



Evaluation of WRF-Chem Simulations of Carbonaceous and Inorganic Aerosols

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• Utilize extensive measurements collected during the CalNex and CARES campaigns to evaluate regional predictions ($\Delta x = 4 \text{ km}$) of aerosol mass, composition, size distribution, and optical properties

clean' conditions observed

challenging for models to accurately represent 'clean' conditions

- Understand the sources of uncertainties associated with simulated aerosol radiative forcing over California
- Determine the relative importance of long-range transport and local emission sources on aerosol loading and radiative forcing



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Weather Research Forecasting (WRF) Model

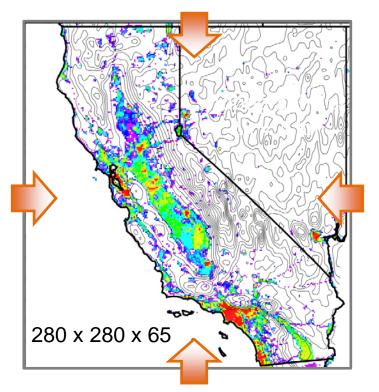
Meteorology:

- Boundary Layer: MYJ
- Land Surface: Noah
- Radiation: RRTMG
- Microphysics: Morrison
- Convection: new Kain-Fritsch
- IC/BC: GFS + analysis nudging

Chemistry:

- Trace Gases: SAPRC
- Photolysis: FTUV
- Aerosols: **MOSAIC**, 8 size bins,
- Volatility Basis Set approach for SOA
- Direct effect on, indirect effect off
- Wet Scavenging: off
- ► IC/BC: MOZART

Simulation Period: May 1 – June 30, 2010



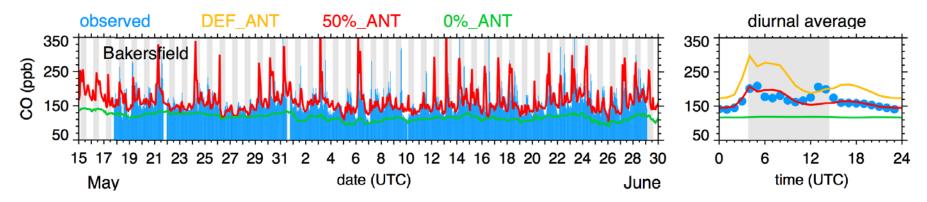
Effects of Long-Range Transport of Trace Gases and aerosols from Global Model



Four WRF-Chem Simulations



- Default: CARB 2008 inventory
 Sensitivity 1) 50% CARB inventory (except SO₂ and NH₃)
 Sensitivity 2) no anthropogenic emissions
 Sensitivity 3) same as 1), but MOZART aerosol for 50%_LBC boundary conditions reduced by 50%
- Initial simulations (as well as other modeling studies) suggest that CARB 2008 inventory is too high, especially for CO, NOx, POA, and BC
 - For example at Bakersfield:



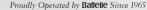


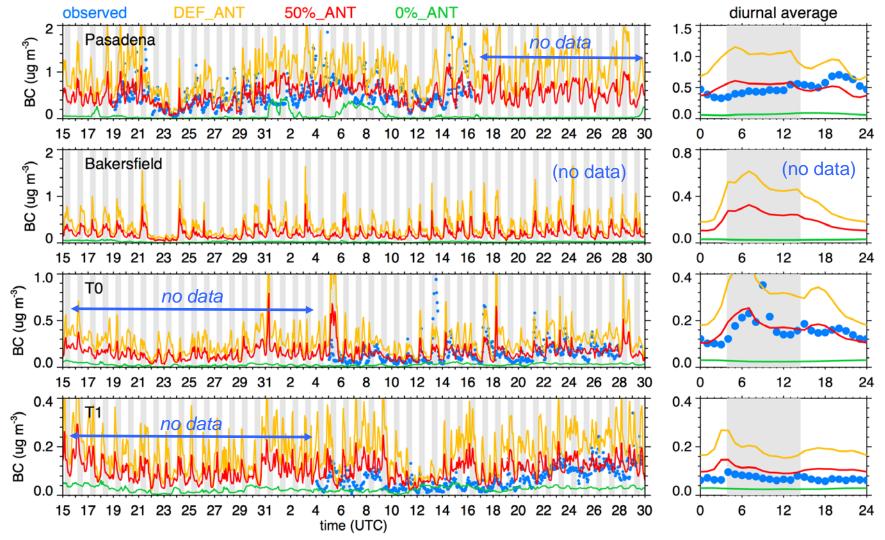
Proudly Operated by Battelle Since 1965

Results: Aerosols

Black Carbon at the Supersites





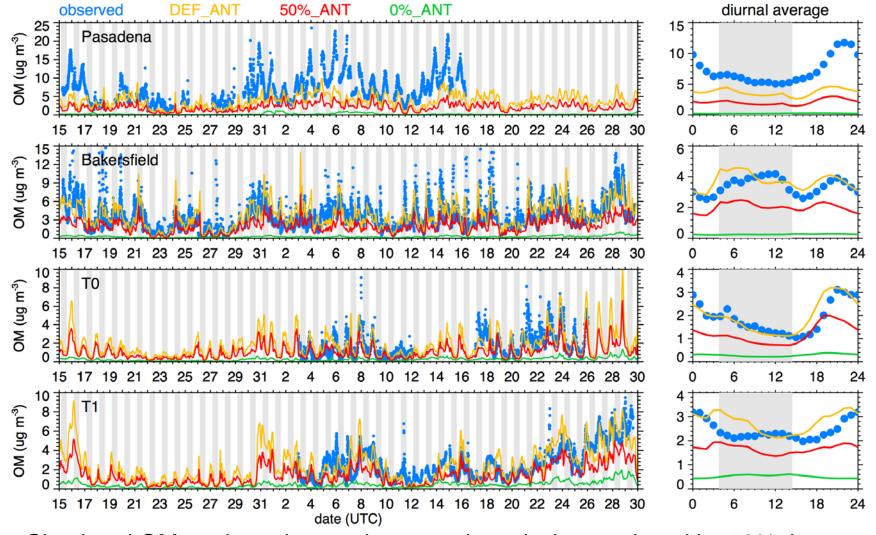


Simulated BC improved when anthropogenic emissions reduced by 50%, but correlation could be improved

Organic Matter at the Supersites



Proudly Operated by Battelle Since 1965

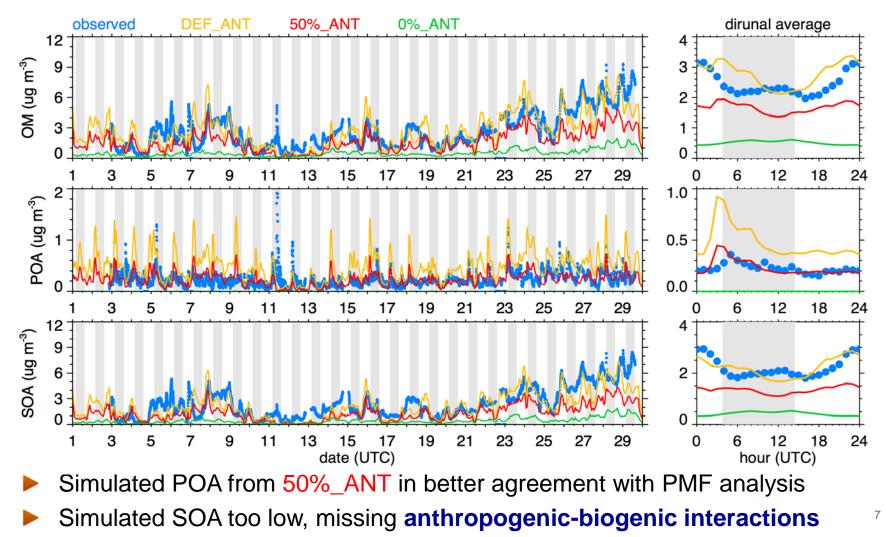


Simulated OM too low when anthropogenic emissions reduced by 50%, but need to examine components of organic aerosols ...

