# Chemical composition and optical properties of wildland and agricultural biomass burning particles measured downwind during BBOP study

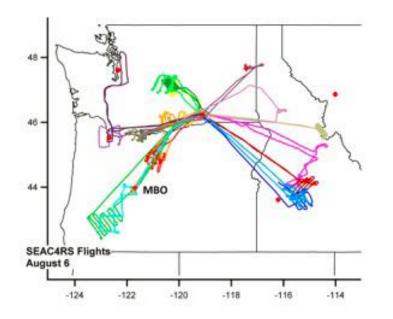
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DOE ASR and ARM funding

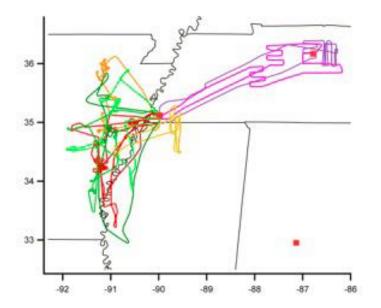
#### **Biomass Burn Observation Project (BBOP)**

A Department of Energy (DOE) sponsored study to measure wildland fires in the Pacific Northwest and prescribed agricultural burns in the Central Southeastern US from the DOE Gulfstream-1 aircraft platform over a four month period in 2013.



Wildland Fires (17):

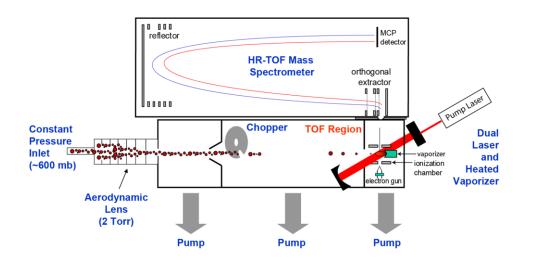
Shrub, Forest **Urban**: Seattle (3), Portland (2), Spokane (2) **MBO** (3) **SEAC4RS:** Joint mission Aug., 6



Prescribed Agricultural burns (24): rice, soybean, sorghum Urban: Nashville (2), Memphis (2)

# Chemical and Optical Measurements

Chemical & Physical Measurements	
NR-PM:	SP-AMS, TEM
rBC:	SP2, SP-AMS, TEM
Size:	UHSAS, PCASP, FIMS



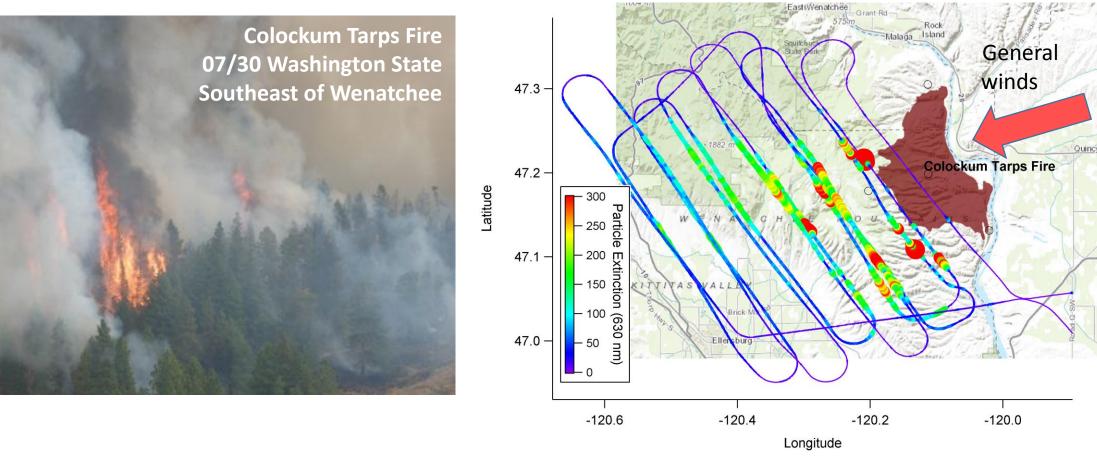
Soot Particle Aerosol Mass Spectrometer (SP-AMS)

