

Seasonal and interannual variability of upper tropospheric aerosols: Sources and the role of Asian monsoon transport

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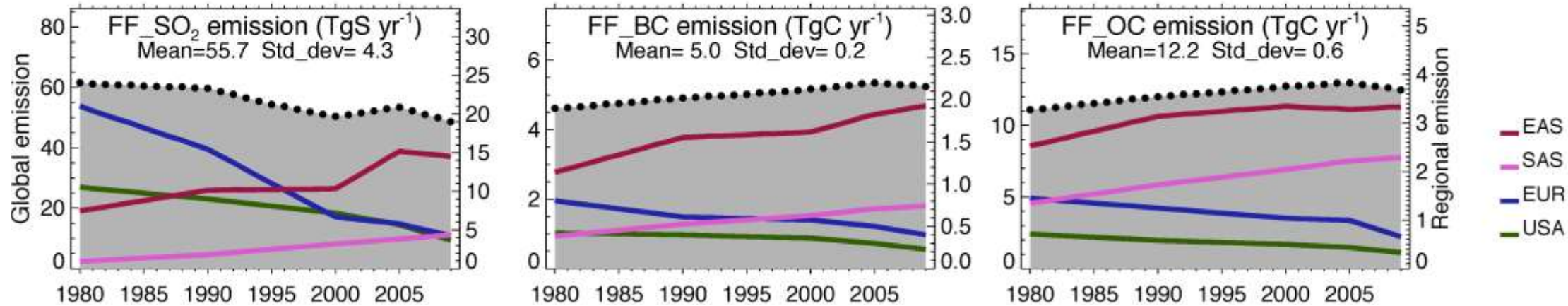
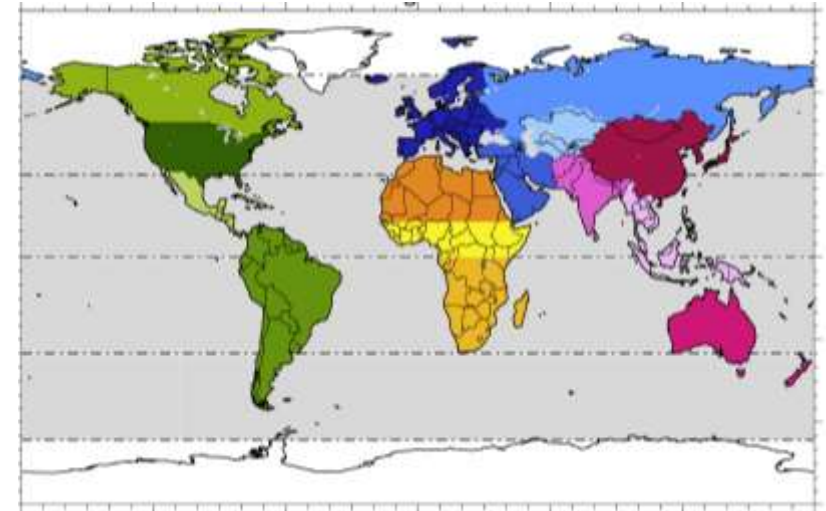


Introduction

- The origin and variability of stratospheric aerosol have drawn considerable attention because the change of such aerosol could have long-term climate effects
- Recent observations seem to suggest that the stratospheric aerosol has been increasing in the past decade without major volcanic eruptions
 - Is the increase due to the Asian anthropogenic emission?
 - Or volcanoes?
- This work uses a global model to estimate the aerosol sources in the UTLS region and to elucidate the role of Asian monsoon transport

