Initial Study of the Roles of Chemical Composition and Meteorology on Aerosol Radiative Effects in the SE U.S.- Results from a Regionally-Representative Site

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Motivation for Aerosol Research in SE U.S. APP Monitoring Site

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III. Dependence of aerosol properties on meteorology and source region

IV. Conclusions

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# Introduction



#### Trenberth, et al., 2007

Southeastern US: one of a few regions in world not to have warmed in 20<sup>th</sup> century, but is beginning to warm in recent decades







Boone, NC, 36.2° N, 81.7° W, 1076 m



NASA - AERONET aerosol monitoring site Appalachian



Tower height 34 m



NOAA-ESRL aerosol monitoring site



Data products	Measurement techniques
Aerosol light absorption at 467 nm, 530 nm, 660 nm wavelengths (size-resolved, sub-μm, sub-10 μm)	Radiance Research Particle Soot Absorption Photometer (PSAP)
Aerosol total light scattering and hemispheric backscattering at 450 nm, 550 nm, and 700 nm (size-resolved, sub-µm, sub-10 µm)	TSI 3563 Nephelometer
Aerosol hygroscopic growth: total light scattering & hemispheric backscattering	TSI 3563 Nephelometer operating at a reference RH (< 40%) in series with a TSI 3563 scanning a higher RH range (up to 85%)
Aerosol number concentration	TSI 3010 Condensation Particle Counter (CPC)
Aerosol size distributions Spectral aerosol optical depth (AOD) at eight wavelengths between 340-1020 nm)	CIMEL 318 Solar-Tracking Sun/Sky Radiometer
Aerosol chemical composition (size- resolved, sub- µm)	Aerodyne Aerosol Mass Spectrometer (AMS)
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#### Initial Studies of Relationships between Aerosol Radiative Properties, Aerosol Chemistry, Meteorology, and Source Region

- Lower tropospheric aerosol properties have been measured continuously since June 2009
- Column-averaged and vertically-resolved aerosol properties have been measured near-continuously since August 2010 and April 2011, respectively
- This <u>initial study</u> will focus on a 7-week period (June 4-July 21, 2012), when the aerosol chemical composition measurements were initiated. This period will be referred to as 'Summer 2012' in the presentation
- The upcoming summer 2013 field campaign and subsequent measurements will provide opportunity to expand upon this study





Summer 2012 statistics of non-refractory aerosol chemical composition



Ammonium sulfate: ~7.5  $\mu$ g/m<sup>3</sup>, Particle organic matter: ~ 4.0  $\mu$ g/m<sup>3</sup> IMPROVE 2005-2008 regional monthly mean PM <sub>2.5</sub> in June in Southern Appalachian mountain region) [*Hand et al.*, 2012]

 $(NH_4)SO_4$ : 54% Org.: 36% in  $PM_{2.5}$  in summer at rural sites in southeastern US. (*Zhang et al.,* 2012)





## **Chemical composition relationships**



### Dependence of aerosol properties on synoptic meteorology













Direct Radiative Forcing Efficiency at 550nm (6/4/2012-7/21/2012)





Scattering Hygroscopic Growth Factor (6/4/2012-7/21/2012)





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# Dependence of aerosol properties on source region

Source Region Code	Туре	# Hours	% Hours	-120 110 -100 -90 y -90 y
0	Back trajectories did not clustered	483		
1	Long range transport from West/Southw est	0	49%	a solution of the solution of
2	Long range transport from north	48	27%	
3	Southeast/loc al	168	24%	12 24 36 48 60 72













# Conclusions

- Organics dominated the non-refractory aerosol mass loading during the summer 2012 period, consistent with other recent studies in the region (Goldstein, 2008).
- Aerosol single scattering albedo and hygroscopic growth factor increased with sulfate fraction. Aerosol hygroscopic growth factor decreased with increased organic fraction.
- Aerosol organic fraction, mass loading and extinction coefficient increased with ambient temperature due to the increased photochemical formation of secondary organic aerosol and highly temperature-dependent BVOC emissions from forests.
- Aerosol originating in the SE U.S. (local) had relatively high organic fraction consistent with the forested and rural environment. Aerosol transported from the west and southwest corresponded to high sulphate fraction. Aerosol transported from the north had anthropogenic influences with relatively lower single scattering albedo.
- The results are based on ~7 week measurements. Long term observations are needed to improve the understanding of the relationships among aerosol properties, source and meteorology.



