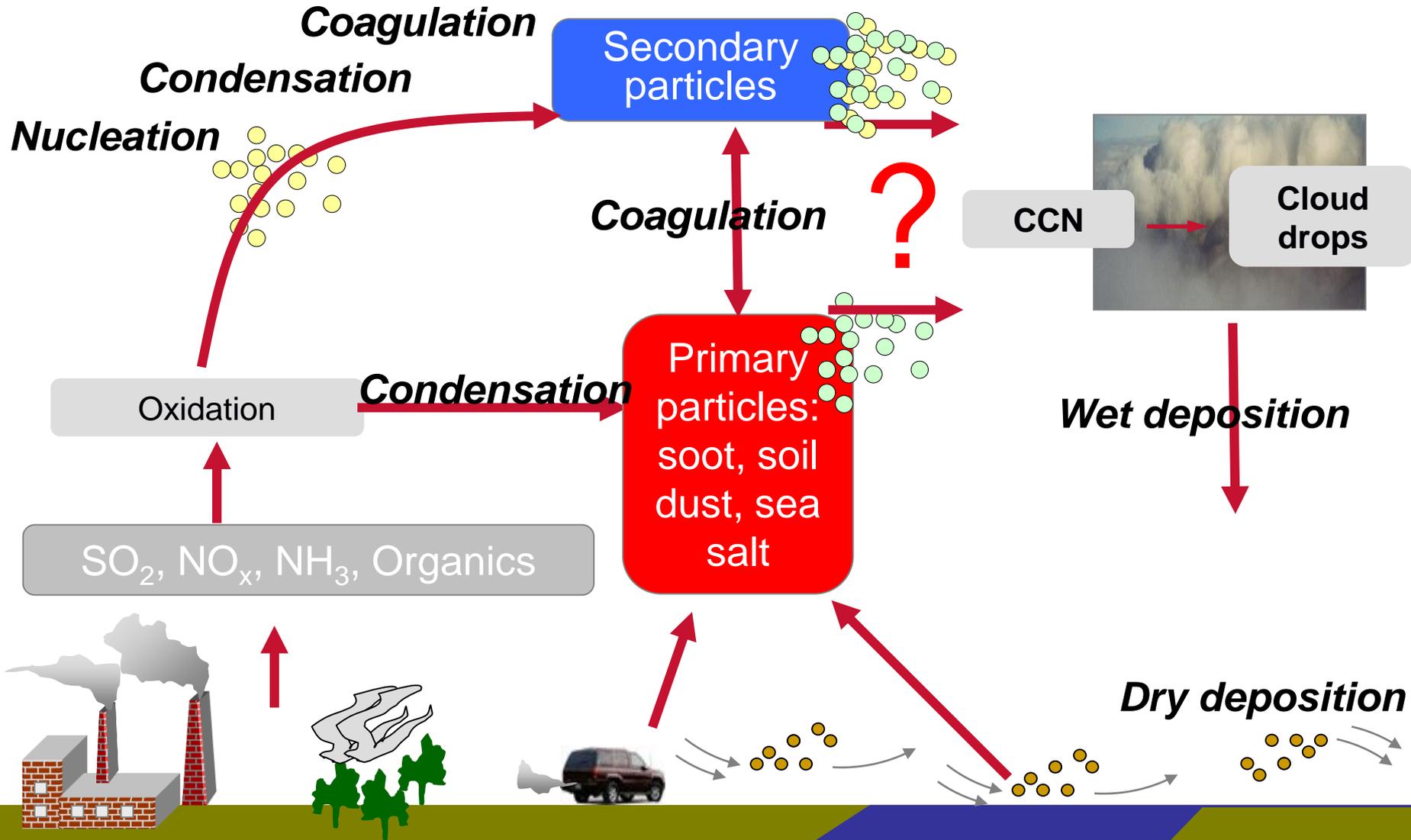


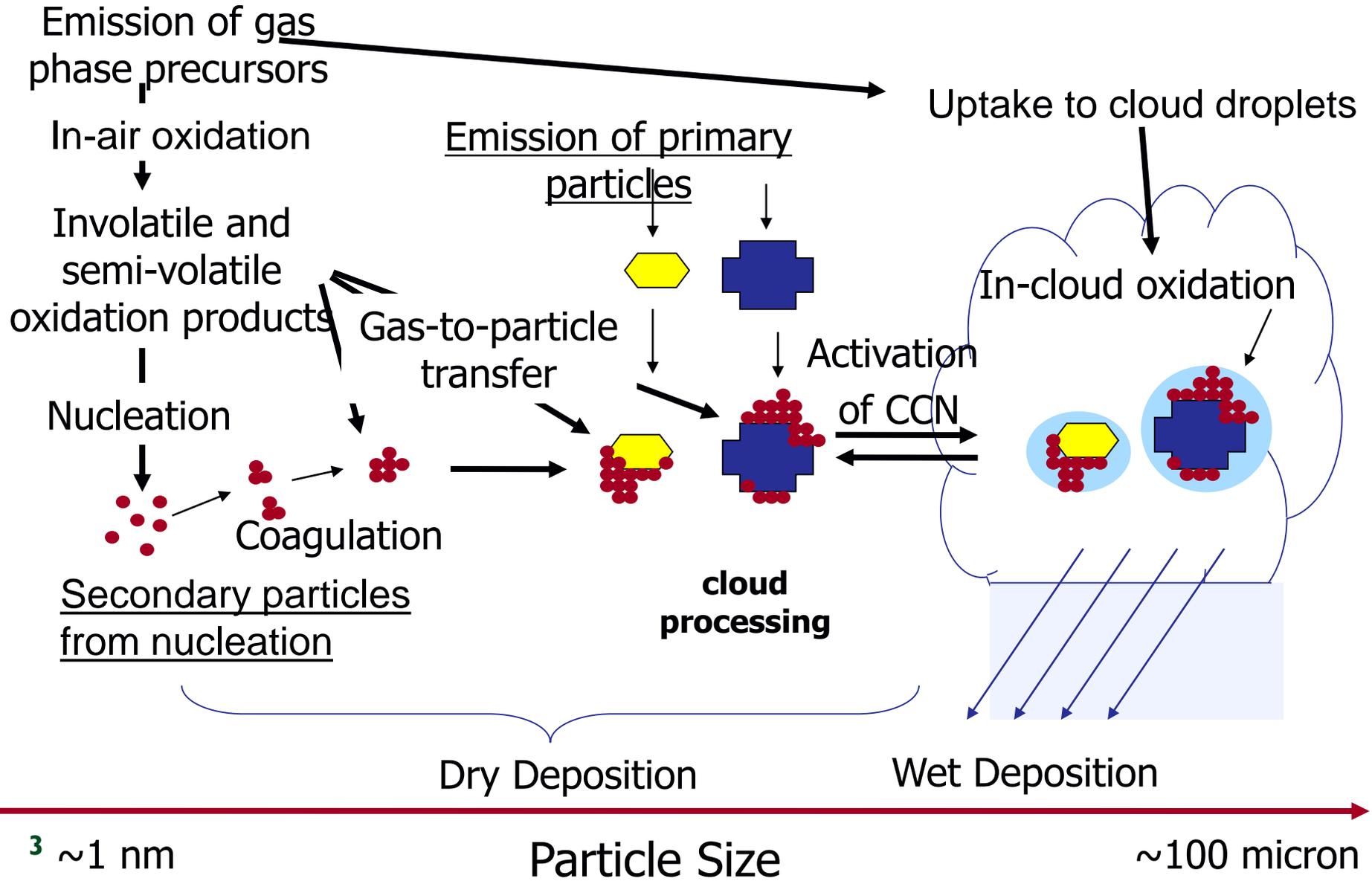


Sources of aerosol particles – a complex mix





Processes control particle size & composition



³ $\sim 1 \text{ nm}$

Particle Size

$\sim 100 \text{ micron}$

- Collaboration between NCAS & UK Met Office Hadley Centre since 2005. Universities of Leeds, Cambridge and Oxford are main NCAS partners
- Aerosol-chemistry sub-model in Met Office Unified Model environment for a range of applications (climate, Air Quality, Earth System science etc.)
- Tropospheric and stratospheric chemistry schemes. Aerosol precursor extension to UKCA chemistry schemes so that climate model simulated aerosol is coupled to atmospheric chemistry.
- Improved representation of aerosol in UK climate model simulations
 - new particle formation & growth using GLOMAP aerosol microphysics
 - internally mixed aerosol (e.g. BC & sulphate) affect optical properties
 - biogenic secondary organic aerosol from monoterpene oxidation
- UKCA interactive ozone, methane and aerosol (direct/indirect) radiative effects for fully coupled composition-climate simulations.
- Enhances UK capability in aerosol-climate-earth system modeling and provides integration for NCAS and Met Office initiatives.



UKCA aerosol scheme is GLOMAP-mode

UK Chemistry and Aerosol project has developed aerosol-chemistry sub-model of the Met Office Unified Model suitable for climate, air-quality & aerosol effects on NWP

Aerosol scheme is called GLOMAP-mode (sometimes MODE or UKCA-MODE in UM)

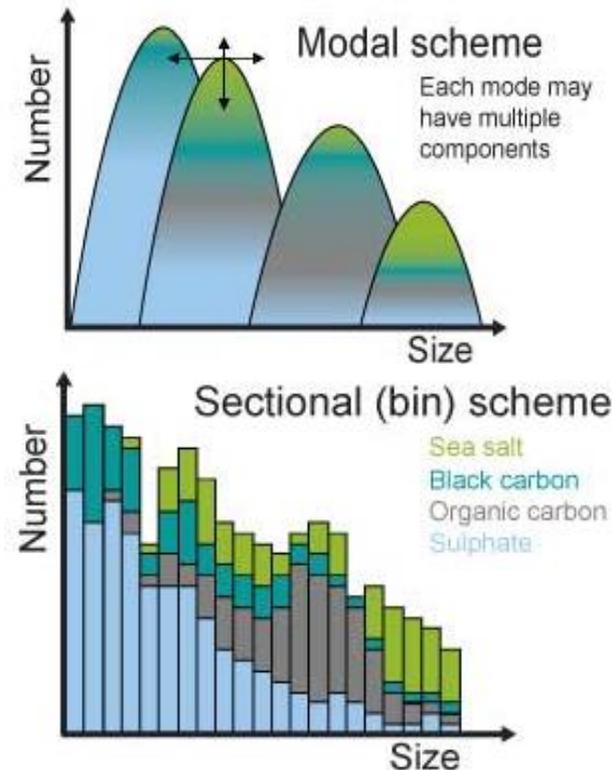
Transported tracers are size-resolved component mass & particle number mixing ratios

Uses size-resolved aerosol microphysics from GLOMAP-bin scheme but using log-normal size modes rather than bins:

- Primary emissions (sulphate, BC/OC, sea-salt, dust)
- Secondary sulphate particle formation (nucleation)
- Coagulation (within modes, between modes).
- Condensation (of sulphuric acid, bulk condensible organic)
- Cloud processing
- Dry deposition and sedimentation
- In-cloud and below-cloud scavenging

Benchmark MODE scheme vs detailed BIN scheme in CTM.

Simulates size-resolved multi-component aerosol lifecycle including new particle formation & growth.



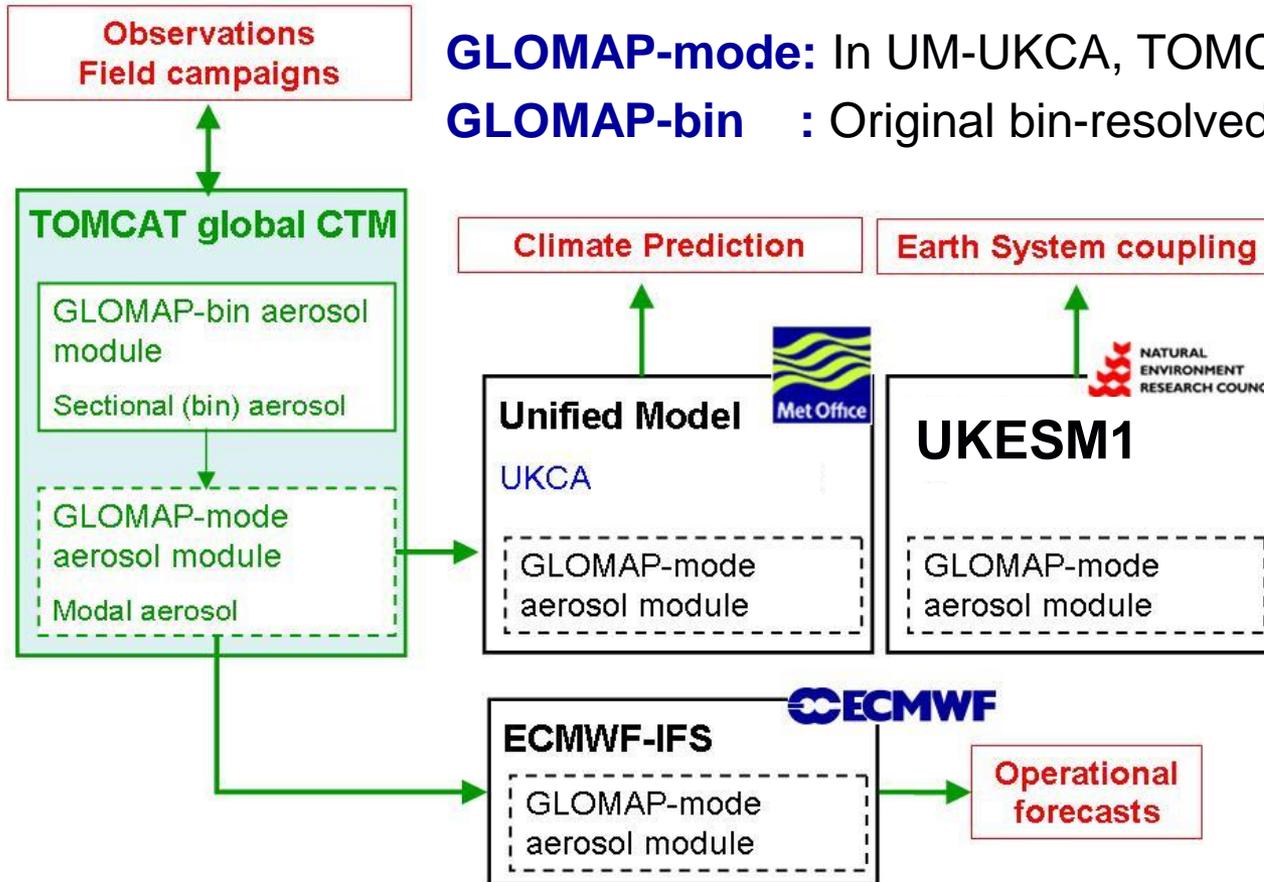
GLOMAP-bin : Spracklen et al. (ACP 2005a,b, 2006, 2007), Korhonen et al (JGR, 2008),
 GLOMAP-mode: Manktelow et al (GRL, 2007), Woodhouse et al (Atmos Env, 2008; ACP, 2010),
 Schmidt et al (ACP,2010), Mann et al, (GMD,2010), Mann et al (ACP, 2012)



UKCA Aerosol Strategy

UM-UKCA is one of several model frameworks which include GLOMAP.

GLOMAP-mode: In UM-UKCA, TOMCAT CTM & ECMWF-IFS
GLOMAP-bin : Original bin-resolved version in TOMCAT CTM



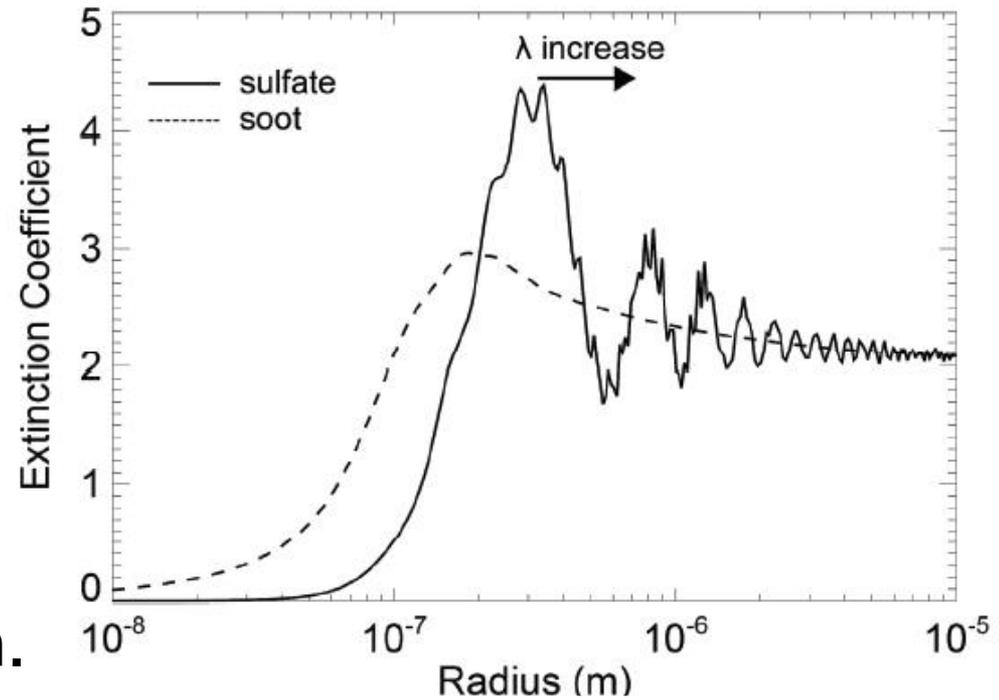
Strong connection between the future development of UM-UKCA and improvements and developments to GLOMAP in the TOMCAT CTM and ECMWF-IFS

UM-UKCA release job at v8.4 with GLOMAP & CheST stratosphere-troposphere chemistry with extension for aerosol precursor chemistry

Importance of size distribution: AOD

Differences in particle size distribution strongly affect the extinction.

Constant mass extinction efficiency will not capture variability from changes to the particle size distribution.

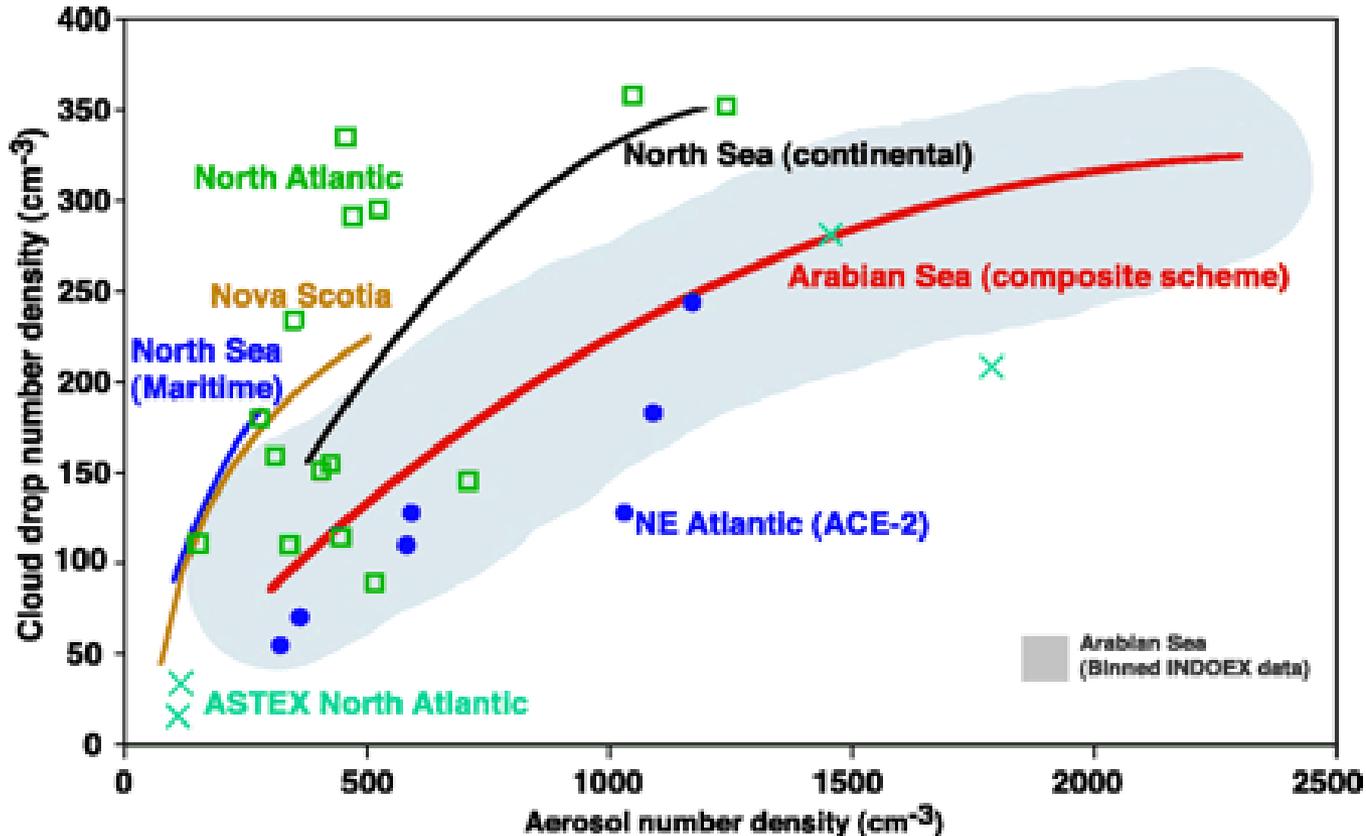


$$b_{ext} = \pi \sum_{i=0}^n Q_{ext} R_i^2 \frac{dN_i}{d \ln R_i} \Delta \ln R_i \quad \tau_{\lambda} = \int_0^{z_{top}} b_{ext}(\lambda) dz ,$$

Same mass distributed in different sizes gives different AOD



Cloud droplet vs aerosol from observations

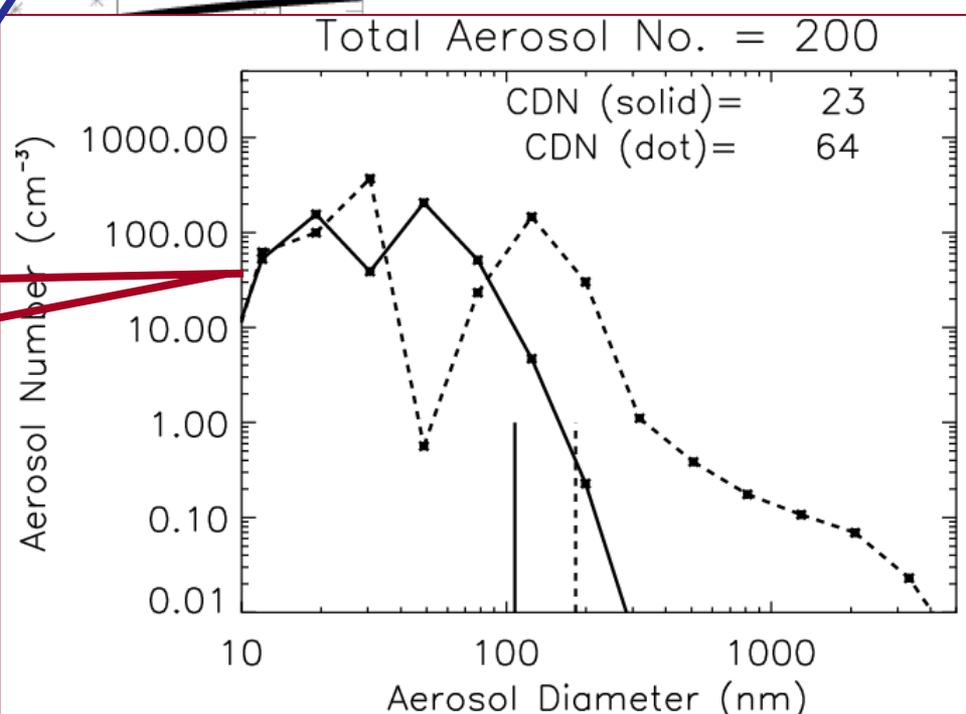
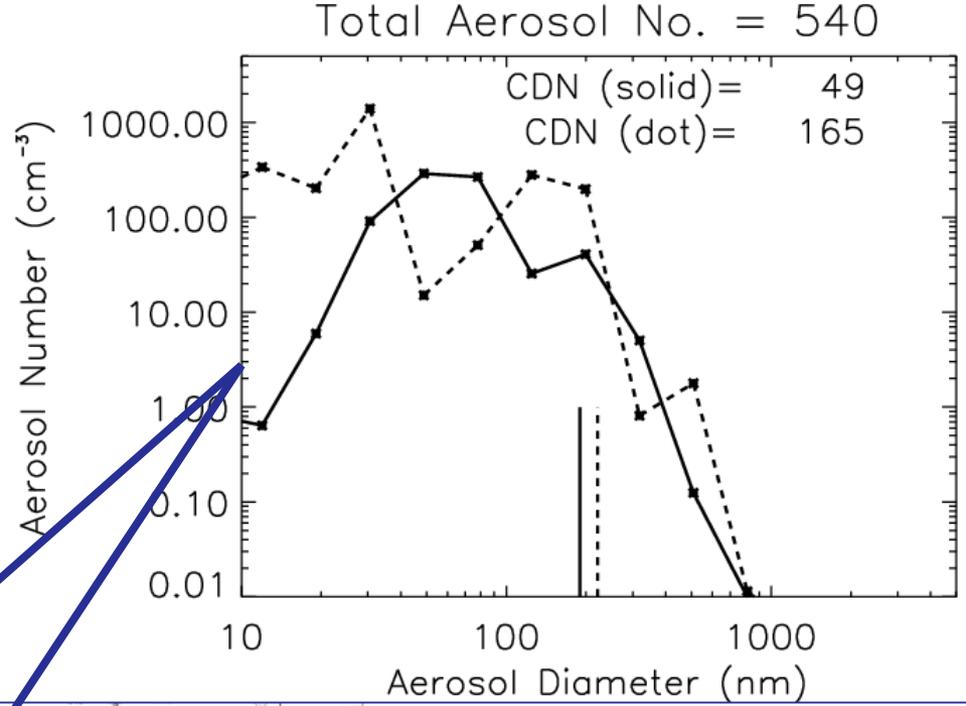
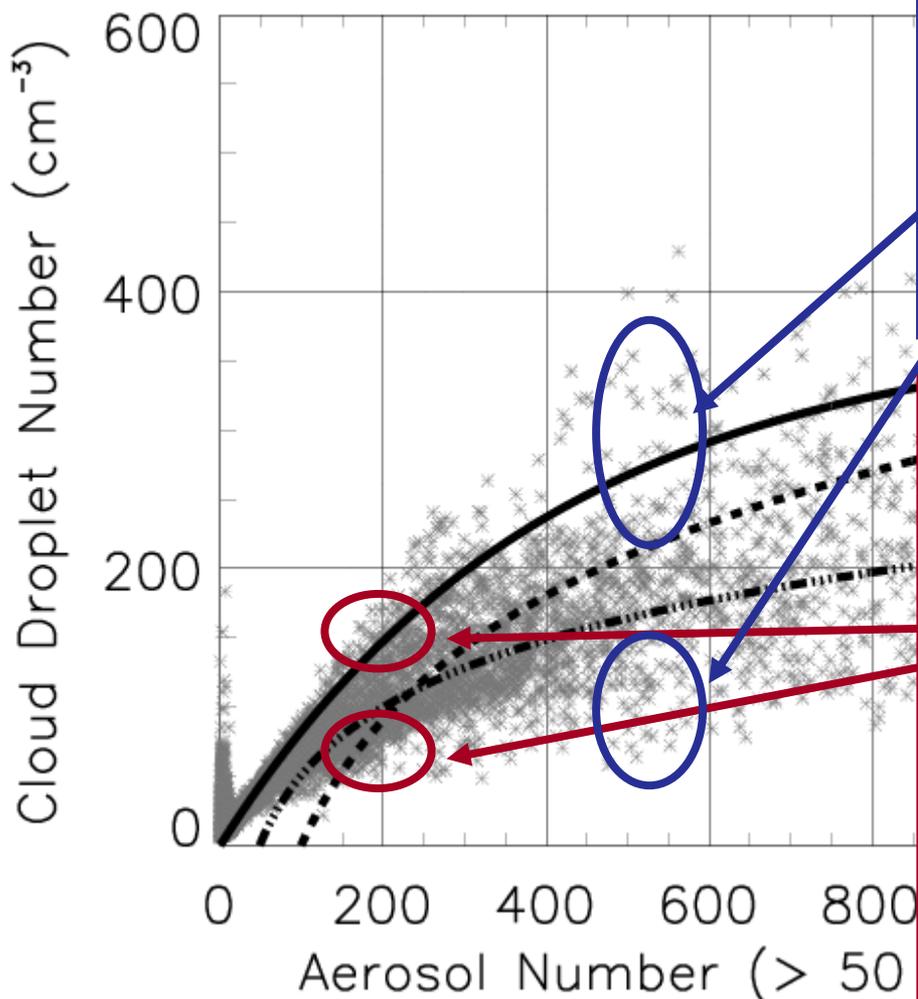