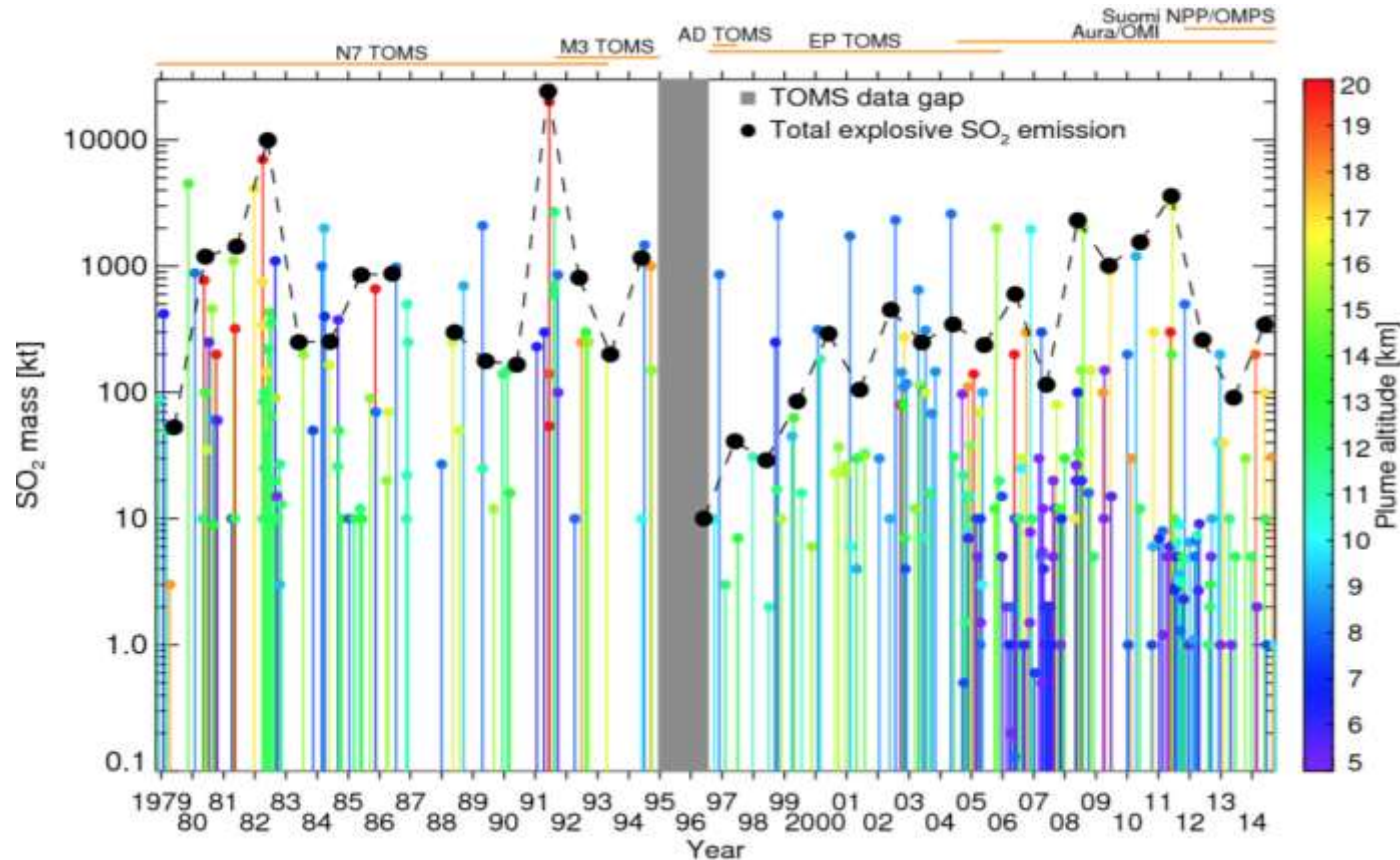
Anthropogenic emission

- Anthropogenic SO₂ (and other pollutants as well) emissions in East Asia and South Asia have increased significantly in the last decade
- EAS emission is much higher than SAS
- The question is: How efficient the transport is to lift surface pollution to the UTLS?



(Figures from Chin et al., 2014)

