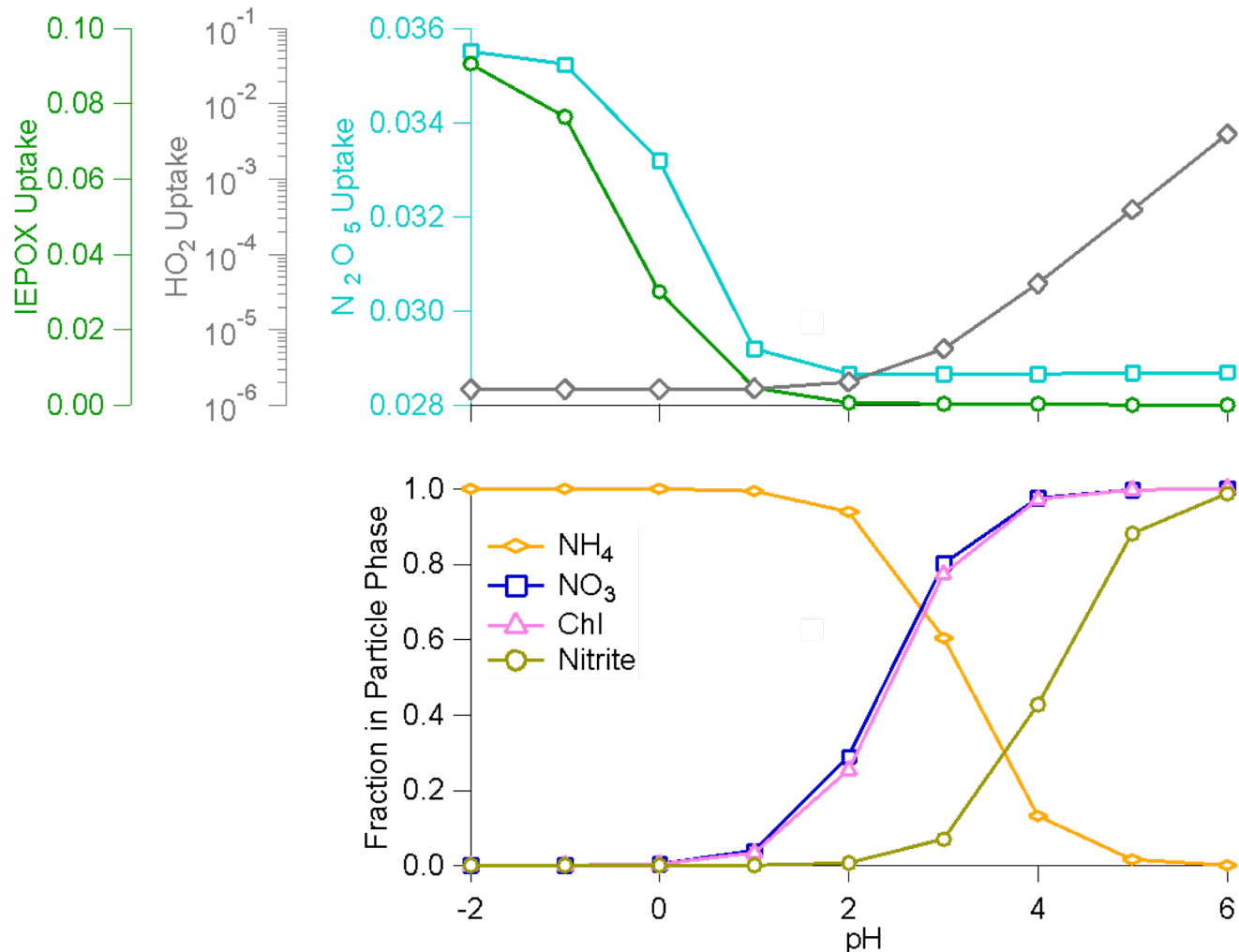


# Aerosol pH is important



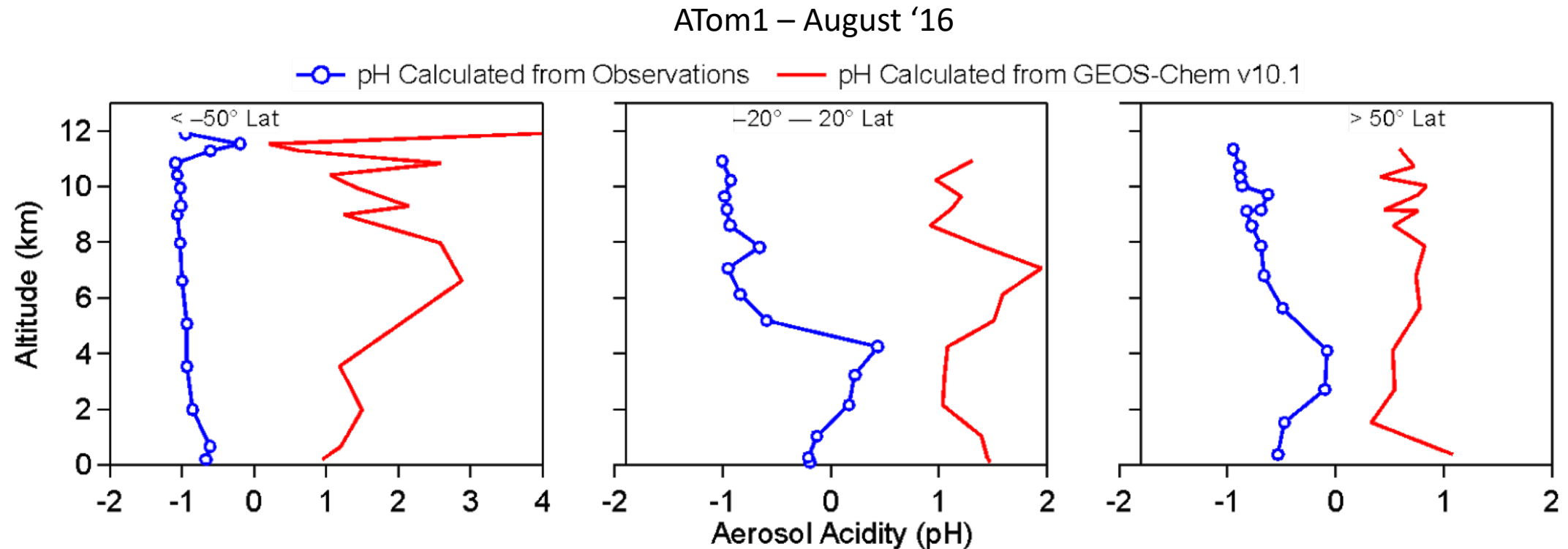
Various aerosol processes depend on pH, including uptake and partitioning