Optical Measurements	
Extinction:	1- $\lambda$ CAPS PMex (630 nm)
Scattering:	$3-\lambda$ Nephelometer (450, 550, 700 nm)
Absorption:	1-λ PAS (355 nm)
	1-λ PTI (532 nm)
	3-λ PSAP (462, 523, 648 nm)

SSA's derived from SCAT/EXT

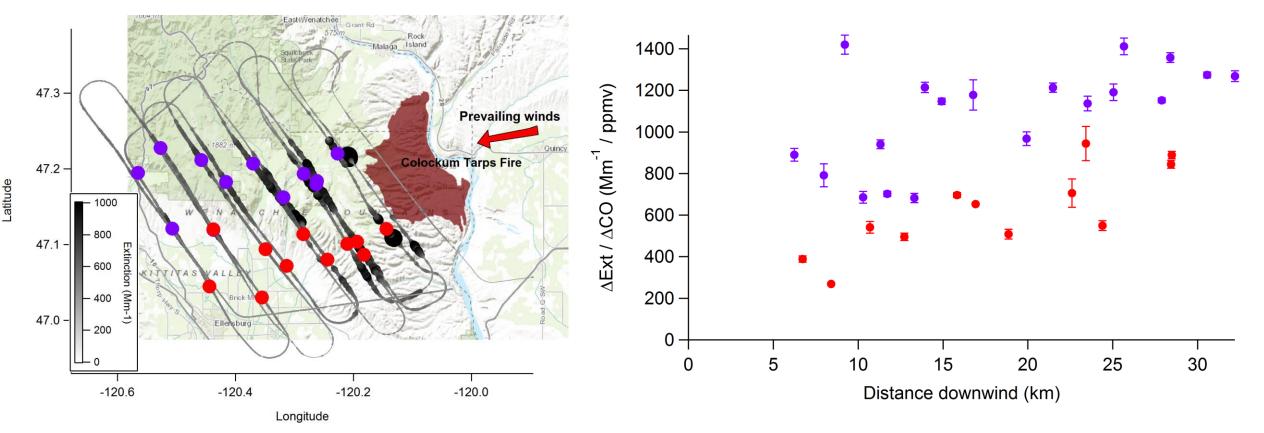
- EXT: CAPS PMex (630 nm)
- SCAT: NEPH (550 and 700 nm)
  - Interpolated at 630 nm using Ångström coefficient

#### Colockum Tarps Fire (2<sup>nd</sup> flight July 30) Extinction



• Downwind transects across fire front tracking plumes

#### $\Delta$ Extinction / $\Delta$ CO downwind



- Extinction relative to CO is increasing downwind
- Different burn plumes from same fire
- Average MCE ~ 0.92 in both plumes (red ≥ purple)

#### $\Delta$ Extinction / $\Delta$ CO

#### Flight 730b (31 plumes)

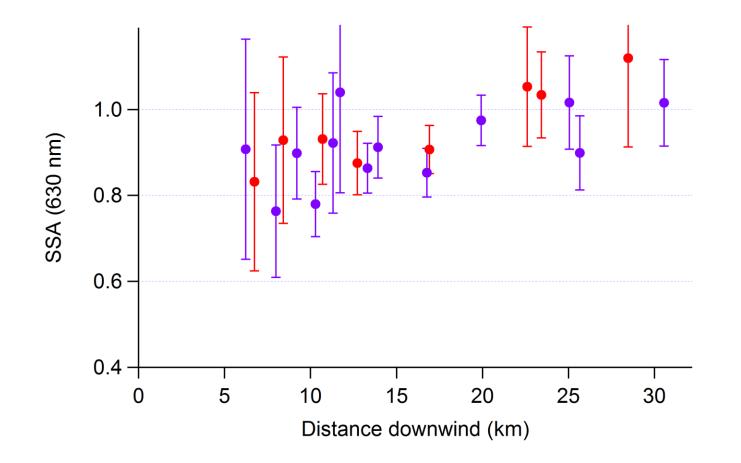
#### 35 -5 -30 · 4 25 mass influenced Plume # Plume # 3 -20 15· 2 -10 -Jrban 1 5 -0 -0 500 1000 1500 0 500 1000 1500 0 $\Delta Ext / \Delta CO (Mm^{-1} / ppmv)$ $\Delta Ext / \Delta CO (Mm^{-1} / ppmv)$