Ramanathan et al.
(2001, Science)

No single relationship

Different particle types, compositions, size distributions, etc

Exploring the scatter in CD

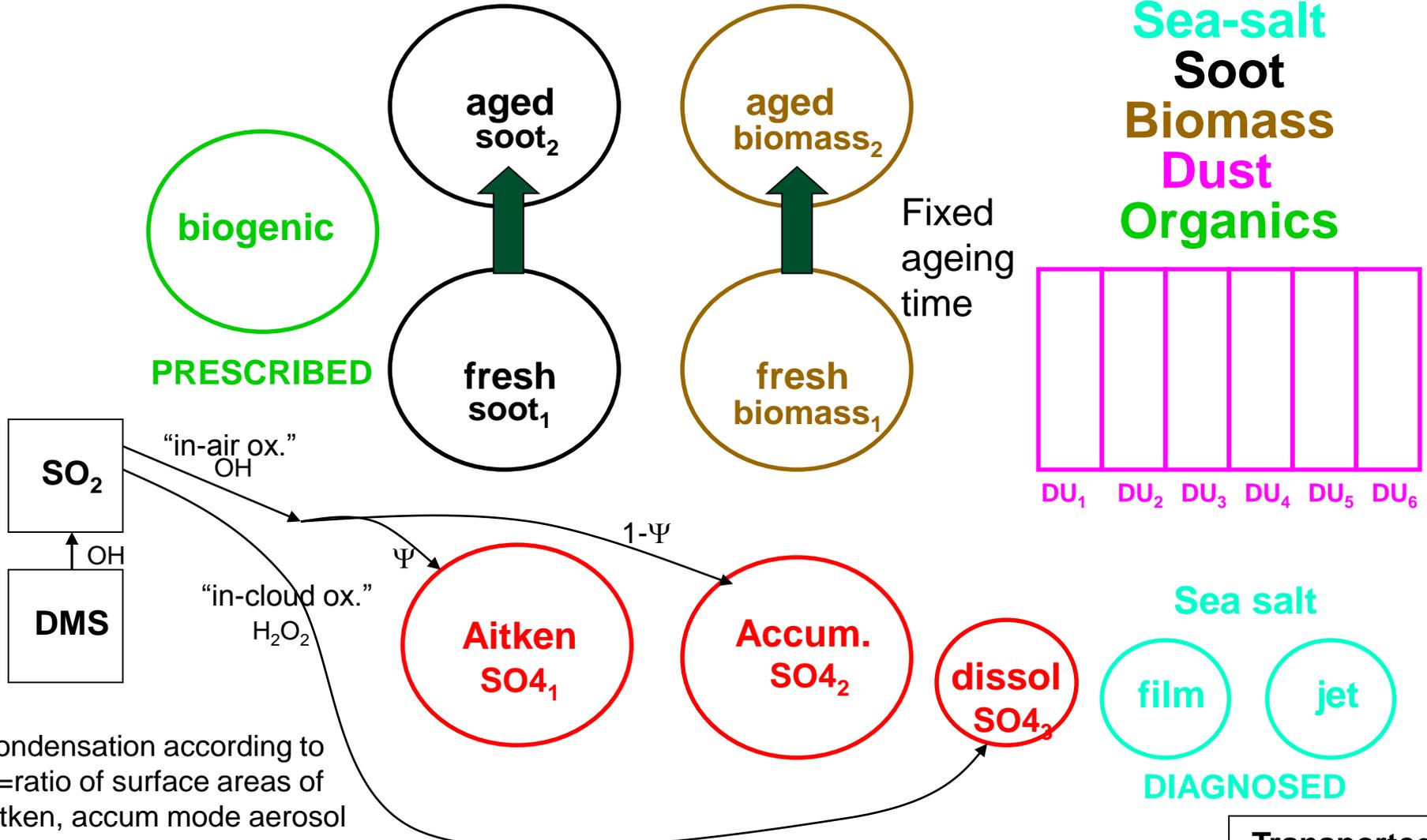




Mass-based UM aerosol scheme (CLASSIC)

Has aerosol mass as "types" of fixed size (externally mixed)

Sulphate
Sea-salt
Soot
Biomass
Dust
Organics



Condensation according to Ψ =ratio of surface areas of Aitken, accum mode aerosol

Aerosol optical properties & CCN derived based on fixed size distribution

Transported tracers=13



GLOMAP 7-mode aerosol configuration

Has aerosol mass as “components” in internally mixed modes

Sulphate, **sea salt**, **black carbon**, **organic carbon**, **dust**

MONOTER

SEC_ORG

PRIMARY EMISSIONS

Wildfire & industrial BC/OC, mineral dust

In-cloud produced S(VI) to mode according to ratio of accum, coarse mode N.

H₂SO₄

Condensation

SO₂

DMS

Insoluble Aitken N₅

insoluble accum N₆

insoluble coarse N₇

Soluble nucln N₁

soluble Aitken N₂

soluble accum N₃

soluble coarse N₄

component mass conc.

SO₄ OC

SO₄ OC
BC Cl

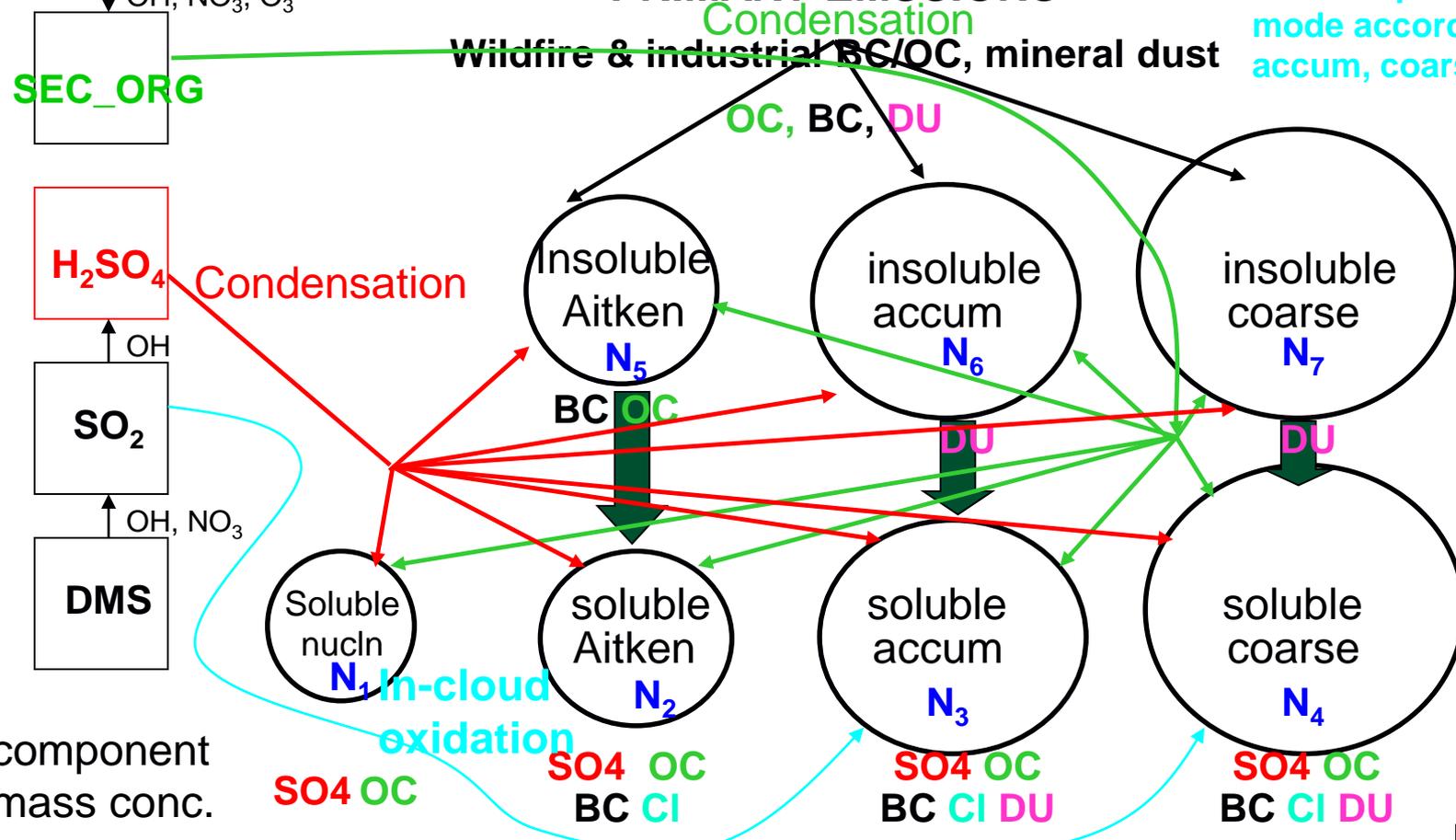
SO₄ OC
BC Cl DU

SO₄ OC
BC Cl DU

20 mass
7 number

AOD, CCN determined by size-resolved emission, processes & removal.

Transported tracers=27

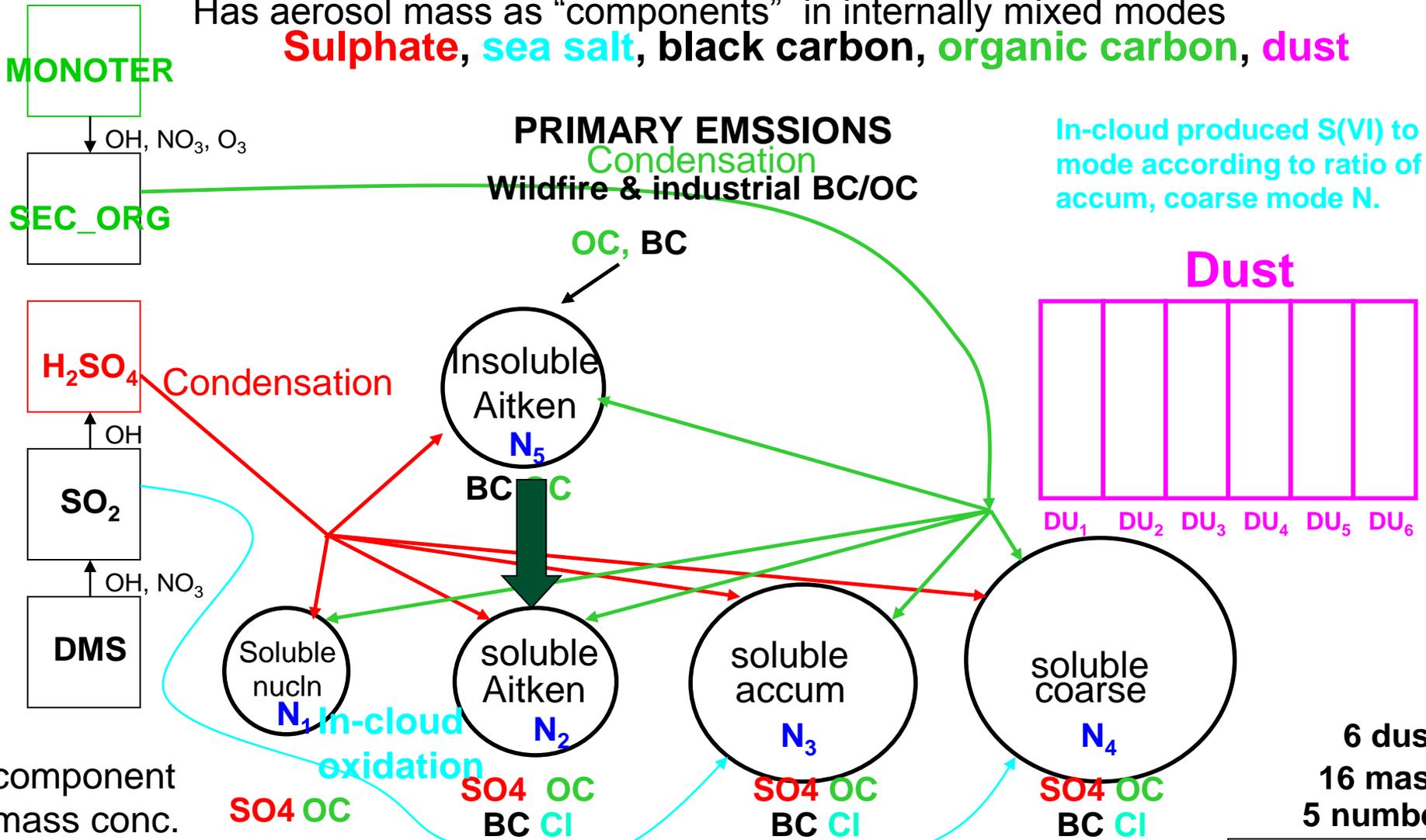




UM-UKCA uses 5-mode GLOMAP plus 6-bin dust

Has aerosol mass as “components” in internally mixed modes

Sulphate, **sea salt**, **black carbon**, **organic carbon**, **dust**



AOD, CCN determined by size-resolved emission, processes & removal.

CLASSIC and UKCA compared



	CLASSIC	UKCA
Transported particle types	Associated with emissions (sulphate, biomass, etc)	Defined by microphysics (Aitken, accumulation, etc)
Size distribution	Prognostic m Fixed size N derived from m and size	Prognostic N, m Variable size Log-normal modes
Mixed composition	No	Yes
Chemistry	Simple S-chem driven by offline oxidants or UKCA	Coupled chemistry
Cloud drop number	From mass	From size, N, mixed composition
Particulate tracers	13	27 for full 7 modes (21 for 5-mode setup)

“Standard” UKCA Aerosol Scheme



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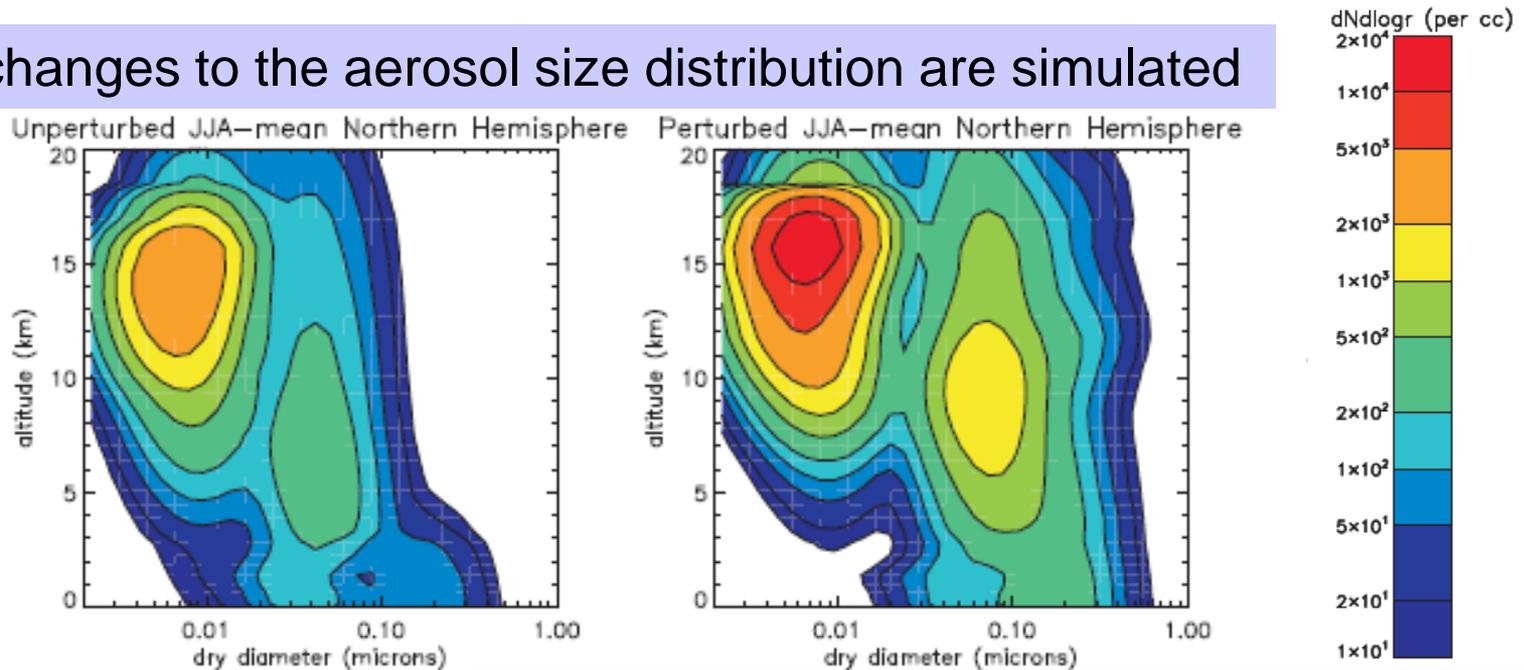
Mode name	Mean rad range nm	Composition	Production	Comments
nucleation soluble	< 5	SO ₄	nucleation	Currently only binary H ₂ SO ₄ -H ₂ O, later BLN
Aitken insoluble	5 – 50	BC, OC	primary BC/OC emissions	Separation to handle BC/OC ageing (necessary if timescale for ageing > lifetime in grid box)
Aitken soluble	5 – 50	SO ₄ , BC, OC	growth of nucl. soluble, condensation (sol,ins Ait), primary SO ₄ ems, coag.	
Accum. insoluble	50 – 500	DU	primary DU emissions	Separation to handle dust ageing (as above) Primary accum. mode BC/OC ageing?
Accum. soluble	50 – 500	SO ₄ , SS, DU, BC, OC	growth of Aitken soluble, condensation (sol,ins acc) primary SS ems, coag.	
Coarse insoluble	>500	DU	primary DU emissions	Separation to handle dust ageing
Coarse soluble	>500	SO ₄ , SS, DU, BC, OC	growth of accum soluble, primary SS emission	

UKCA-aerosol applications : Volcanic aerosol: impacts on climate



- Simulate aerosol from 1783 Laki eruption of 120 TgS over 8 months.
- Revisits Stevenson et al (2003) and Highwood & Stevenson (2003) studies with GLOMAP-mode aerosol microphysics (in CTM with coupled-chemistry)

Enormous changes to the aerosol size distribution are simulated

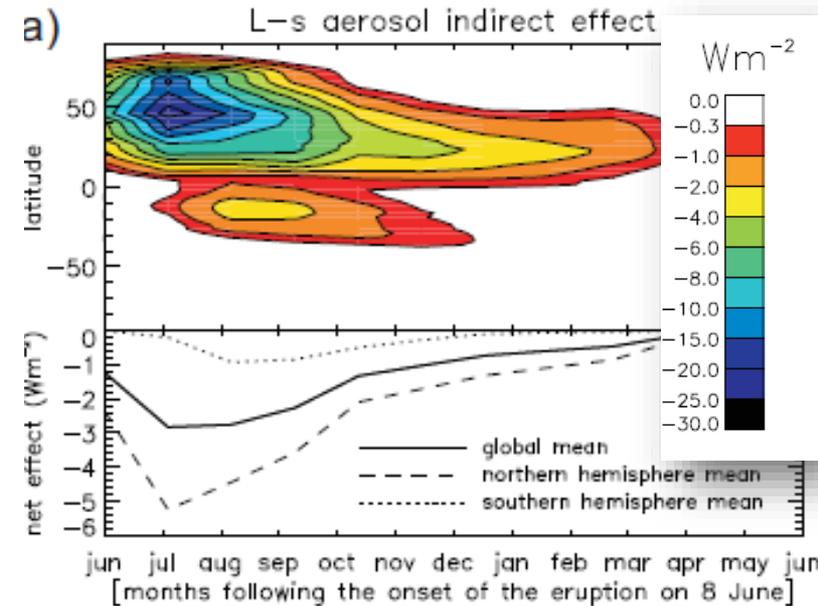
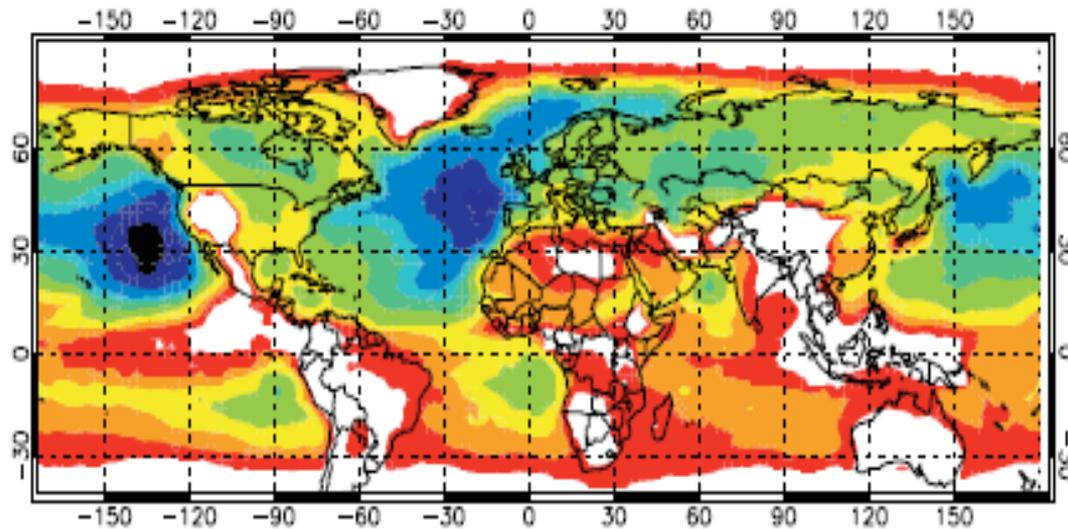


Schmidt et al (2010, ACP)

Eruption leads to stronger UT nucleation & growth → strong accum mode ~100nm

1783 Laki eruption: aerosol indirect climate effects

L-s JAS-mean aerosol indirect effect

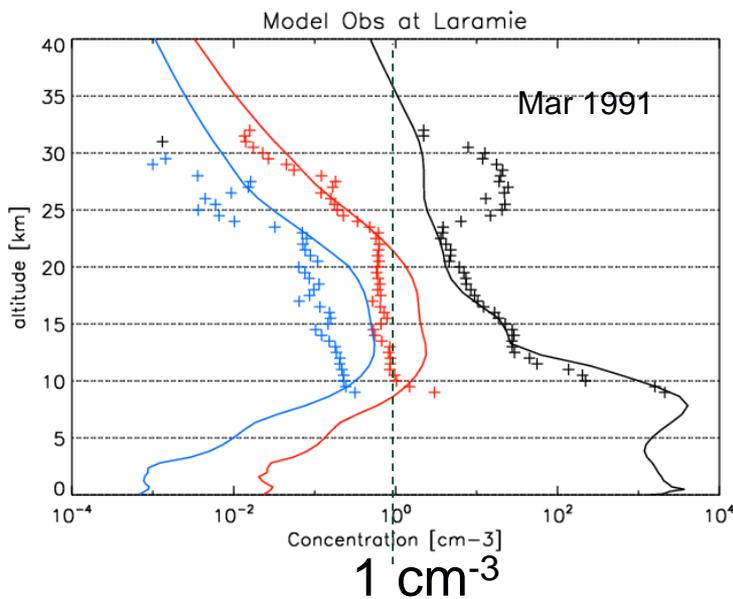


1st indirect forcing as large as the direct forcing,
but spatially much more widespread (due to
nucleation and growth)

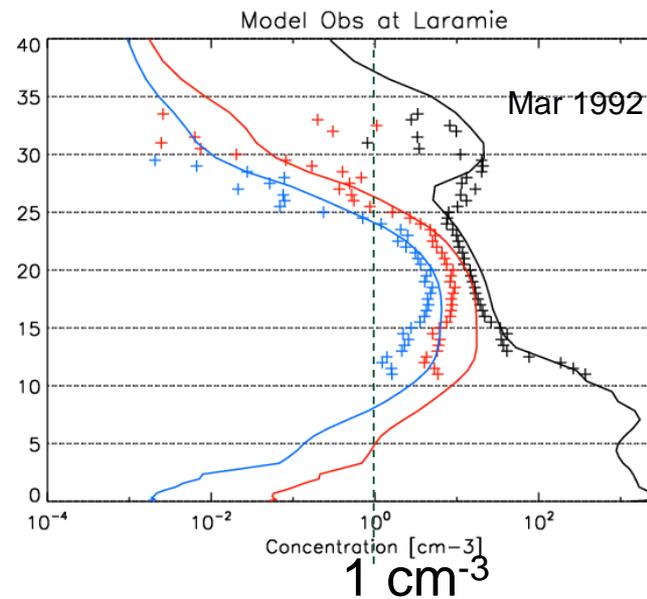
Schmidt (PhD Thesis, 2011)

Why UKCA?