# Aerosol pH is important



Various aerosol processes depend on pH, including uptake and partitioning

# pH from observations is lower than predicted by models

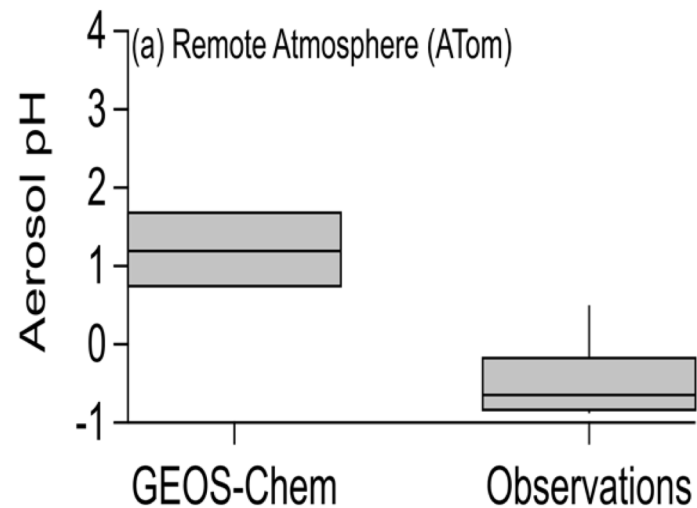


Uncertainty on pH calculated from observation  $\pm 0.5$

# Models fails to capture decrease in pH between urban and remote environments

**GEOS-Chem = 1.19**

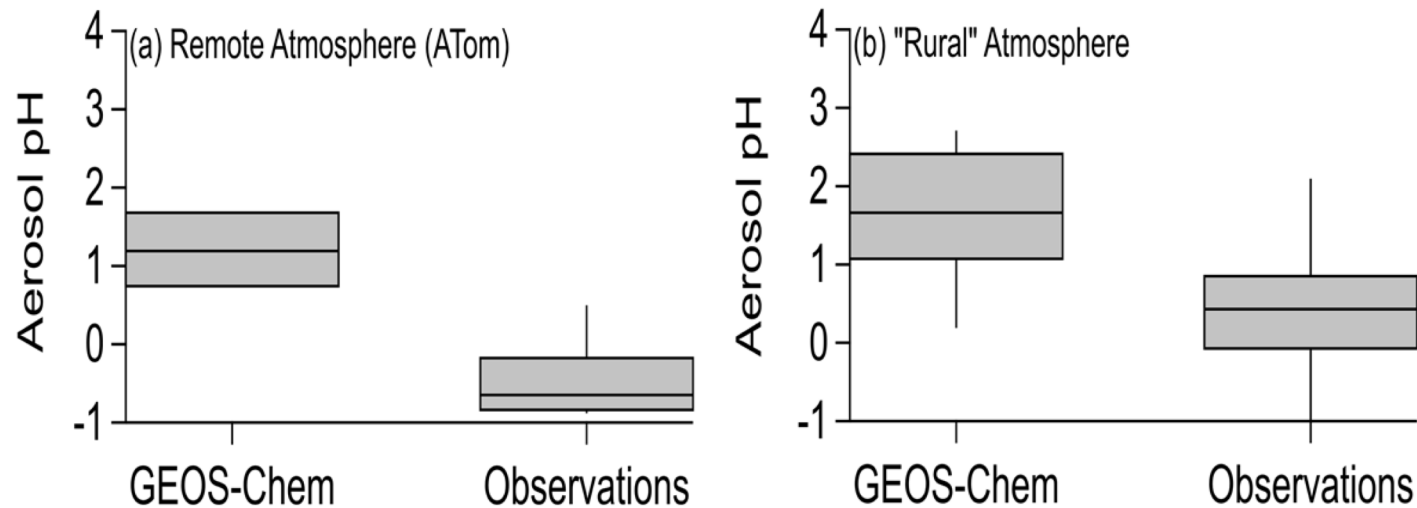
**Observations = -0.65**



# Models fails to capture decrease in pH between urban and remote environments

**GEOS-Chem = 1.19**  
**Observations = -0.65**

**GEOS-Chem = 1.66**  
**Observations = 0.43**



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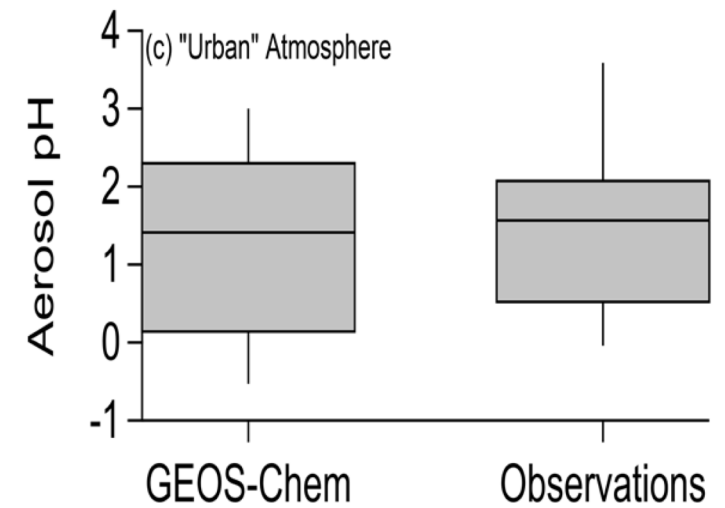
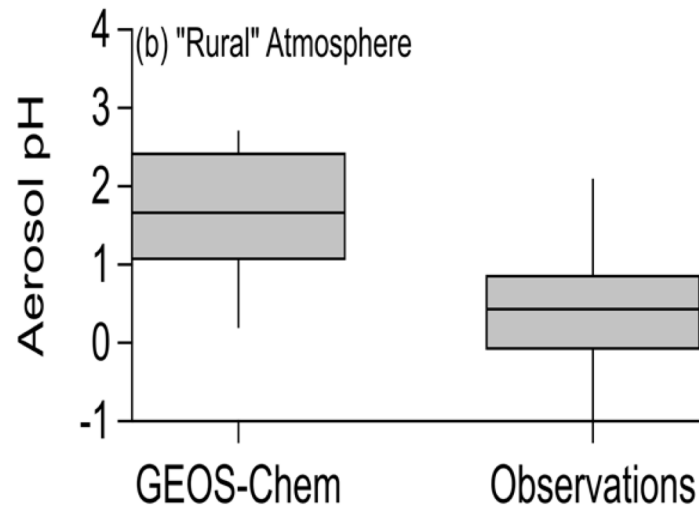
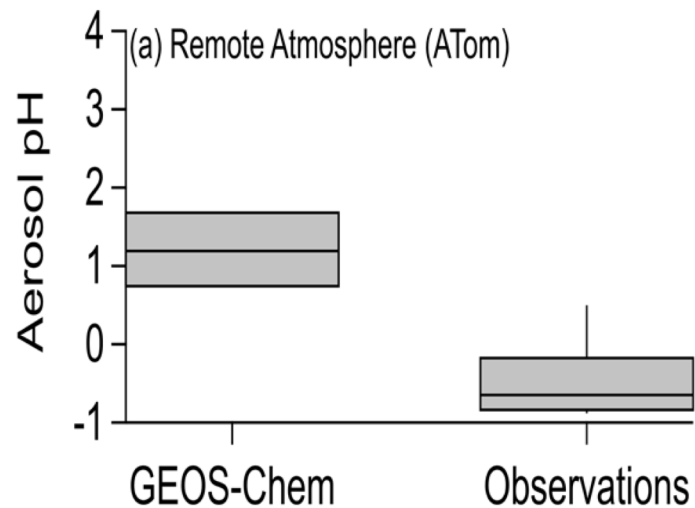
**GEOS-Chem = 1.66**

