

# The influence of meteoric smoke particles (MSPs) on stratospheric aerosol properties

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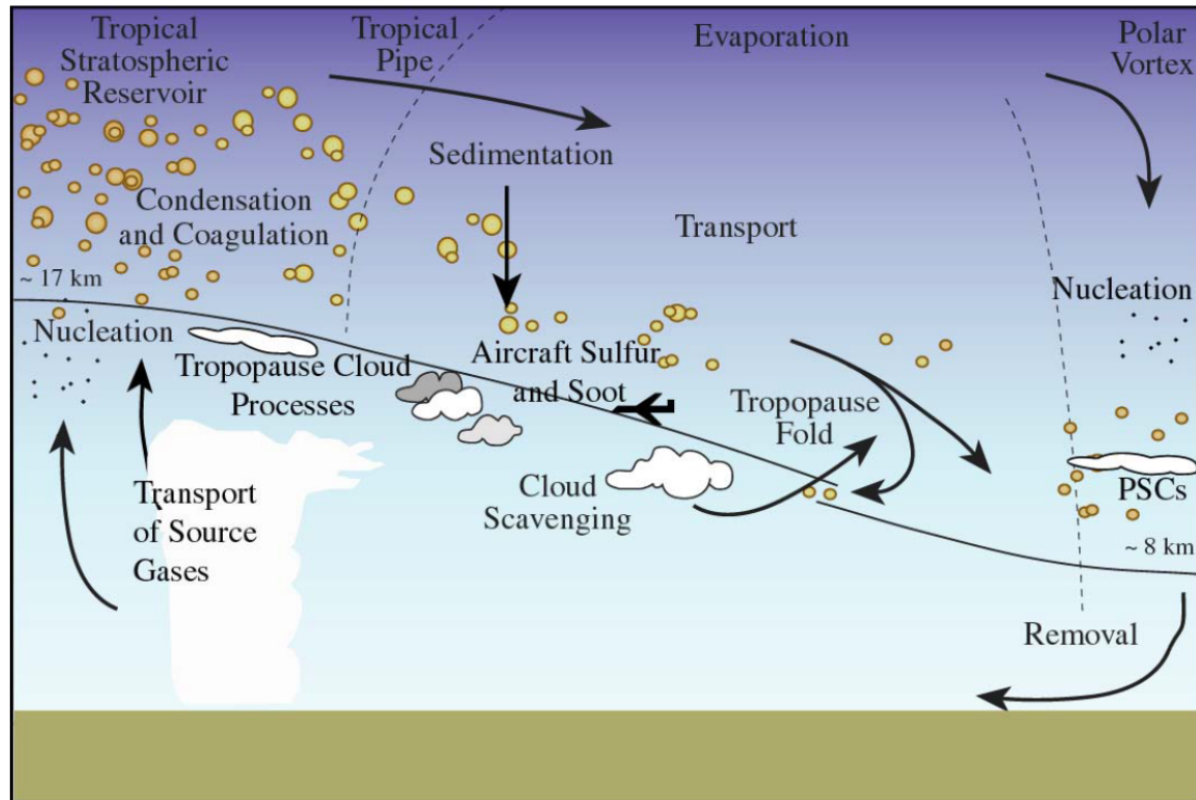
# Meteoric Smoke Particles (MSP)

- Ablation of micrometeorites in the mesosphere results in the formation of MSPs -- initially at nm sizes (Hunten et al., 1980) but at number conc'ns  $\sim 1000 \text{ cm}^{-3}$  in the upper stratosphere,
- Particle concentrations decrease and particle sizes increase during descent to the lower stratosphere and MSPs have the potential to strongly influence the Junge layer (Turco et al. 1982)
- The main way the MSPs influence the Junge layer is to provide additional condensation nuclei which compete with homogeneously-nucleated particles for available  $\text{H}_2\text{SO}_4$  vapour
- Reactive uptake of gas phase sulphuric acid to MSP surfaces also occurs above 35km where  $\text{H}_2\text{SO}_4$ - $\text{H}_2\text{O}$  particles evaporate.
- MSPs are most prevalent in the wintertime polar vortex  
COPAS measurements (Curtius et al. 2005) show  $\sim 12$ - $45\%$  of particles contain refractory cores outside of polar vortex
- Aircraft measurements (Murphy et al., 1998; Cziczo et al., 2001)  
<sup>2</sup>  $\sim 50\%$  strat aerosol particles contain  $\sim 1$ - $2 \text{ wt}\%$  meteoric material.

# MSPs & stratospheric aerosol

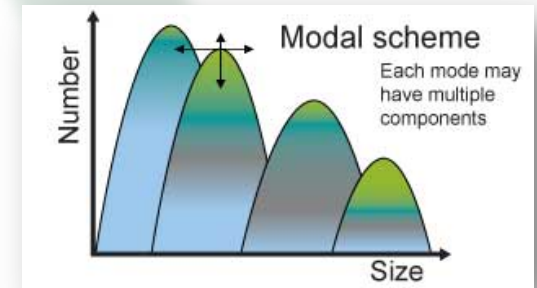
- Meteoric smoke particles form in mesosphere & transported into the polar stratosphere but interactive stratospheric aerosol models include only limited representation of their effects (if any)