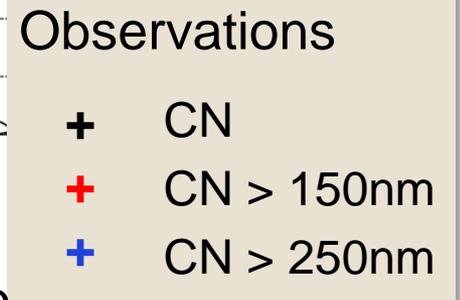
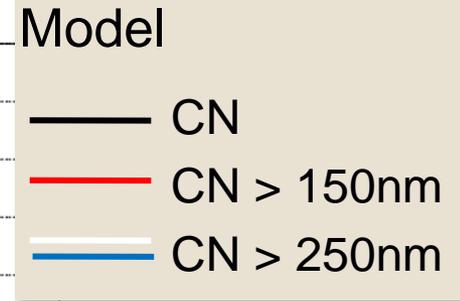
- Particle size distribution of the perturbed stratospheric sulphate aerosol is a major factor in predicting direct radiative forcing from stratospheric injection of SO₂ (affects growth, sedimentation, particle lifetime)
- In volcanically quiescent conditions, stratospheric aerosol particles are much smaller than following volcanic eruptions
- Climate forcing of very large volcanic eruptions is greatly reduced when the growth of sulphate droplets is accounted for (Timmreck et al., 2009, 2010)
- Applying UM-UKCA in coupled-AO to simulate stratospheric injection of SO₂ will maximise realism of predicted radiative forcings and climate responses



3 months before Pinatubo



9 months after Pinatubo

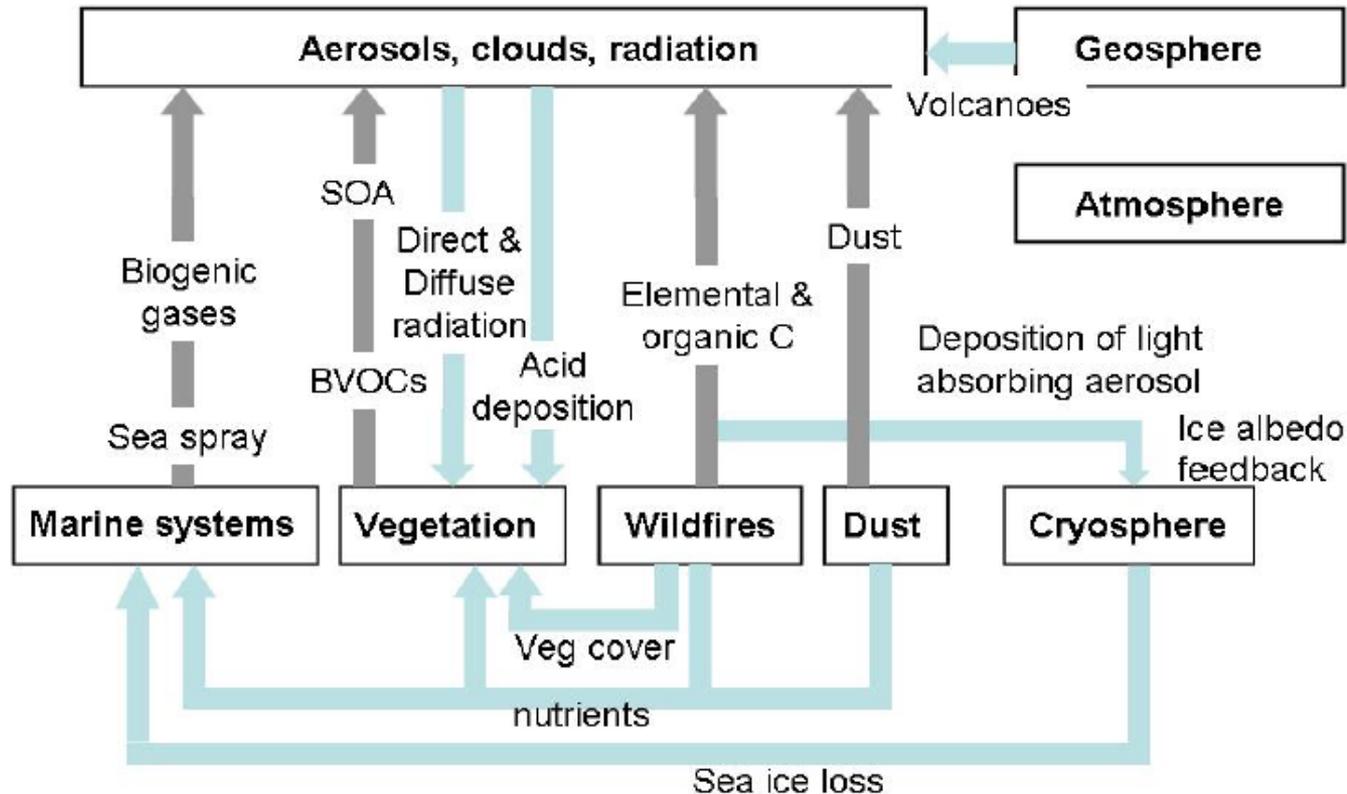


Balloon measurements of number concentration at Laramie, Wyoming (Deshler et al, 2003)

See Dhomse et al. (2014, ACP)

UKCA-aerosol applications: Role of natural aerosol in the Earth System

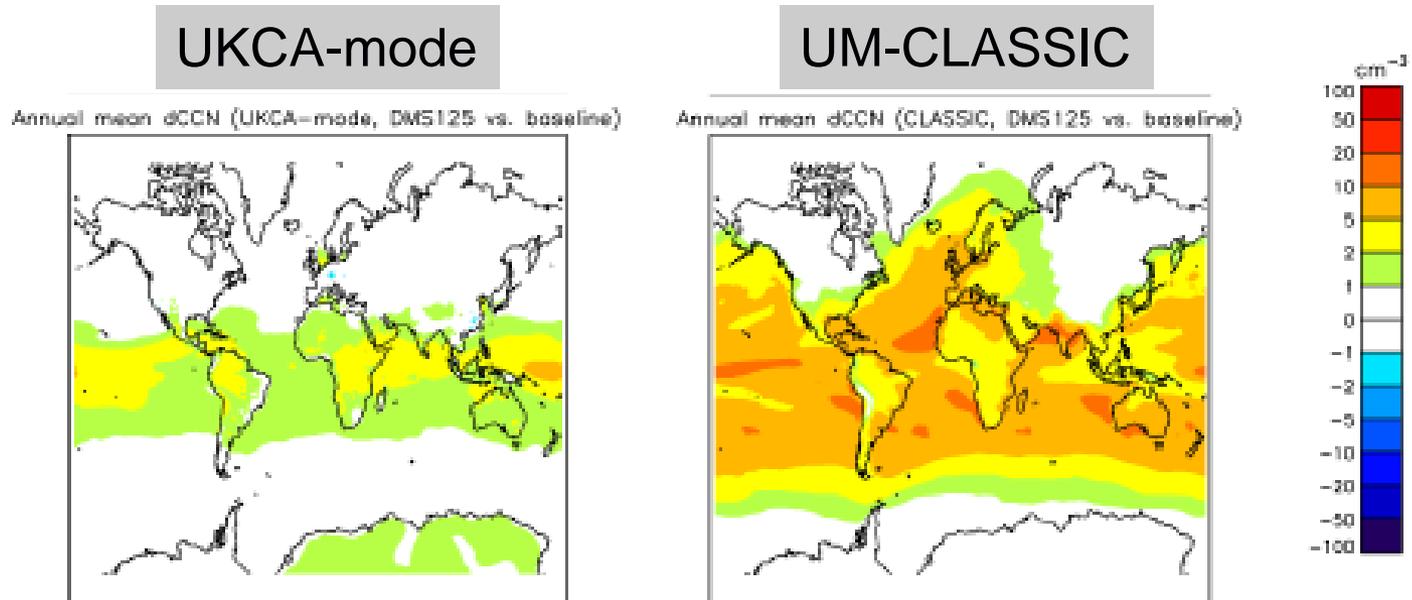
Carslaw et al (2010) review of *Natural aerosol interactions in the Earth System*



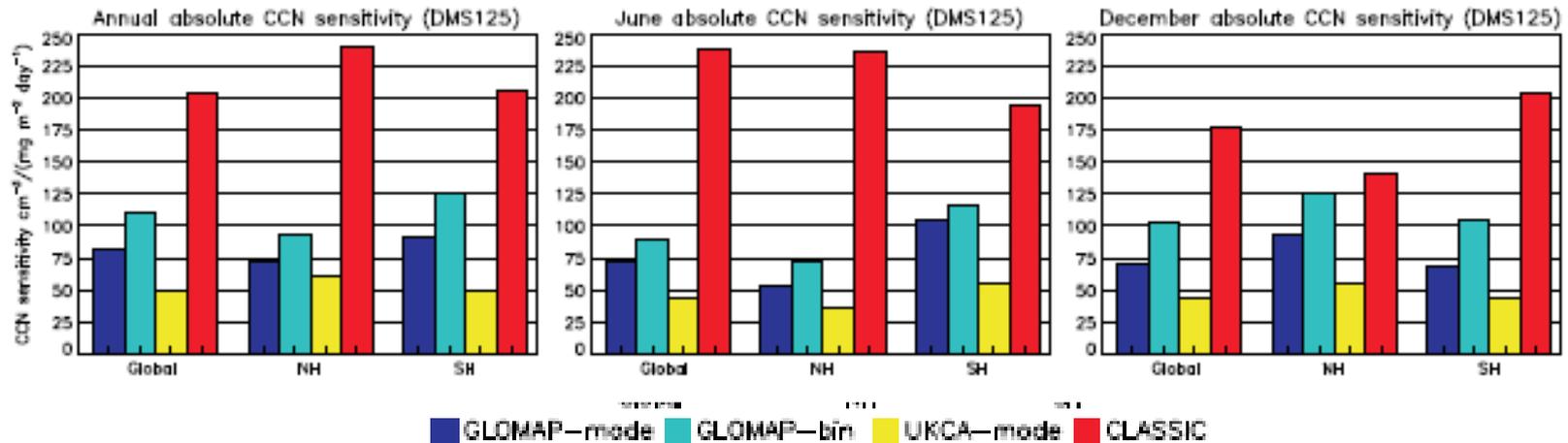
Offline CTM simulations have examined the role of these sources. UKCA chemistry & size-resolved aerosol microphysics will allow state-of-the-science studies of these within the new UKESM1 AO-GCM.

Change in CCN due to 25% increase in DMS emissions

- UKCA-mode predicts a much smaller response of CCN



Matt Woodhouse (Leeds) : from PhD thesis 2010.



CLASSIC mass-based aerosol scheme (in HadGEM2-ES for IPCC AR5) overestimates CLAW feedback because increase in aerosol mass (e.g. from in-cloud oxidation of SO_2) increases cloud drops

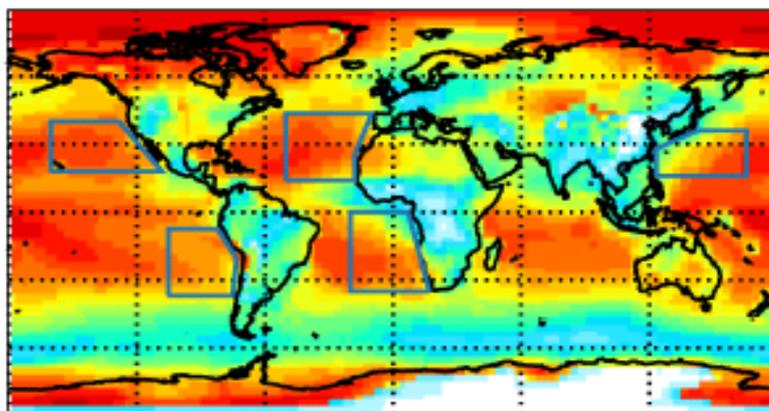
GLOMAP simulates particle growth (conserves number), hence changes to CCN and indirect climate effects more realistic.

Matt Woodhouse (Leeds) : from PhD thesis 2010.

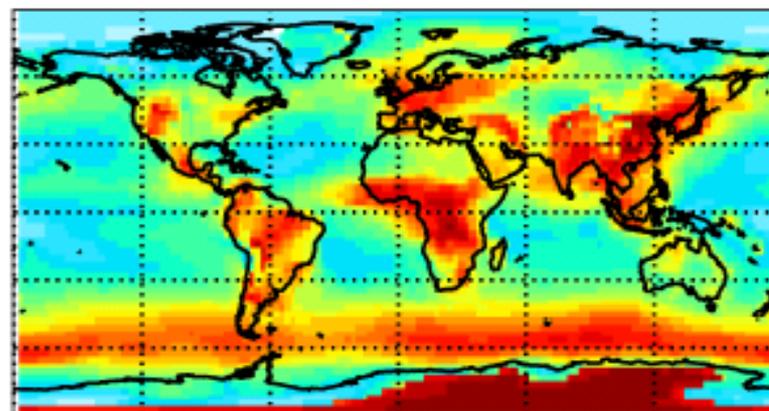
Percentage of primary and nucleated CCN

Nucleated particles important for aerosol indirect effects.
Growth to CCN sizes via coagulation and gas to particle transfer.

CCN from nucleation



CCN from primary emissions



Globally:

39% of low cloud-level CCN are from nucleation, 61% from primary particles

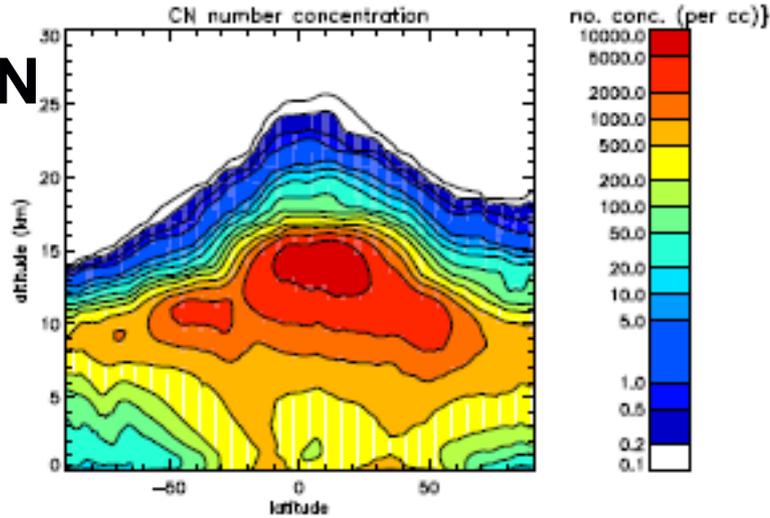
Nucleated CCN dominate in cleaner regions.

Merikanto et al (2009, ACP)

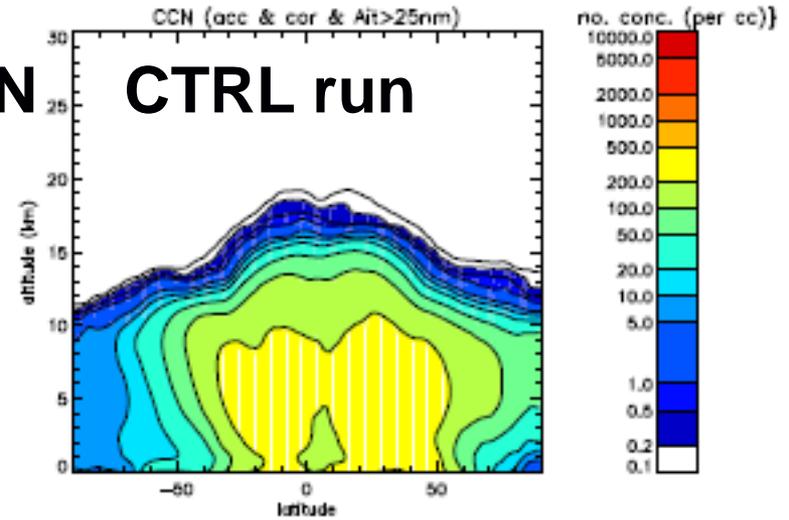


UT nucleation contribution to CN, CCN in UKCA

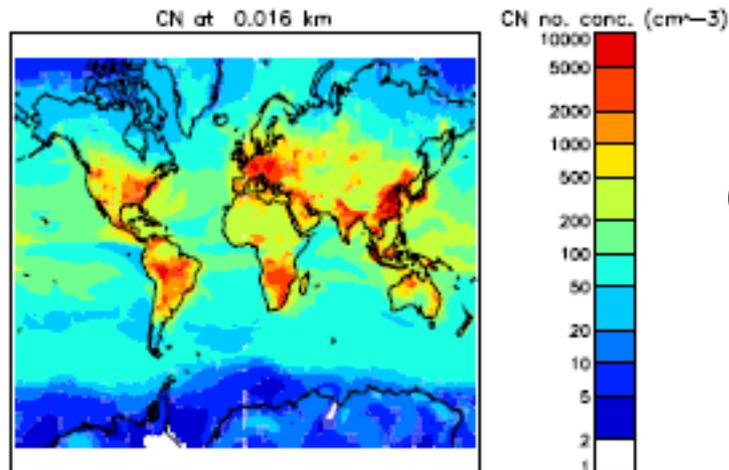
CN



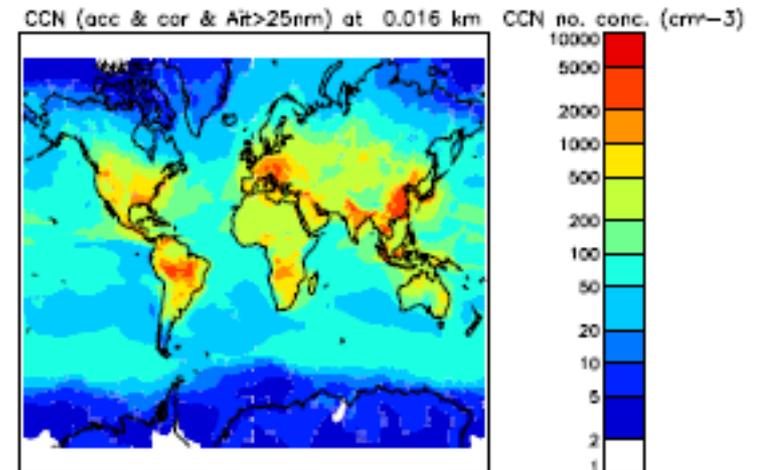
CCN CTRL run



CN

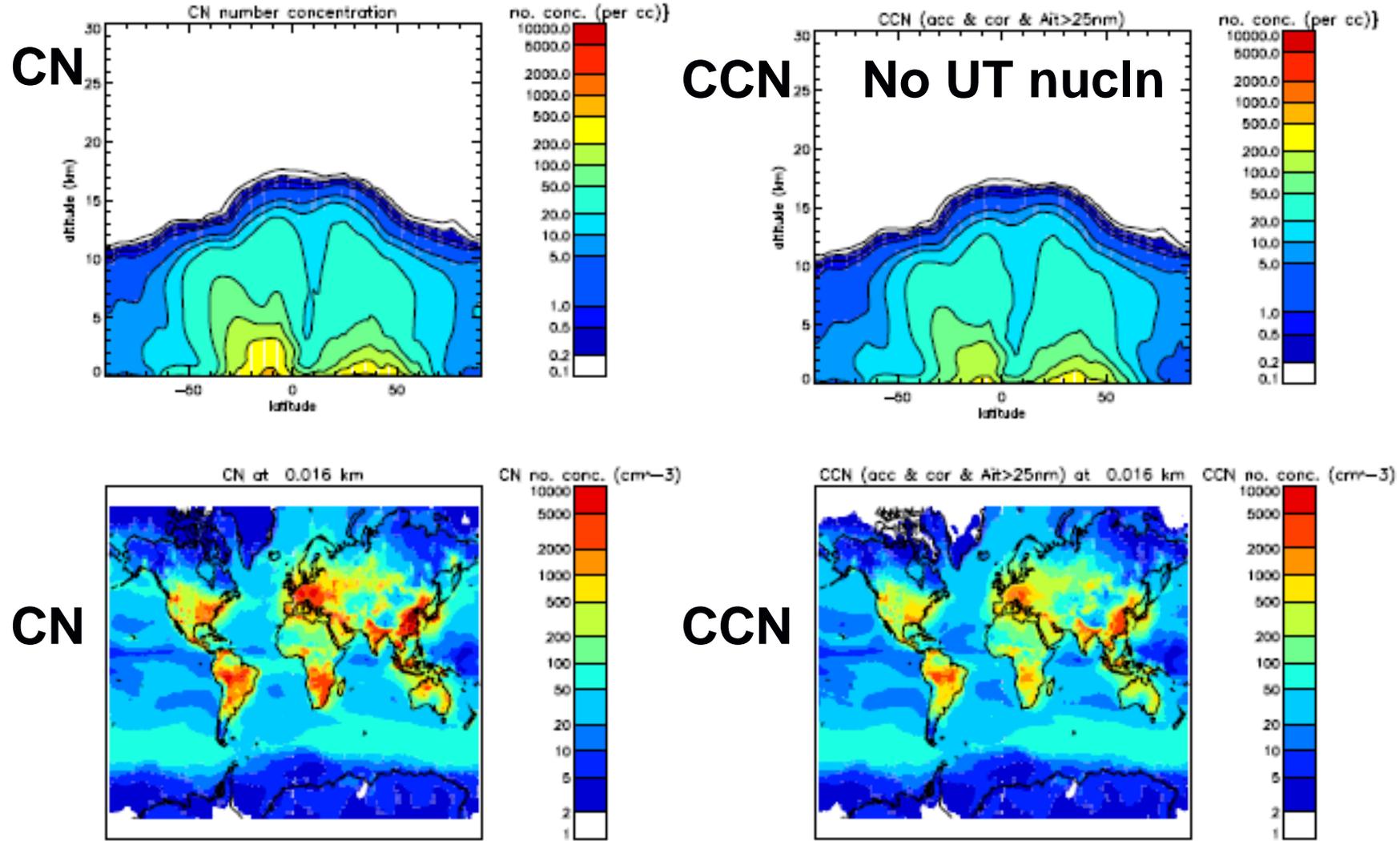


CCN





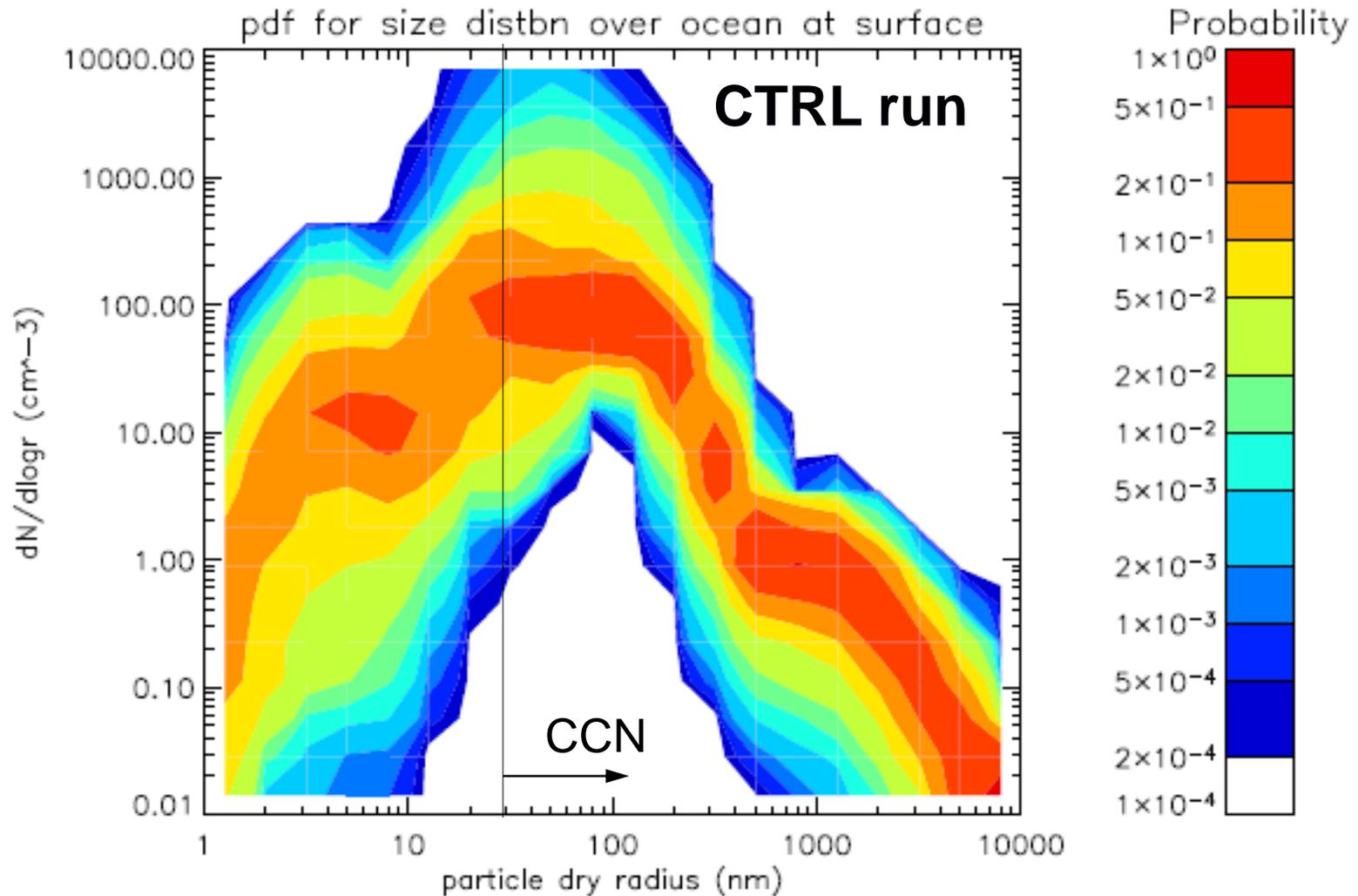
UT nucleation contribution to CN, CCN in UKCA



Large proportion of marine CN & CCN from secondary (FT nucleated) particles.
Need to consider nucleation for realistic aerosol indirect effects on climate

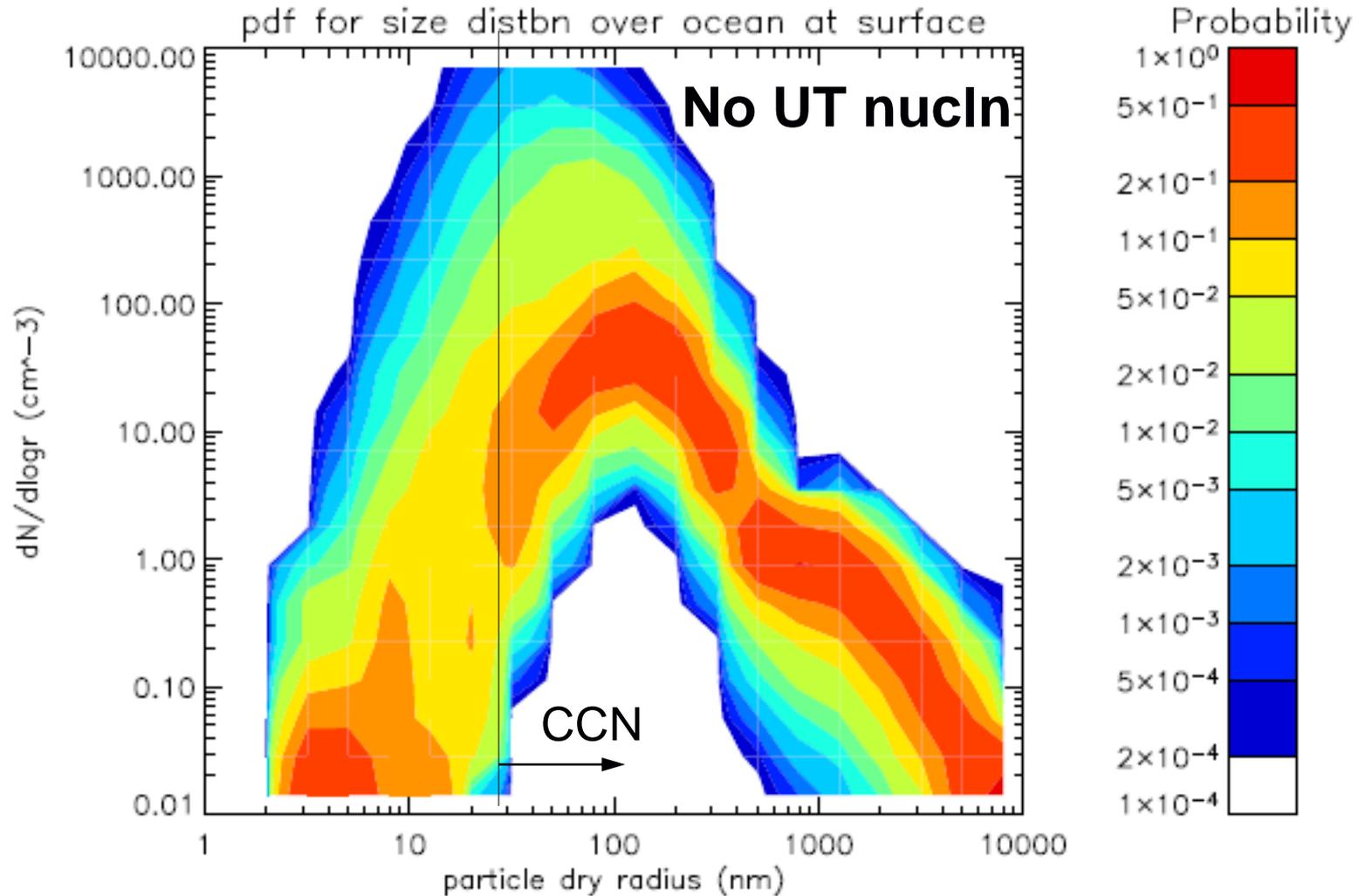


Marine BL particle size distribution in UKCA





Marine BL particle size distribution in UKCA

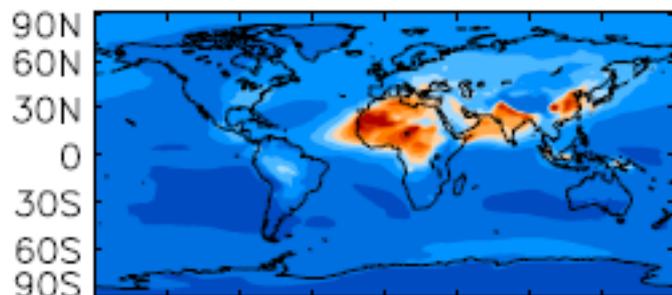


UKCA-aerosol applications: Nucleation effects on climate



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$\tau_{0.55}$ PD
UKCA-MODE BLN Off