**GEOS-Chem = 1.41**

**Observations = -0.65**

**Observations = 0.43**

**Observations = 1.57**



# Models fails to capture decrease in pH between urban and remote environments

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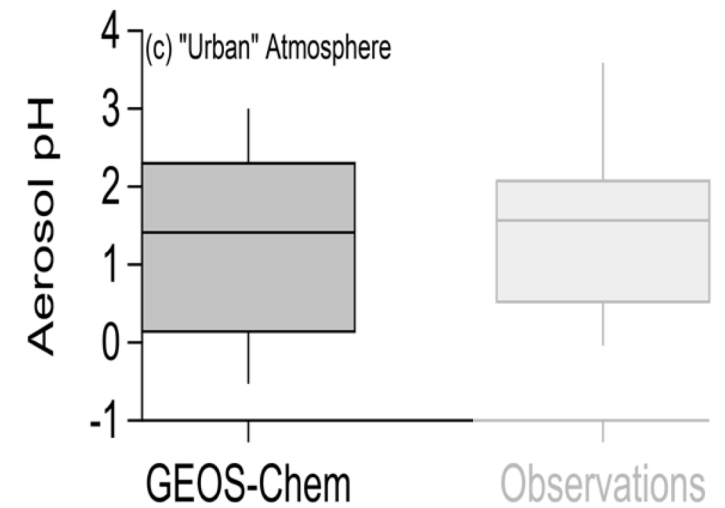
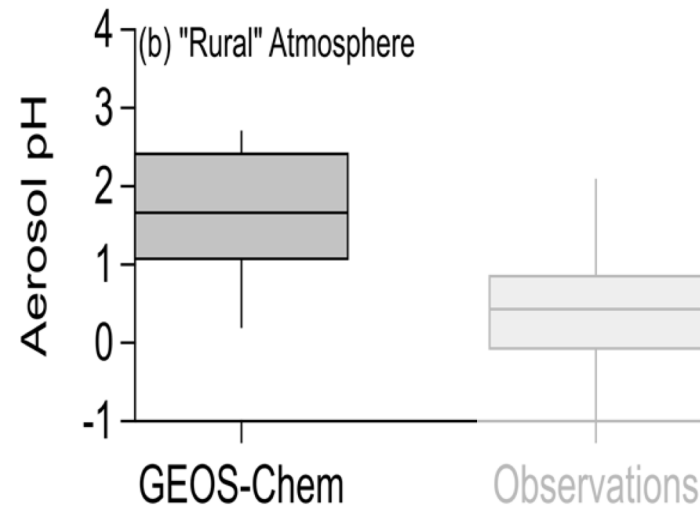
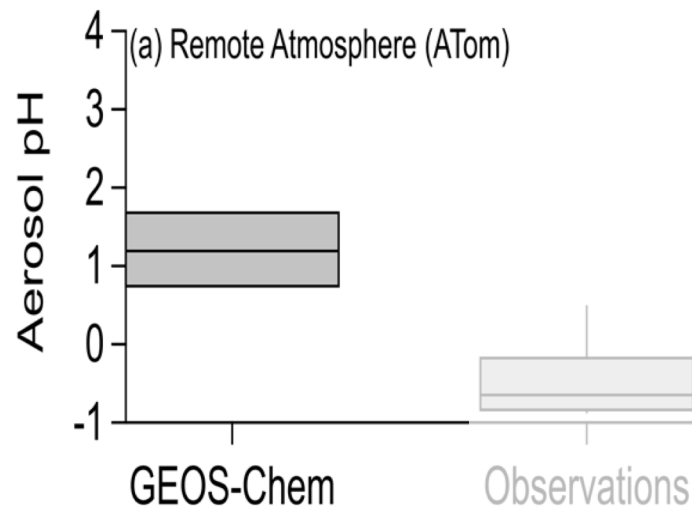
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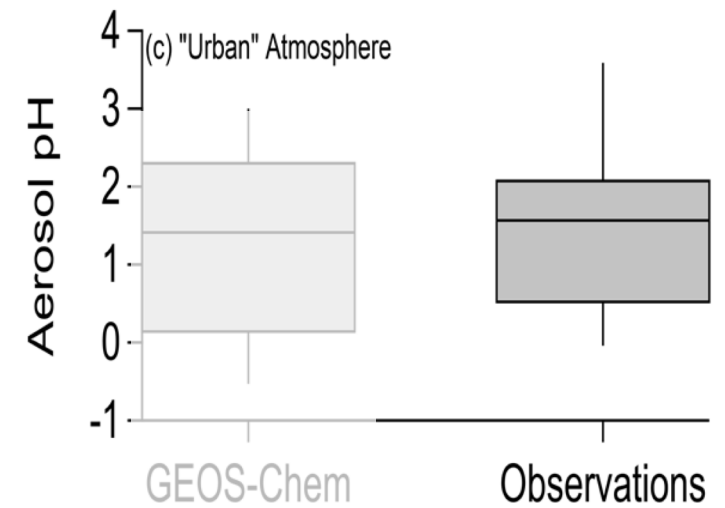
