

# WHY

Aerosol optical depth (aot) comparisons to data from ground and space are preferred ways to demonstrate the skill of aerosol modules in global modeling. Comparisons among aerosol module detail demonstrate strong differences at sub-components, which may go unnoticed when looking at integrated properties. Specifically we have to wonder: *Are 'good' aot totals skillful, just luck (off-setting errors) or a matter of tuning?* Investigations of detailed aerosol output of control experiments as proposed in AEROCOM will tell.



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# Simulated aerosol components global fields of yearly averages and evaluations

Models	Resolution	Simulation	authors
• LO LOA	3.8/2.5deg	yr 2000	Reddy / Boucher
• LS LSCE	3.8/2.5deg	yr 2000	Schulz / Balkanski
• UL ULAQ	22.5/10deg	yr 2000	Pitari / Montenaro
• SP SPRINTARS	1.1/1.1deg	yr 2000	Takemura
• CT CANADA	2.8/2.8deg	yr 2000	Gong
• MI MIRAGE	2.5/2.0deg	1yr avg	Ghan / Easter
• EH ECHAM5 HAM	1.9/1.8deg	3yr avg	Stier / Feichter
• NF NCAR MATCH	1.9/1.9deg	yr 2000	Fillmore / Collins
• OC OSLO-CTM	2.8/2.8deg	yr 1996	Myhre / Isaksen
• OG OSLO-GCM	2.8/2.8deg	3yr avg	Iversen et al.
• IM IMPACT	2.5/2.0deg	yr 2000	Liu / Penner
• GM GFDL MOZART	2.5/2.0deg	yr 2000	Ginoux / Horowitz
• GO GOCART	2.0/2.5deg	yr 2000	Chin / Diehl
• GI GISS	4.0/5.0deg	yr 2000	Koch / Bauer
• TM TMS	4.0/6.0deg	yr 2000	Krol / Dentener
• EM ECHAM4 MADE	3.8/3.8deg	10yr avg	Lauer / Hendricks
• GR GRANTOUR	5.0/5.0deg	1yr avg	Herzog / Penner
• NM NCAR MOZART	1.9/1.9deg	1yr avg	Tie / Brasseur
• NC NCAR CAM	2.8/2.8deg	1yr avg	Mahonwald
• EL ECHAM4	3.8/3.8deg	3yr avg	Lohmann / Feichter
• HA HADAM4	3.8/2.5deg	3yr avg	Roberts / Jones

## RESULTS

- better mode agreement on source location, but differences in strength
- large differences in simulated transport (incl. removal processes)
- large differences in modeled aerosol water impacts  $m \rightarrow aot$  conversions
- simulated (total) aot seems low over tropical oceans and tropical biomass