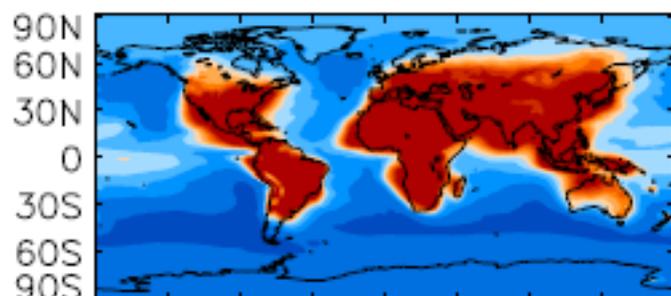


180 135W 90W 45W 0 45E 90E 135E 180
Mean: 0.108017



0 0.1 0.2 0.3 0.4 0.5

CDCN (1 km, cm^{-3}) PD
UKCA-MODE BLN Off

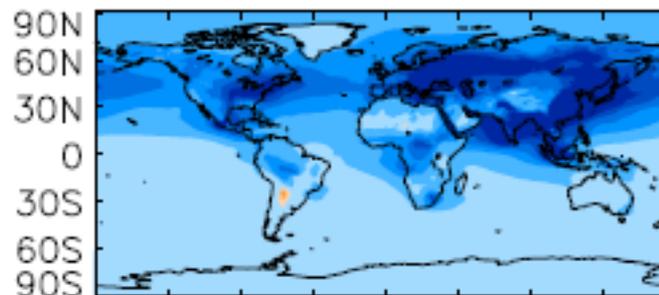


180 135W 90W 45W 0 45E 90E 135E 180
Mean: 60.5613



0 20 40 60 80 100

Clear-sky DRF PD w.r.t PI TOA
UKCA-MODE BLN Off

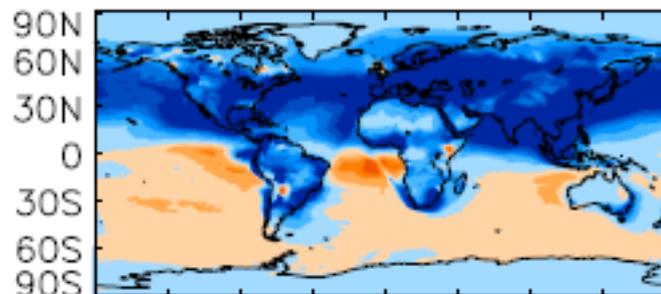


180 135W 90W 45W 0 45E 90E 135E 180
Mean: -0.966487



-2 -1 0 1 2

All-sky D+1IRF PD w.r.t PI TOA
UKCA-MODE BLN Off



180 135W 90W 45W 0 45E 90E 135E 180
Mean: -1.45930

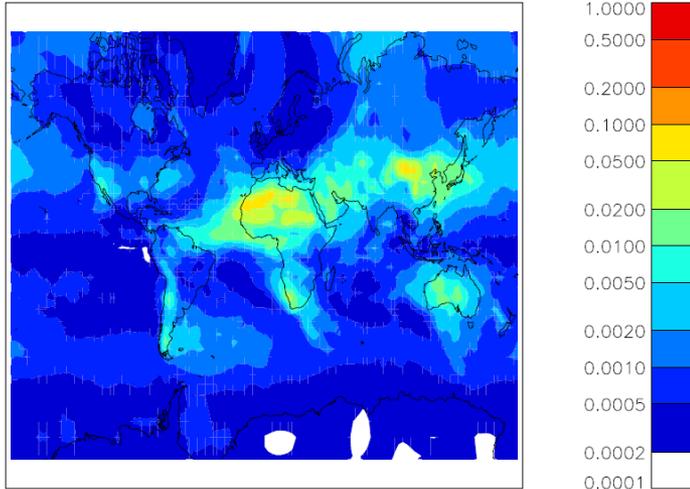


-2 -1 0 1 2

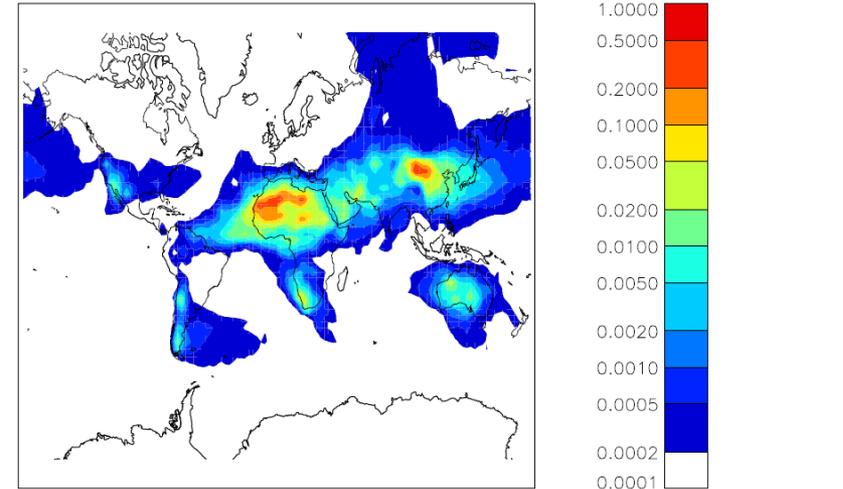
Nicolas
Bellouin
(Met Office)

UKCA aerosol applications: Nutrient availability to marine phytoplankton

Soluble Fe deposition (BJ06 solubility)

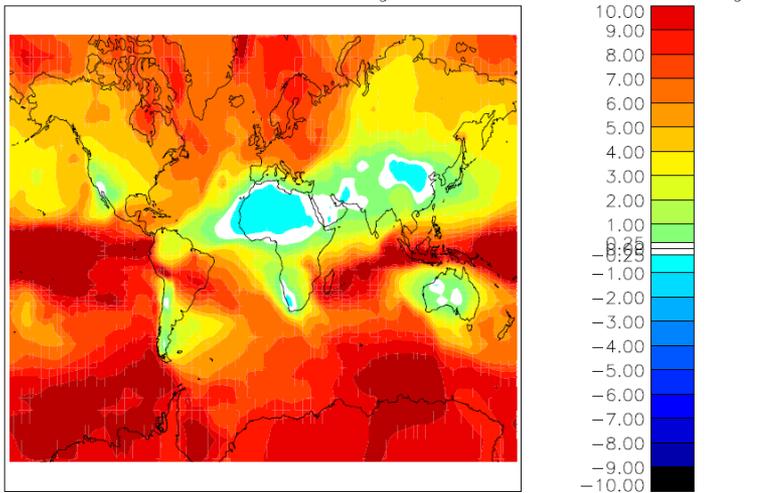
Solubility as in Baker & Jickells (2006) solFe deprn (mg/m²/da:

Soluble Fe deposition (1% solubility)

Constant solubility = 1% solFe deprn (mg/m²/day)

Difference in soluble Fe deposition simulated with BJ06 relative to 1%

Difference relative to 1% when using BJ06 relation relative change



GLOMAP dust deposition → Fe to marine biota

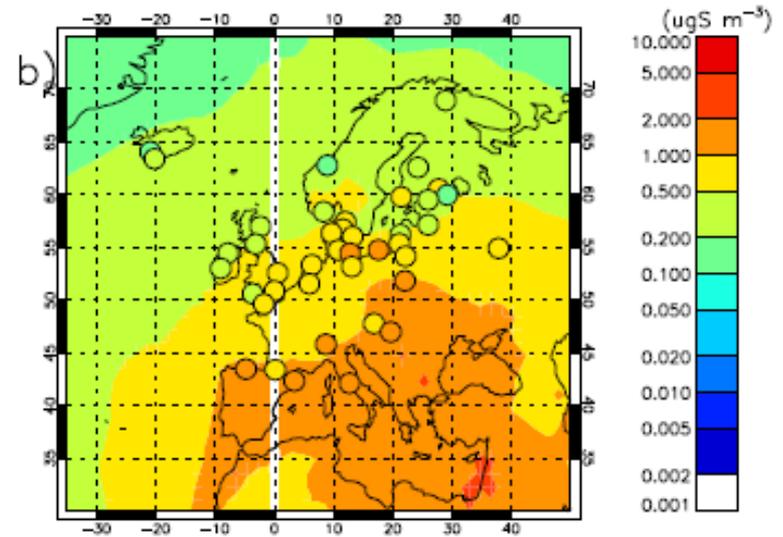
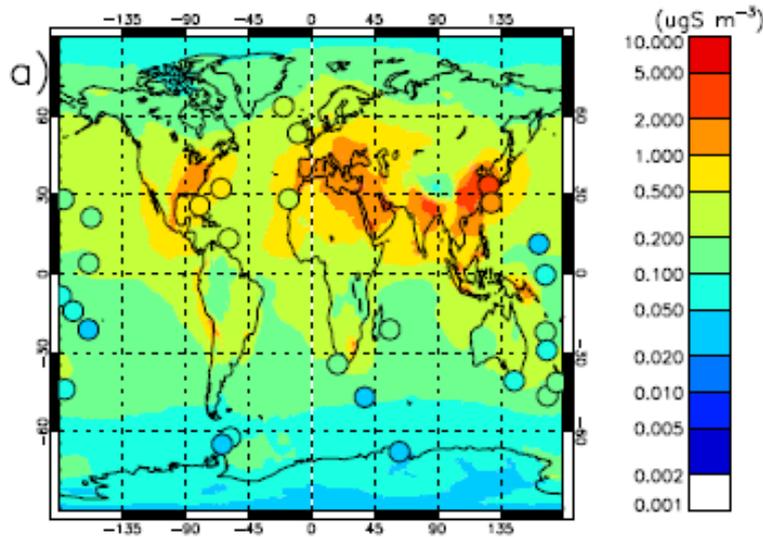
Soluble Fe deposition simulated by UKCA will perturb nutrient availability for phytoplankton in ocean biogeochemistry model in UKESM1.

Observations suggest far-field deposited dust much more soluble than that near source.

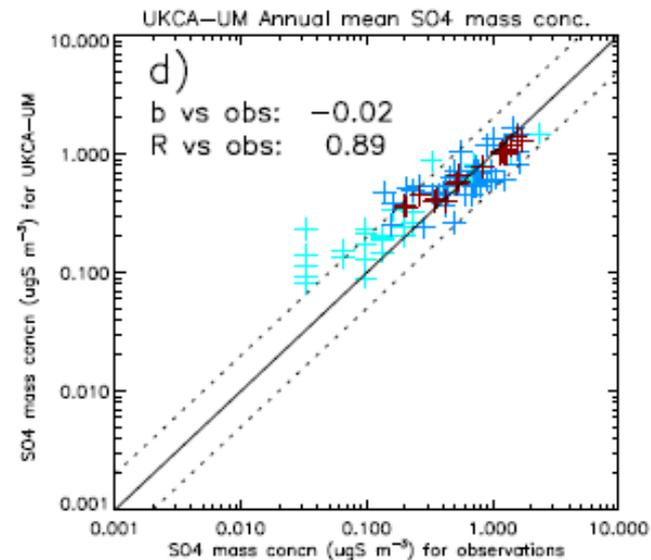
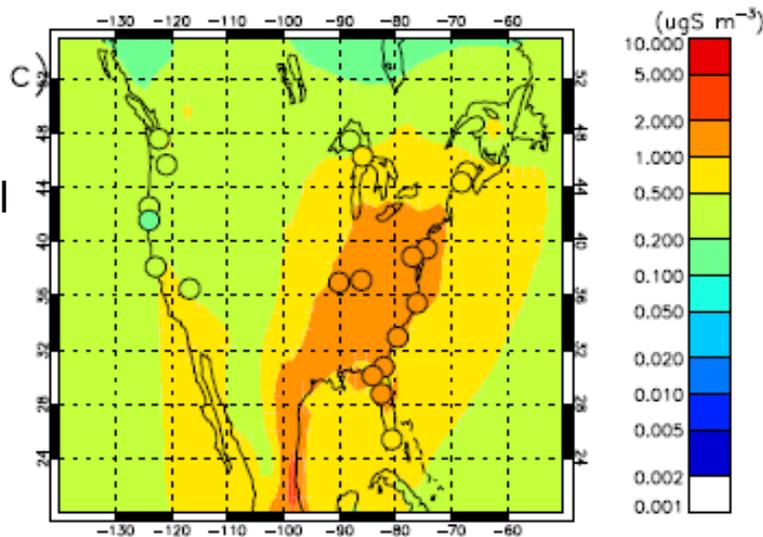
Having fresh and aged dust modes in UKESM allows variation in Fe solubility due to size and atmospheric processing (ageing) to be resolved.

UM-UKCA v8.4 GA4 CheST+GLOMAP RC6.2(xkjj): Evaluation vs Sulphate mass measurements
Europe (EMEP : Loevblad et al., 2004) & N. America (IMPROVE network; Malm et al., 2002)
and from marine sites (U. Miami network, Savoie et al., 2001)

Annual
mean

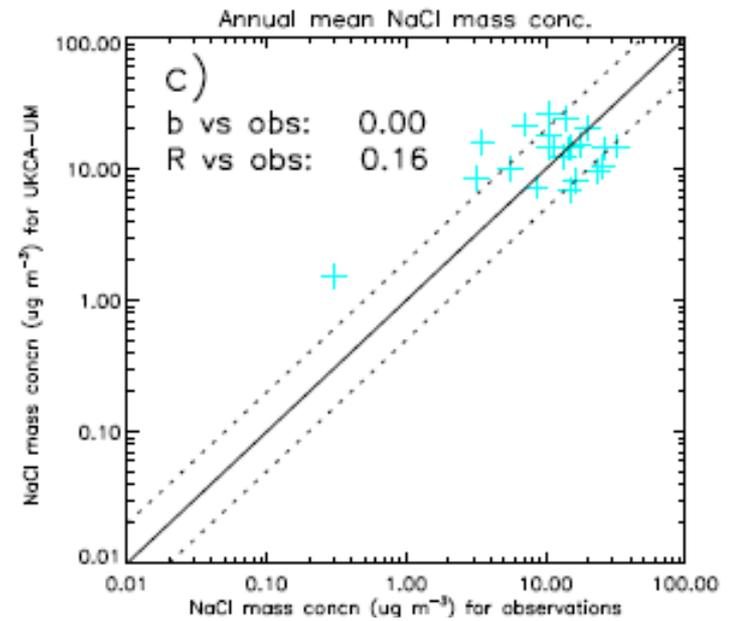
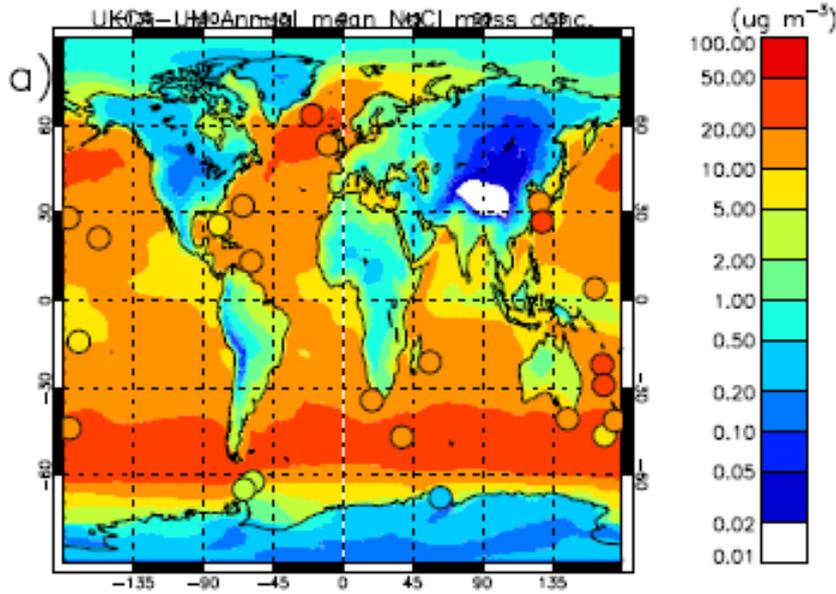


Annual
mean

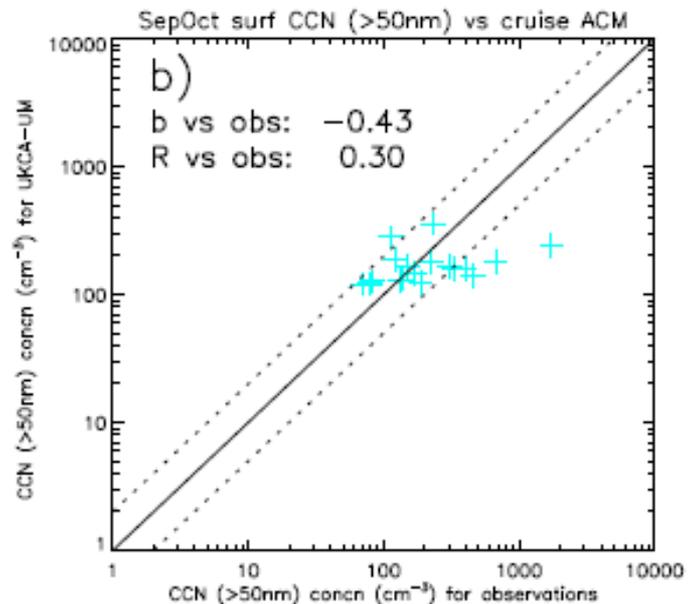
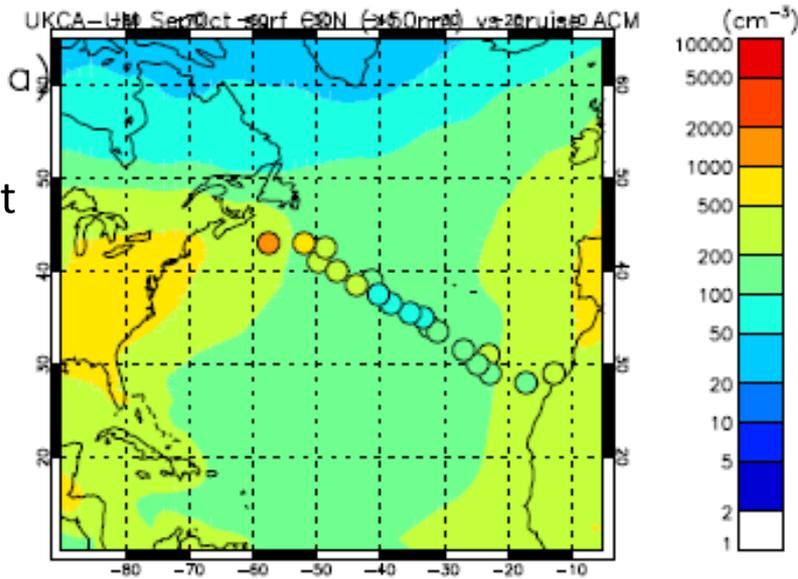


UM-UKCA v8.4 GA4 CheST+GLOMAP RC6.2(xkjj): Evaluation vs sea-salt mass measurements (U. Miami network, Savoie et al., 2001) and cruise CCN conc'ns (Van Dingenen et al., 1995)

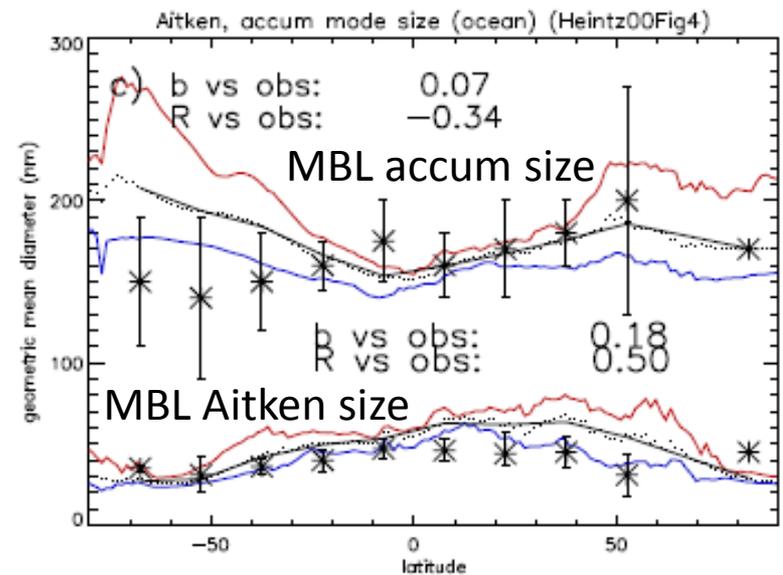
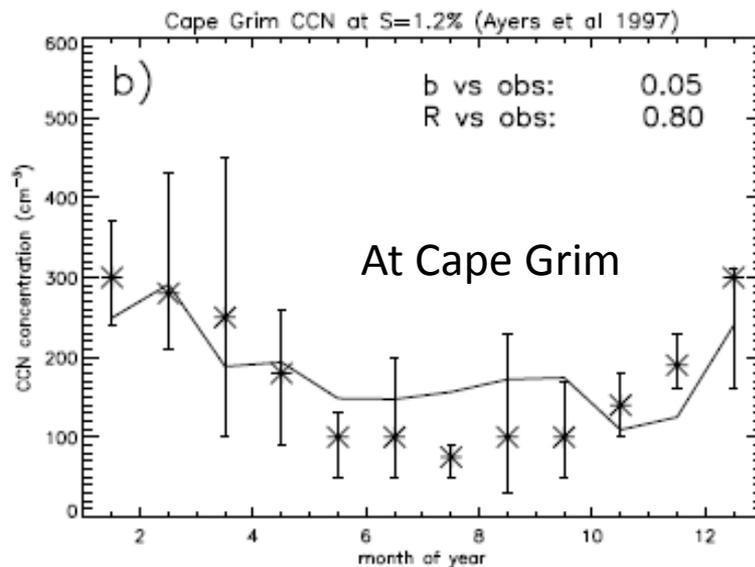
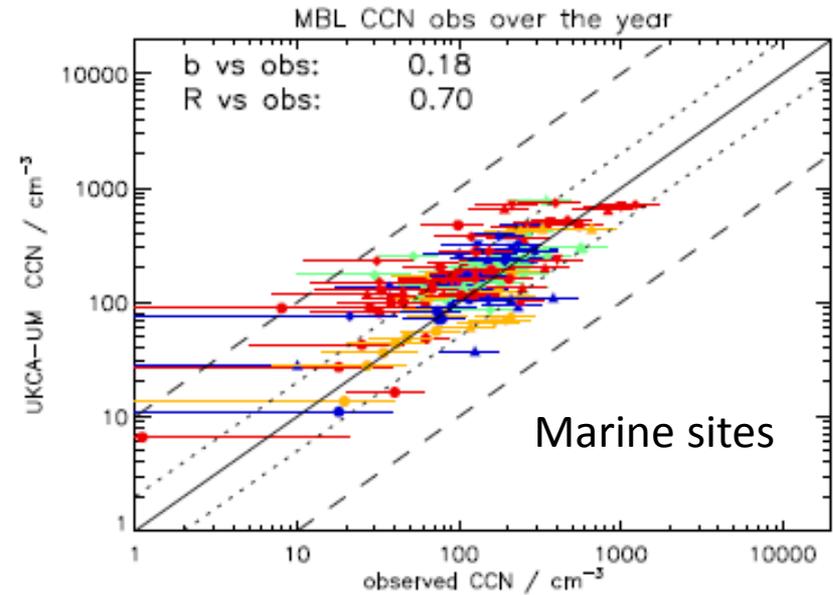
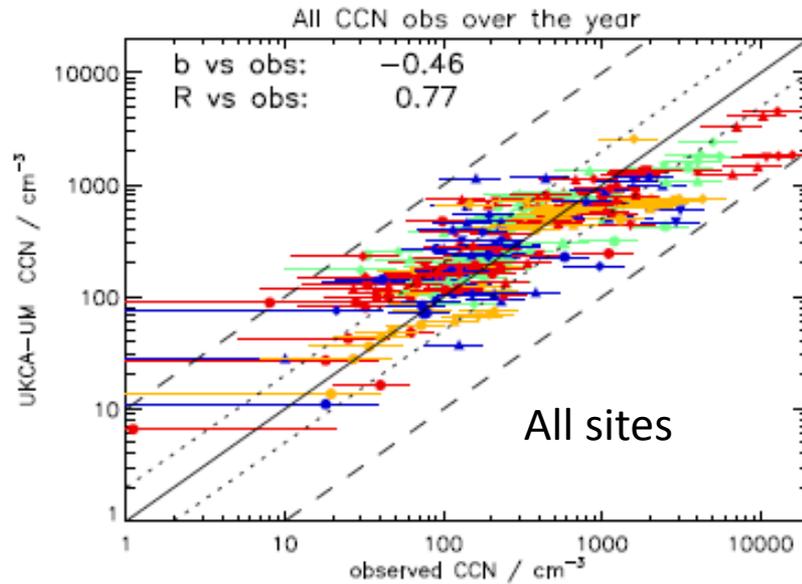
Annual mean



SepOct mean

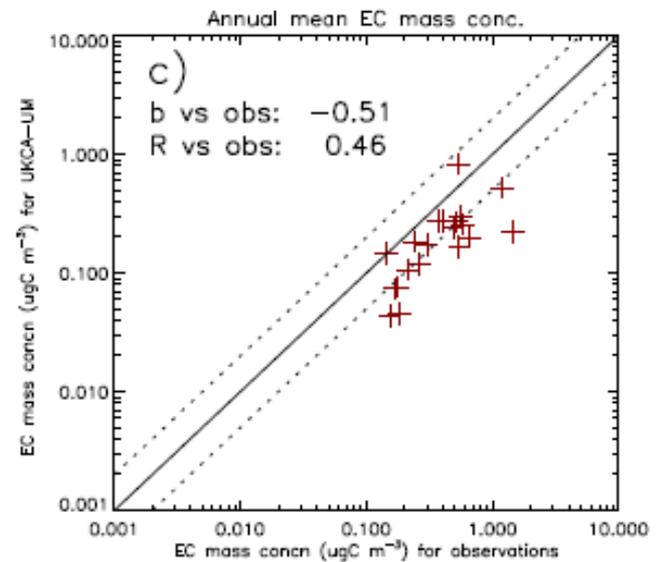
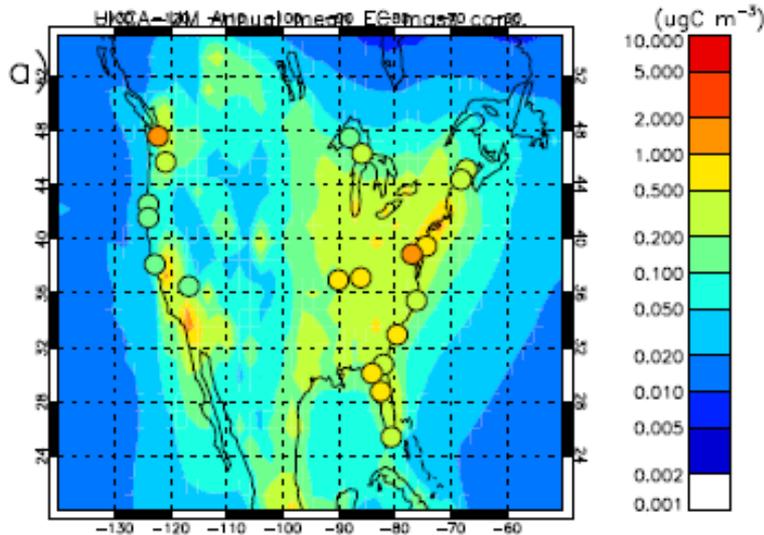


UM-UKCA v8.4 GA4 CheST+GLOMAP RC6.2(xkjj): Evaluation vs compilation of CCN concn measurements (Spracklen et al., 2011) and Aitken/accum mode size vs Heintzenberg (2000)

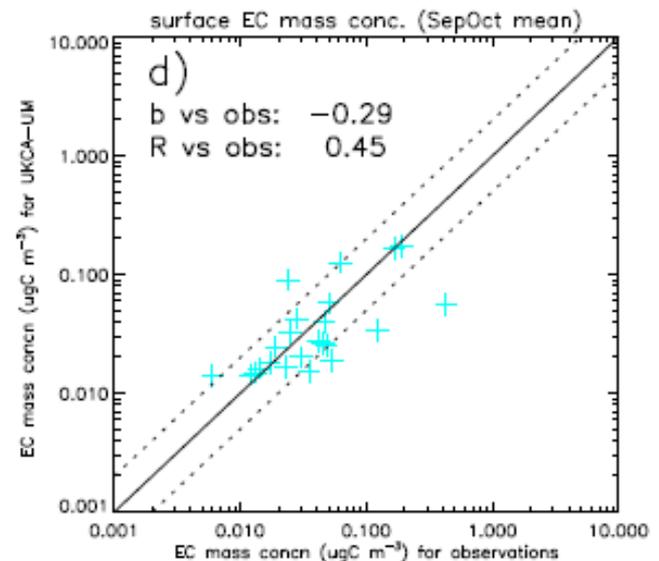
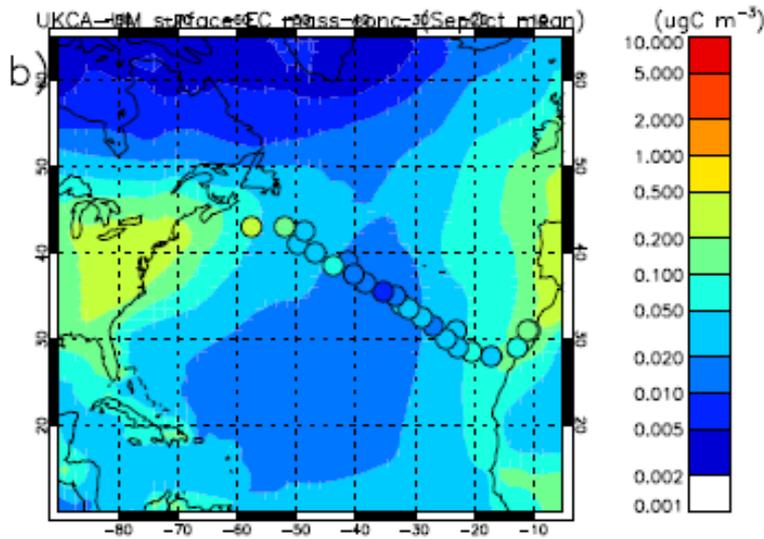