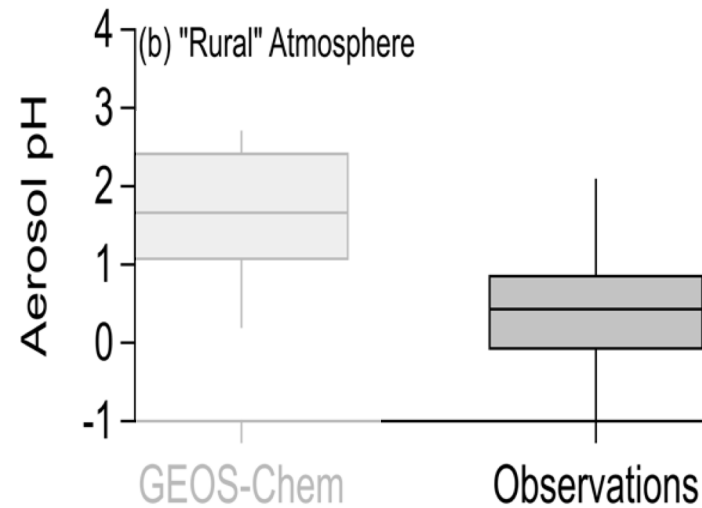
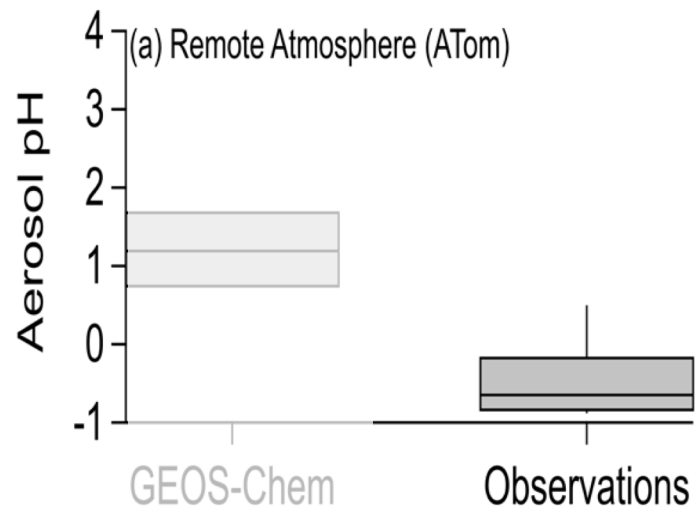
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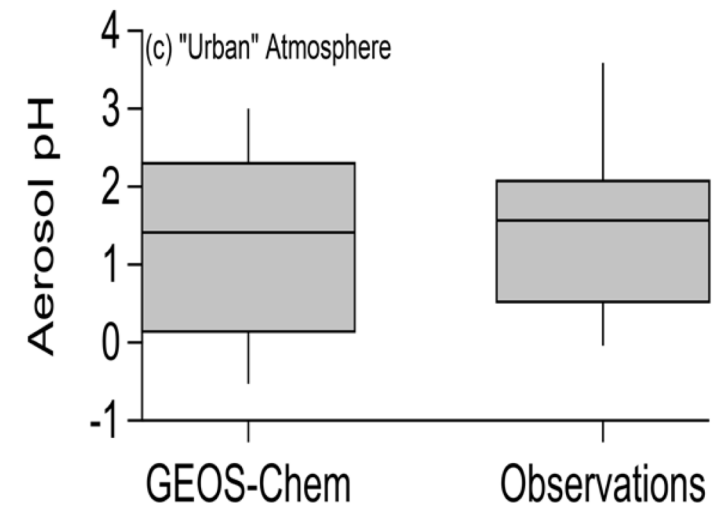
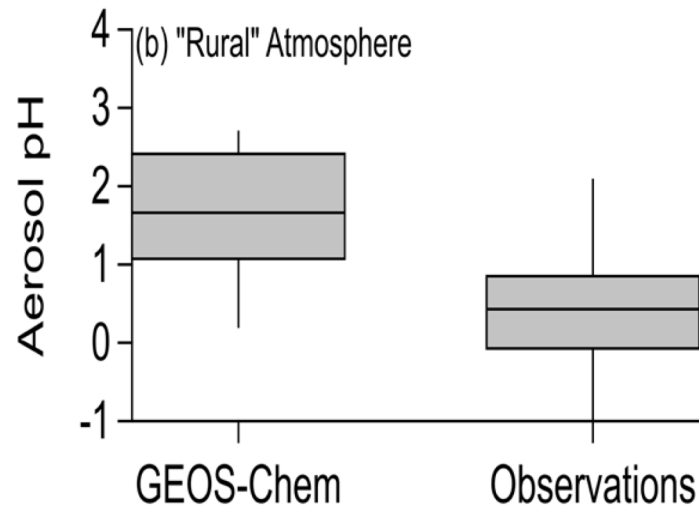
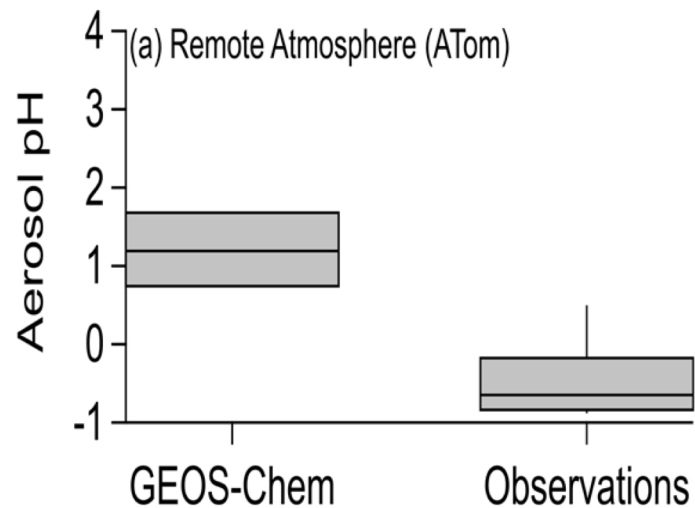
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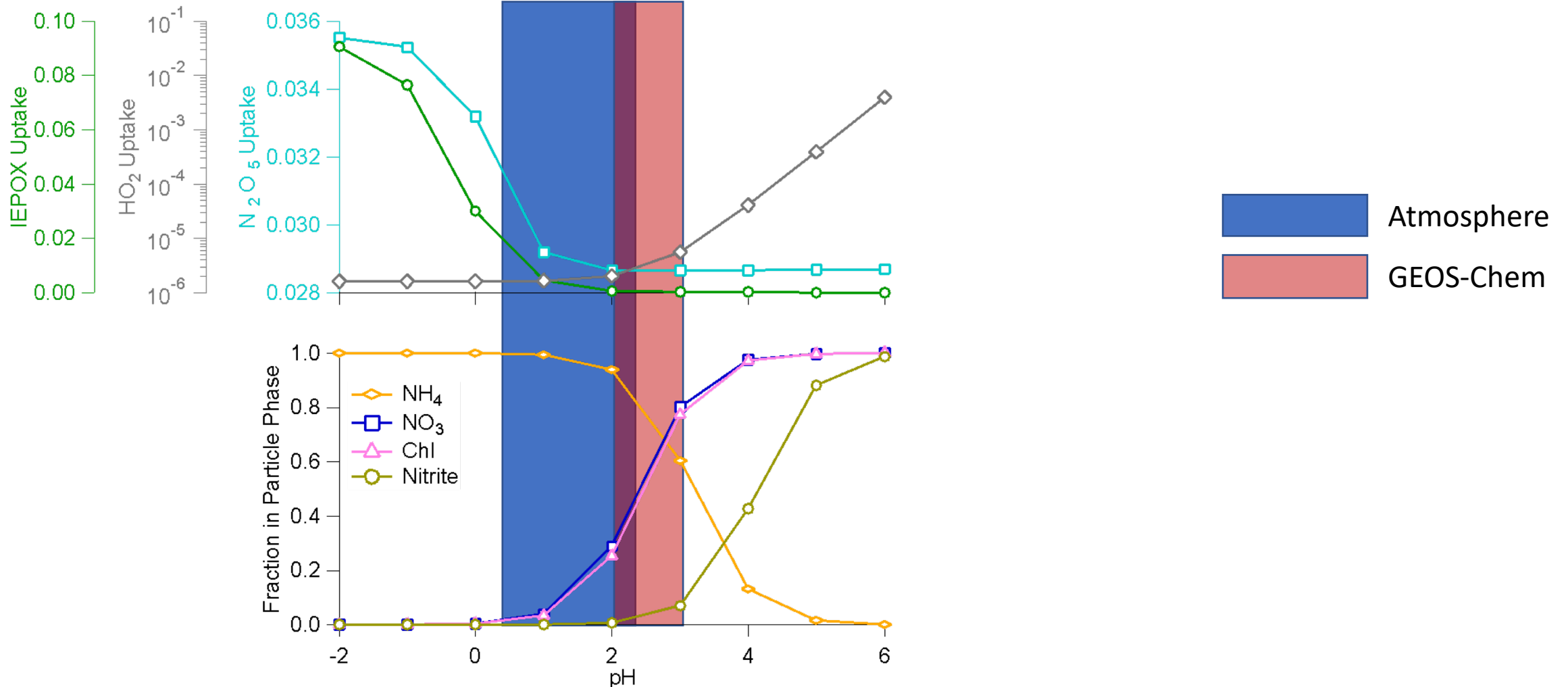
Observations = -0.65