Volcanic SO₂ emission



- Volcanic emissions release SO₂ usually at higher altitudes than anthropogenic emissions to have a more direct influence in the UTLS

SO₂ emission from eruptive volcanoes from 1979 to 2014. Data source: Carn et al., 2015).

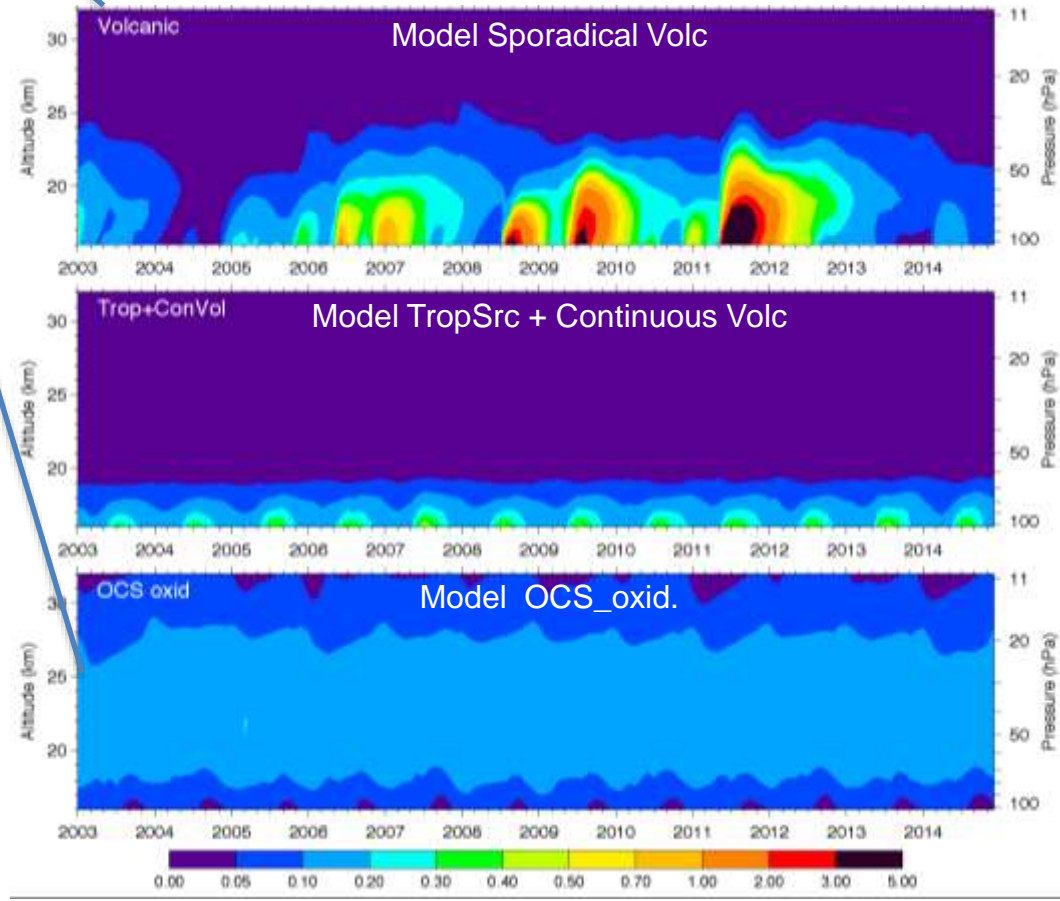
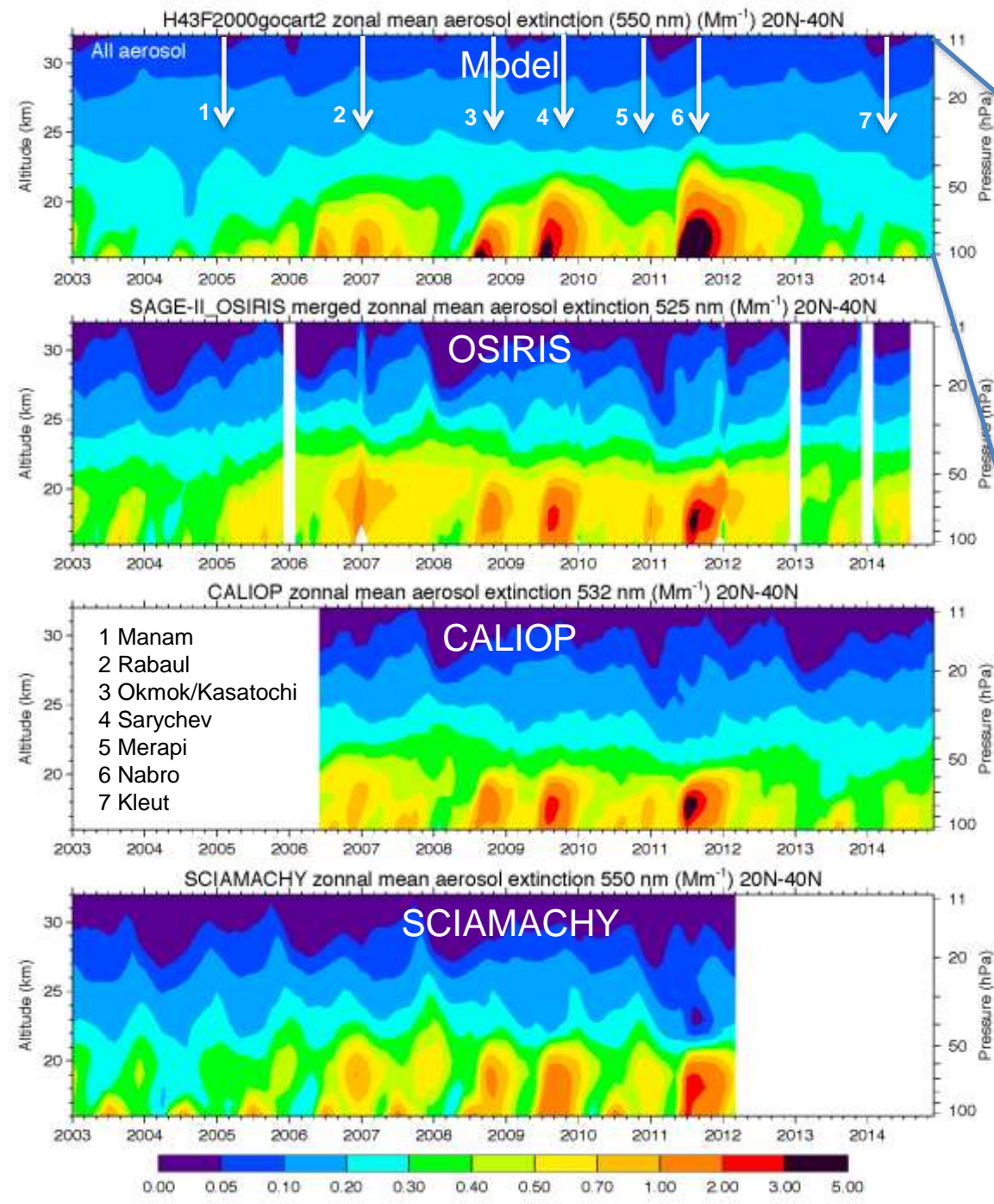
Model simulations