## RESULTS

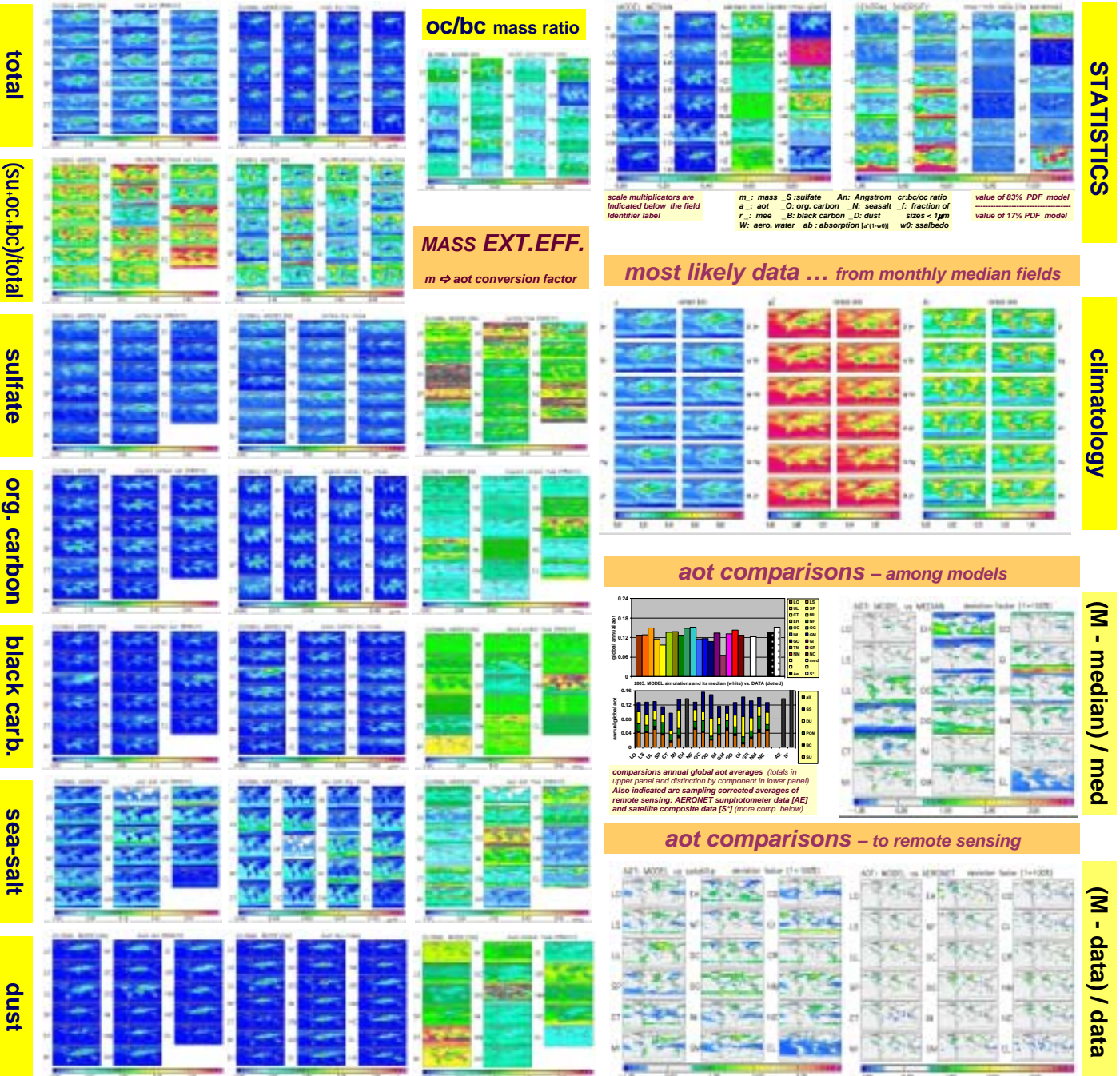
Human activity has increased atmospheric concentrations of greenhouse gases and aerosol. Our understanding of associated climatic impacts is largely based on global modeling. With respect to aerosol, though, uncertainties remain large. For an improved representation, new aerosol modules in recent years started to distinguish between sulfate, organic carbon, black carbon, dust and sea-salt components. Here, simulations of 21 modules are presented (most of them participate in the **AeroCom** model diagnosis). Here simulated global fields of mass (m) and optical depth (aot) are compared (both are intermediate products on the way to radiative forcing – quantifying the climatic impact). **Model diversity for totals (m, aot) is smaller (!) than for almost all sub-components.** Contributing factors are also differences for  $m \rightarrow aot$  conversion factors (mass ext. eff.), which rely on assumptions (size, humidification, available water) that need to be further investigated. Thus, (dis-) agreement of aot totals among models and to measurements from remote sensing (samples are below) are insufficient for model evaluations. Moreover, as different component contributions are expected to increase model diversity for aerosol absorption, the diversity of simulated radiative forcing (influenced by both aot and absorption) is a poor measure for the real diversity in modeling.

### OPTICAL DEPTH

### MASS

### Median Fields

### Diversity-Fields



## next

**AEROCOM project**  
 $\rightarrow$  detailed evaluations

- to understand reasons for differences in mass to optical depth conversions among models: **identical year, identical water uptake**
- to identify major causes for differences in mass distribution, including transport: **identical inventories (sources), identical meteorology**
- to understand observed seasonal and regional patterns of aerosol/chemistry: **satellite data, field studies, long-term monitoring**