Observations = 0.43

Observations = 1.57



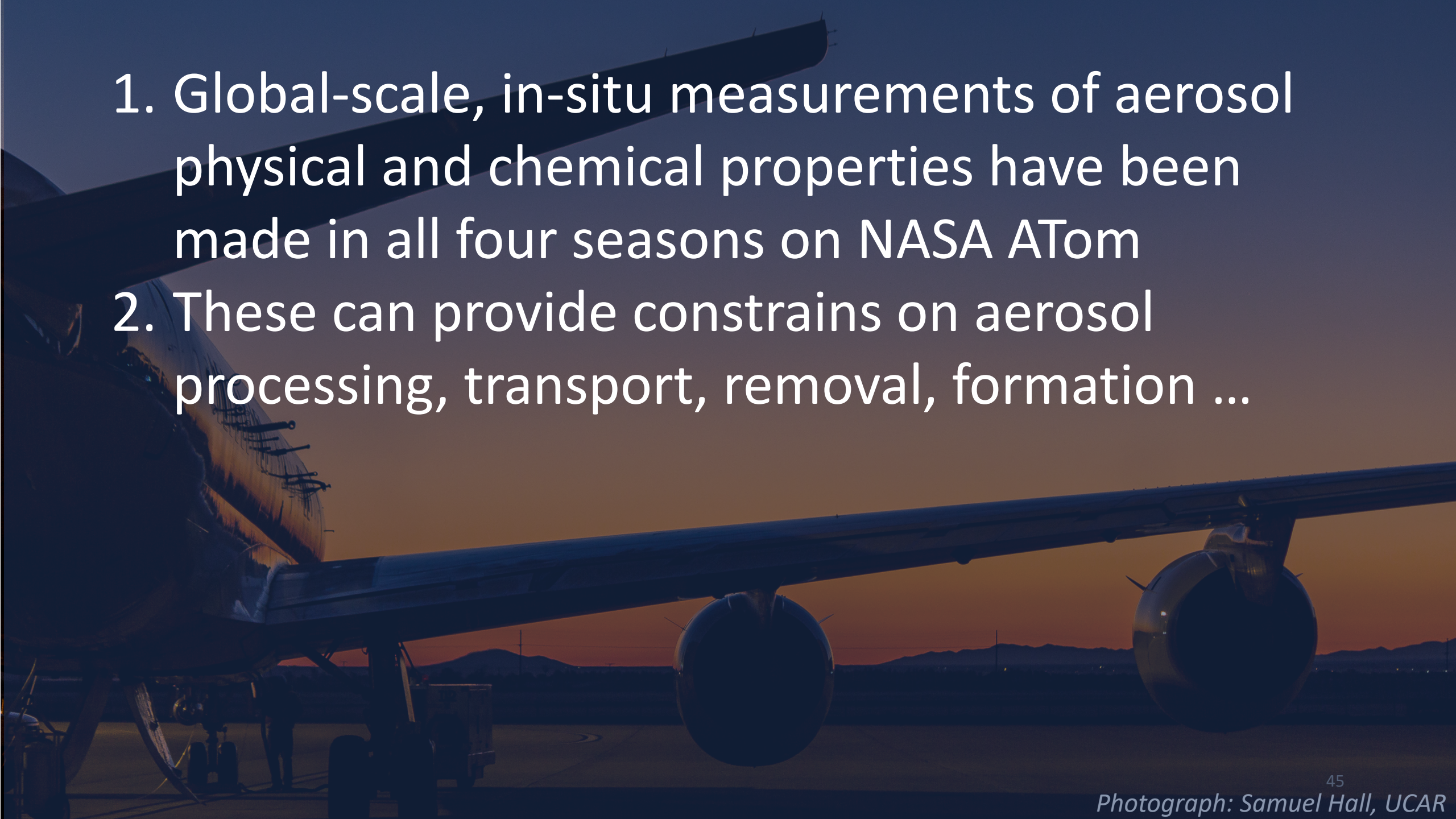
# Incorrect aerosol pH in models will affect many processes