Dingle et al., 2016 ACPD

•  $\Delta Ext/\Delta CO$  for a single fire spans nearly the full range of observed wildfire plumes

#### **BBOP wildfires (173 plumes)**

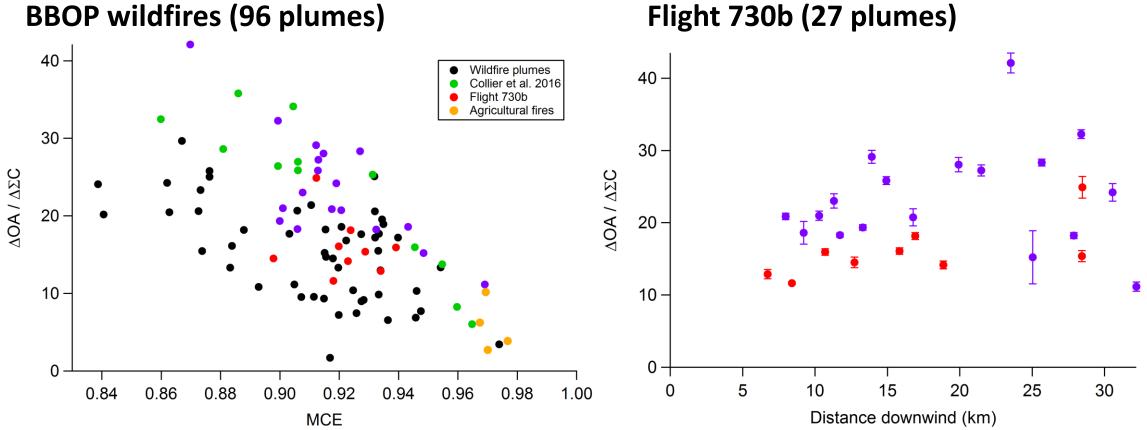
#### SSA (630 nm) measurements downwind



- No apparent differences in SSA with burn plume
- Increasing SSA (630 nm) with downwind transport indicating changes in particle scattering > particle absorption
- Reach SSA ~ 1 within ~3 hours down wind

Collier, S. et al. Environ. Sci. Technol. acs.est.6b01617 (2016)

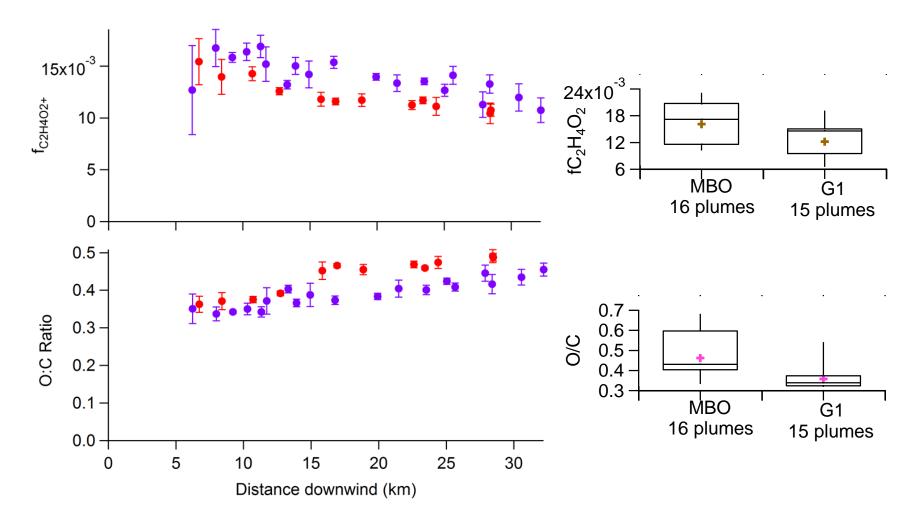
#### Biomass burning PM chemical composition



