Chemical and Physical Properties of Aerosols in Regional Background and Free Tropospheric Air Masses in the Western U.S.

Shan Zhou¹, Sonya Collier¹, Daniel A. Jaffe², Qi Zhang^{1*} (dkwzhang@ucdavis.edu)

¹Department of Environmental Toxicology, University of California, Davis, CA, USA ²School of Science, Technology, Engineering, and Mathematics, University of Washington, Bothell, WA, USA

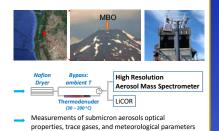


Introduction

- o Atmospheric aerosols affect the Earth's radiative budget directly and indirectly.
- Aerosols are mostly produced and concentrated in the planetary boundary layer (BL), but can be transported into the free troposphere (FT), where they are subjected to fewer removal processes and therefore have longer lifetime and climate effects.
- Understanding the sources, evolution, and physicochemical properties of aerosols in regional background air masses and in the FT is crucial for constraining the climate impacts of aerosols on a global scale.
- High-altitude mountaintop observatories provide the opportunity to study aerosol properties in remote areas, especially the FT, without the added expense and difficulty of making airborne measurements.

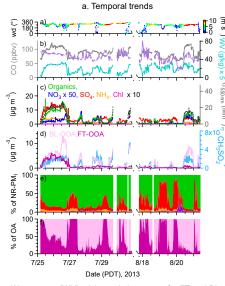
Methodology

- The Mt. Bachelor Observatory (MBO; 2763 m a.s.l.) is a regional background site in the western US.
- The location allows for FT air masses to be sampled during the night and air coming from the BL during daytime.
- Real-time aerosol measurements were conducted at MBO during the Biomass Burning Observation Period (BBOP) in July 25 – August 25, 2013.

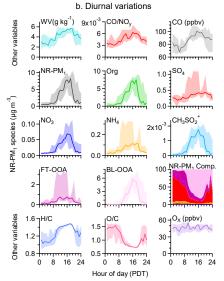


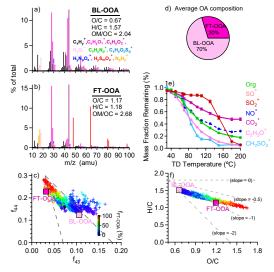
Results

Aerosol concentration and composition at MBO during periods representing regional background and free tropospheric conditions



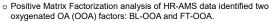
o Water vapor (WV) mixing ratio is a tracer for FT and BL air
 o Aerosol concentration and composition differ significantly between FT and BL air masses.





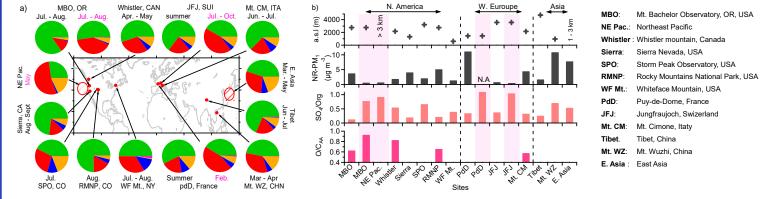
c. Composition and volatility of organic aerosol (OA)

 Major aerosol components and primary gaseous pollutants show clear diurnal cycles driven by the BL.



o BL-OOA and FT-OOA have different compositions and volatilities

Global observations of aerosol composition in the regional background air masses at high-altitude



A summary of measurements of regional background aerosols at multiple high-altitude sites and aircraft. Pink labels in (a) and shades in (b) indicate FT air and the rest represent mixed BL and FT air.
 Compared to aerosols in mixed air masses, FT aerosols have much lower aerosol concentrations with substantially higher mass fractions of sulfate and sulfate-to-organic ratios.

Conclusions

Chemical and physical properties of aerosols in regional background and free tropospheric air masses in the western US were studied in summer 2013 at MBO:

Low aerosol loading (average NR-PM₁ = 3.75 μg m⁻³) was observed.

• Main aerosol components showed clear diurnal variations driven by boundary layer dynamics with higher concentrations occurring during daytime and lower concentrations at night.

- Aerosols in the free troposphere tended to be more acidic and contained a higher mass fraction of ammonium sulfate (up to 90% of NR-PM₁ mass) compared to air masses from the boundary layer.
 Organic aerosol was highly oxidized (O/C = 0.84) and dominated aerosol composition (85% of NR-PM₁ mass):
 - BL-OOA (O/C = 0.67; 70% of OA mass) representing biogenically-influenced SOA formed in the BL
 - FT-OOA (O/C = 1.17) representing highly oxidized low-volatility organics in the FT.

Acknowledgements: This work was funded in part by US Department of Energy Atmospheric System Research Program, Grant No. DESC0014620 and DE-SC0007178. Shan Zhou acknowledges funding from the Chinese Scholarship Council (CSC), Donald G. Crosby Fellowship, Matsumura Memorial Fellowship, and Coulson ASGG Representative Travel Award.