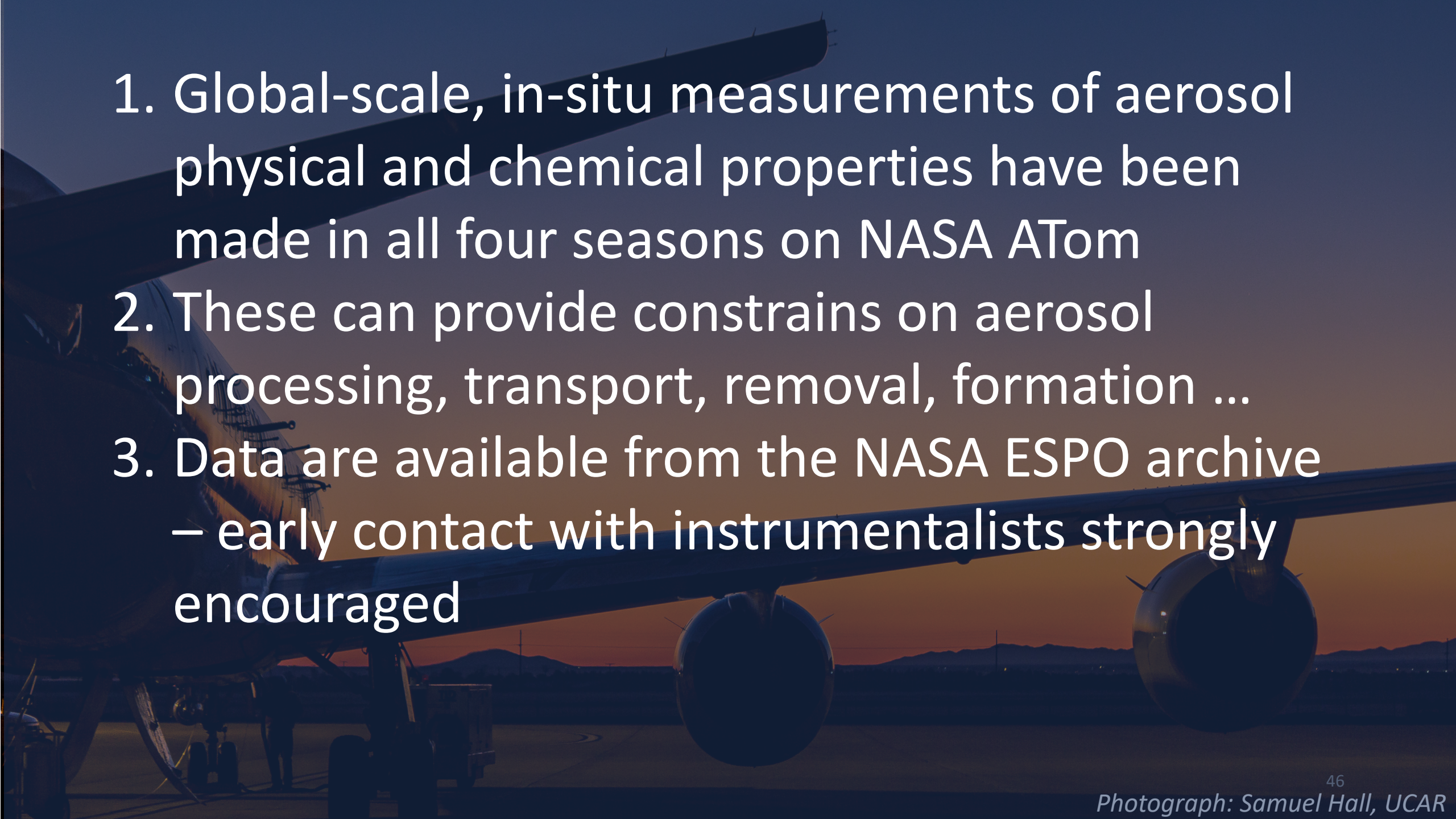


# Summary



1. Global-scale, in-situ measurements of aerosol physical and chemical properties have been made in all four seasons on NASA ATom

- 
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  2. These can provide constrains on aerosol processing, transport, removal, formation ...

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  2. These can provide constrains on aerosol processing, transport, removal, formation ...
  3. Data are available from the NASA ESPO archive  
– early contact with instrumentallists strongly encouraged

# Example Aerosol Products

- Concentration by number, surface area and volume
  - Nucleation , Aitken, Accumulation and Coarse Modes
- Aerosol Scattering, Absorption and Extinction
- Aerosol Mass
  - Organic            - Ammonium            - Sulfate
  - Dust- Biomass Burning       - Sea Salt
  - Black Carbon    - Brown Carbon       ...
- Number fraction from
  - Dust - Biomass Burning       - Sea Salt       ...
- Calculated pH
- Cloud phase
- Cloud droplet number
- ...



# Contact us:



## **Size Distributions (AMP):**

christina.williamson@noaa.gov, agnieszka.kupc@univie.ac.at , charles.a.brock@noaa.gov

## **Single Particle Composition (PALMS):**

karl.froyd@noaa.gov , gregory.schill@noaa.gov , daniel.m.murphy@noaa.gov

## **Black Carbon (SP2):**

joseph.m.katich@noaa.gov, joshua.p.schwarz@noaa.gov

## **Bulk Composition (AMS):**

jose.jimenez@colorado.edu , pedro.campuzanojost-1@colorado.edu, benjamin.nault@colorado.edu