***Q: How does the presence of the MSPs alongside the homogeneously nucleated particles affect the Junge layer?***



# UK Chemistry and Aerosol project

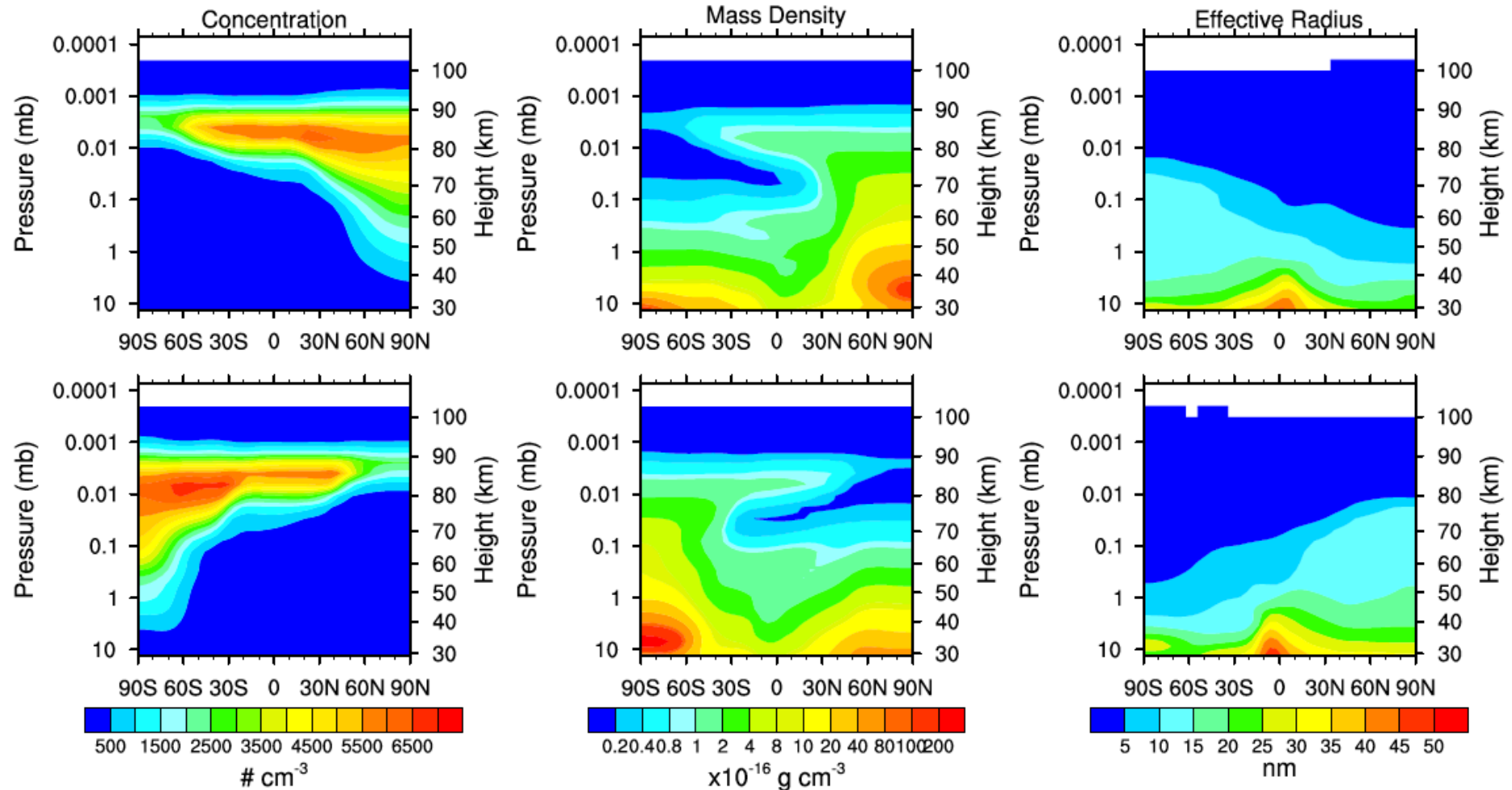
- Collaboration between UK National Centre for Atmospheric Science (Leeds, Cambridge, Oxford) & UK Met Office since 2005
- Has built aerosol-chemistry sub-model in the UK Met Office Unified Model, being applied for a range of applications (climate, air quality, Earth system science, weather)
- Chemistry schemes & aerosol configurations including for stratosphere-troposphere
- Multi-component aerosol microphysics scheme (GLOMAP)
- Global variations in particle size distribution → sedimentation and SW & LW radiative effects
- UK Earth System model in development (UKESM) will simulate strat-trop ozone and strat-trop aerosol interactively and radiatively coupled with composition-dynamics interactions