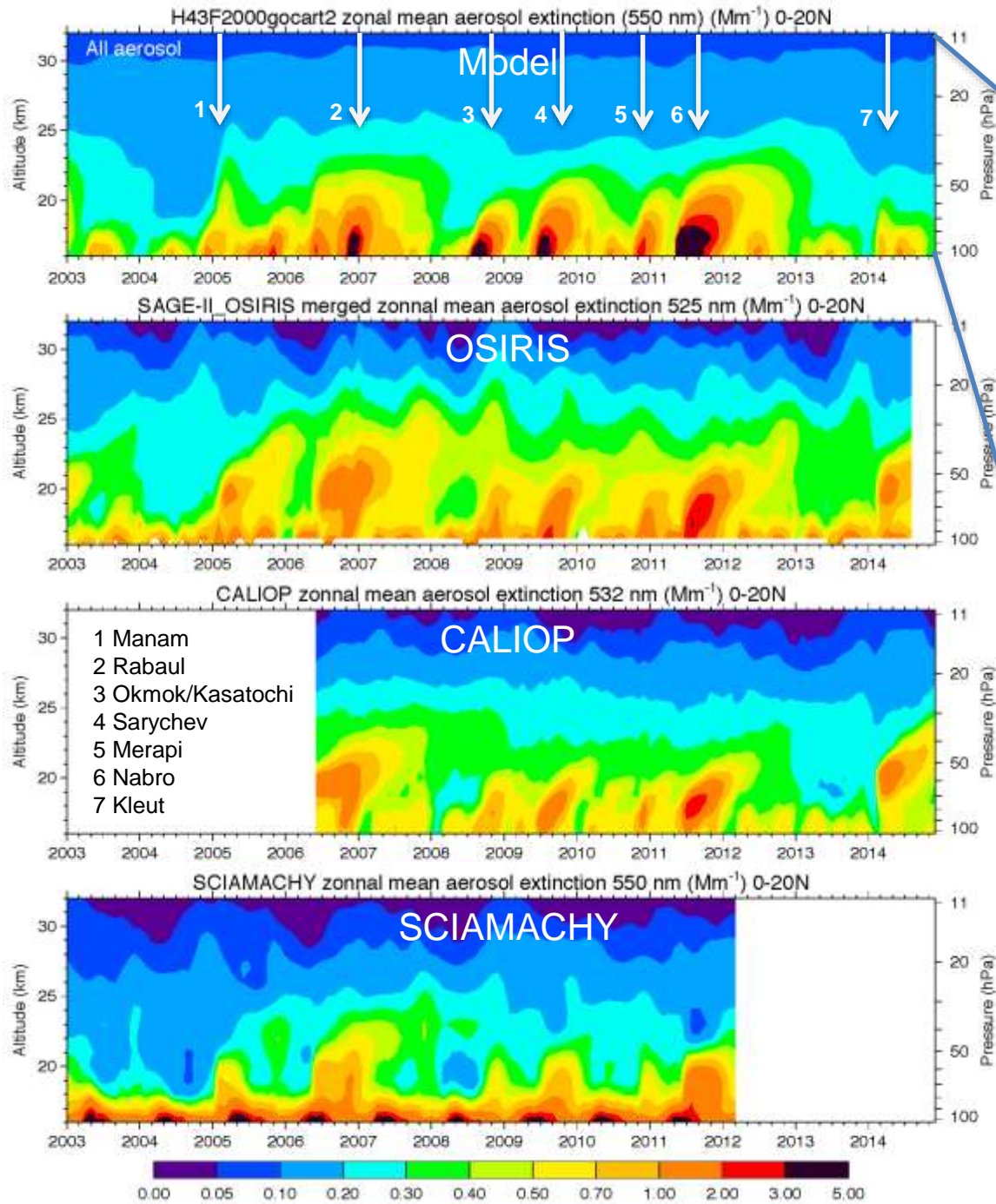
- Model simulations:
 - GEOS-5 AGCM/GOCART, 2.5°lon x 2° lat horizontal resolution, 72 vertical layers
 - Anthropogenic and biomass burning emission: ACCMIP (Granier et al., 2011)
 - Volcanic emission: OMI-based sporadically erupting volcanic emission (Carn et al., 2015) + continuously erupting volcanic emission (Andres and Kasgnoc, 1998)
 - Sulfate from OCS oxidation included
 - Aerosols identified from (1) sporadically erupting volcanic source, (2) stratospheric background source (OCS oxidation), and (3) other (anthropogenic + biomass burning + non-volcanic natural + continuously erupting volcanic sources)
- Time period of this study: 2003-2014