Organic Aerosol Components at T1



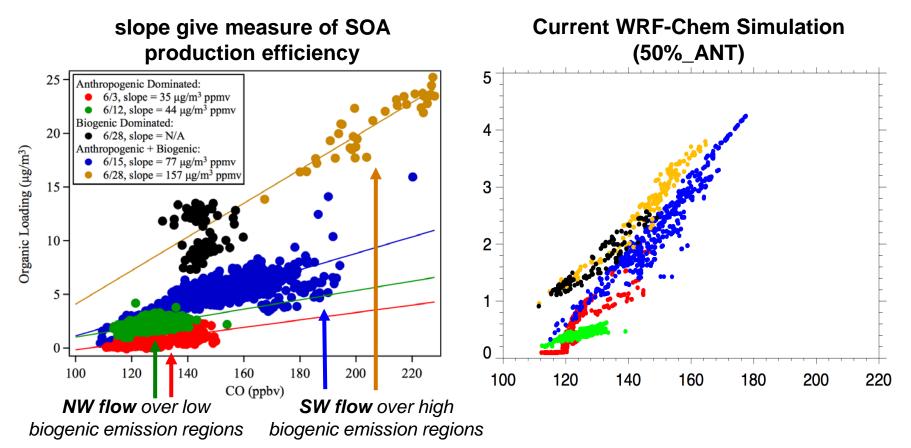
Use of Postive Matrix Factorization to Determine POA and SOA from Aerosol Mass Spectrometer (AMS) data



Role of Anthropogenic-Biogenic Interactions



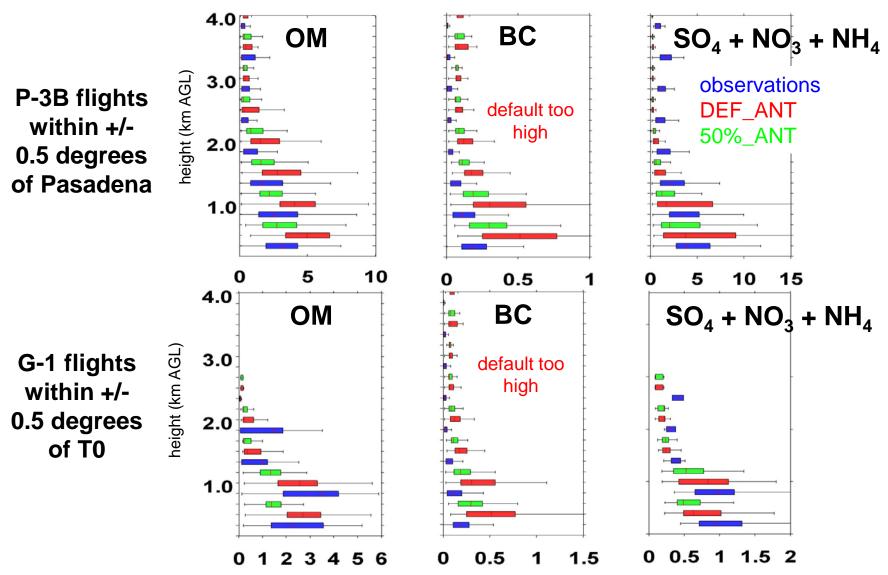
 Shilling et al., ACP (2013) use G-1 AMS and PTR-MS data to show that OM is enhanced when anthropogenic and biogenic emissions mix together



Chemistry associated with this enhancement is not well known, and therefore not represented by treatments of SOA formation