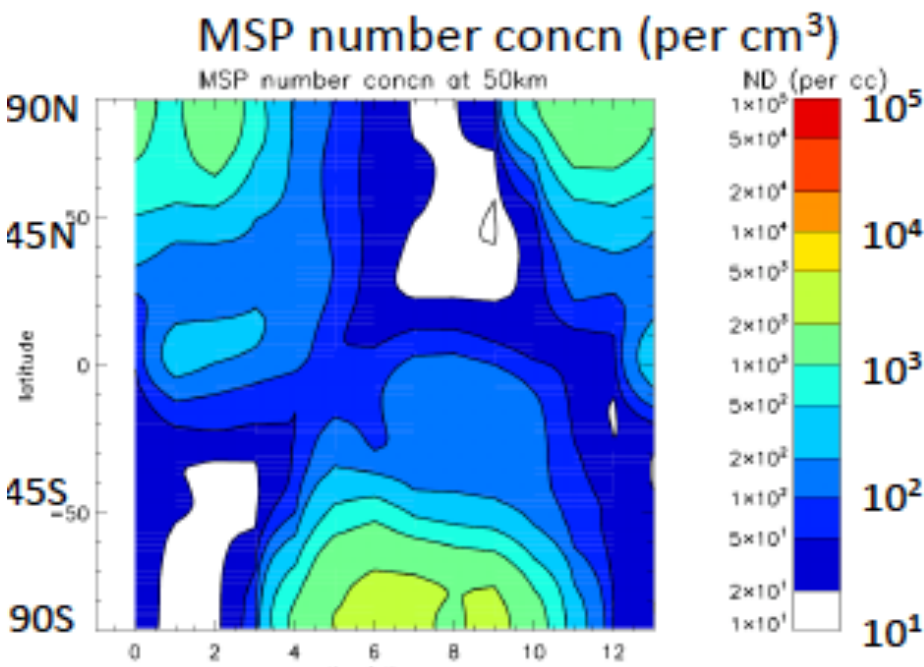


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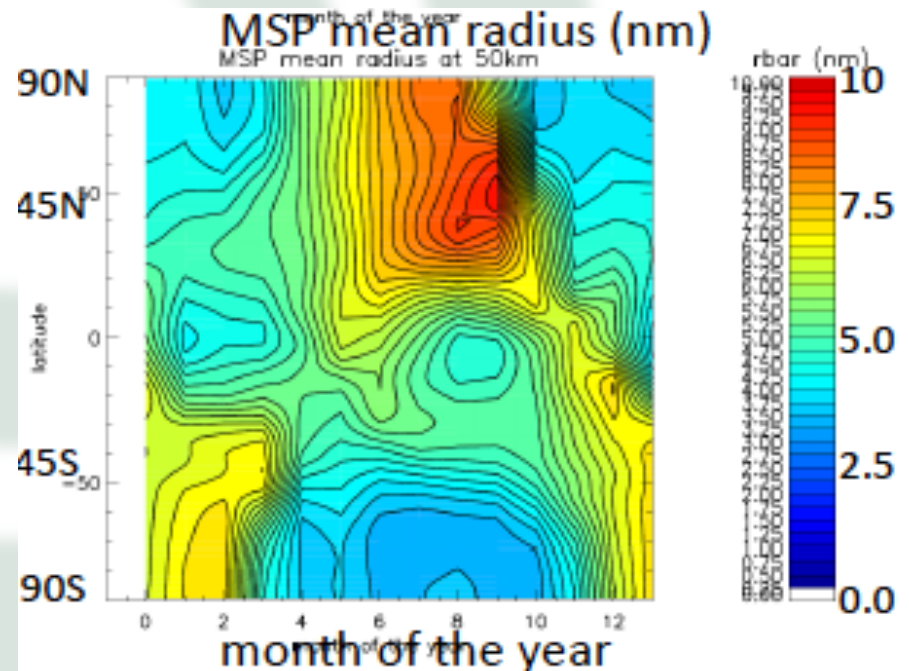
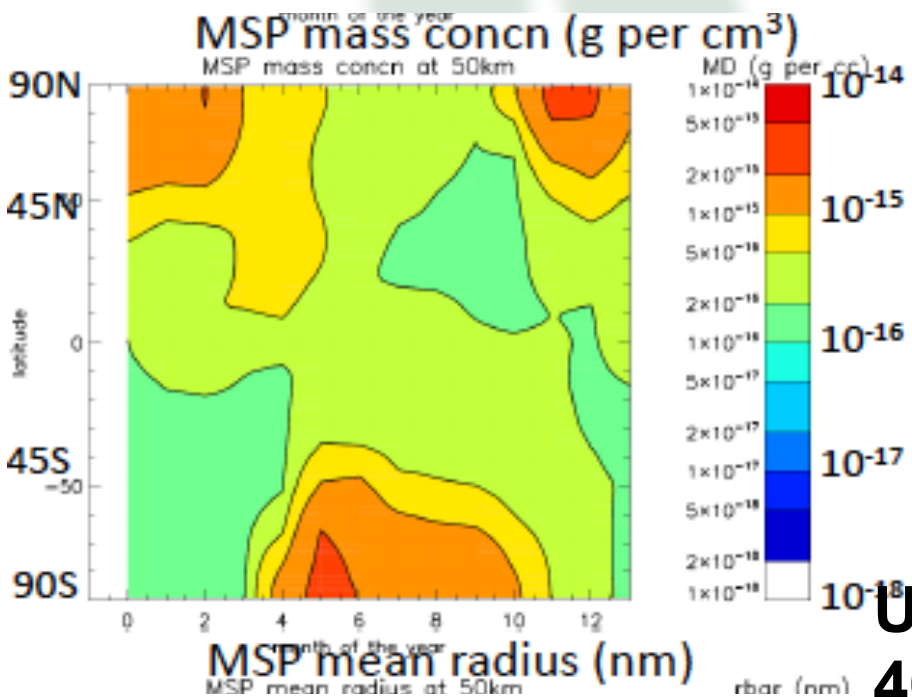
# Distribution of MSP simulated by WACCM-CARMA (120km top, bin-resolved aerosol scheme)





**Introduce zonal-mean MSP number & mass mixing ratio from WACCM-CARMA into UM-UKCA**

**Over-write UM-UKCA tracer values for  $p < 0.2$ hPa**

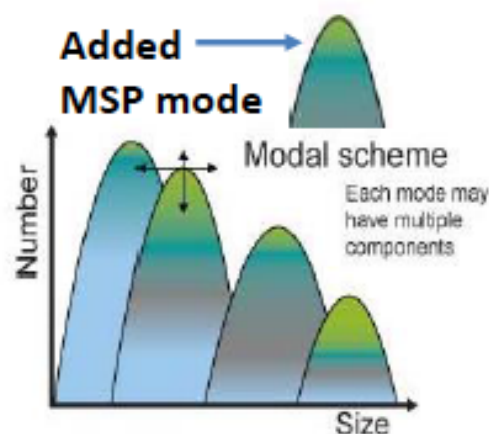


**Upper limit MSP input flux of 40 kt/yr (Love & Brownlee, 1993)**

### 3. MSP added to UM-UKCA within GLOMAP-mode

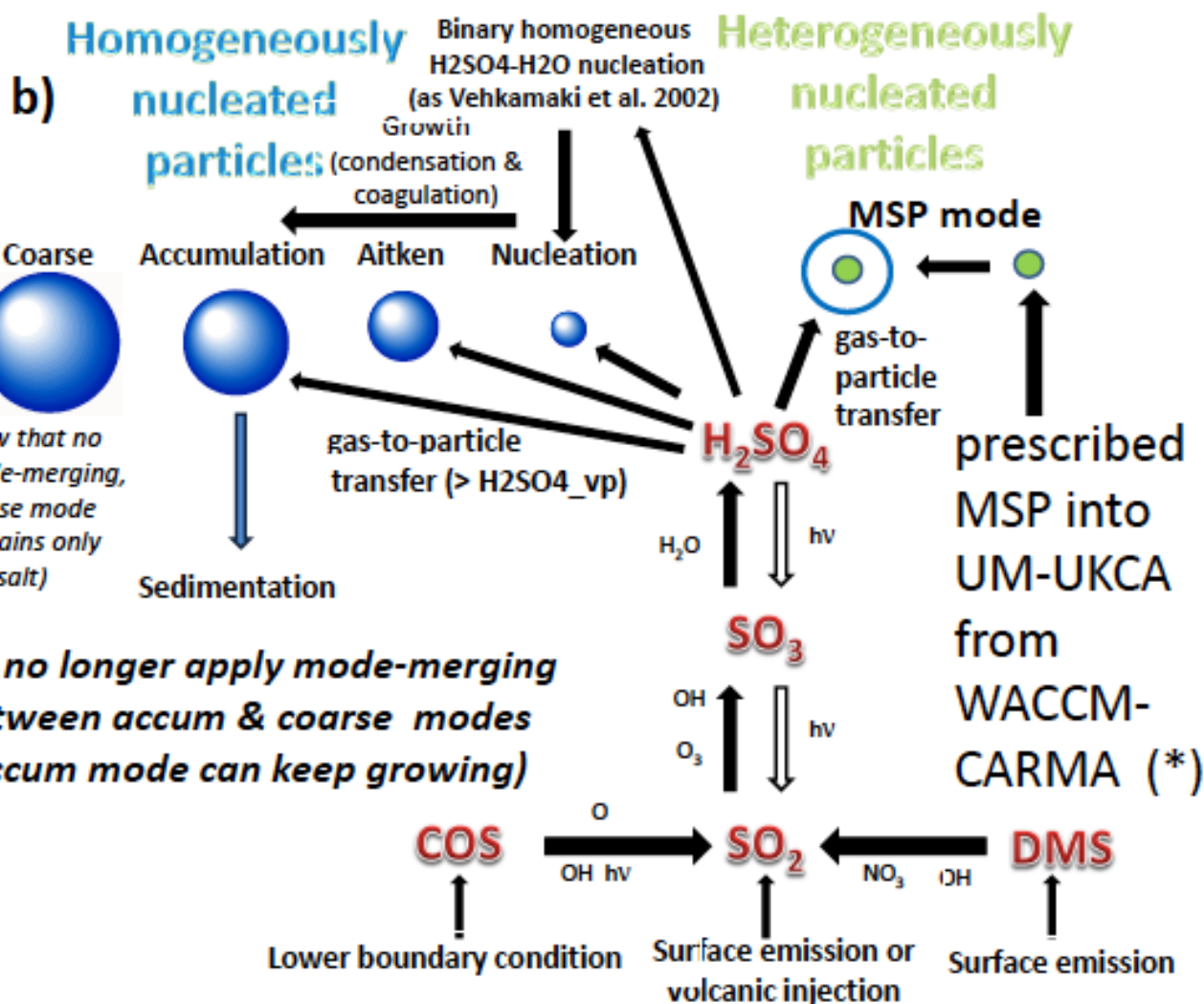
Include MSPs within existing accumulation insoluble mode (but size restrictions removed to allow to cover terrestrial and extra-terrestrial dust.

