

Improving aerosol size representation in the UKMO Hadley Centre climate model

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Motivation

- Aerosol optical properties are very sensitive to their assumed size distribution.
- Accurate direct aerosol radiative forcing calculation requires adequate characterisation of size distribution.
- The number of aerosol particles that can be activated as CCN depends critically on their size distribution.
- Accurate indirect aerosol radiative forcing calculation also requires adequate characterisation of size distribution





Aerosol in existing UKMO Hadley Centre scheme

- Sulfate aerosol exists in Aitken, accumulation or dissolved modes.
- Only aerosol mass carried for each mode.
- Fixed geometric mean radius and standard deviation for each mode.

• Jones et al. (2001) estimated indirect effect using

$$N_{d} = \max \{ 3.75 \times 10^{8} [1 - \exp(-2.5 \times 10^{-9} N_{tot})], N_{min} \}$$

$$N_{tot} = N_{Ait} + N_{acc} + N_{dis} + N_{jet} + N_{film} [N_{jet} + N_{film} \text{ from O' Dowd et al (1999)}]$$

$$N_{min} = 5.0 \times 10^{6} \text{ over water and } 3.5 \times 10^{7} \text{ over land}$$

$$N_{Ait}, N_{acc}, N_{dis} \text{ calculated from mass assuming } r_{med}, \sigma_{g}$$



Processes in existing Hadley Centre sulfur cycle

- Gas phase oxidation of SO2 to Aitken & accum mode according to ratio of surface areas
- In-cloud aqueous phase oxidation of SO2 to dissolved mode
- Activation of accumulation mode to dissolved mode
- Evaporation of dissolved mode to accumulation mode
- Coagulation of Aitken mode with accum mode to more accum mode
- Diffusion of Aitken mode aerosol to dissolved mode
- Dry deposition of aerosol
- In cloud scavenging included and assumed to dominate over below cloud scavenging (BCS neglected).

For details see Jones et al., (2001)





A new UK Chemistry and Aerosol model (UKCA)

- Collaboration between University of Leeds, Cambridge University and UKMO Hadley Centre to develop next generation aerosol-chemistry module for use in century-scale climate GCM simulations.
- Implemented existing Hadley Centre scheme within an aerosol-chemistry transport model framework.
- Comparing size distributions using existing scheme with those from microphysical sectional GLOMAP scheme (in same framework) and measured size distributions – which processes/aspects need particular improvement?
- GLOMAP is described in Spracklen et al. (2004, ACPD) [single-component sulfate-seasalt version]





TOMCAT

•3D Offline CTM •Forced by ECMWF Winds

- •Convective transport
- Convective and resolved rain

GLOMAP

Aerosol size spectrum (~ 1 nm – 24 μm) Two-moment sectional scheme 20 bins in ptcl number & average mass





UGAMP 2004





GLOMAP vs Hadley Comparison Experiment

- Spin-up from an aerosol-free atmosphere, October December 1995 at T42 (2.8° x 2.8°) horizontal resolution, 31 vertical levels
- GLOMAP --- Sulfate and sea salt aerosol only, treated in the same distribution. 20 bins of numbers and mass.
- Hadley --- Aitken, accum, dissolved mode sulfate aerosol scheme used. Only mass carried, number deduced from r_{med} , σ_q
- Equivalent emissions of SO2, DMS & sea-salt, oxidant (OH, H2O2, NO3)
 & cloud fields, precipitation rates and gas phase chemistry schemes
- More sophisticated aqueous phase oxidation scheme in GLOMAP (cloud droplet spectrum produced for assumed cloud water content)
- No nucleation in Hadley centre scheme, condensation parameterized by Aitken/accum surface area ratio and coagulation treatment very basic.
- •Both runs have aerosol dry deposition although aerosol wet removal only by in-cloud scavenging (no BCS) in Hadley Centre run.



Hadley Centre Aerosol Modes in FT (at ~2.5km)



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Hadley vs GLOMAP sectional at surface (Dp < 10 nm)



Hadley centre







Hadley vs GLOMAP sectional at surface (Dp 10-100 nm)



Hadley centre







Hadley vs GLOMAP sectional [surface] (Dp 100-1000 nm)



Hadley centre







Hadley vs GLOMAP sectional [surface] (Dp > 1000 nm)



Hadley centre







Number conc. size distributions (surface)

MBL off E coast of U.S.A.

Continental U.S.A.



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Size distributions in volume concentration



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<u>GLOMAP CCN no. conc. (surface) using Kohler theory vs</u> <u>Hadley Centre CCN no. conc. (surface) - Jones et al (2001)</u>





<u>Summary</u>

- GLOMAP model framework will be useful for identifying areas for improvement in Hadley Centre aerosol scheme
- By comparison of Hadley scheme with observations and GLOMAP sectional scheme (and using method of "process reduction") we will identify which processes are the most important to include in a new computationally affordable scheme for runs on centennial timescales
- For example does nucleation in FT and subsequent coagulation and condensation growth need to be included for accurate forcing calculation?
- Does sea salt need to be included as an interactive component?





Plan for next 6 months

- Thorough evaluation of existing Hadley scheme against GLOMAP and observations
- GLOMAP already evaluated against observations
- Investigate:
 - Response to changing emissions
 - Relative change in Aitken/accumulation modes (critical for direct and indirect forcing)
- Identify key model updates/refinements to sulphate/sea salt scheme
- Initial development of new scheme





Plan for longer term

- GLOMAP multi-component version is currently being developed at University of Leeds
- Update dust, black carbon, organic carbon scheme in Hadley centre testing versus multi-component GLOMAP and observations
- Include secondary organic aerosol formation.
- Include improved chemistry scheme (nitrate, ammonium).
- Investigate bio-geochemical feedbacks ---- iron fertilization of oceans from dust deposition ---- increased DMS production ---- increased sulfate aerosol ---- increased cloud albedo ---- investigate feedback on DMS production





Hadley Centre aerosol no. conc. by size (surface level)





GLOMAP aerosol no. conc. by size (surface level)