Comparison with satellite aerosol data

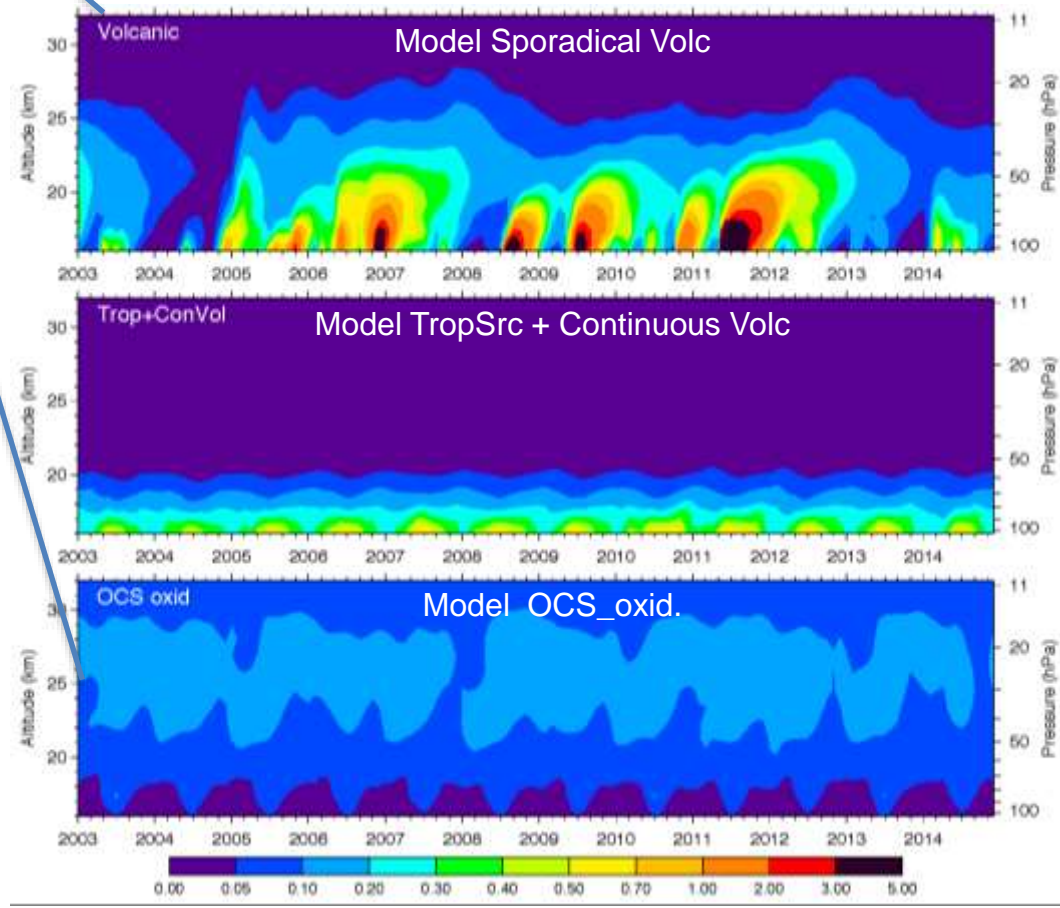
- OSIRIS:
 - V5-07 level-3 monthly zonal averages at 5° latitude resolution and 1-km vertical resolution from 0-40 km (provided by U. Saskatchewan group, POC: Landon Rieger)
 - Merged SAGE-II and OSIRIS: extinction at 525 nm
- CALIOP:
 - Stratospheric aerosol V6, monthly zonal average extinction at 532 nm at 5° latitude resolution, 8 to 40 km (provided by Jean-Paul Vernier, LaRC)
- SCIAMACHY:
 - V1.1. level 3 monthly averages at 5°x5° horizontal resolution and 1-km vertical resolution from 9-40 km (provided by U. Bremen group, POC: Alexei Pozanov)
 - 550 nm extinction was interpolated from 470 and 750 nm using the Angstrom Exponent

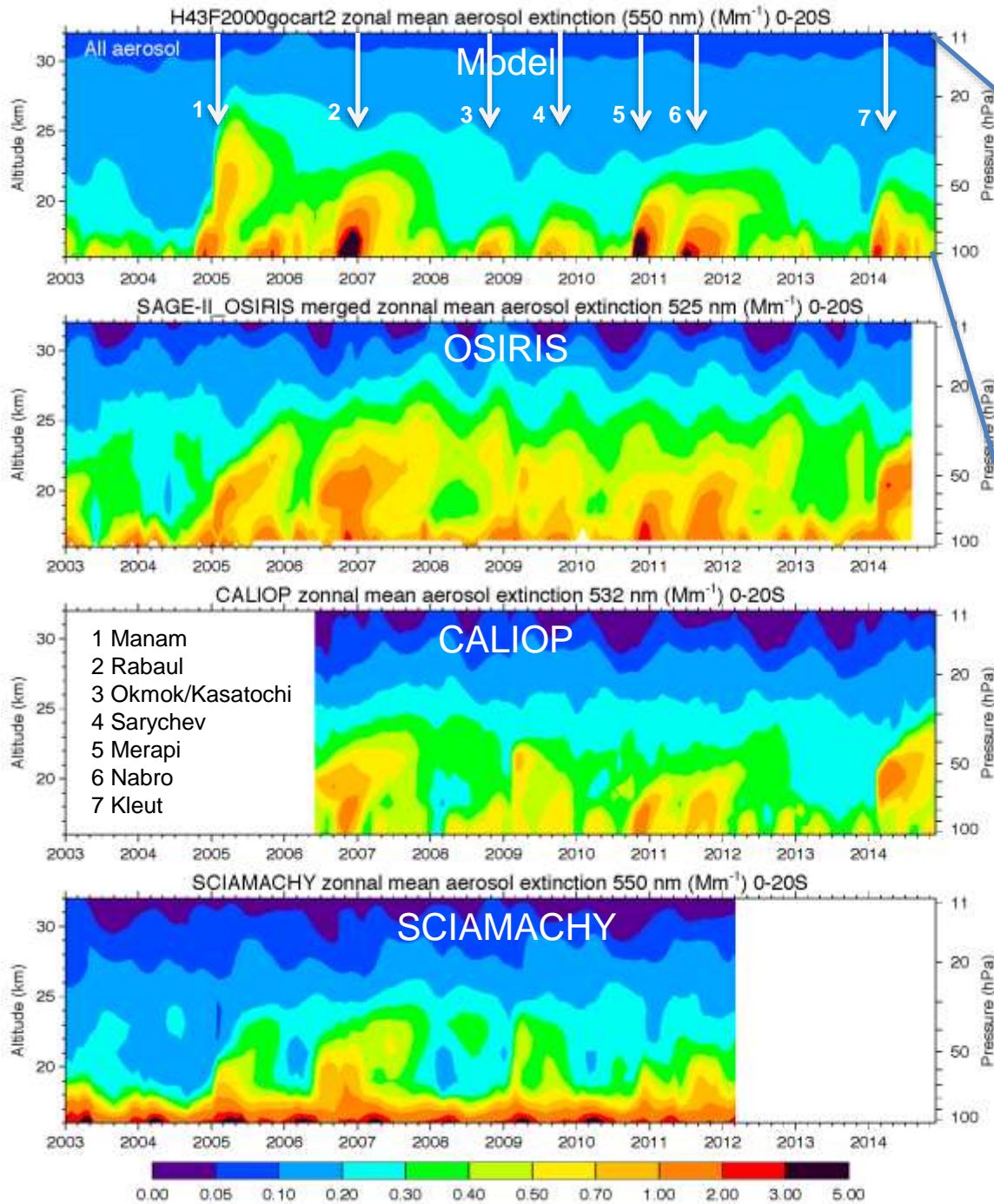
Zonal mean aerosol extinction at 550 nm (Mm^{-1}), 20N-40N