UM-UKCA v8.4 GA4 CheST+GLOMAP RC6.2 (xkjj) --- Evaluation vs EC/BC mass measurements
N. America (IMPROVE network) and from N. Atlantic cruise (Van Dingenen, 1995)

Annual
mean

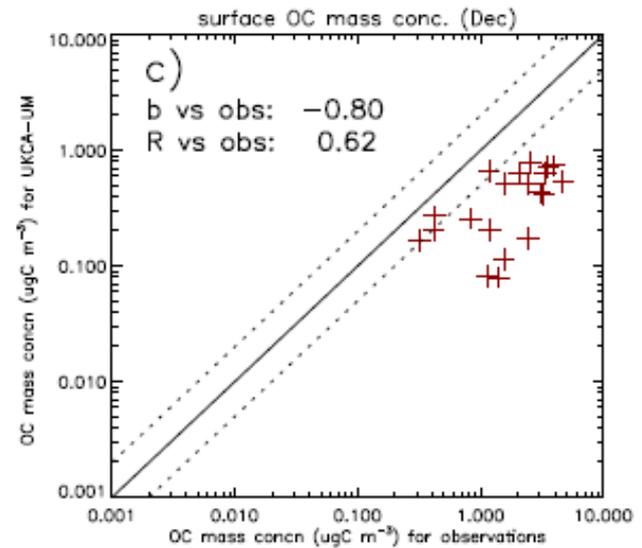
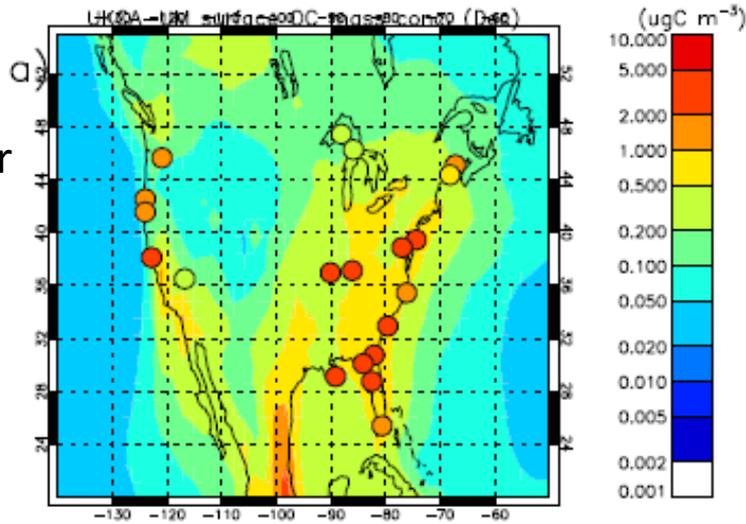


SepOct
mean

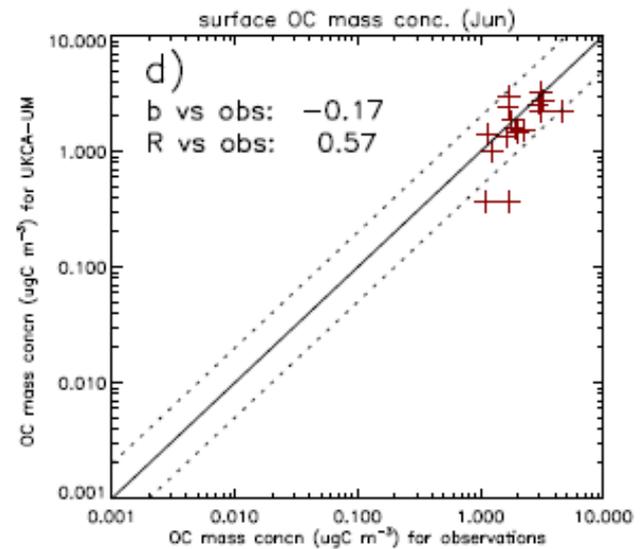
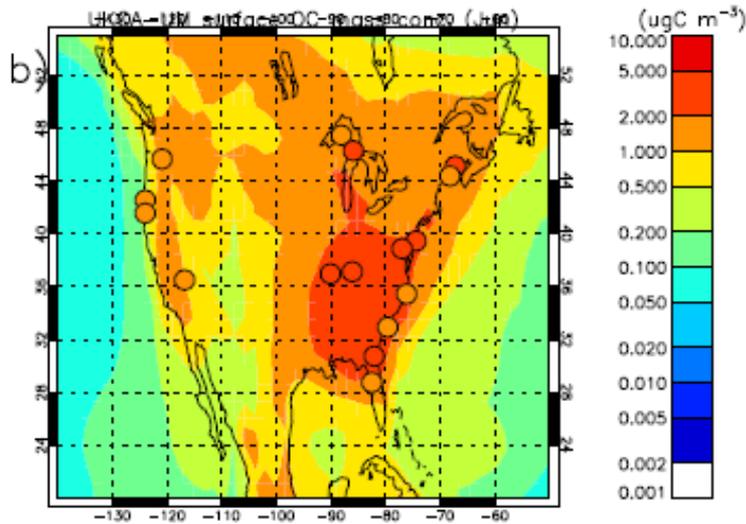


UM-UKCA v8.4 GA4 CheST+GLOMAP RC6.2(xkjgj): Evaluation vs Organic Carbon measurements from North America (IMPROVE visibility network; Malm et al., 2002)

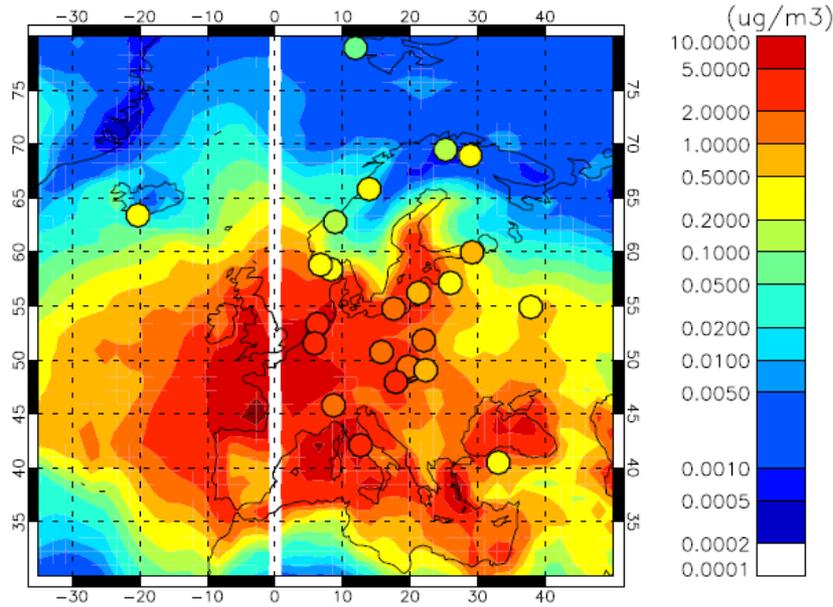
December



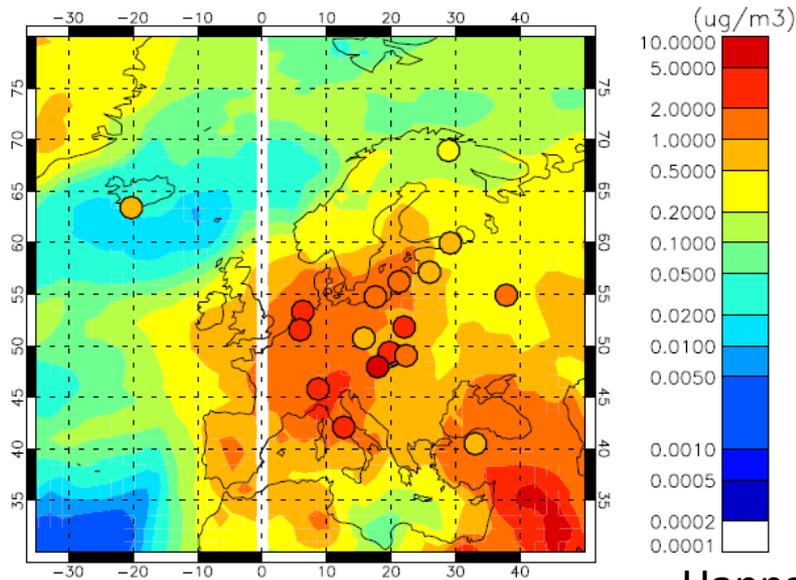
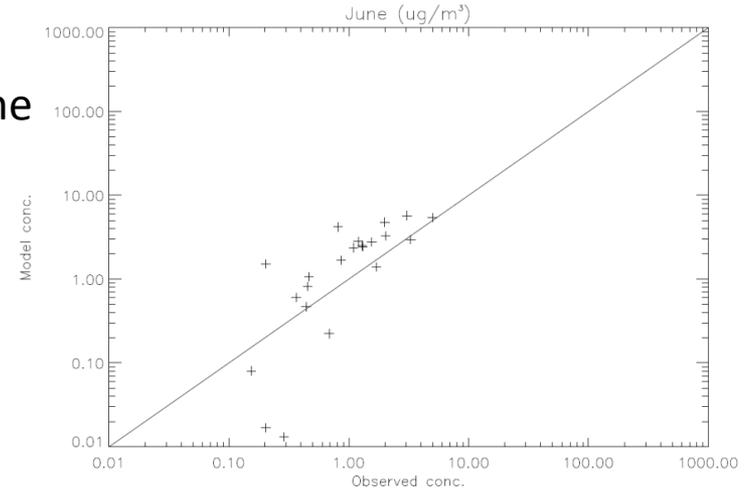
June



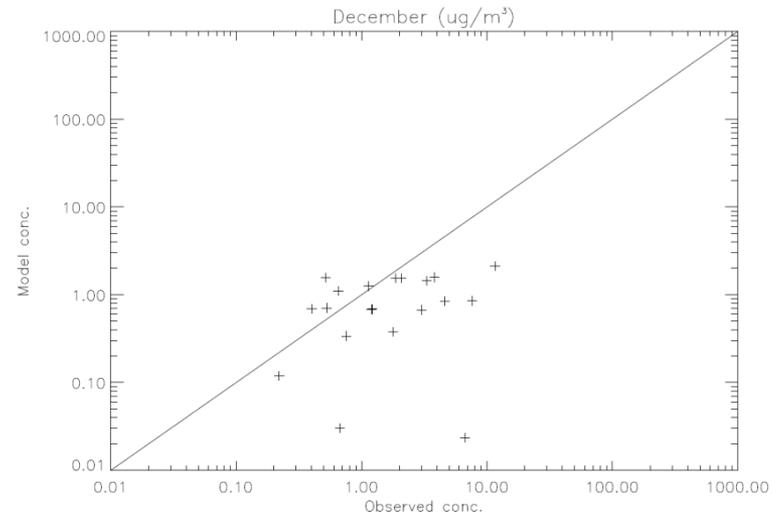
UM-UKCA v7.3 HG3-A-r2 CheT+GLOMAPnit (xiupl) - Evaluation vs nitrate measurements (June and Dec) over Europe (EMEP network; Loevblad et al., 2004)



June

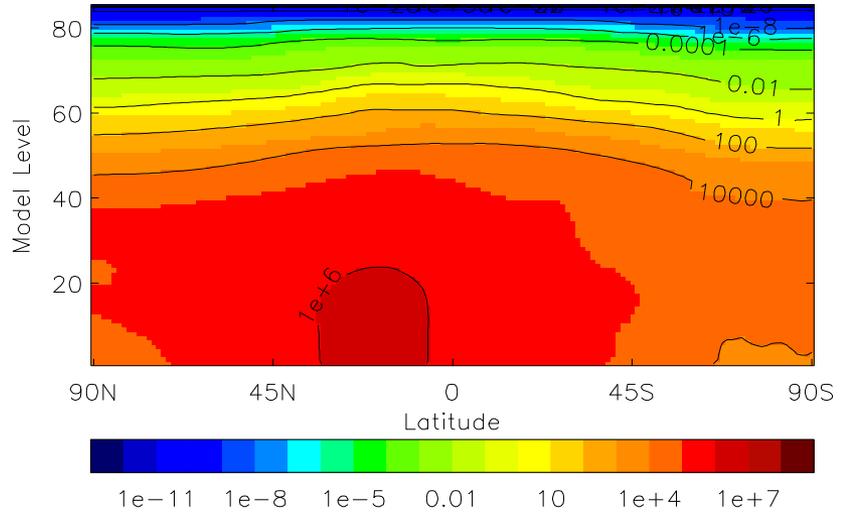


Dec

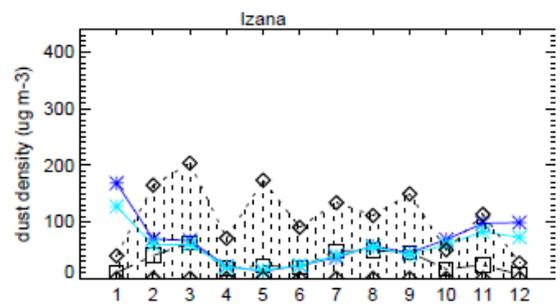
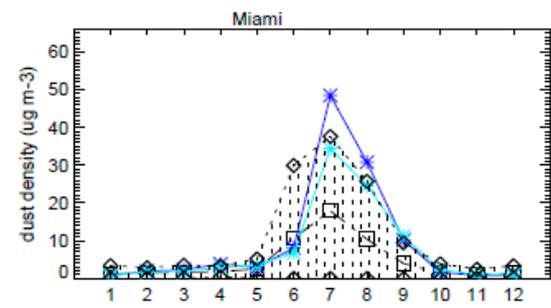
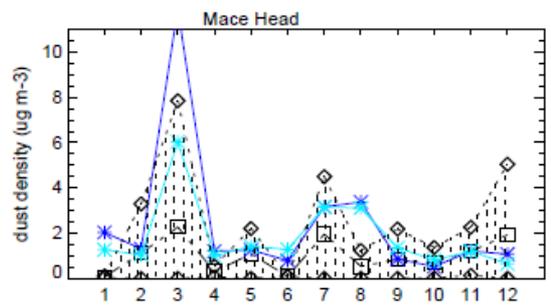
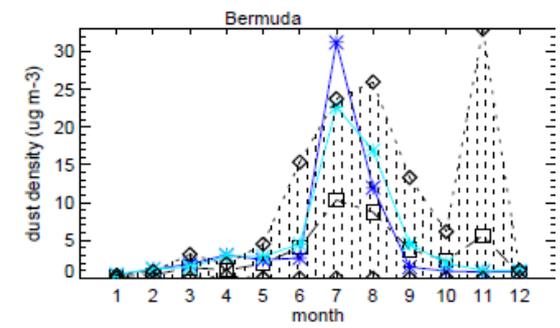
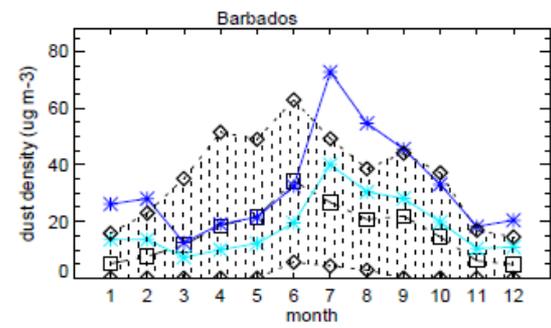
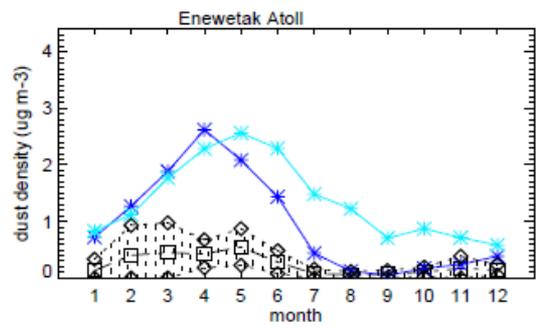
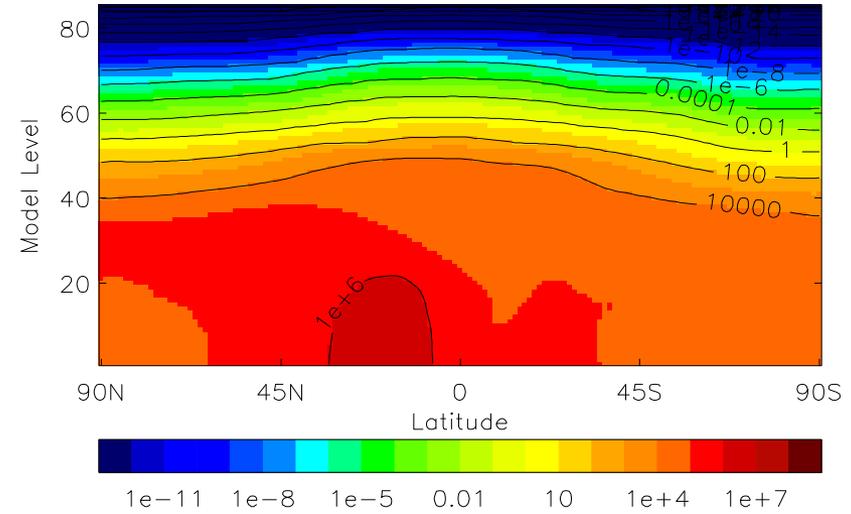


UM-UKCA v8.3 GA4 nochem+GLOMAP vs CLASSIC & Evaluation vs dust measurements (S Steph Woodward, Met Office (U. Miami network, Savoie et al., 2001)

\$LD/anrxqa.py19871201
cse ins number conc N m⁻³



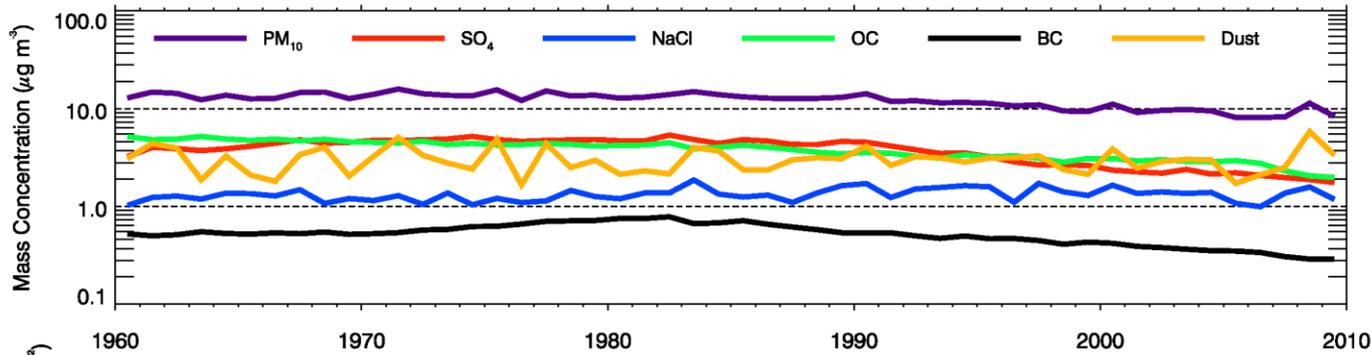
\$LD/anrxqa.py19871201
half of div3 +div4 +div5 number conc N m⁻³



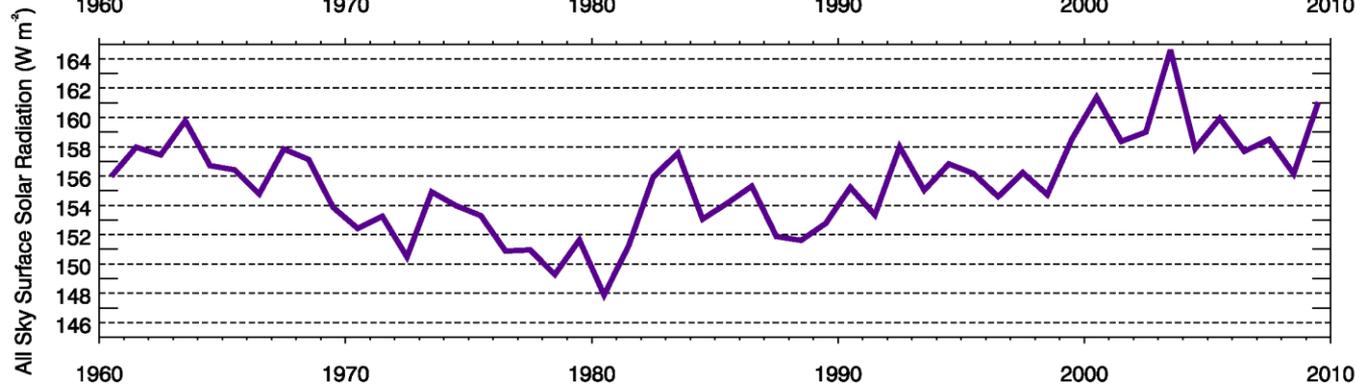
UM-UKCA v7.3 HG3-A-r2 CheT+GLOMAP

Modelled EU Trends in Mass, SSR and AOD

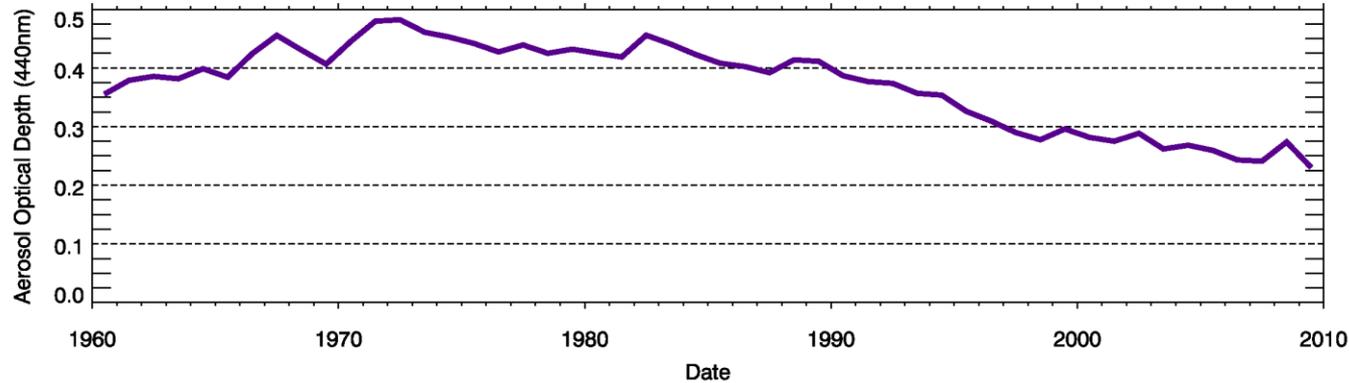
a)



b)

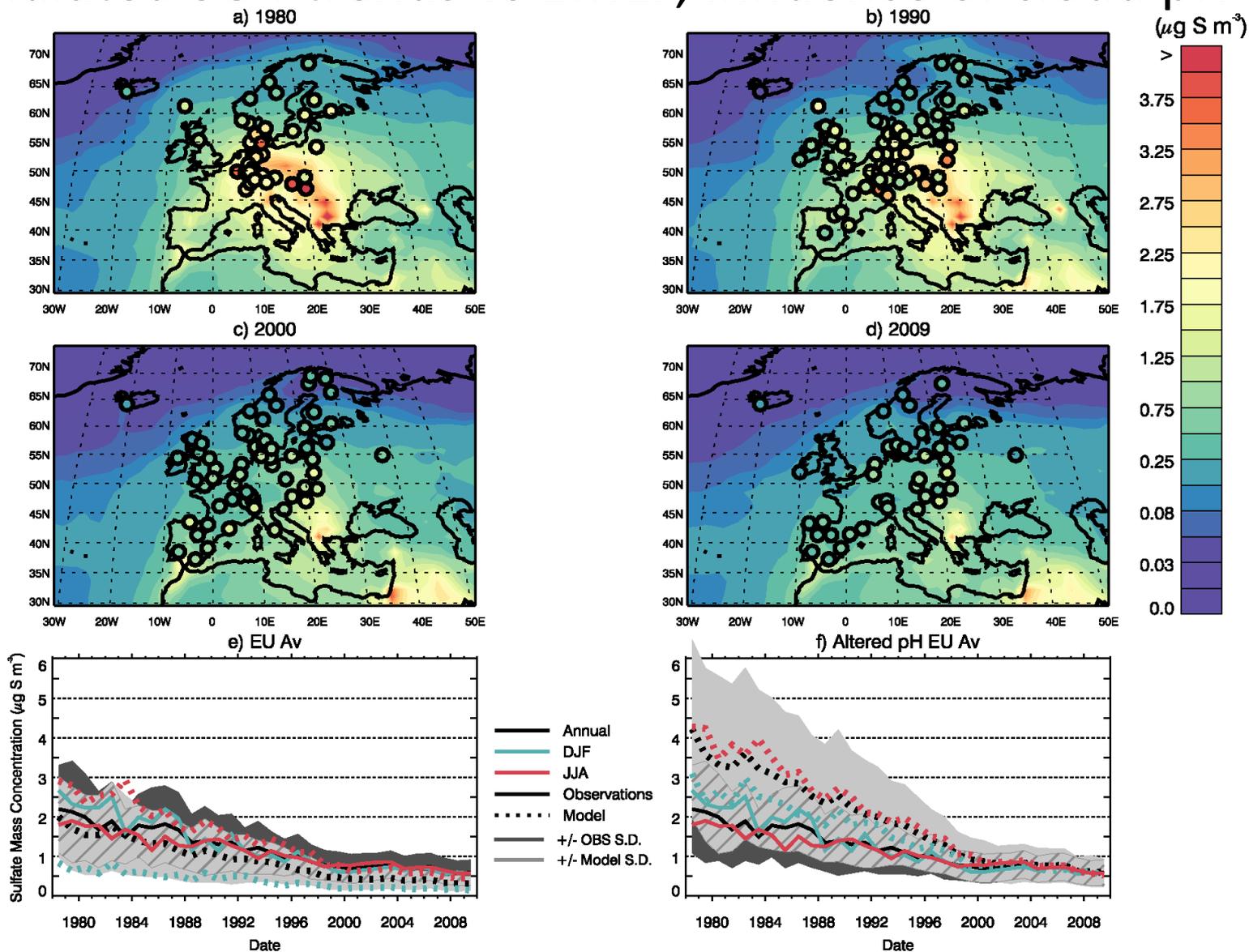


c)