a)



**Fig 2a. Schematic of modal representation GLOMAP-mode aerosol microphysics scheme (Mann et al., 2010)**

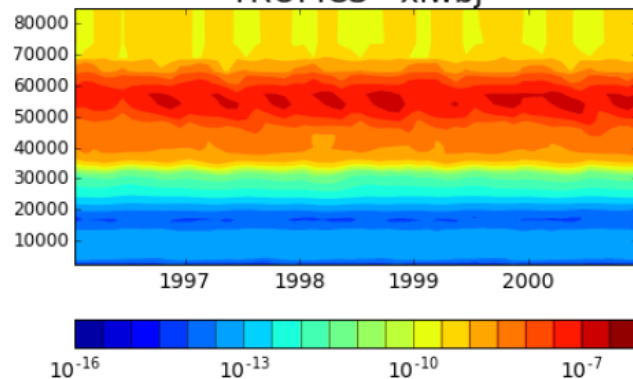
(\*) We prescribe number- and mass- mixing ratios of MSP  $p < 0.2$  hPa based on WACCM-CARMA sectional runs (Bardeen et al., 2008).



**Fig 2b. Schematic of UM-UKCA stratospheric sulphur chemistry and coupling to GLOMAP-mode aerosol microphysics (see Dhomse et al., 2014)**

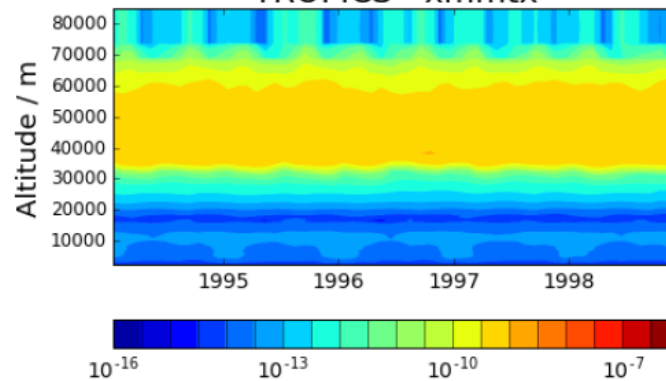
# **a** $\text{H}_2\text{SO}_4(\text{g})$ : no MSP

TROPICS - xlwbj

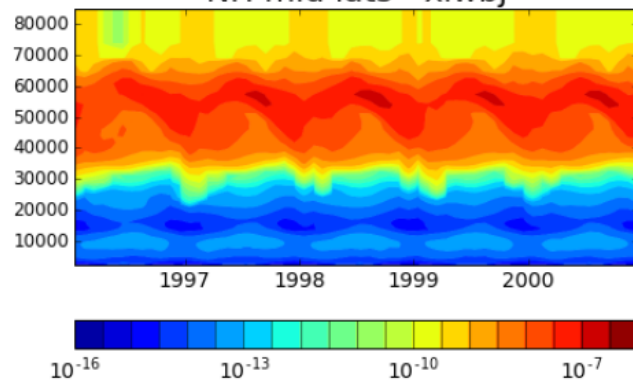


# $\text{H}_2\text{SO}_4(\text{g})$ : withMSP

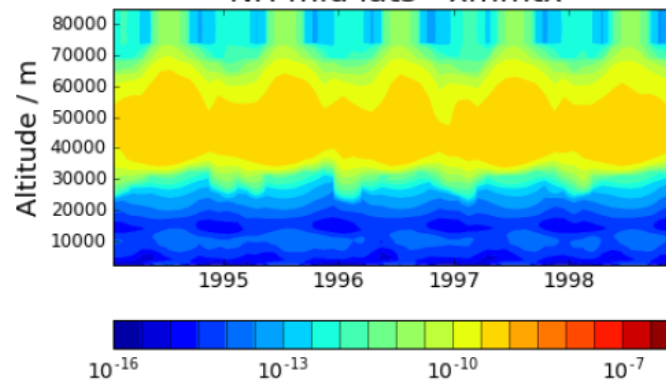
TROPICS - xmmtx



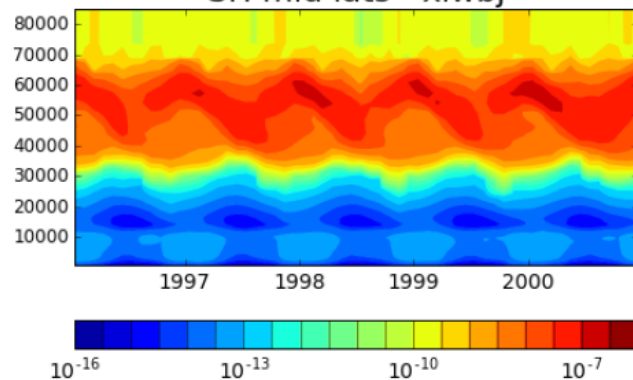
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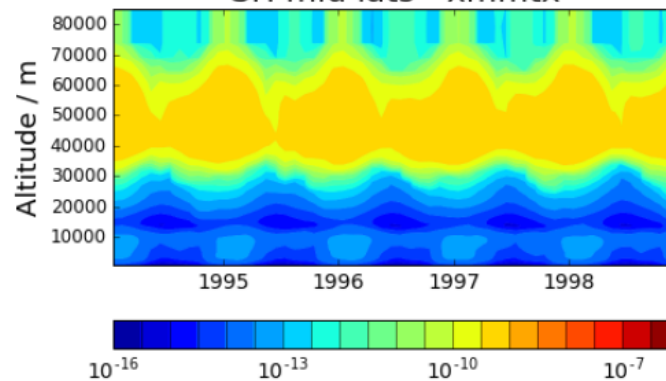
NH mid lats - xmmtx