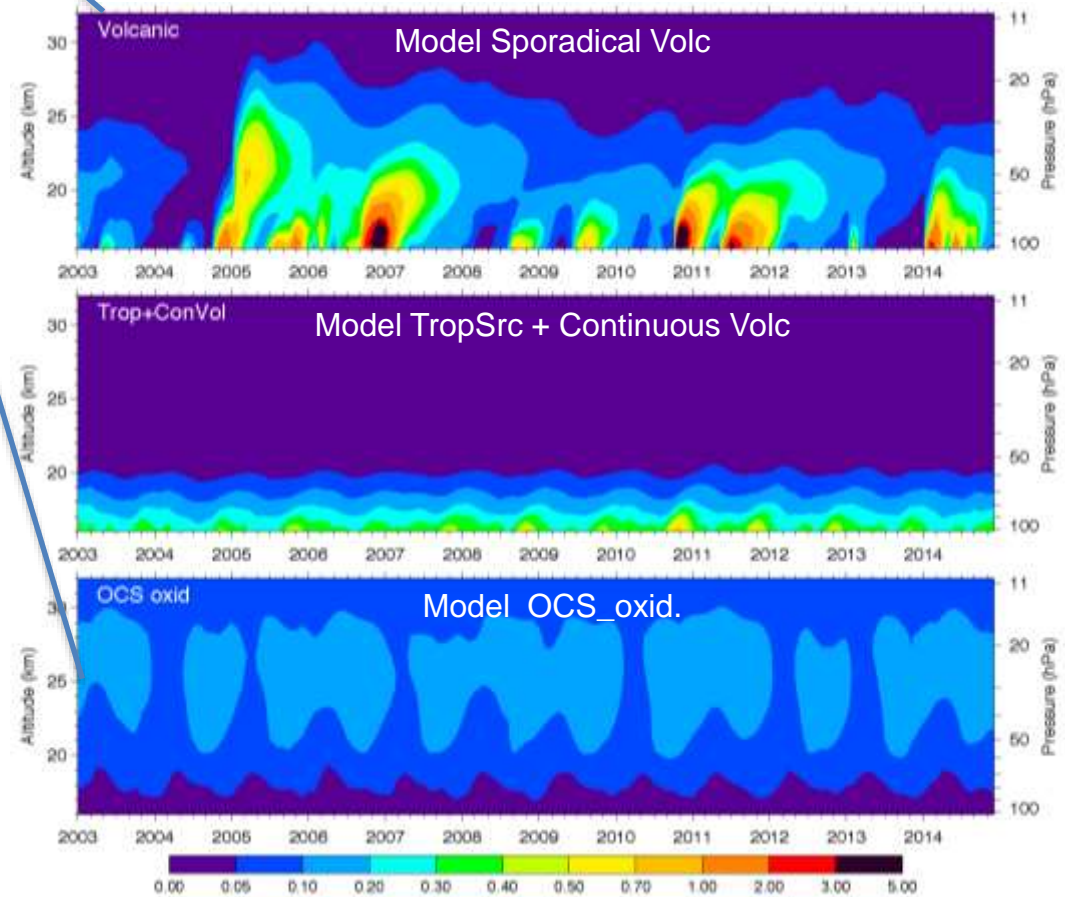


Zonal mean aerosol extinction at 550 nm (Mm^{-1}), 0-20N





Zonal mean aerosol extinction at 550 nm (Mm^{-1}), 0-20S

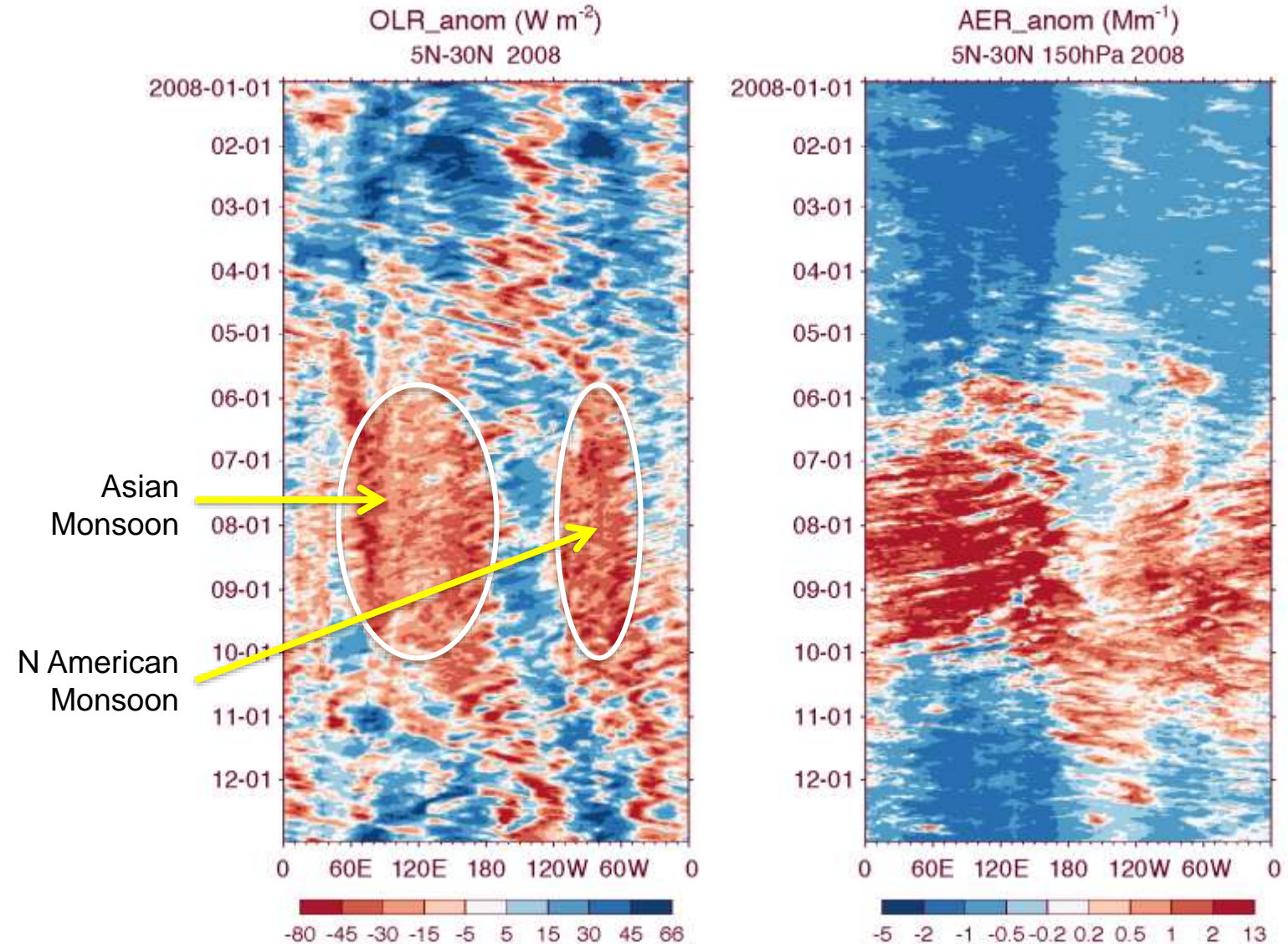


Source attribution – volcanic, anthropogenic, and background

- Overall, the volcanic aerosol dominates the stratospheric aerosol loading even without Pinatubo-scale large eruption (4-5 higher than anthropogenic)
- Near the tropopause, anthropogenic aerosol shows a well organized, repetitive seasonal cycle
- The “background” sulfate aerosol from OCS oxidation is the most important aerosol source >25 km

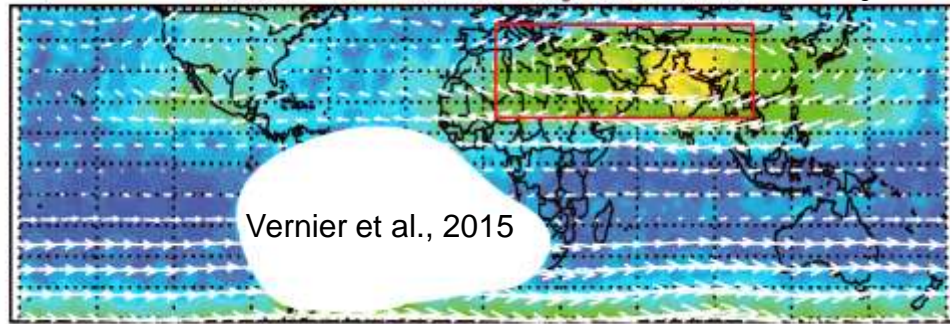
Seasonal variations of deep convective transport: Average OLR and UT aerosol anomaly, 5-30N

- In the subtropical northern hemisphere, the most pronounced convective features are the Asian summer monsoon and the North American summer monsoon
- The monsoon convections pumps aerosols to the upper troposphere, even though the convection always associated with heavy rainfall
- Composition of UT aerosol varies from year to year depending on the variation of aerosol sources