UM-UKCA v7.3 HG3-A-r2 CheT+GLOMAP

Simulated SO₄ trends vs EMEP, influence of cloud pH

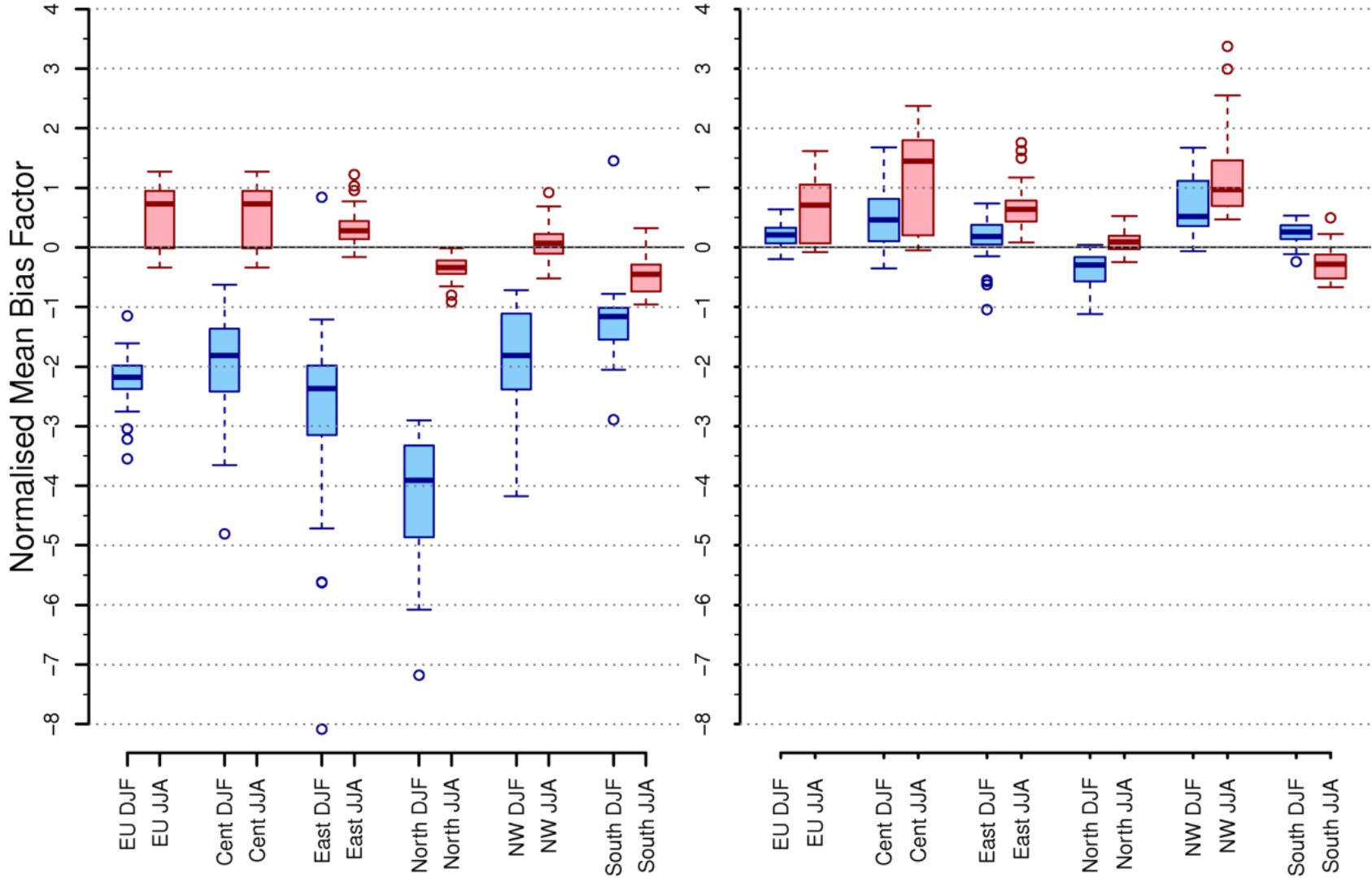


UM-UKCA v7.3 HG3-A-r2 CheT+GLOMAP

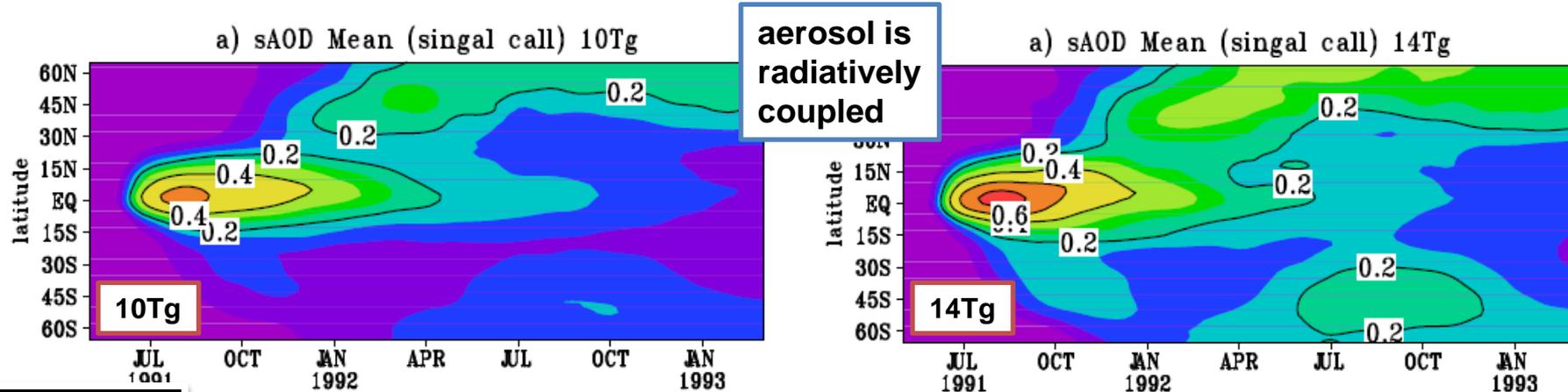
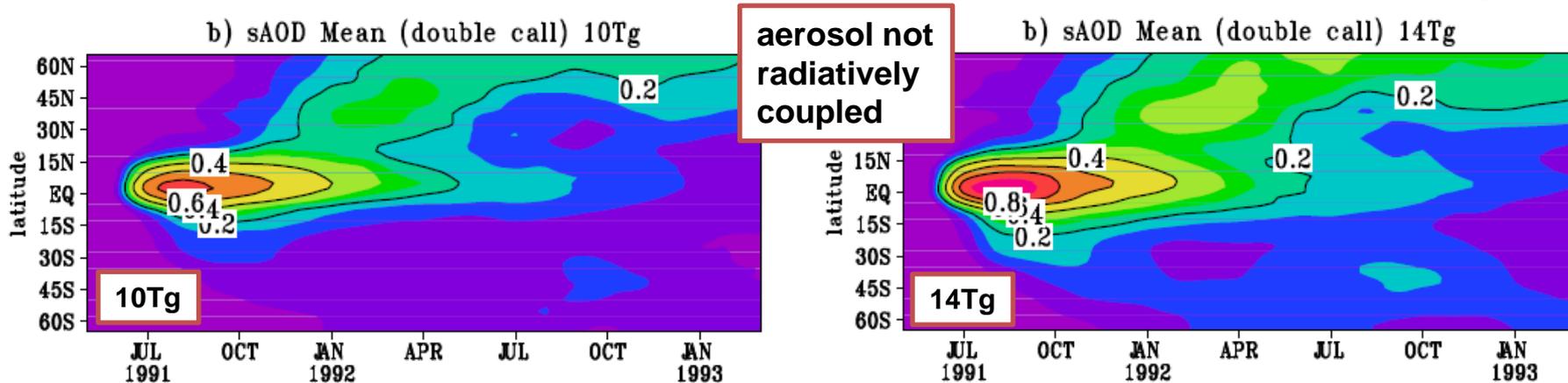
NMBF for SO4 vs EMEP (plus new pH) DJF, JJA

a) Original

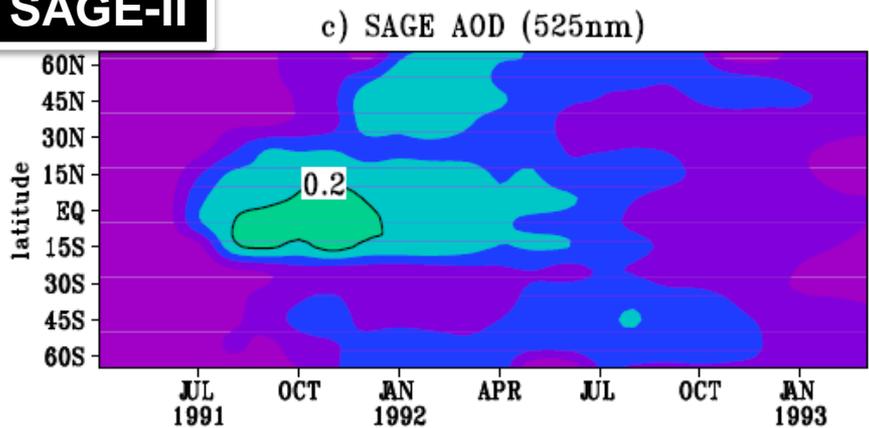
b) Altered Ph



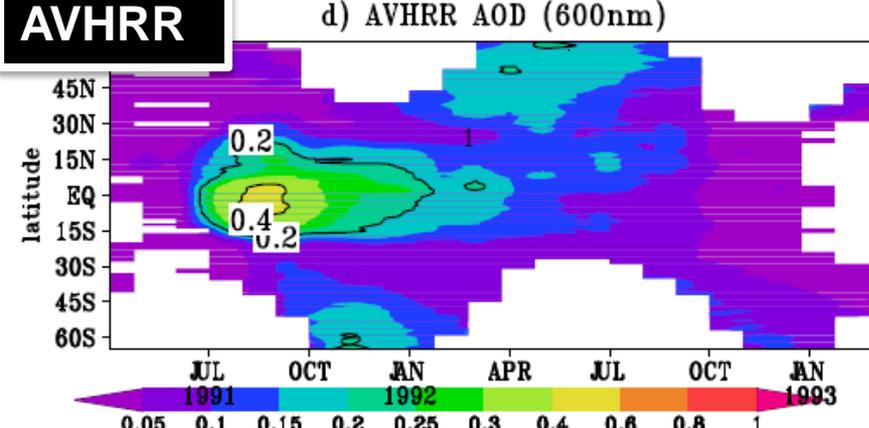
v7.3 CheS+GLOMAP N48L60: sAOD evolution vs SAGE-II sAOD & AVHRR anomaly



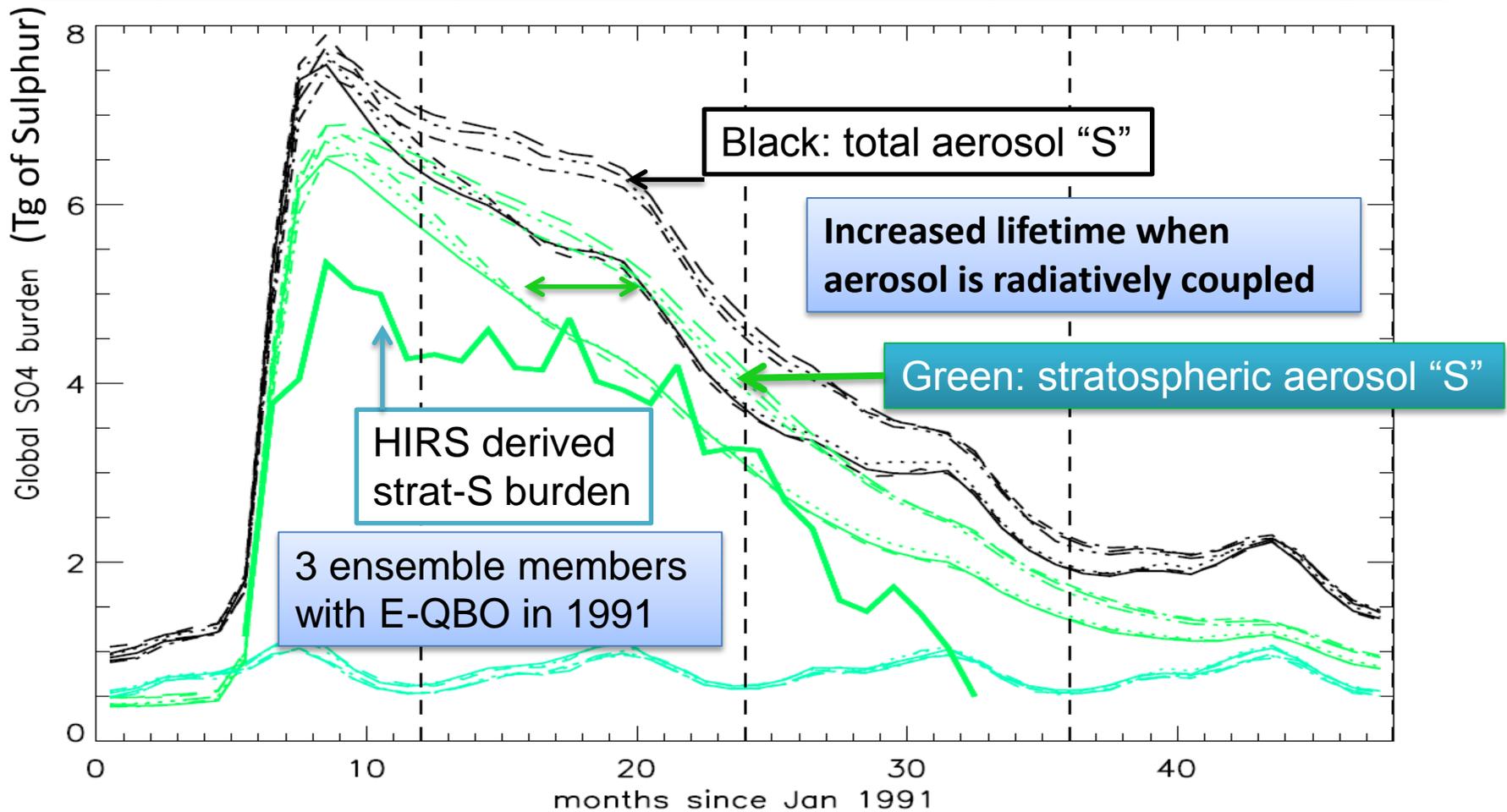
SAGE-II



AVHRR



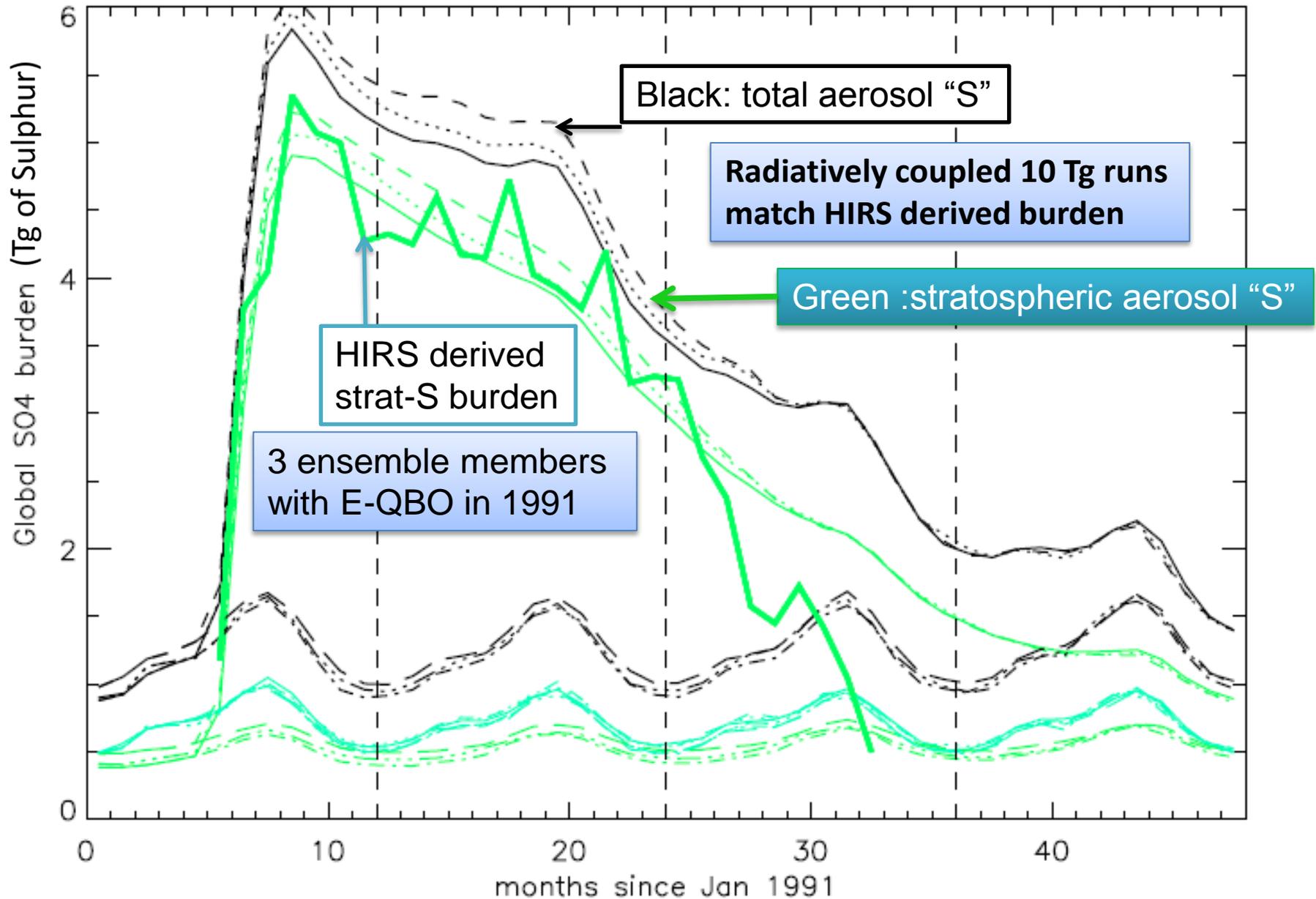
Stratospheric aerosol sulphur burden in UM-UKCA (thin green) vs HIRS (thick green)



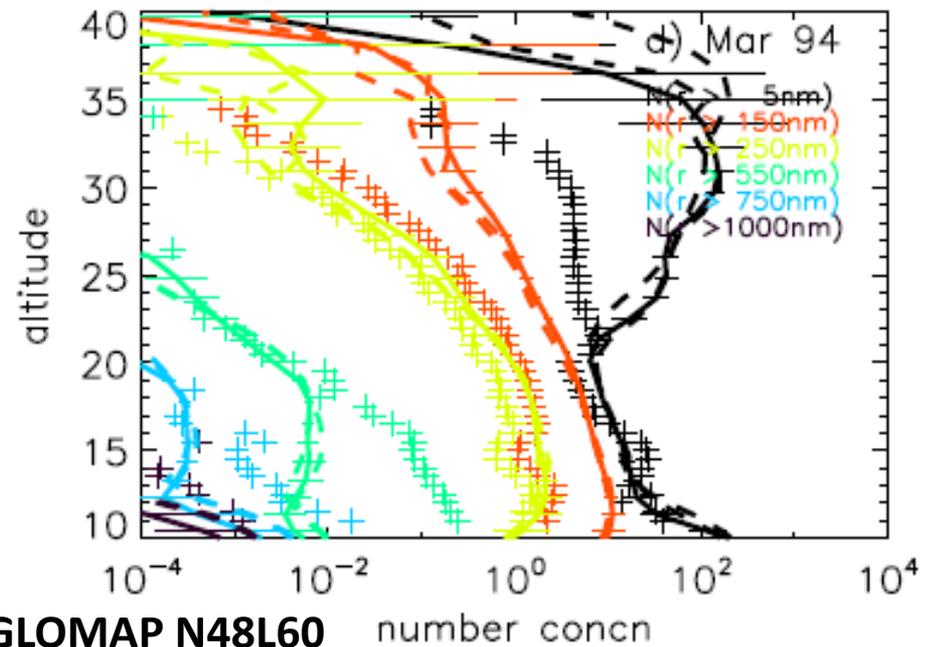
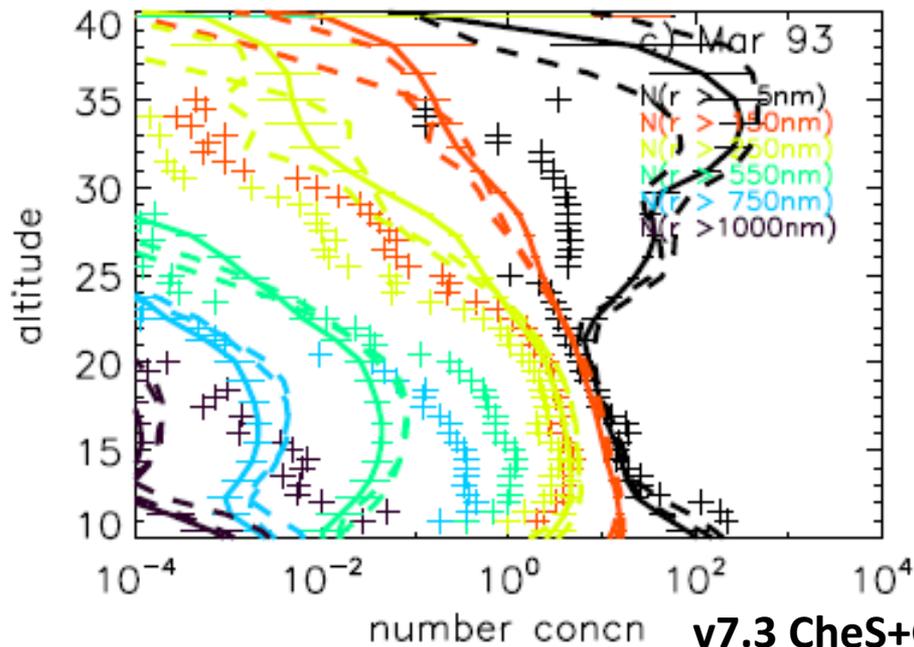
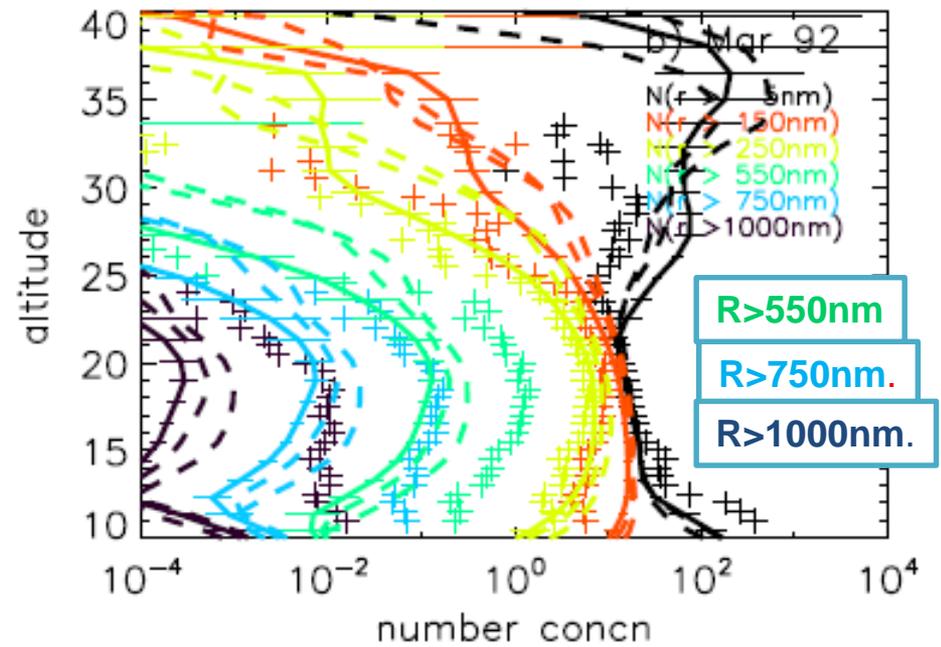
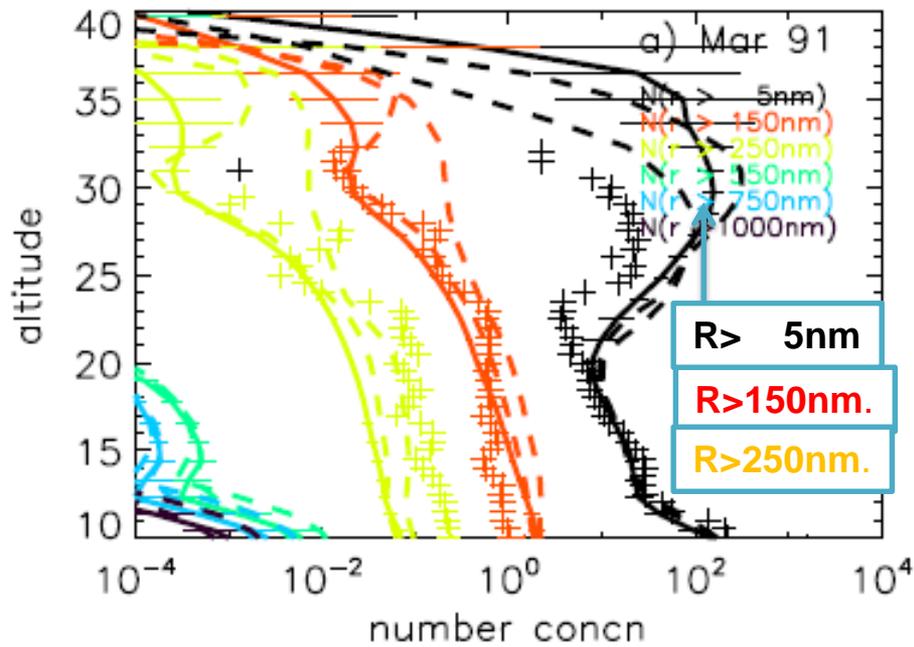
Impact of radiative coupling on lifetime in stratosphere

Mann et al., in preparation

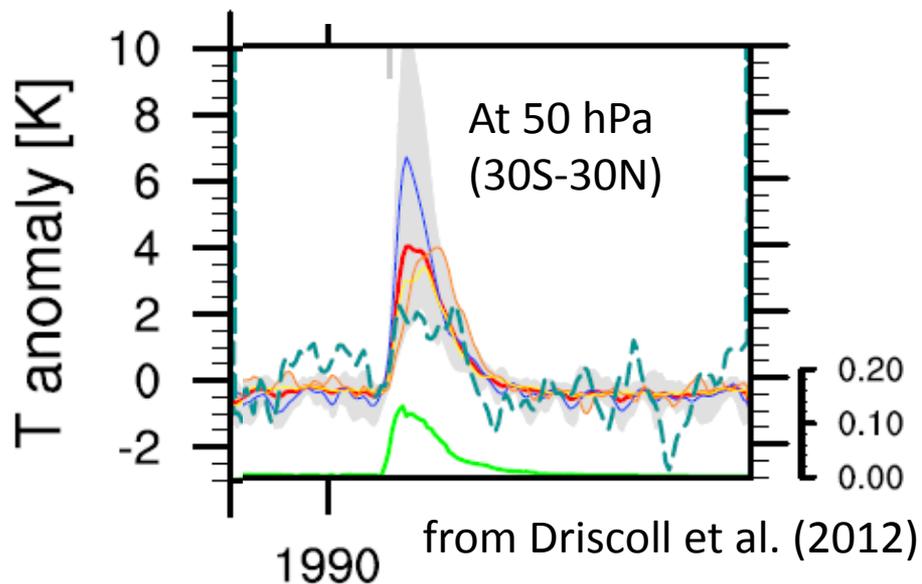
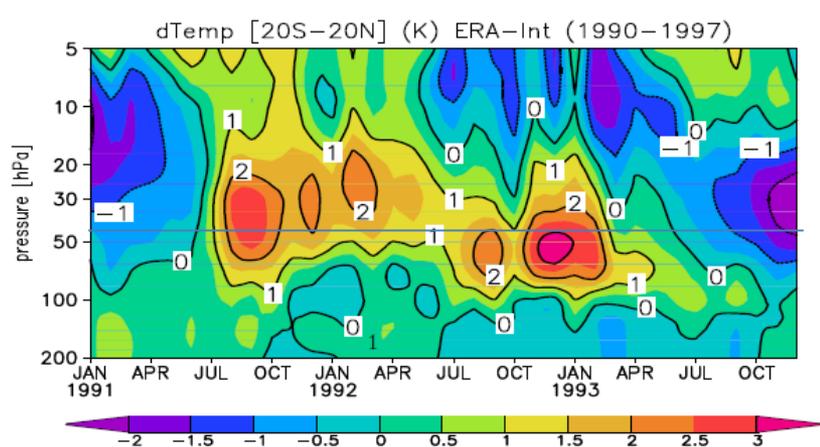
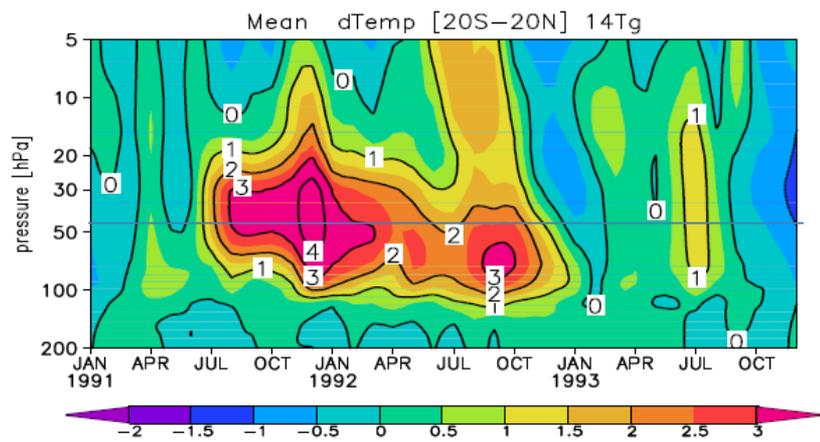
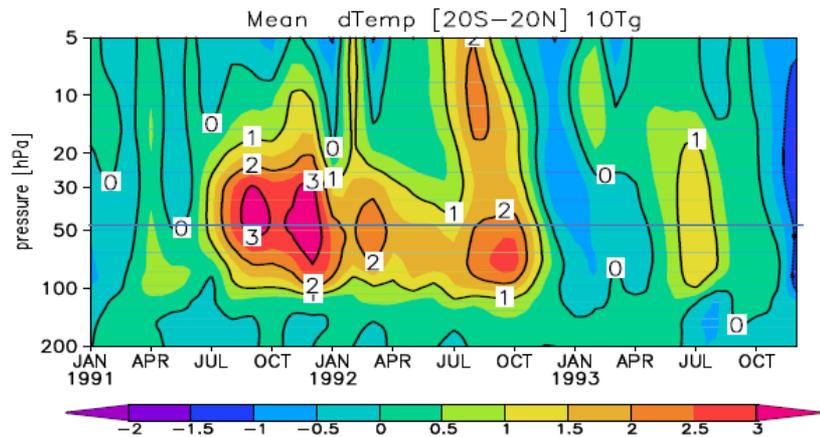
Stratospheric aerosol sulphur burden in UM-UKCA (thin green) vs HIRS (thick green)