# Acknowledgements

<u>Appalachian State University :</u> College of Arts and Sciences, Office of Research and Sponsored Programs, Research Institute for Environment, Energy, and Economics

<u>Undergraduates</u> Ian Krintz, Craig Stewart, Jason Hightower, Matt Karas

<u>Graduate Students</u> Colonel William Bullitt Beuttell John Markham

ASU Faculty Brett Taubman

<u>ASU Staff</u> Mike Hughes Dana Green Robert "Butch"Miller AppalAIR Collaborators Patrick Sheridan (NOAA-ESRL) John Ogren (NOAA-ESRL) Betsy Andrews (NOAA-ESRL) Brent Holben (NASA AERONET)

<u>AMS community</u> John Jayne, Jose Jiminez, James Allan, Alice Delia

<u>HYSPLIT model</u> Air Resources Laboratory (NOAA)



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#### Summer 2012 non-refractory aerosol size distribution





**Figure 1:** Summer 2012 time serials of total non-refractory aerosol mass loading (AMS  $\mu$ m/m<sup>3</sup>), aerosol optical thickness (AOT) at 500 nm, and sub-10 $\mu$ m aerosol scattering ( $\sigma_{so}$ ) and absorption ( $\sigma_{ap}$ ) at 550 nm







Summer 2012 time serials of single-scattering albedo ( $\omega_0$ ), back-scatter fraction (*b*) at 550 nm, Angstrom exponent (a) and sub-1um scattering fraction (R sp)



Aerosol optical depth (AOD) at 500 nm, and sub-10µm aerosol extinction ( $\sigma_{ext}$ ), single scattering albedo ( $\omega_{o}$ ), and back scattering fraction (*b*) at 550 nm





Aerosol optical depth (AOD) at 500 nm, and sub-10µm aerosol extinction ( $\sigma_{ext}$ ), single scattering albedo ( $\omega_{o}$ ), and back scattering fraction (*b*) at 550 nm





 $f(RH) = \frac{\sigma_{bsp}(85\%)}{\sigma_{sp}(40\%)}$ 





Summer 2012 statistics of aerosol optical thickness (AOT) at 500 nm, and sub-10µm aerosol scattering ( $\sigma_{sp}$ ), absorption ( $\sigma_{ap}$ ), extinction ( $\sigma_{ext}$ ), single-scattering albedo ( $\omega$ ), back-scatter fraction (*b*) at 550 nm, and Angstrom exponent (a)

	AOT_50	0 σ <sub>sp</sub>	$\sigma_{ap}$	$\sigma_{ext}$	
Mean	0.21	44.2	3.0	47.1	
Stdev	0.10	25.3	1.3	26.2	
					_
	ω	b	а	R <sub>ap</sub>	R <sub>sp</sub>
Mean	0.927	0.146	0.598	0.90	0.94
Stdev	0.038	0.023	0.264	0.08	0.10

SGP 1996-2000  $\sigma$  ap: 2.8 (June), 2.4 (July)  $\sigma$  ap: 50 (June), 48 (July)  $\Omega$ : 0.92 (June), 0.95 (July) Sheridan et al., 2001







120 Extinction Coefficient 100 80 60 40 20 0 0.1-0.2 0.2-0.3 <0.1 0.3-0.4 >0.4 ALL SO4 Fraction Hemispheric Backscatter Fraction at 550nm (6/4/2012-7/21/2012) 0.22 0.2 0.18 0.16 هـ 0.14 0.12 0.1 <0.1 0.1-0.2 0.2-0.3 0.3-0.4 ALL >0.4 SO4 Fraction

Aerosol Extinction Coefficient at 550nm (6/4/2012-7/21/2012)

140

#### 155

#### Meteorological relationships

























#### Case study





# **Time series in Summer 2012**

# Chemical composition



## Size distribution

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Aerosol optical depth (AOD) at 500 nm, and sub-10µm aerosol extinction ( $\sigma_{ext}$ ), single scattering albedo ( $\omega_{o}$ ), and back scattering fraction (*b*) at 550 nm





Aerosol hygroscopic growth factor (f(RH)) and directive radiative forcing efficiency (DRFE)







