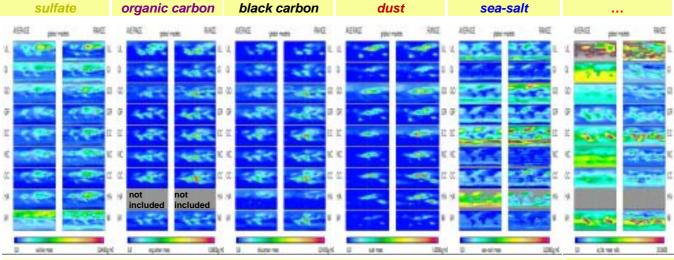
There are more differences among component models than a comparison of total aots would suggest WHY New aerosol components in global (climate) models distinguish between sulfate, organic carbon, black carbon, dust and sea-salt. The sum of all aerosol types provide the totals. Are good totals for optical depth skillful or just luck?

## Simulated aerosol components

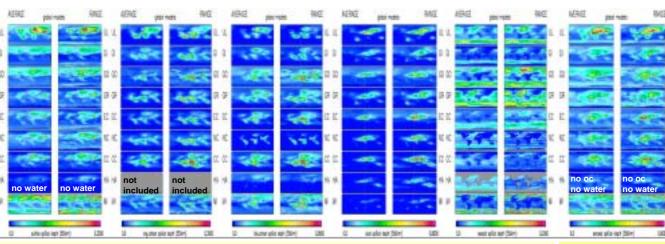
global fields of yearly averages and seasonality\*

seasonality = range of 3 month running means

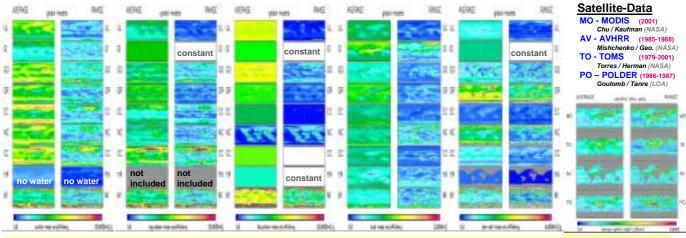
Models Resolution Authors Simulation Message + ULAQ (GCM) 10/22.5deg 3yr avg Pitari + GISS (GCM) 4.0/5.0deg Koch / Tegen 3yr avg + GOCART (CTM) 2.0/2.5deg (90, 96, 97) Chin / Ginoux • Grantour (GCM) 5.0/5.0deg Herzog / Penner 1yr avg + ECHAM4 (GCM) 3.8/3.8deg Feichter / Lohmann /Schulz 3yr avg 1.9/1.8deg size assumptions Collins / Rasch NCAR (GCM) (95-00) ⇔ humidification assumptions + CCSR (CTM) 2.8/2.8deg Takemura / Nakaiima (90)ambient relative humidity used + HadAM4 (GCM) 2.5/3.8deg Roberts / Jones 5vr avg MIRAGE (GC/TM) (6/94-5/95) 2.8/2.8deg Ghan / Easter







## AEROSOL mid-visible OPTICAL DEPTH



## AEROSOL mid-visible MASS EXTINCTION EFFICIENCY

next

Add. Comparisons: - 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* AEROCOM project - to understand observed seasonal and regional patterns of aerosol/chemistry: *satellite data, field studies, long-term monitoring* 

overall agreement for source location, but difference in strength large differences in simulated transport (and/or removal rates) Iarge differences in conversion (of mass into optical depth) due to

extra comparisons needed to identify/ remove poor assumptions

OC / bC ratio

total aot

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satellite aot