Aerosol Composition Aloft

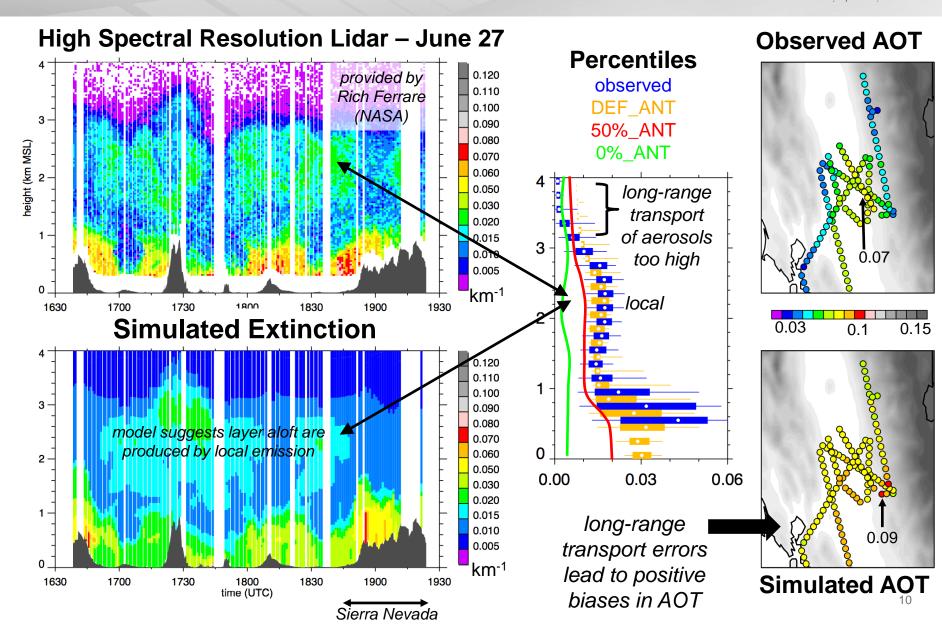




mass concentration (µg m⁻³)

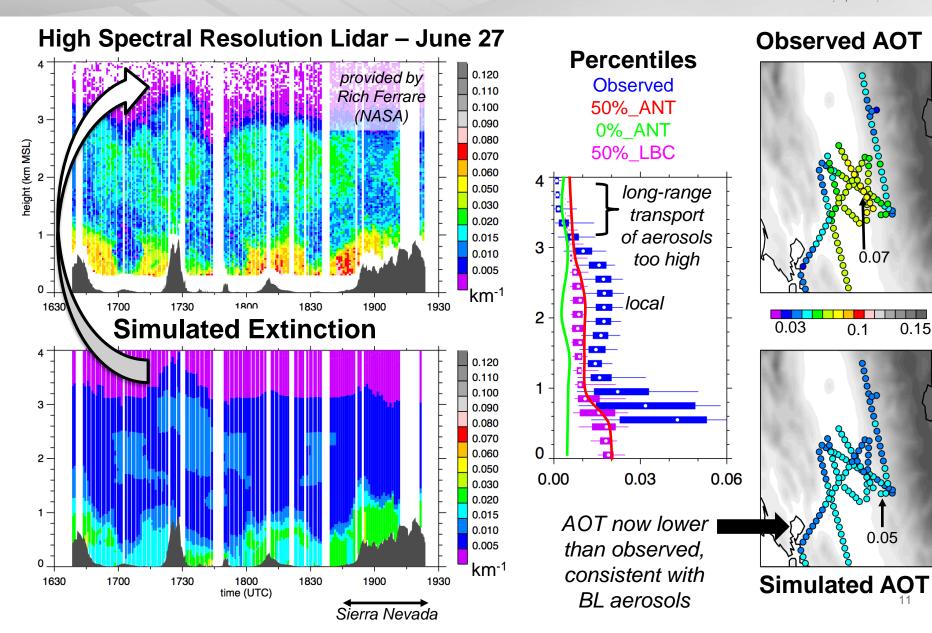
Extinction Profiles – Sacramento Valley





Extinction Profiles – Sacramento Valley









- Model is able to capture general spatial and temporal variations in aerosols reasonably well; however, simulated ...
 - **BC** too high unless emission rates reduced
 - **OM** too low (uncertainties in SOA), especially in southern CA
 - **SO₄** too low in southern CA, better represented in northern CA
 - **NO₃** too low everywhere, but better represented in southern CA
- Spatial variations in extinction and AOT agree reasonably well with Lidar, errors in magnitude consistent with errors seen with in-situ data
- Emission inventories are not the only source of error
 - Long-range transport of aerosols likely too high (both dust and anthropogenic)
 - Adjustments in emissions likely depends on region
- High bias in long-range transport leads to a positive bias in AOT and thus total radiative forcing (bias in transport offset to some extent by low bias in local aerosol sources)