Biomass burning plumes were > 92 % Organic

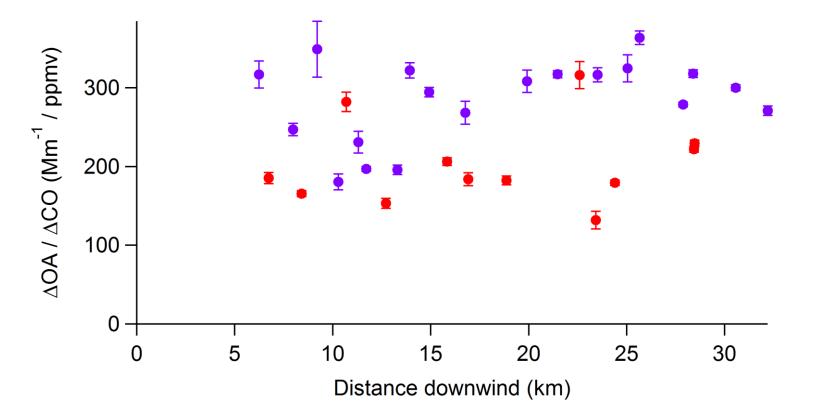
 While MCE appears to be strong driver, atmospheric changes may provide secondary variances

### OA chemical changes downwind

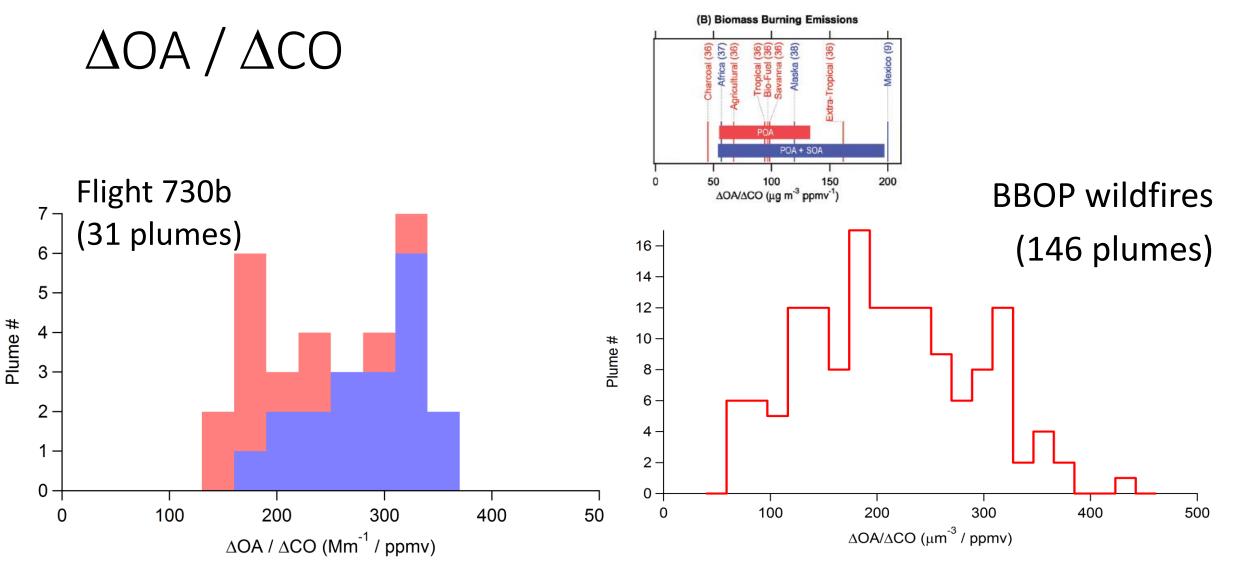


- f60 decreases with downwind distance
- O:C ratio increases with downwind distance
- G1 organics less oxidized than MBO, further down wind
- Observations are consistent with loss of primary OA and/or gain of secondary OA due to photochemical processes

## $\Delta \text{OA}$ / $\Delta \text{CO}$ downwind



- Differences in  $\Delta OA/\Delta CO$  with burn plume