SH mid lats - xlwbj

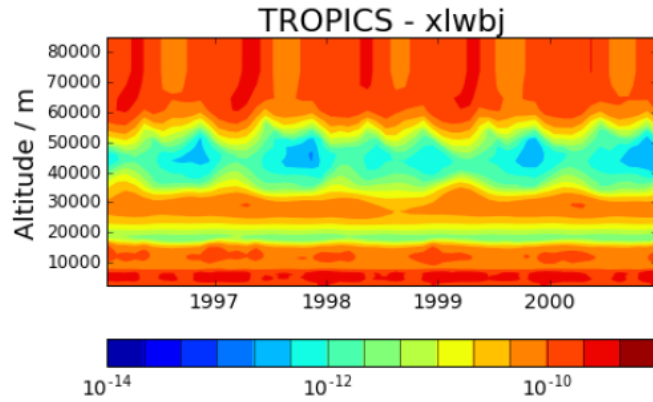


SH mid lats - xmmtx

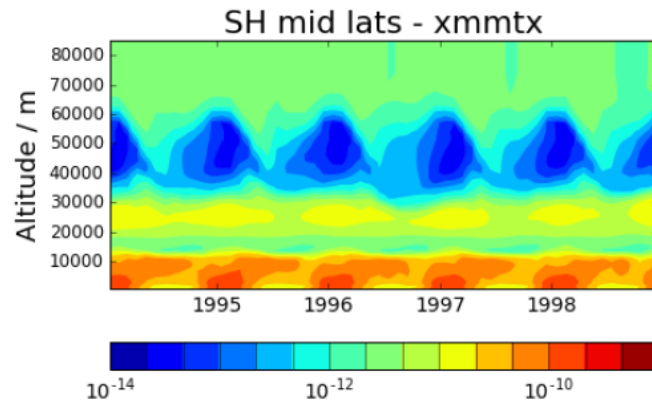
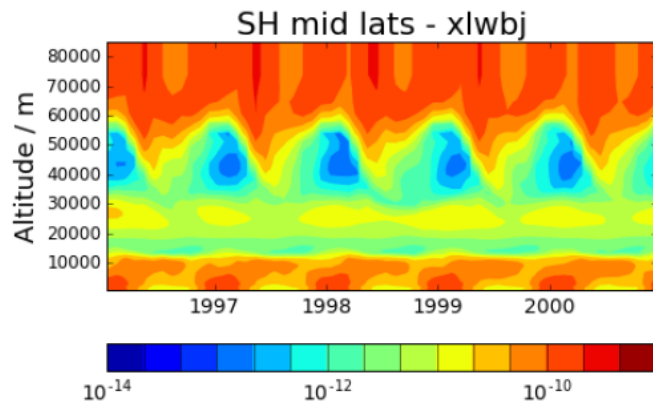
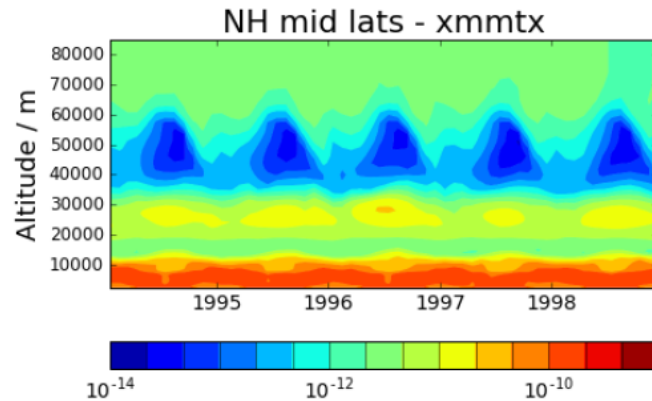
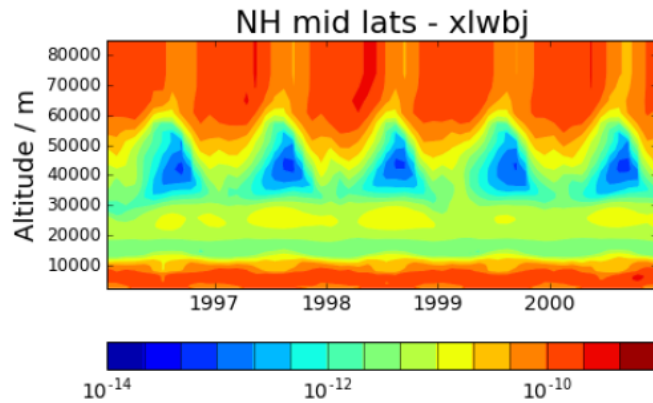
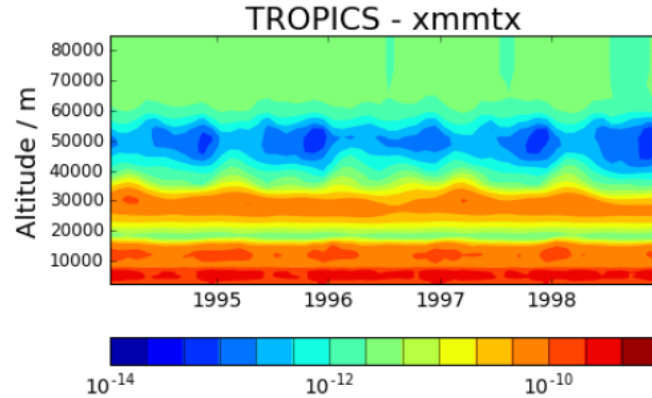