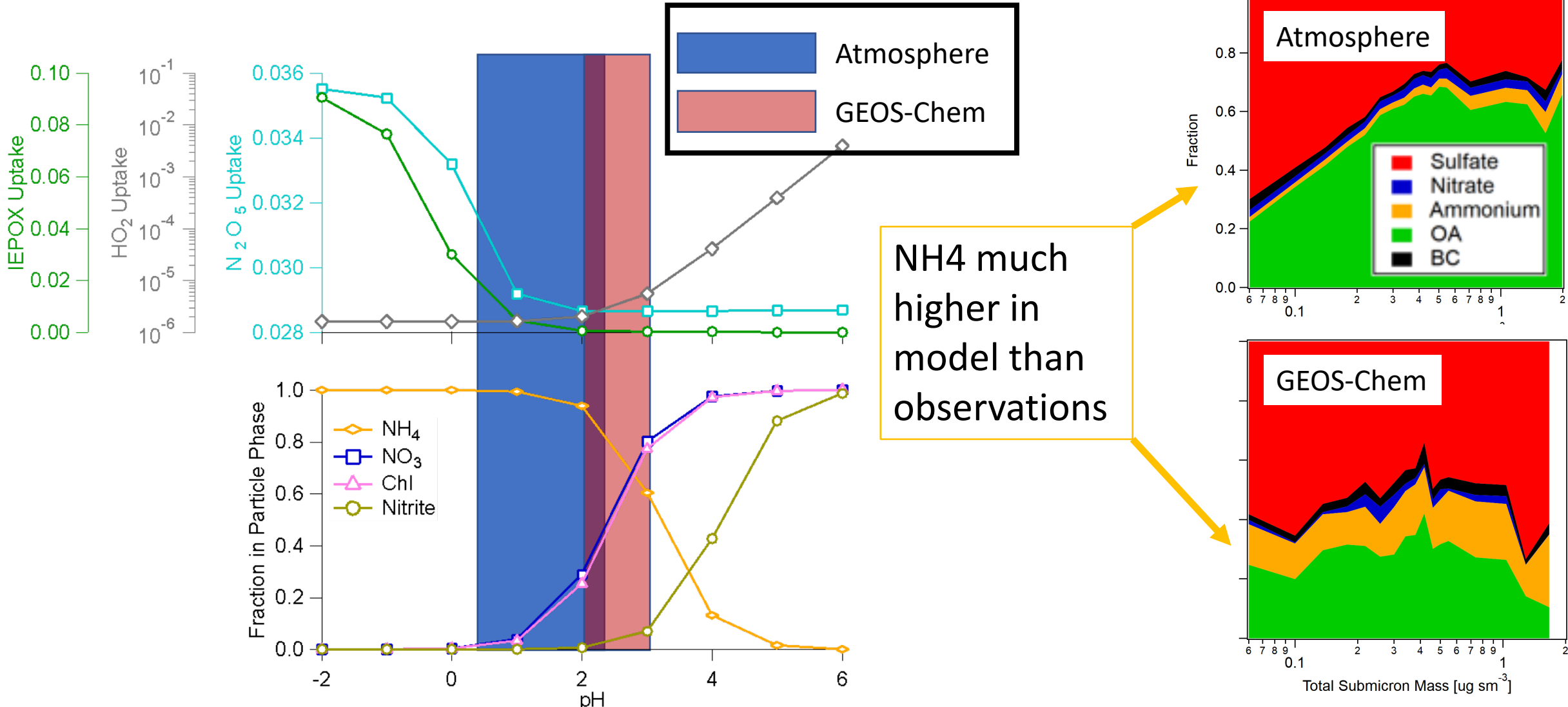
## **Cloud Properties (CAPS):**

bernadett.weinzierl@univie.ac.at , maximilian.dollner@univie.ac.at



backup

# Excess ammonia in models increases aerosol pH



# How submicron aerosol pH is estimated with a thermodynamic model

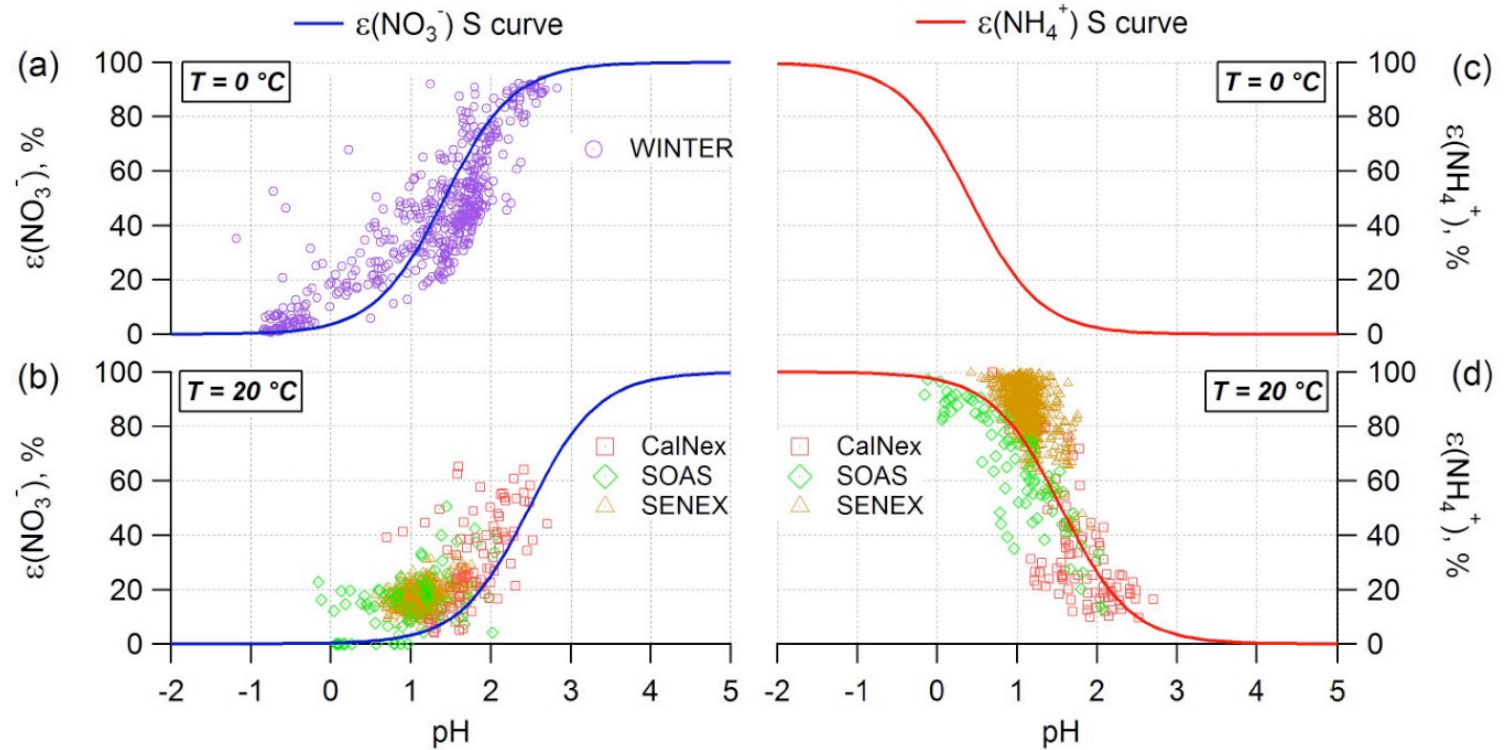
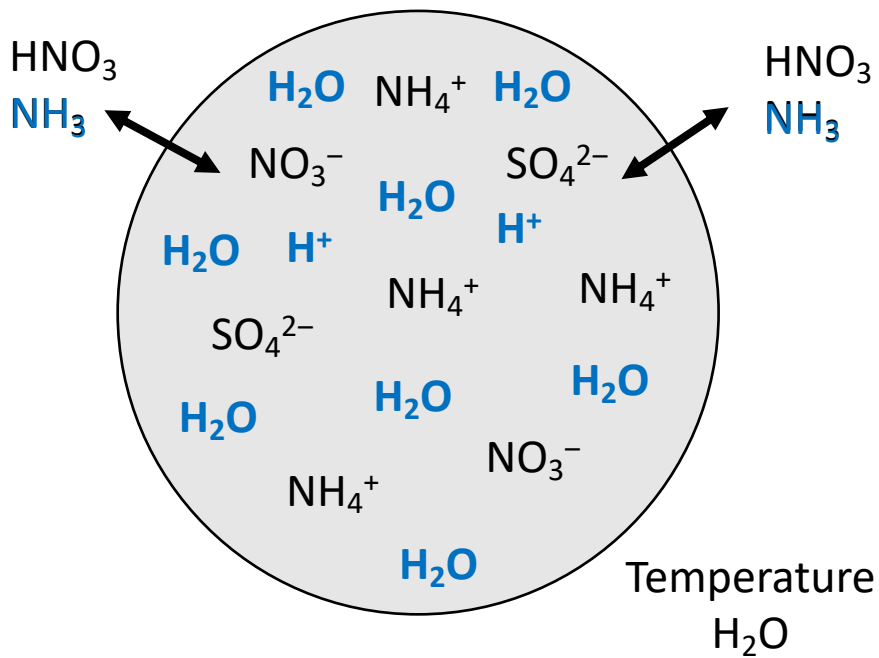
## Model Input:

Total  $\text{NO}_3$  or aerosol  $\text{NO}_3^-$

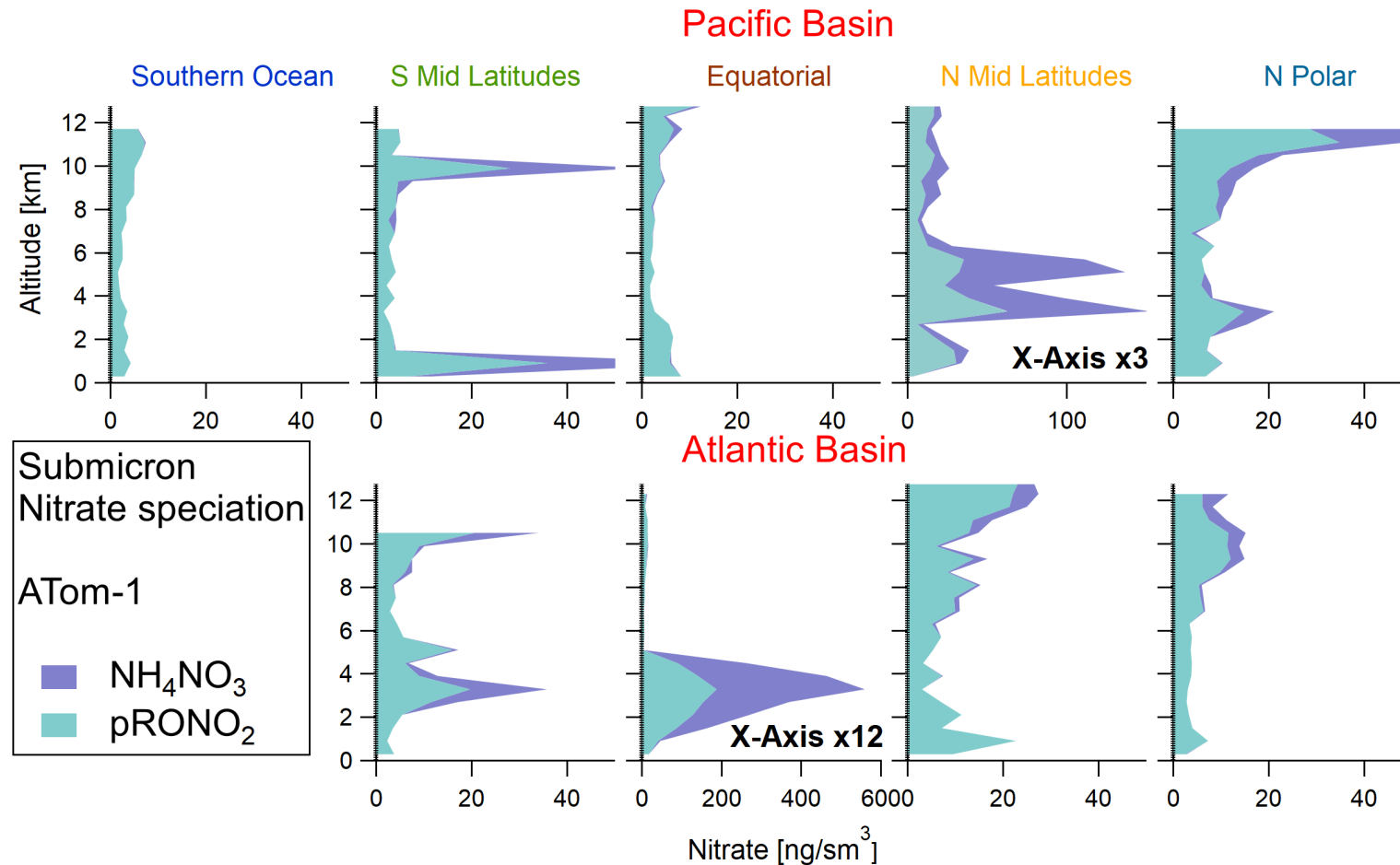
Total  $\text{NH}_x$  or aerosol  $\text{NH}_4^+$

$\text{SO}_4$

Temperature & relative humidity



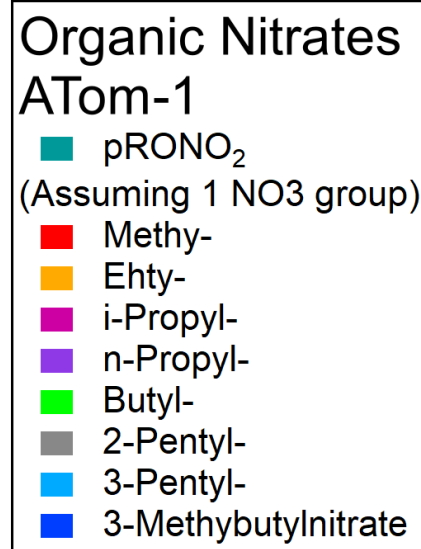
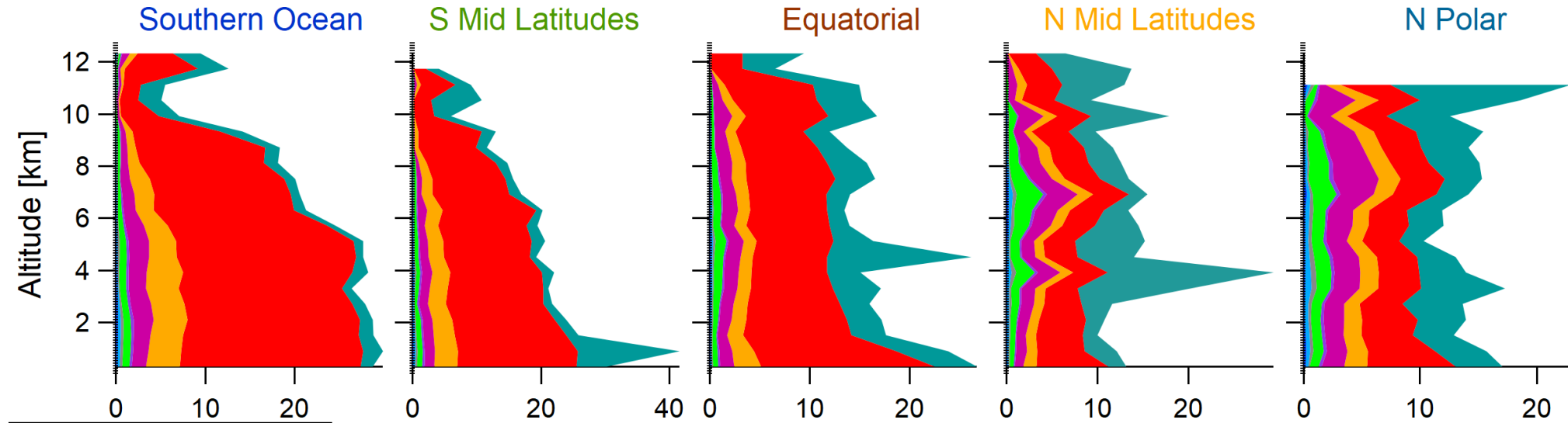
# Most submicron nitrate found in ATom is organic



- Highly acidic aerosol keeps inorganic nitrate in the gas-phase (except in neutralized BB plumes)
- There is a persistent, yet very low background of (likely long-lived) particulate organic nitrate

# pRONO<sub>2</sub> contributes significantly to total RONO<sub>2</sub>

## Pacific Basin



## Atlantic Basin