- Constant △OA/△CO with downwind plume transport time



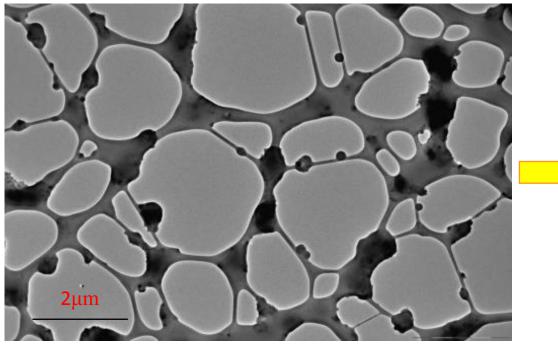
De Gouw et al. Environ. Sci. Technol. 2009, 43, 7614–7618

- $\Delta OA/\Delta CO$  for a single fire spans nearly the full range of observed wildfire plumes
- Near field measurements are higher than previous measurements

#### TEM observations downwind

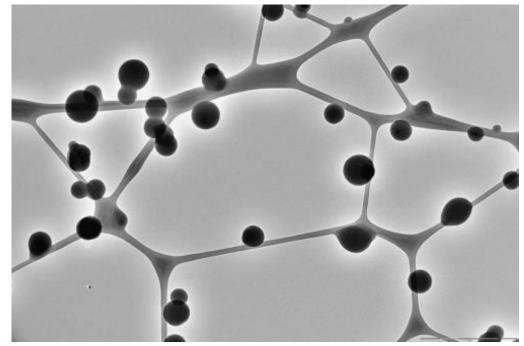
Fresh aerosol particles spread over the substrate

~0 hours



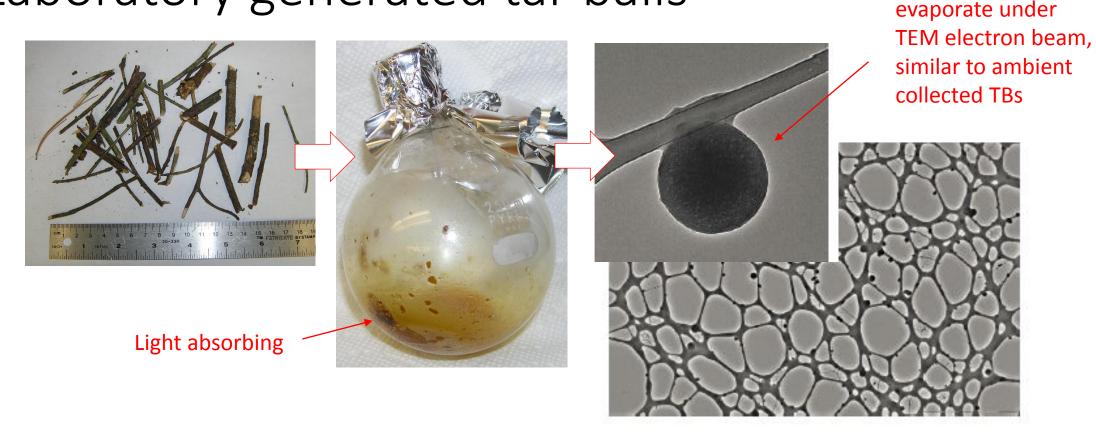
Aging results in highly spherical particles: Tar Balls