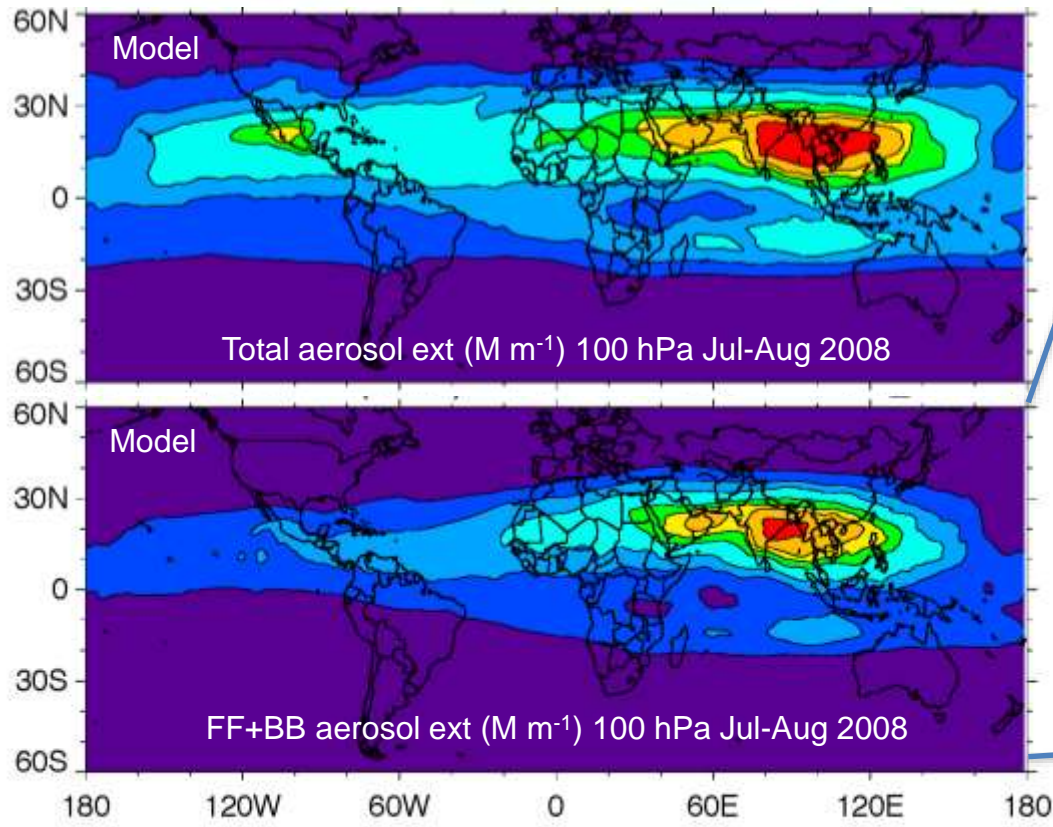


Maximum aerosol in ATAL

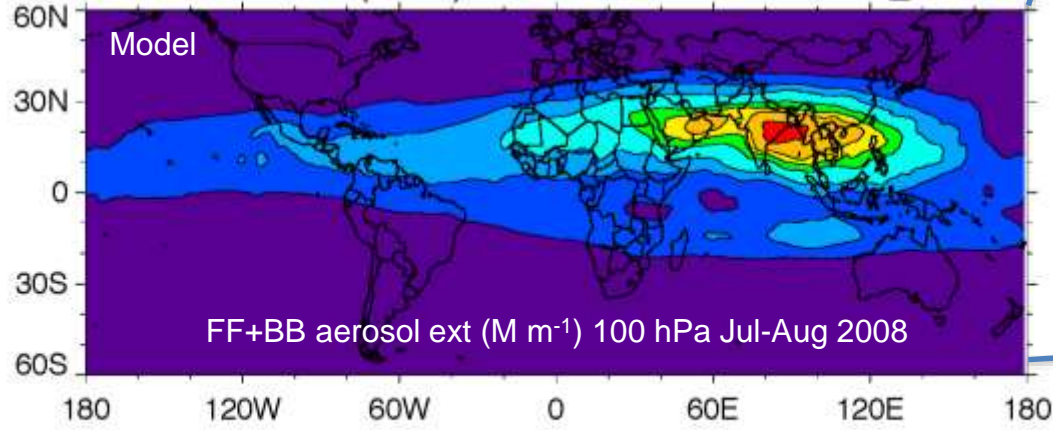
b) CALIOP 15–17km Jul–Aug 2006–2013 Scattering Ratio



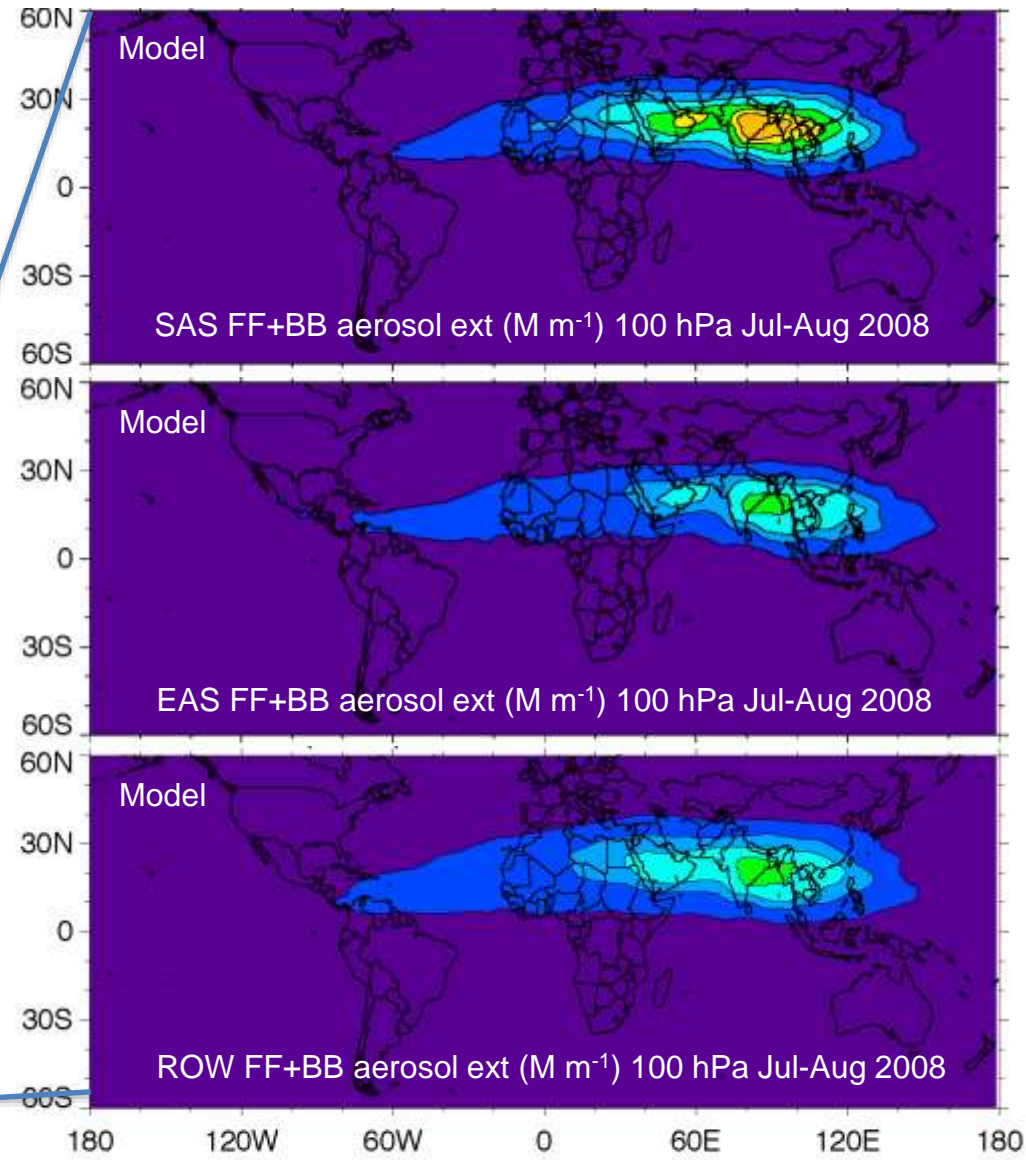
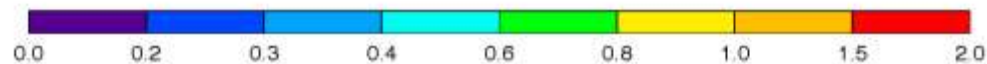
Vernier et al., 2015