Stratospheric particle concentration profiles in UM-UKCA vs Laramie CPC/OPC (pluses)



Post-eruption warming evolution in tropical stratosphere in UM-UKCA vs ERA-interim

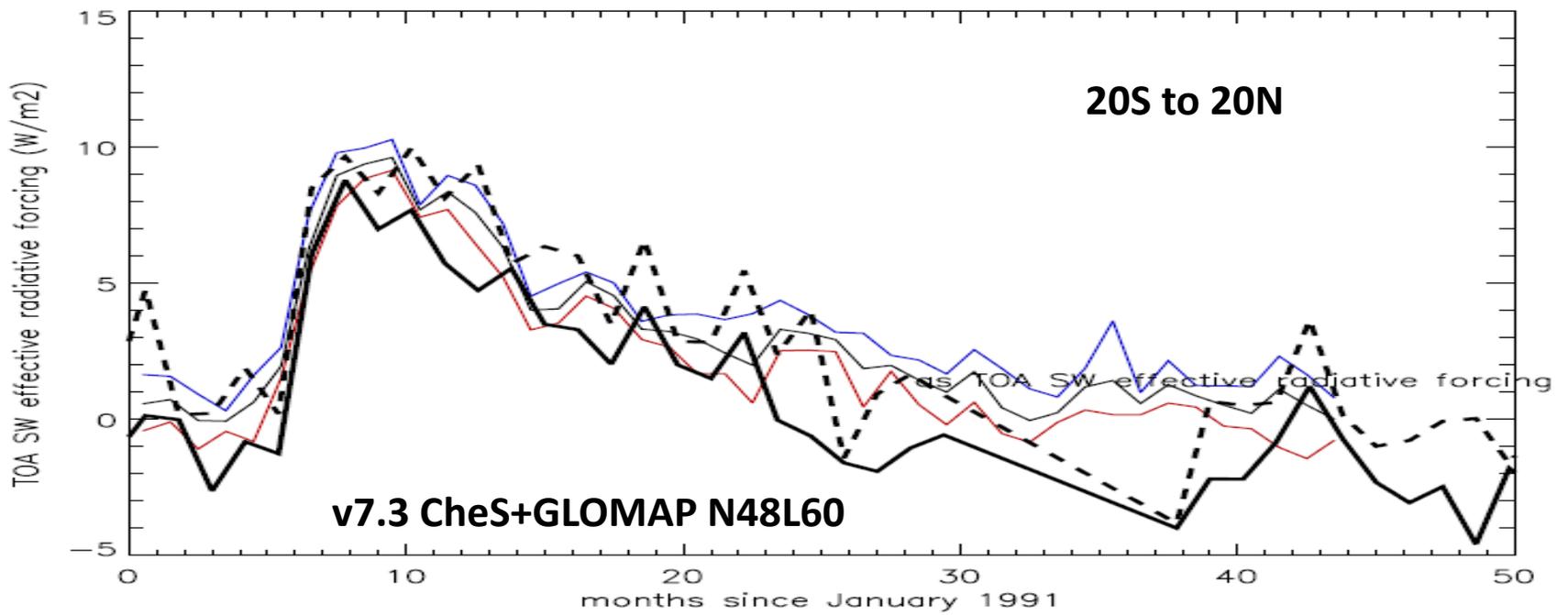
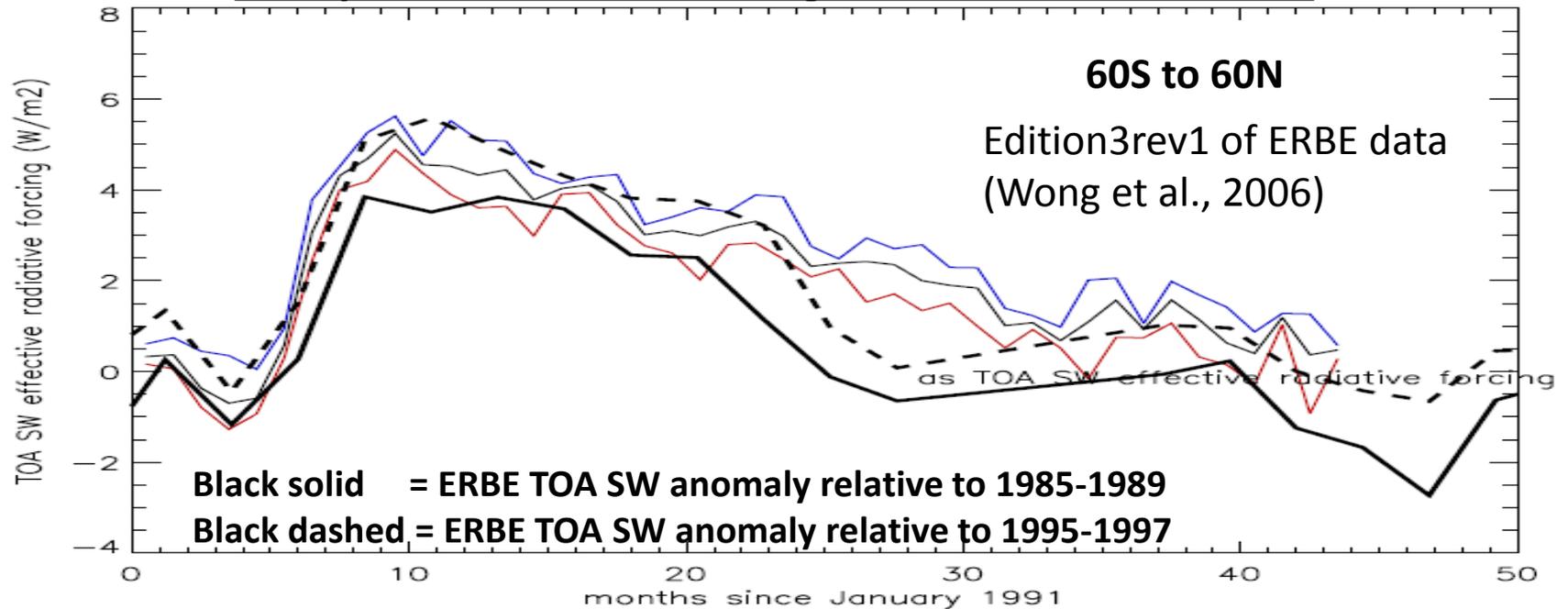


Stratospheric warming from 10 Tg Pinatubo simulation captures approximately the peak seen in the ERA-interim re-analyses.

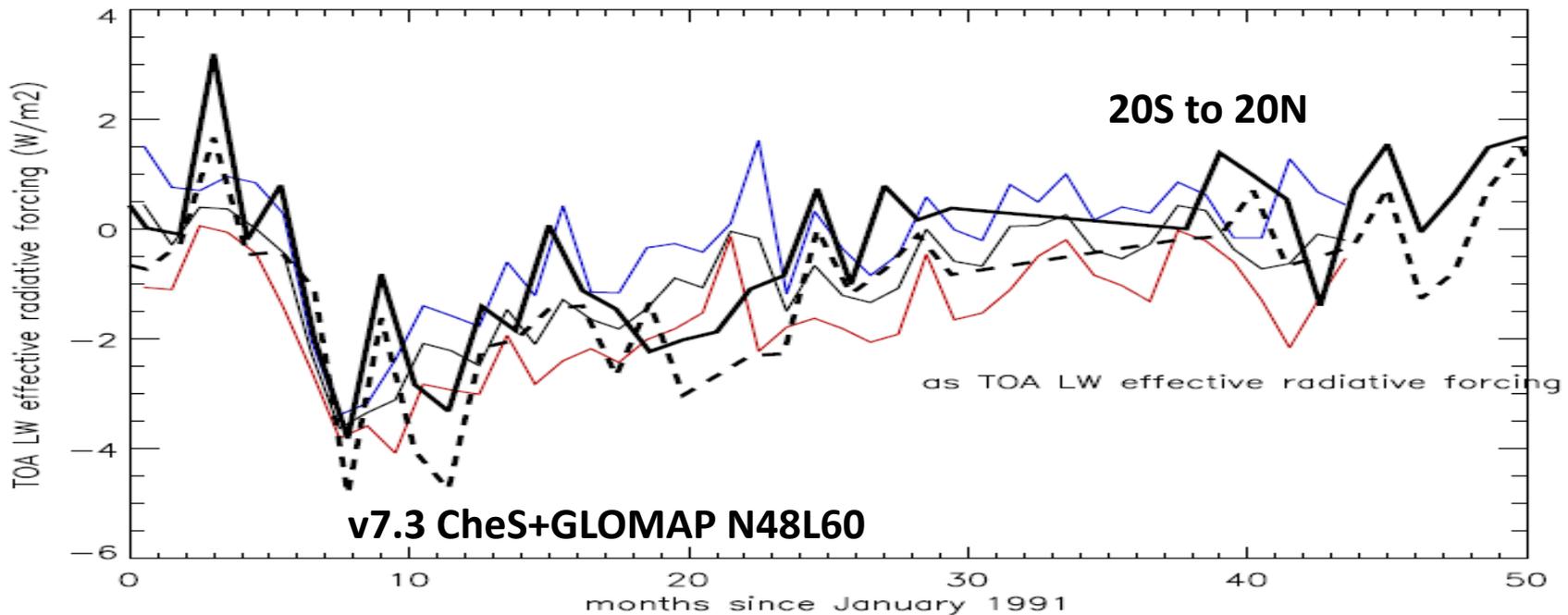
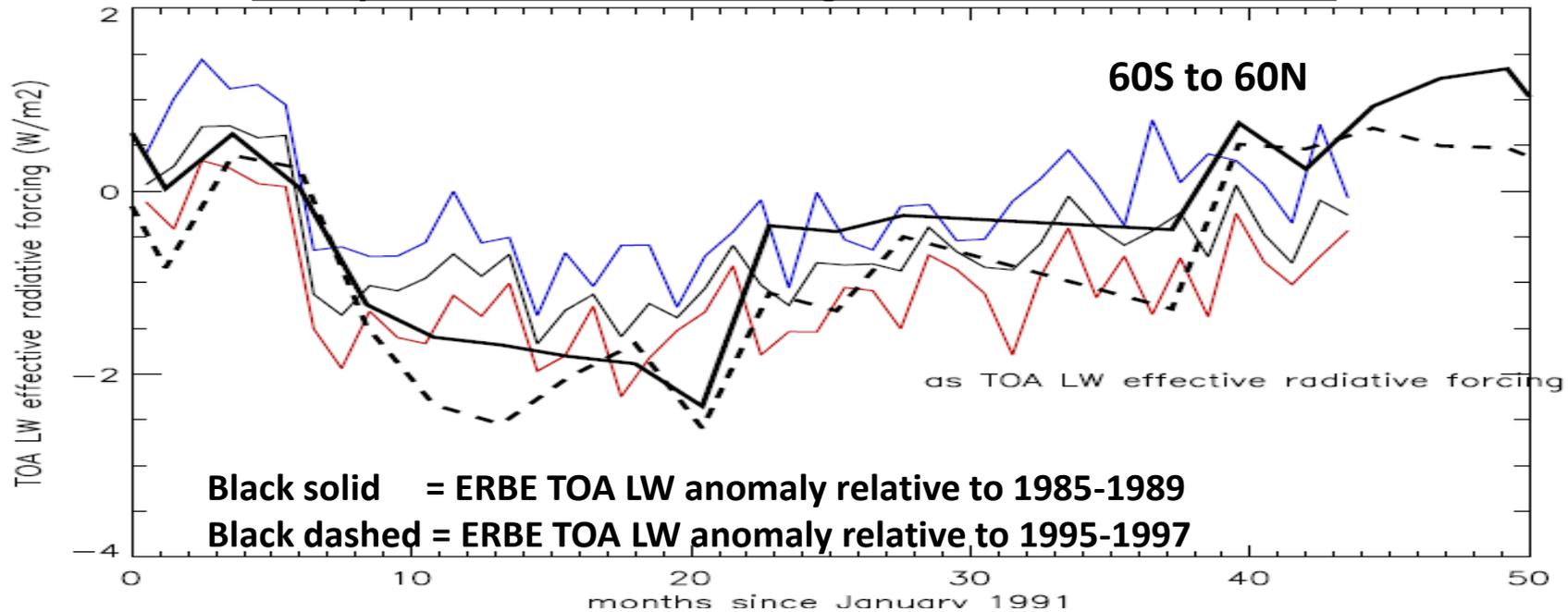
14 Tg Pinatubo injection over-estimates the warming.

Note the delta-Ts include any offset from cooling of the tropical lower stratosphere that results from any chemical and dynamical ozone losses.

All-sky TOA SW radiative forcing timeseries from UM-UKCA



All-sky TOA LW radiative forcing timeseries from UM-UKCA





Summary of UKCA evaluation

- UKCA evaluation suite includes comparisons to sulphate, BC, POM, sea-salt at surface sites. Comparisons to dust, nitrate also carried out.
- Also wider evaluation against CCN, size-resolved particle concentrations
- Trends in sulphate, PM, AOD, surface solar radiation across Europe
- Modal dust scheme now operating well compared to bin scheme
- Evaluation of stratospheric aerosol properties against range of datasets covering both quiescent and volcanically perturbed conditions
- Simulated SW and LW Pinatubo forcings in excellent agreement with observed anomalies from ERBE for 10 Tg injection of SO₂
- Observed global variation and evolution of strat-AOD, strat-S-burden, stratospheric warming, and TOA radiation anomalies all consistent with simulations with 10 Tg SO₂ injection
- Stronger efficacy of Pinatubo forcing per unit mass of aerosol sulphur



UM-UKCA next steps

First stage of UKCA is complete whereby UKCA aerosol scheme using GLOMAP aerosol microphysics is fully integrated within the UM at GA7

Next steps in UKCA development and science:

- Full GLOMAP configuration with modal dust for Earth System science
- Aerosol Optical Depth assimilation for aerosol re-analyses via implementation of GLOMAP into ECMWF Integrated Forecasting System (IFS) (EU-funded MACC projects have built on capability from GEMS project).
- Fully coupled aerosol-climate and aerosol-ES simulations in UKCA and QUEST Earth System Model – explore aerosol-related ES interactions.
- Extended UKCA inorganic composition simulating size-resolved NH_4 & NO_3 Scheme developed by Francois Benduhn to be incorporated into UKCA.
- Secondary organic aerosol chemistry scheme to more realistically simulate production of biogenic and anthropogenic SOA in UKCA
- Stratospheric aerosol and geoengineering simulations

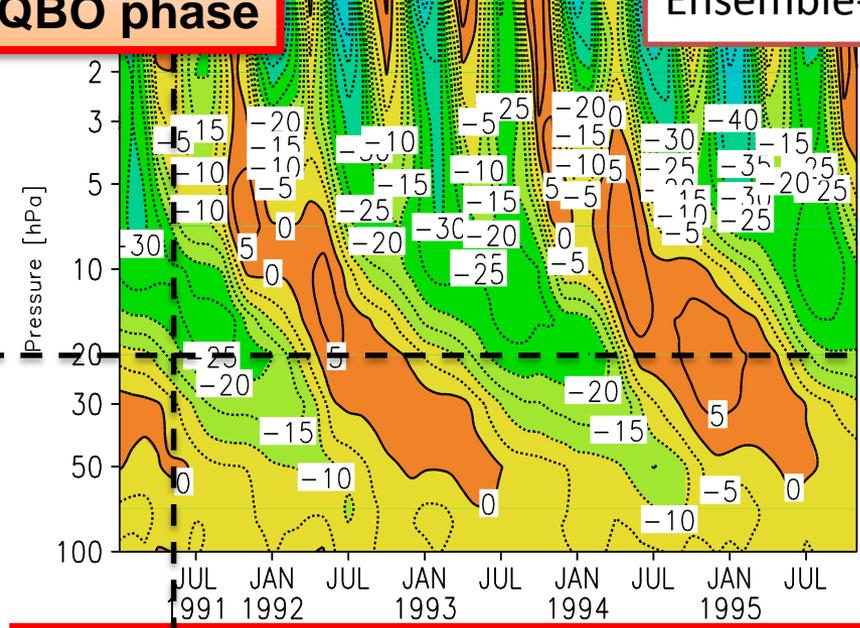
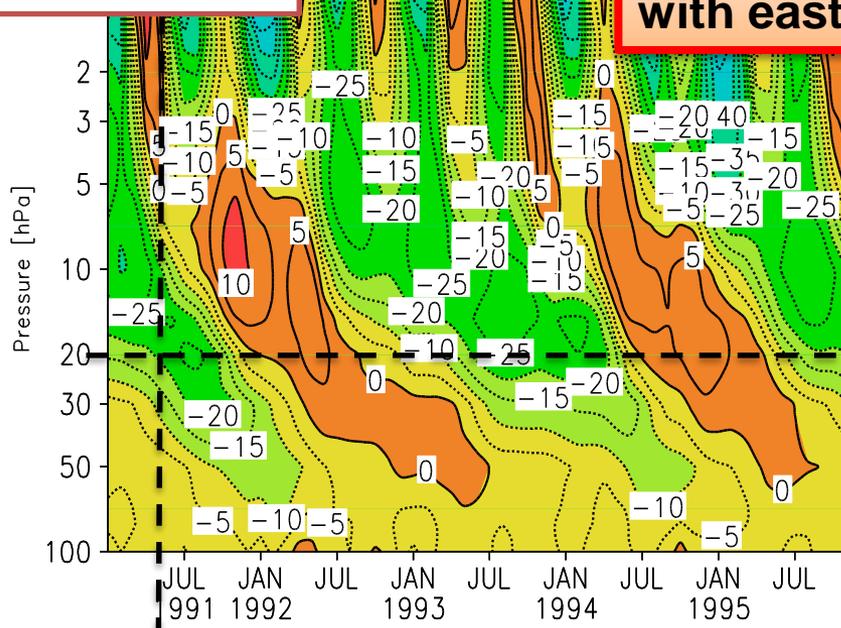
Ensemble-1

Tropical [15SN] Uwind

3 ensemble members with easterly QBO phase

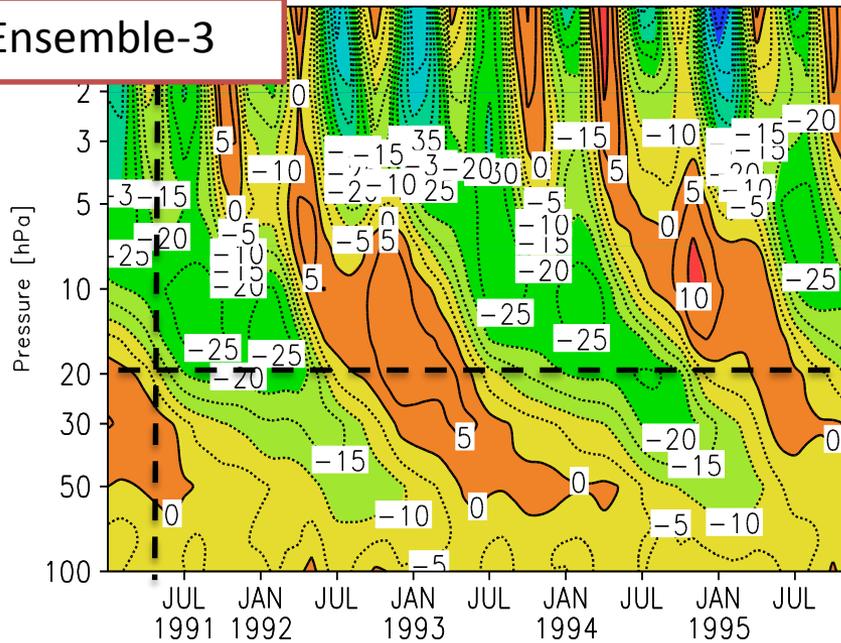
Tropical [15SN] Uwind (°C)

Ensemble-2



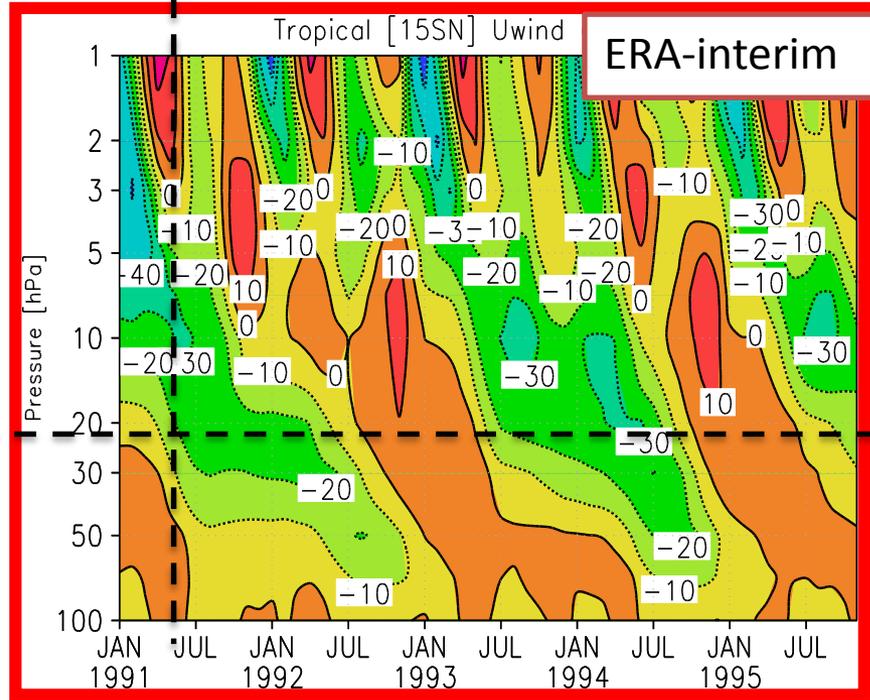
Ensemble-3

Tropical [15SN] Uwind (C)

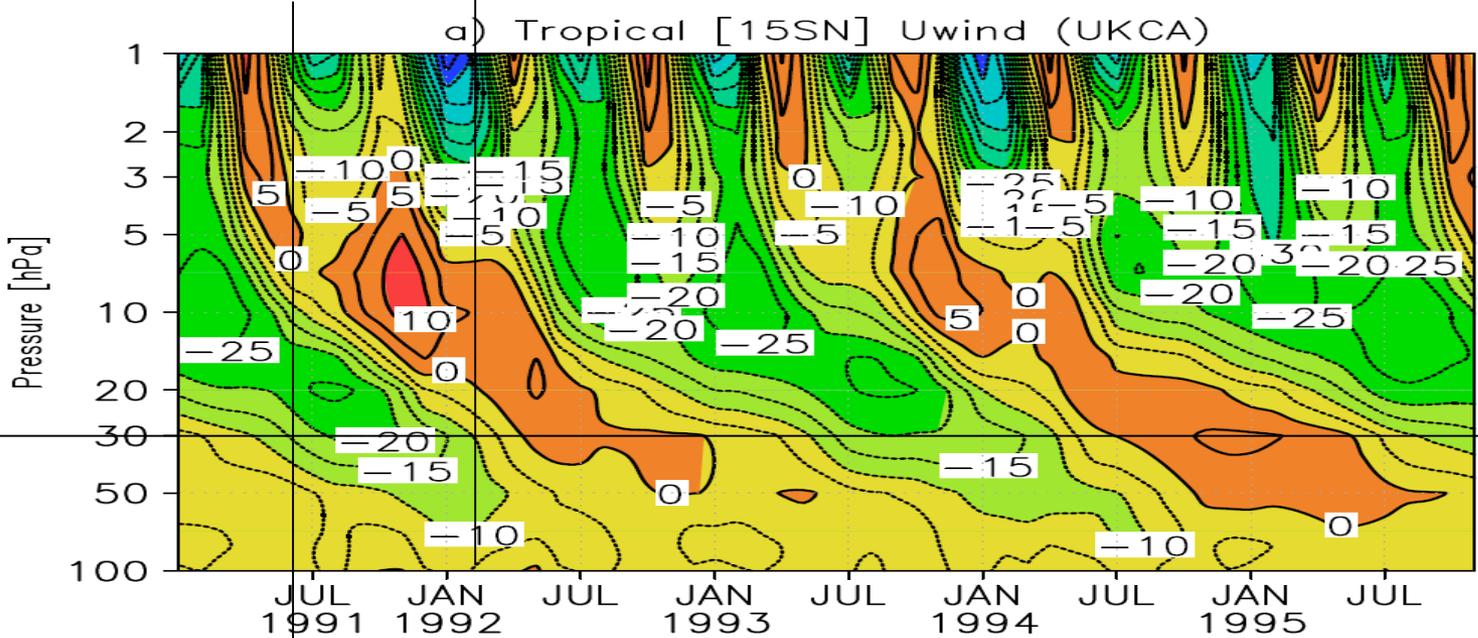


Tropical [15SN] Uwind

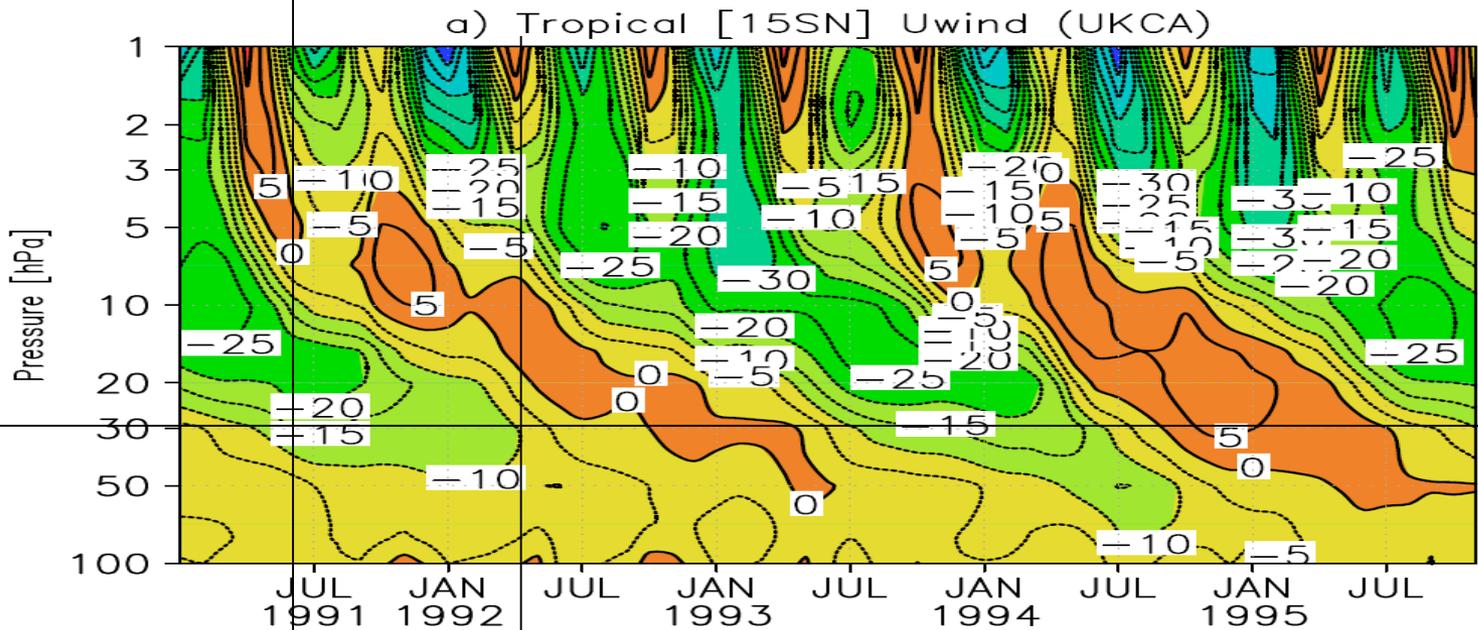
ERA-interim



Impact of radiative coupling: strat. dynamics



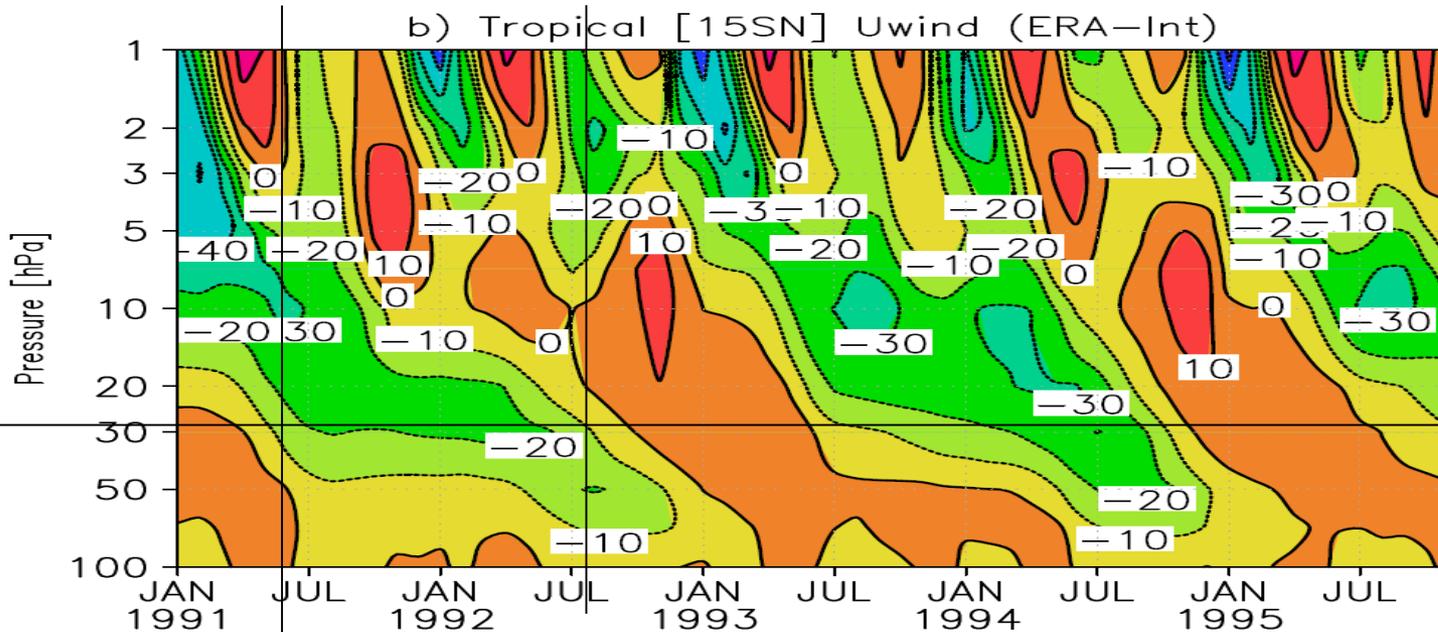
**Tropical
zonal mean
zonal wind for
“no Pinatubo”
run**