~2.5 hours



Loss and/or conversion of low viscous materials

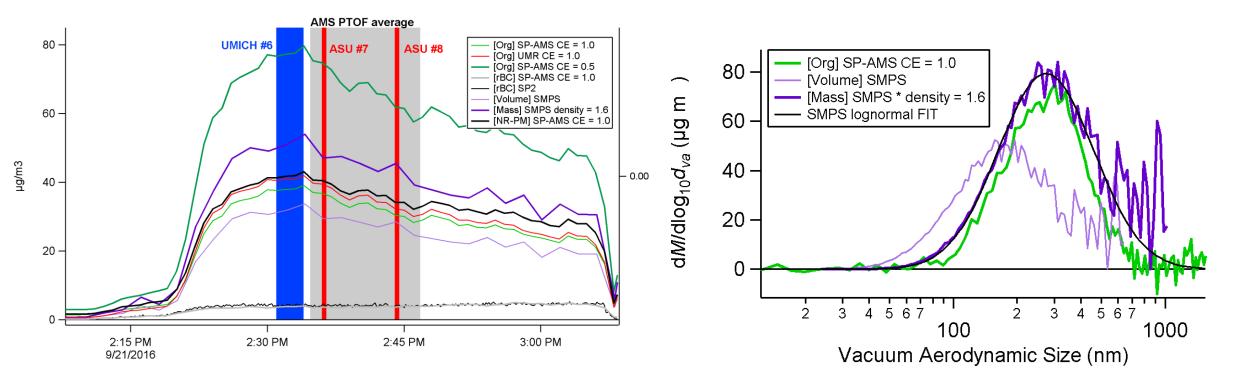
#### Laboratory generated tar balls



TBs slow to

 Tar ball formation and evolution were clearly observed in BBOP samples examined using electron microscopy. We initiated a laboratory study of tar balls to examine chemical and optical properties to augment field observations.

#### Characterized by SP-AMS, SMPS, SP2



- Observed AMS collection efficiencies (CE) of ~0.8, indicating that AMS is sensitive to TBs, but TBs are more refractory than liquid organics (less refractory than AS)
- SP2 and SP-AMS observed minor rBC component in lab TBs (<10%)
- Investigating chemical signatures for potential identification in field observations and correlations with optical properties

## Summary

- Rapid physical, chemical and optical changes in biomass burning particles measured downwind (< 3 hours) from wildland fires in Pacific Northwest.
- $\Delta EXT/\Delta CO$  and SSA increase downwind, indicating scattering increases faster than absorption.
- The chemical composition of OA is rapidly changing, with the O:C and OM:OC increasing and primary components decreasing. Governed by MCE and atmospheric chemistry.
- Organic aerosol loadings ([Org]/[CO]) appear to be relatively constant with time downwind suggesting that the competing evaporation of primary and condensation of secondary particulate material may be of similar order within the first few hours.
- Variability in different burn plumes from one wildland fire similar to the variability in chemical and optical properties for all sampled wildland fires
- TEM's observe formation of tar balls in field. Generated laboratory tar balls to investigate chemical/physical properties and real-time instrument sensitivities.

#### Acknowledgements

- Thanks to the G1 Flight Crew and Ground Crew and the other researchers on board for their help.
- Thanks to DOE Atmospheric System Research and ARM Climate Research Facility for funding and logistical support.





#### Discussion slide

- The DOE 2013 BBOP study is an important project with the correct methodology for providing needed information on biomass burning emissions and plume evolutions.
- Highly suggest that DOE ARM/ASR conduct a second BBOP project in concert with other biomass burning studies already scheduled in years 2018-2019 (FIREX, FIRECHEM, FASMEE, etc.) to take advantage of the planned intensives