Total aerosol ext ($M m^{-1}$) 100 hPa Jul-Aug 2008



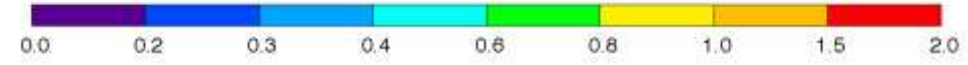
FF+BB aerosol ext ($M m^{-1}$) 100 hPa Jul-Aug 2008



SAS FF+BB aerosol ext ($M m^{-1}$) 100 hPa Jul-Aug 2008

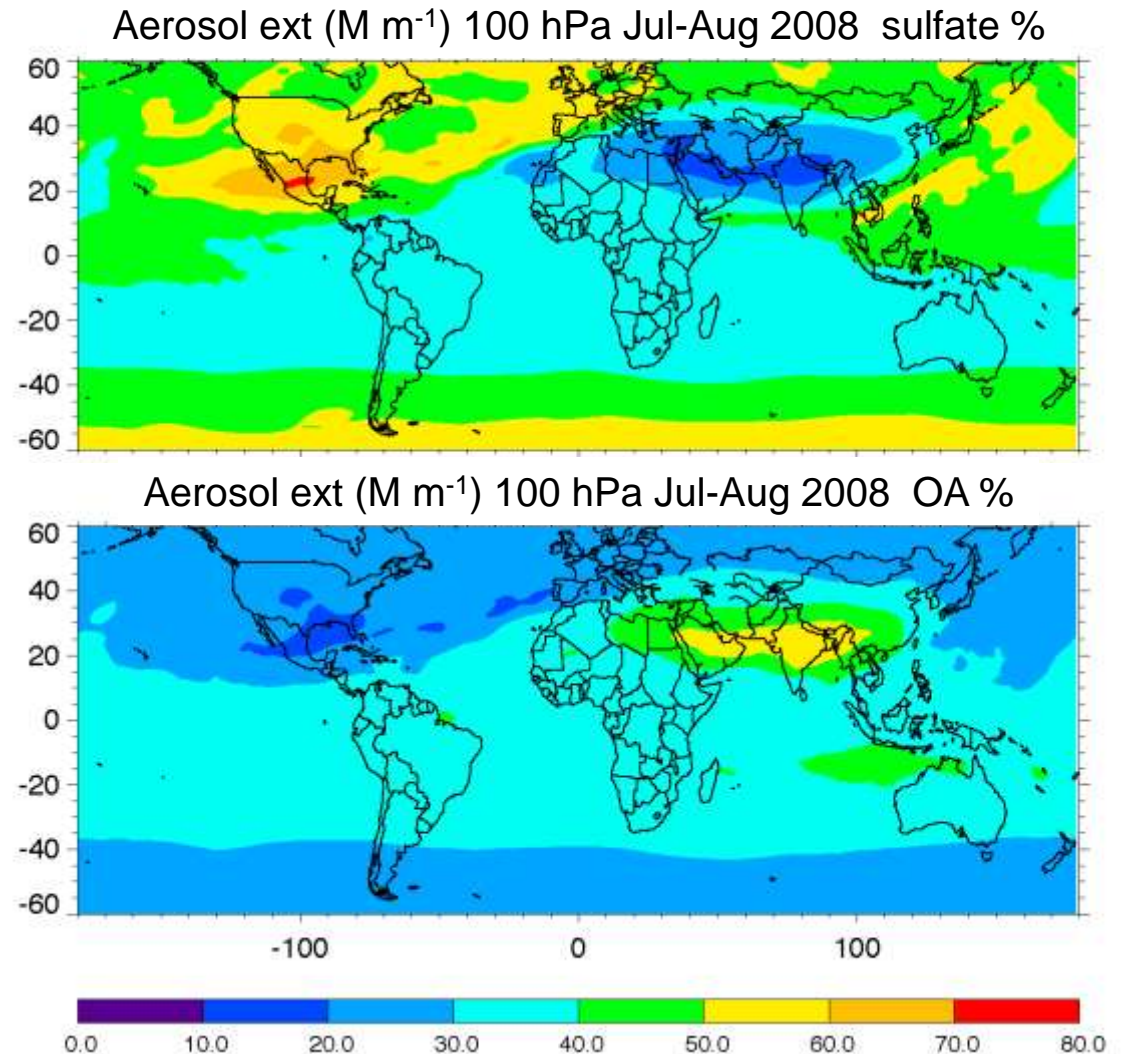
EAS FF+BB aerosol ext ($M m^{-1}$) 100 hPa Jul-Aug 2008

ROW FF+BB aerosol ext ($M m^{-1}$) 100 hPa Jul-Aug 2008



Aerosol composition in the UT monsoon region from GOCART simulation

- GOCART simulations suggest that over the UT Asian monsoon region, organic aerosol accounts for 50-60% of total aerosol extinction at 550 nm while sulfate only 10-30%
- We need measurements to verify aerosol composition
- Further investigation of chemical and physical mechanism determining the aerosol composition in the UT region is in progress



Concluding remarks

- By model experiments separating anthropogenic and natural sources, we have found that
 - volcanic aerosol dominates the total stratospheric aerosol amount even without very large volcanic eruptions like Pinatubo
 - anthropogenic aerosol exhibits well organized seasonal cycle in the tropopause region
- Strong summer monsoon convection in the subtropical northern hemisphere making transport of aerosols to UTLS most effective in the summer
 - SAS anthropogenic aerosol dominates the ATAL aerosol in summer (60-70%), even though EAS anthropogenic emission is much higher than SAS
 - Not all aerosols in ATAL are from Asia – other regions contribute as well
- Aerosol composition from the model suggest that OA is the most component in the UT Asian monsoon region – verification needed!