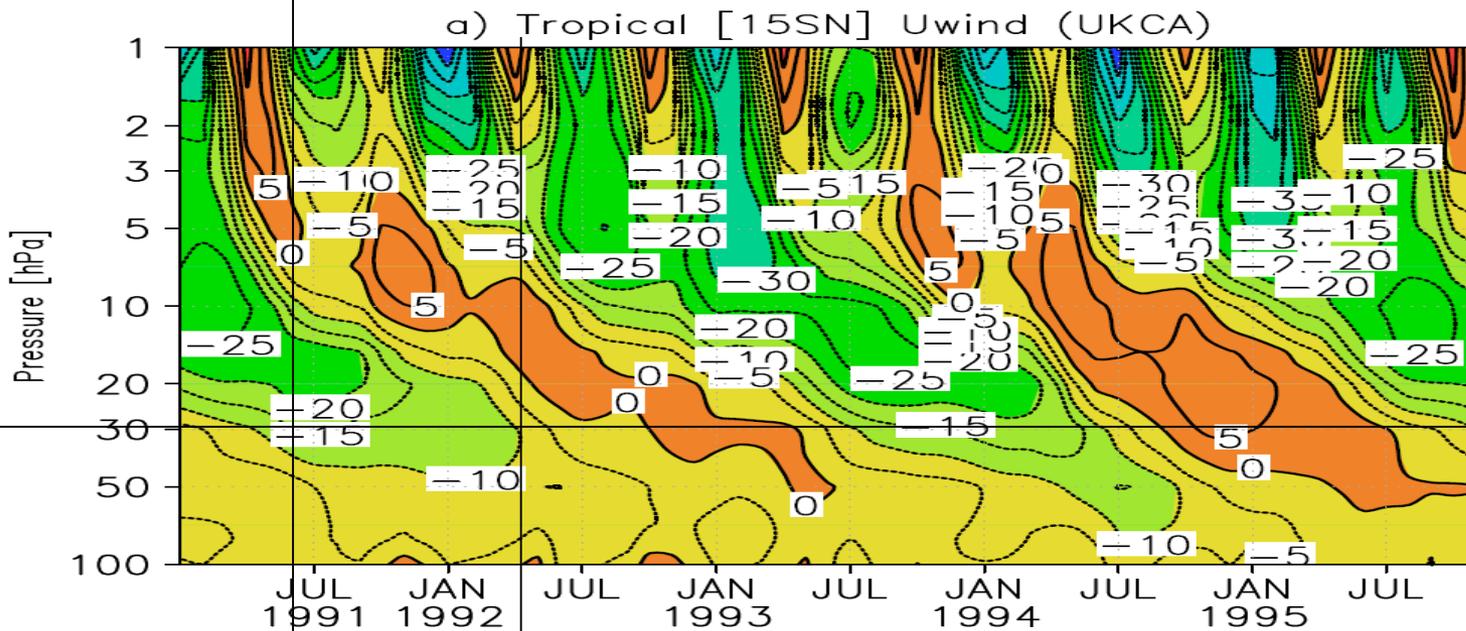
**Tropical
zonal mean
zonal wind for
“Pinatubo20”
run**

Mann et al
(in prep.)

Impact of radiative coupling: strat. dynamics



Tropical zonal mean zonal wind for ERA-interim re-analysis

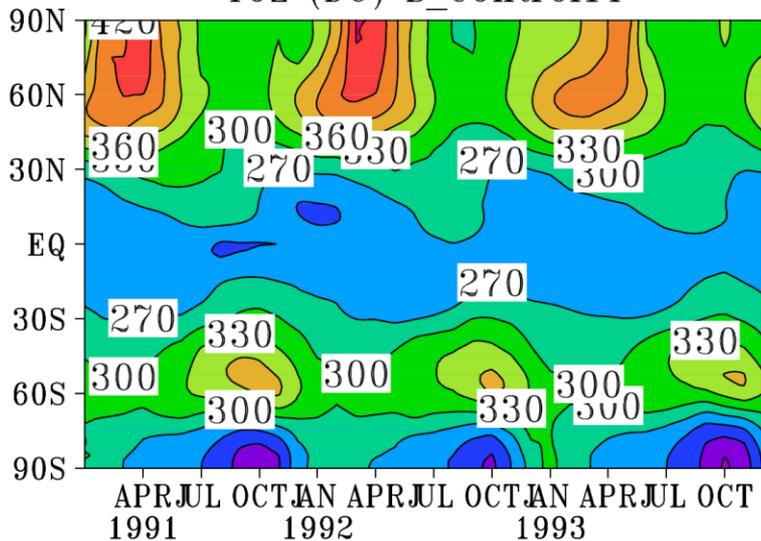


Tropical zonal mean zonal wind for "Pinatubo20" run

Mann et al
(in prep.)

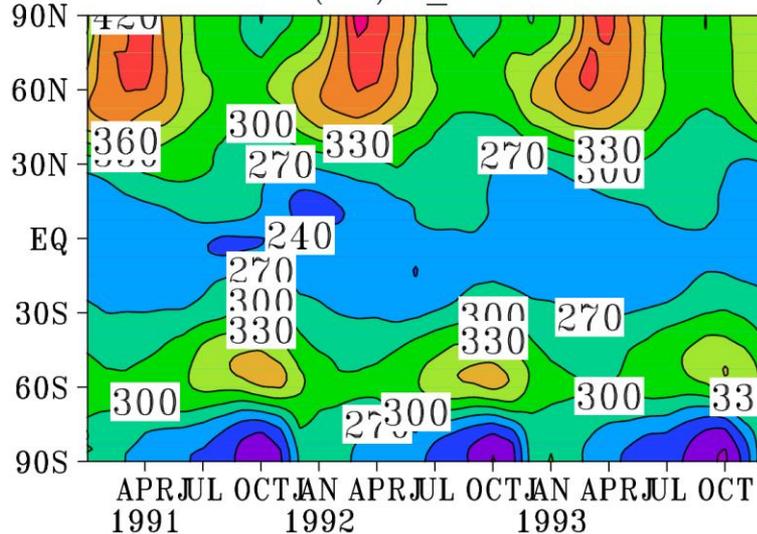
14 Tg SO₂ injection

TOZ (DU) B_Control14

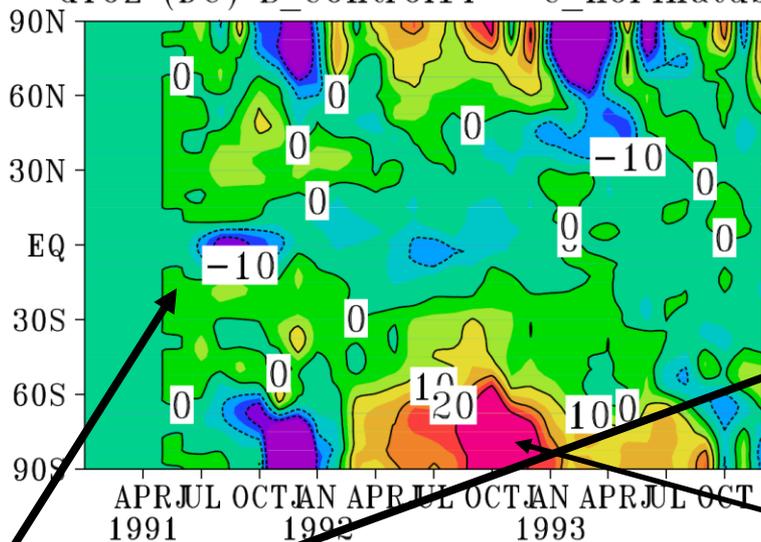


20 Tg SO₂ injection

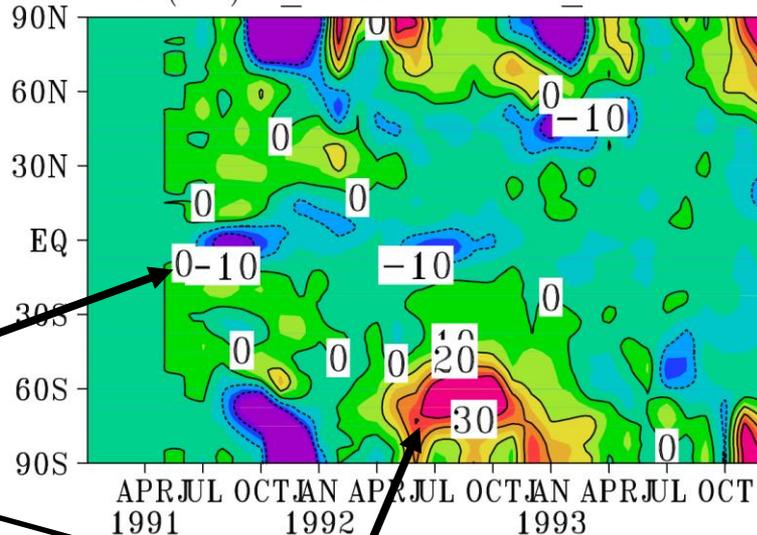
TOZ (DU) A_Control20



dTOZ (DU) B_Control14 - C_noPinatubo



dTOZ (DU) A_Control20 - C_noPinatubo



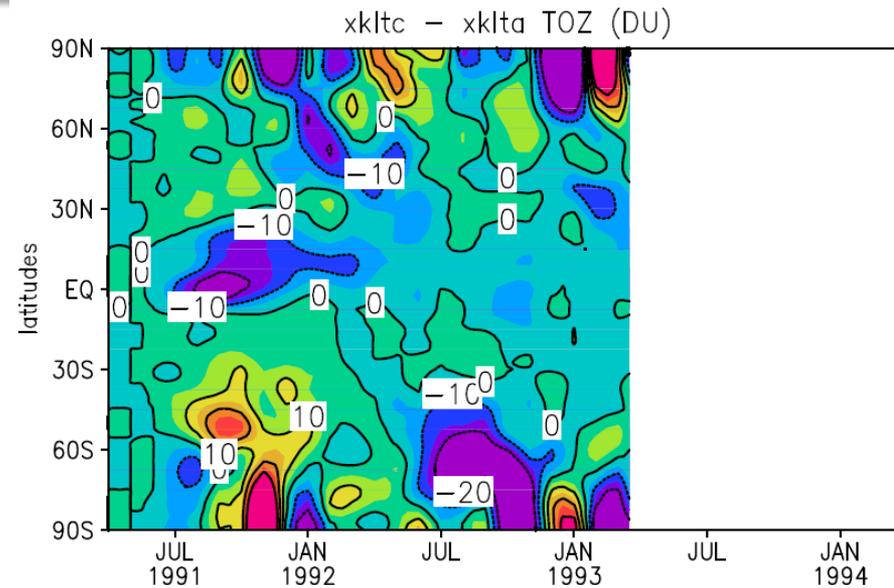
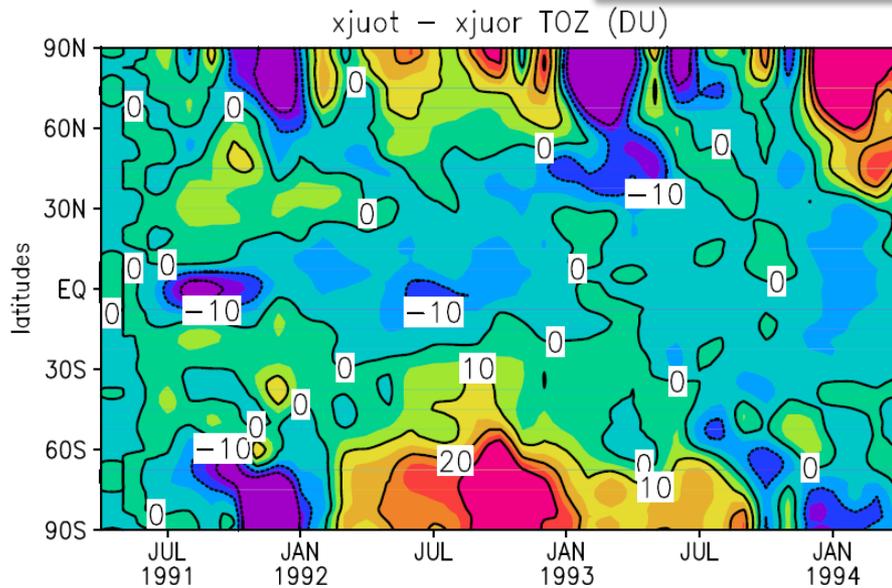
Dynamical O₃ loss in tropics

Dynamical response increases O₃ in SH mid-latitude winter and SH high latitude '92 winter & spring

Ensemble-1

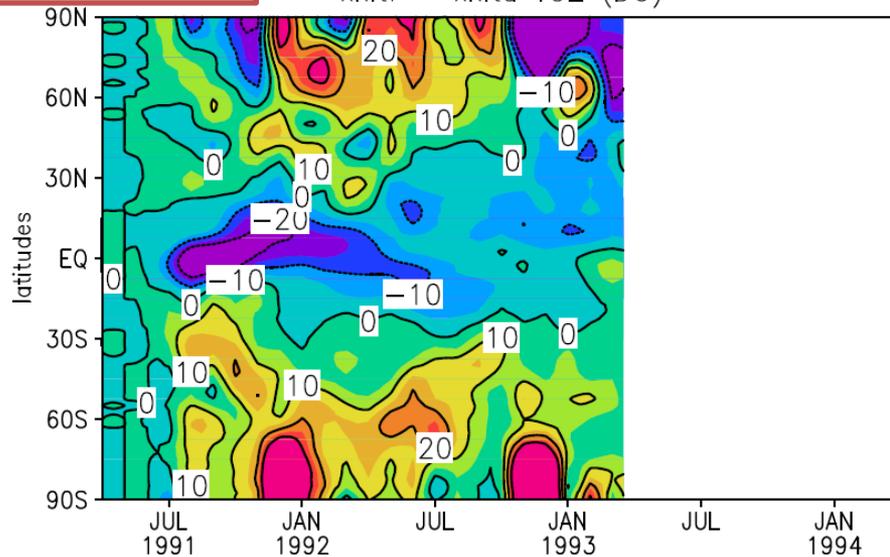
3 14Tg ensemble members with easterly QBO phase

Ensemble-2



Ensemble-3

xkltf - xkltd TOZ (DU)

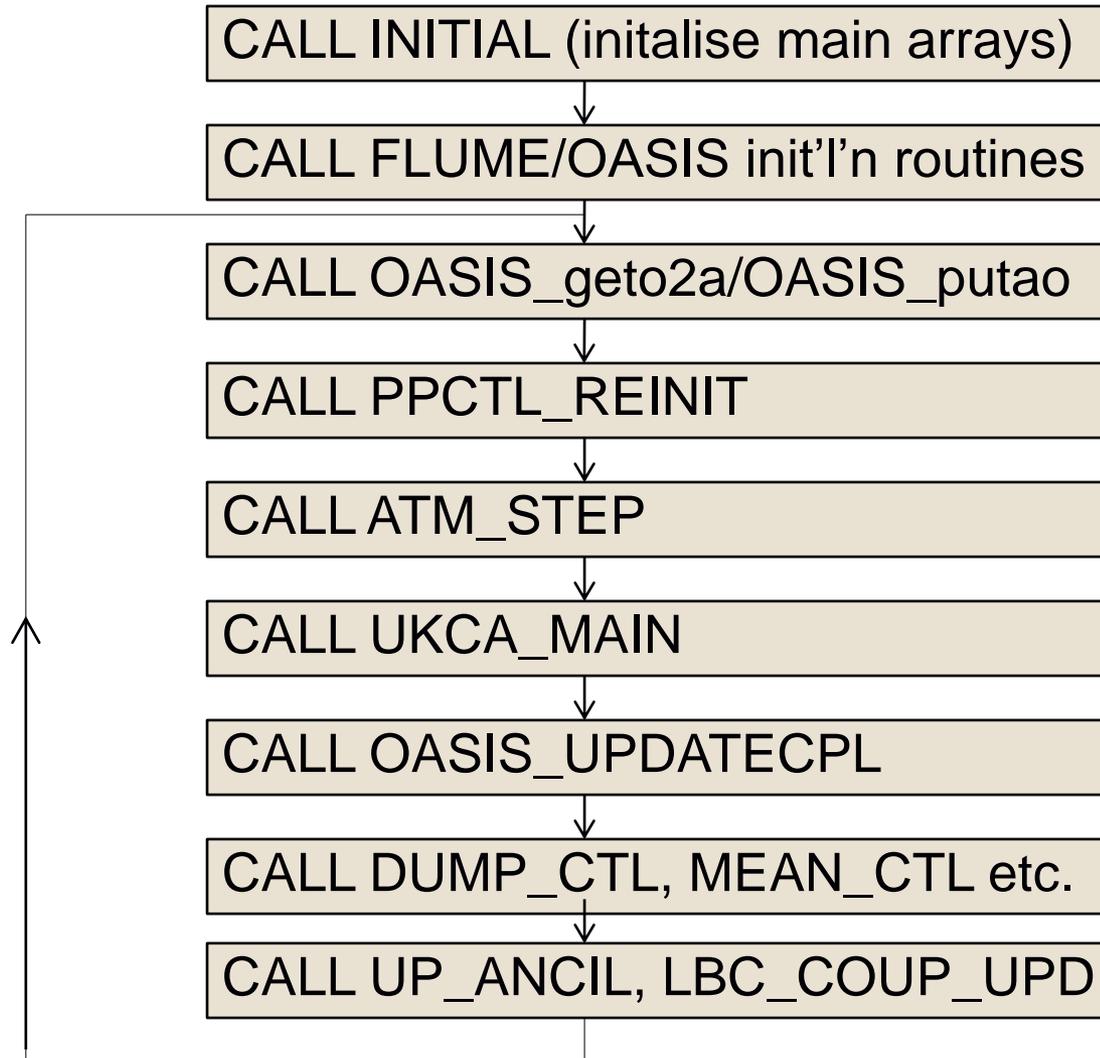


Pinatubo-on and Pinatubo-off runs both use time-varying SAD with volcanic enhancement.

So delta-ozone shown here is that induced from the radiative heating of the stratospheric aerosol layer and the subsequent dynamical response.

UKCA-MODE code in UM (U_MODEL)

UKCA_MAIN is called from U_MODEL after the call to ATM_STEP



UKCA-MODE code in UM (UKCA_MAIN)

UKCA_AERO_CTL called from UKCA_MAIN after UKCA_CHEMISTRY_CTL

Initialise aerosol configuration via subroutines in UKCA_MODE_SETUP and UKCA_SETUP_INDICES)

Set MODE COMPONENT

CALL GETD1FLDS & initialise arrays

CALL UKCA_DUST_CTL & FASTJ

CALL UKCA_CALC_TROPOPAUSE

CALL UKCA_EMISSION_CTL

Update gases for emissions

CALL UKCA_CHEMISTRY_CTL

Update gases for chemistry, BL mix & dry/wet dep

CALL UKCA_AERO_CTL

Update aerosol for emissions, BL mixing, dry/wet dep & microphys.

CALL UKCA_ACTIVATE

CALL PUTD1FLDS



UKCA-MODE code in UM (UKCA_AERO_CTL)

UKCA_AERO_STEP called from UKCA_AERO_CTL

Set input parameters related to primary emissions, dry/wet removal, aqueous SO₄ prodn, cloud processing



Set input params for aero processes (nucleation, condensat'n, coagulat'n)



Copy 3D/2D input UM/UKCA arrays to required 1D arrays for MODE



CALL TR_MIX for aerosol tracers.



Copy emism fluxes, gaseous (S0) & aerosol tracers (ND,MD) → 1D arrays



CALL UKCA_AERO_STEP



Copy updated ND,MD,S0 → tracers

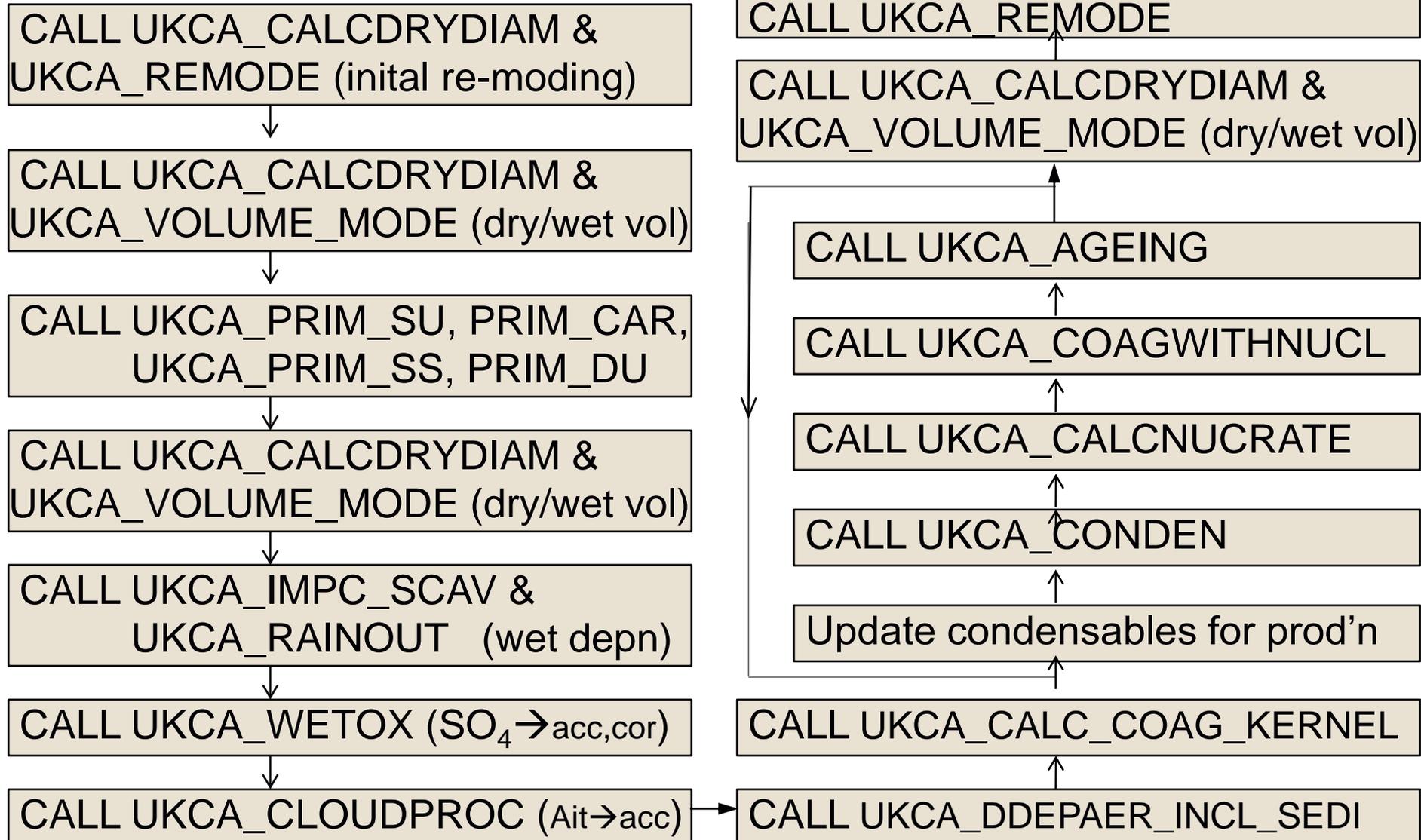
Updates tracers for BL mixing

Updates aerosol tracers for primary ems, dry/wet depn, nucleation, coagulation, condensation (H₂SO₄ & SECORG) and cloud proc.



UKCA-MODE code in UM (UKCA_AERO_STEP)

UKCA_AERO_STEP calls routines for each aerosol process





UKCA-MODE inputs (copied 3D→1D)

- temperature & pressure
(temp→T,pres→PMID,p_bdrs→PUPPER/PLOWER)
- precip. rates (crain/drain → CRAING/DRAING,
CRAING_UP/DRAING_UP)
- autoconversion rates (FCONV_CONV/FCONV_DYN)
and raining fractions (FBOX_CONV/FBOX_DYN)
currently set to constant values)
- moisture (rh3d→RH, q→S)
- cloud (cloud_liquid_wat→LWC,
cloud_frac→LOWCLOUD,[VFAC=1])
- surface layer variables (u_s→USTR,u_10m→US10M,
z0m→ZNOTG)
- sea-ice, land-frac
(sea_ice_frac→SEAICE,land_fraction→LAND_FRAC)
- mass of air and surface area (mass → SM, area → SURF)
- aqueous SO₄ production tendencies (delso2_wet_h2o2→DELSO2
delso2_wet_o3 → DELSO2_2)
- H₂SO₄ vapour prod'n tendency
(delso2_dry_oh→ S0G_DOT_CONDENSABLE)



UKCA-MODE outputs (copied 1D→3D)

- updated number concentration in each mode
(ND → mode_tracers, $ND = tr_rs * AIRD$)
- updated molecules-per-particle for SO₄, BC, POM, NaCl, DU
(MD → mode_tracers, $MD = tr_rs * (MM_DA / MM_AER) * AIRD / ND$)
- diagnosed CN, CCN, CDN concentrations → STASH sec38, 437-441
- diagnosed geometric mean dry/wet diameter
(DRYDP, WETDP → STASH section 38, items 401-411)
- diagnosed aerosol water content (molecular
(MDWAT → STASH section 38, items 412-415)
- diagnosed mode densities
(RHOSOL → STASH section 38, items 430-436)
- diagnosed surface-area concentrations
(SAREA → STASH section 38, items 416-422)
- diagnosed volume concentrations
(VCONC → STASH section 38, items 423-429)
- diagnosed partial volumes for each component and water
(PVOL, PVOL_WAT → STASH section 38, items 442-468)
- diagnosed molecular-fluxes through each process
(BUD_AER_MAS → STASH section 38, items 201-387)