## SO<sub>2</sub>(g) : no MSP

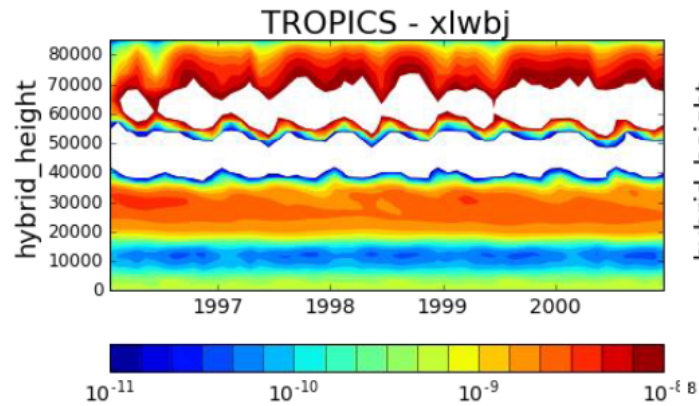


## SO<sub>2</sub> (g) : withMSP

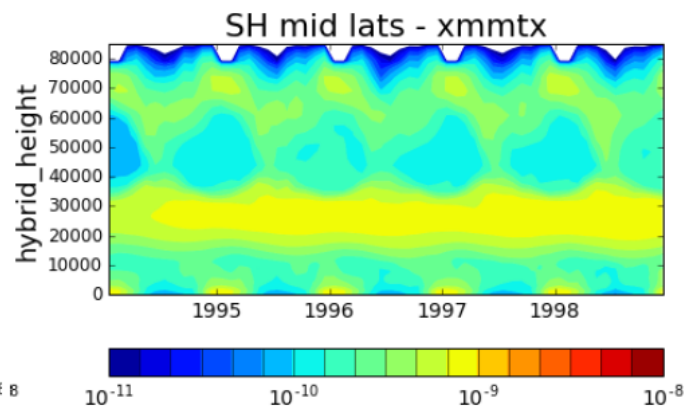
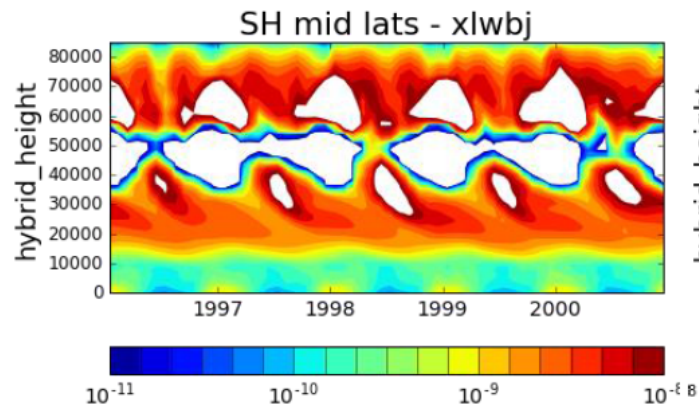
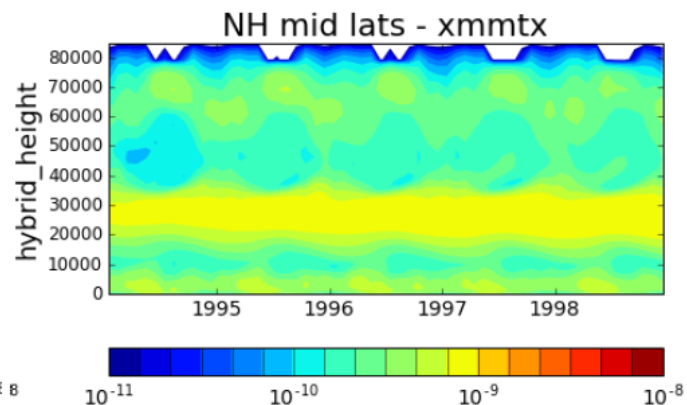
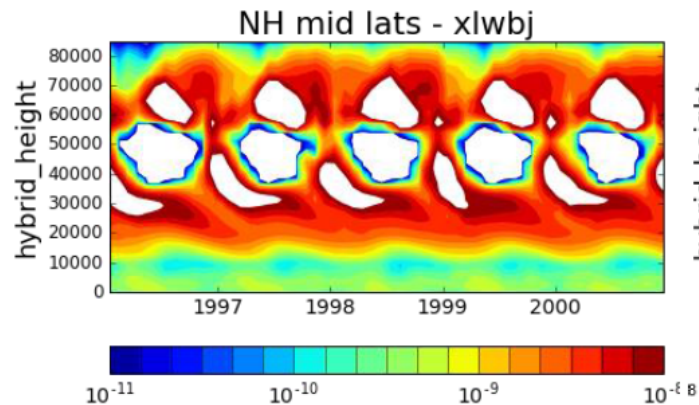
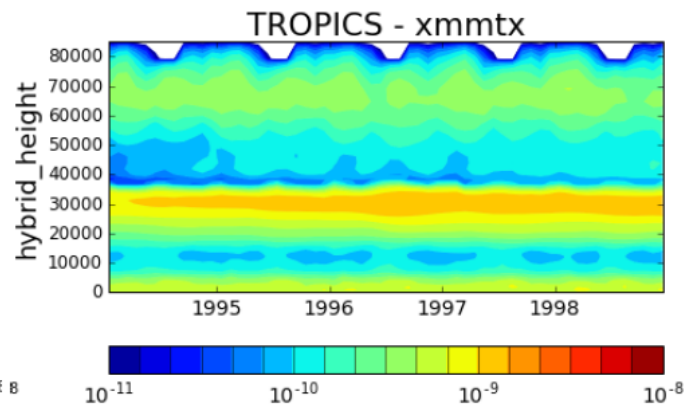




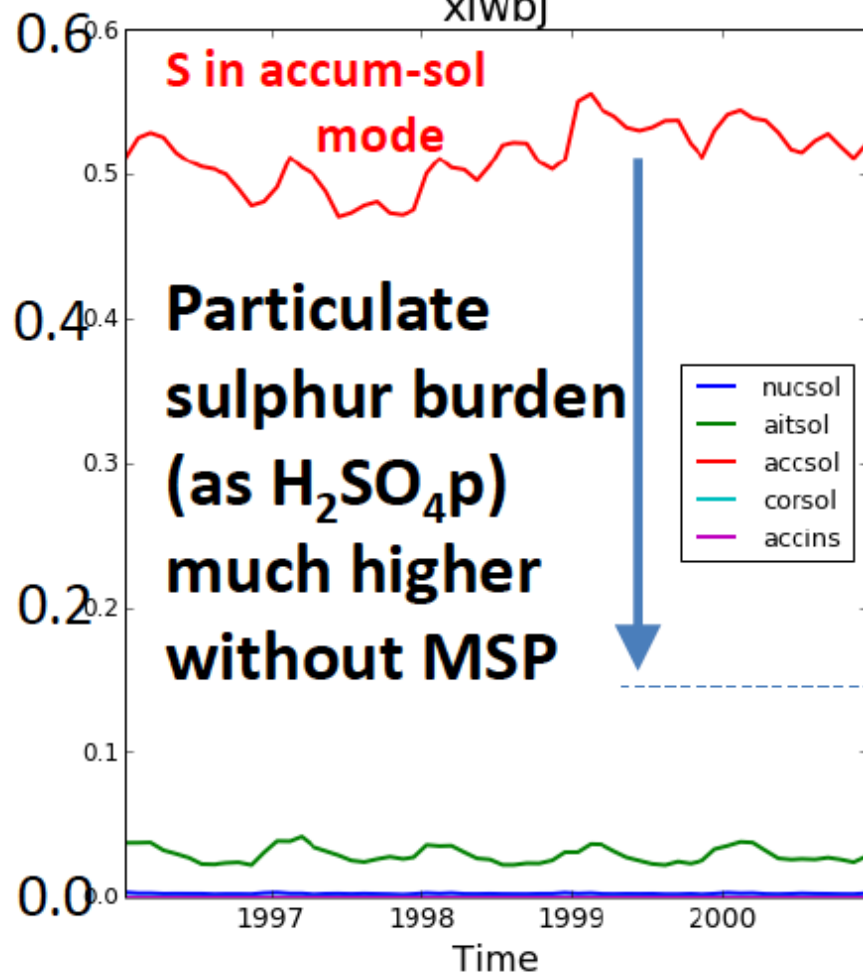
## $\text{H}_2\text{SO}_4(\text{p})$ : no MSP



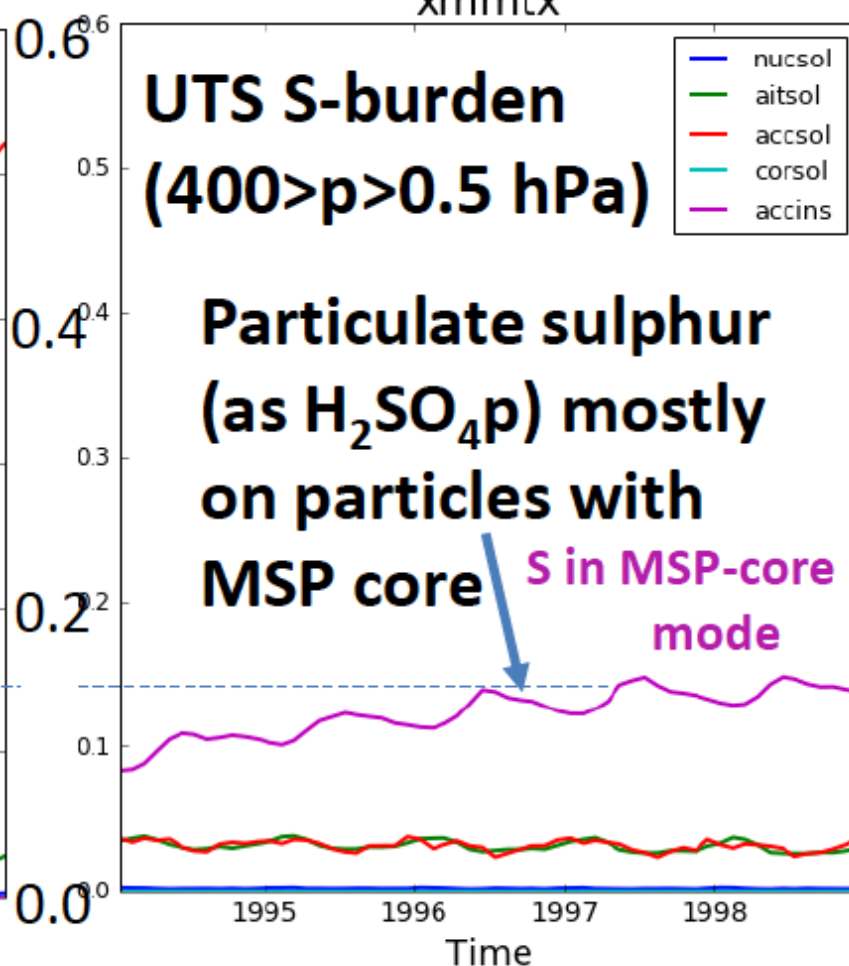
## $\text{H}_2\text{SO}_4(\text{p})$ : withMSP



## Without MSP

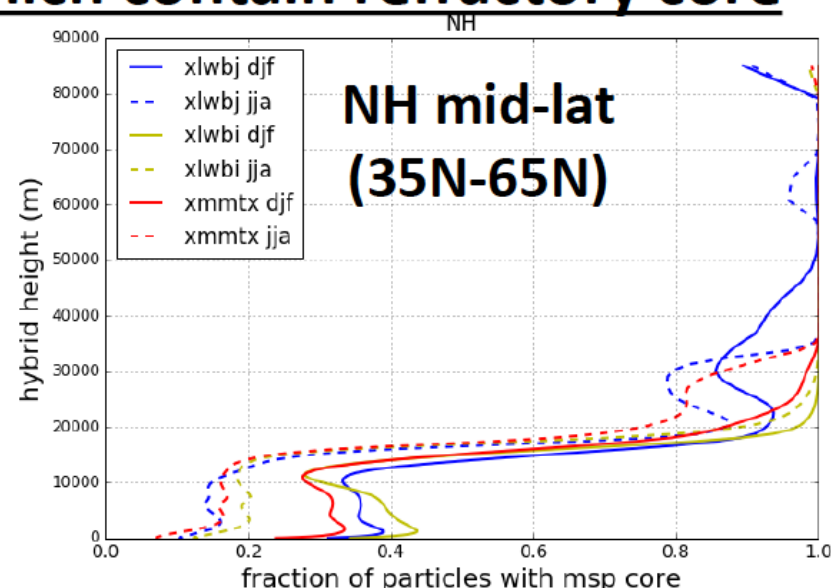
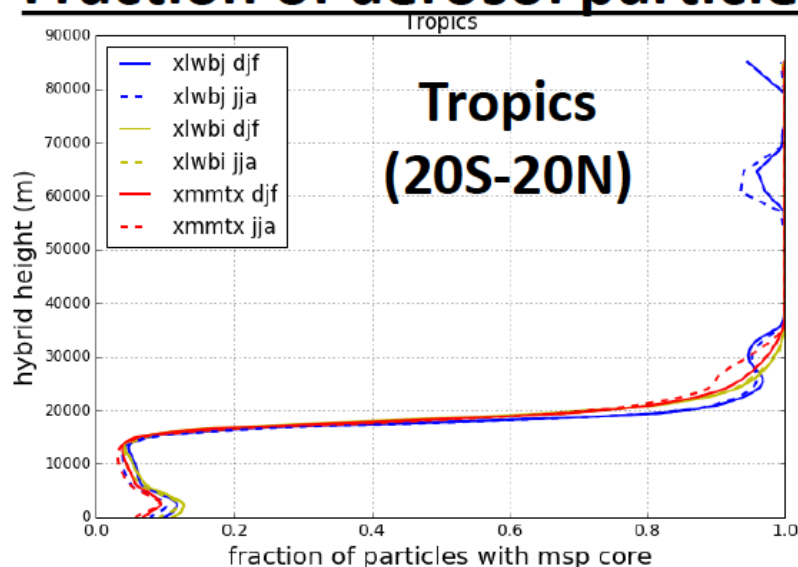


## With MSP interaction



UTS particle sulphur burden decreases as vertical  
 $\text{SO}_4$  distribution shifts towards tropopause

## Fraction of aerosol particles which contain refractory core

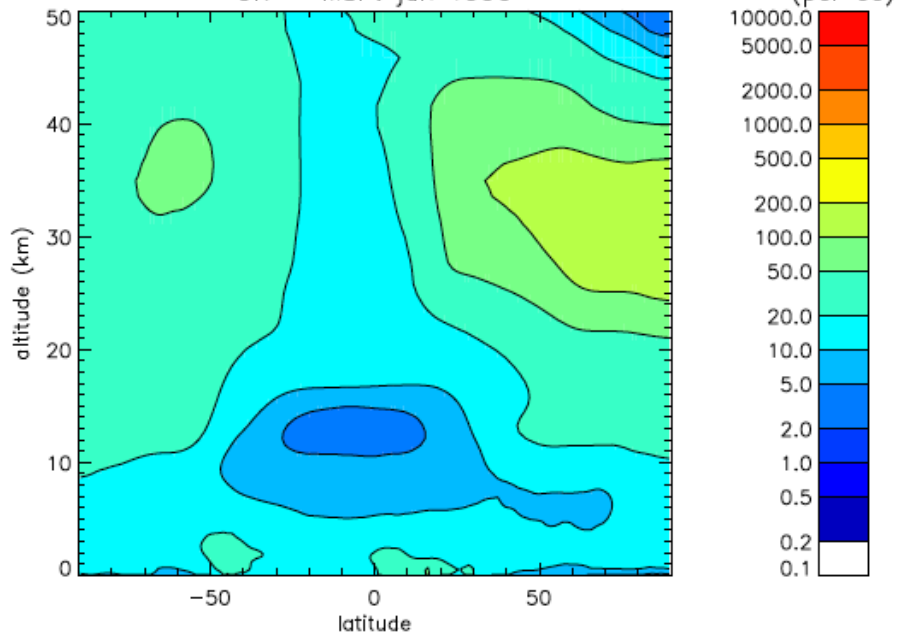


We find that, at least for this upper-limit MSP input flux, the stratospheric aerosol is dominated by MSP-core particles in all but the lowermost stratosphere. *(Note that terrestrial mineral dust is also tracked in this mode. In the troposphere the refractory –core particles will be mostly coated dust.)*

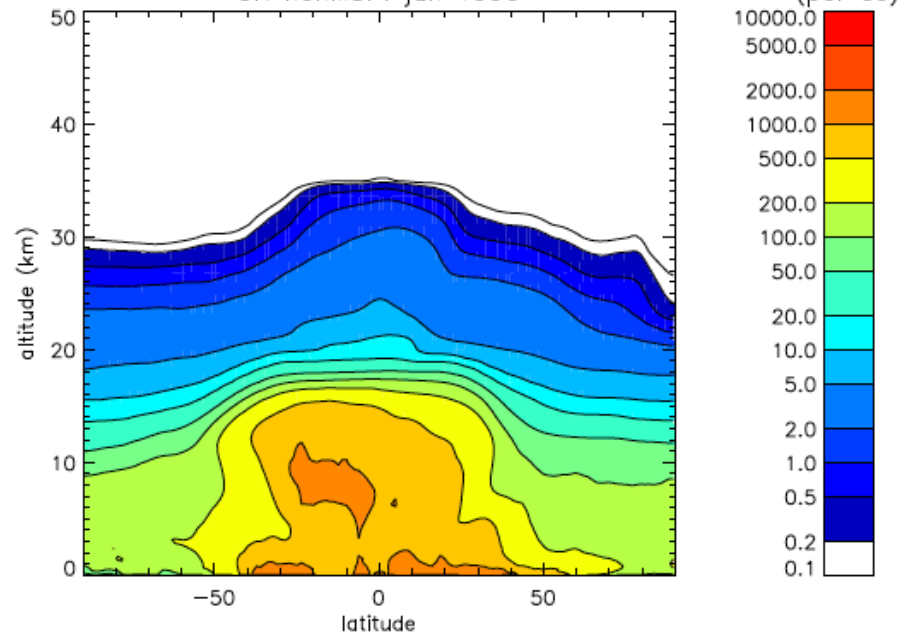
# Summary

- Composition-climate model experiments with interactive tropospheric & stratospheric chemistry and aerosol to assess the influence of MSPs on stratospheric aerosol properties
- MSPs remove  $\text{H}_2\text{SO}_4$  vapour from the gas phase even above the Junge layer since their surfaces are chemically reactive
- MSP reduces gas phase  $\text{H}_2\text{SO}_4$  &  $\text{SO}_2$  concentrations  $> 35\text{km}$
- Sulphur burden reduced due to change in vertical distribution
- Results suggest MSP-core (heterogeneously nucleated) particles dominate aerosol particle population throughout the Junge layer
- Have used “upper-limit-estimate” of MSP input flux (40 kt/yr)
- Simulations using a new “best-estimate” input flux of 5 kt/yr being spun-up → initial results suggest a factor  $\sim 20$  lower MSP number concentrations in lower stratosphere with reduced particle size.

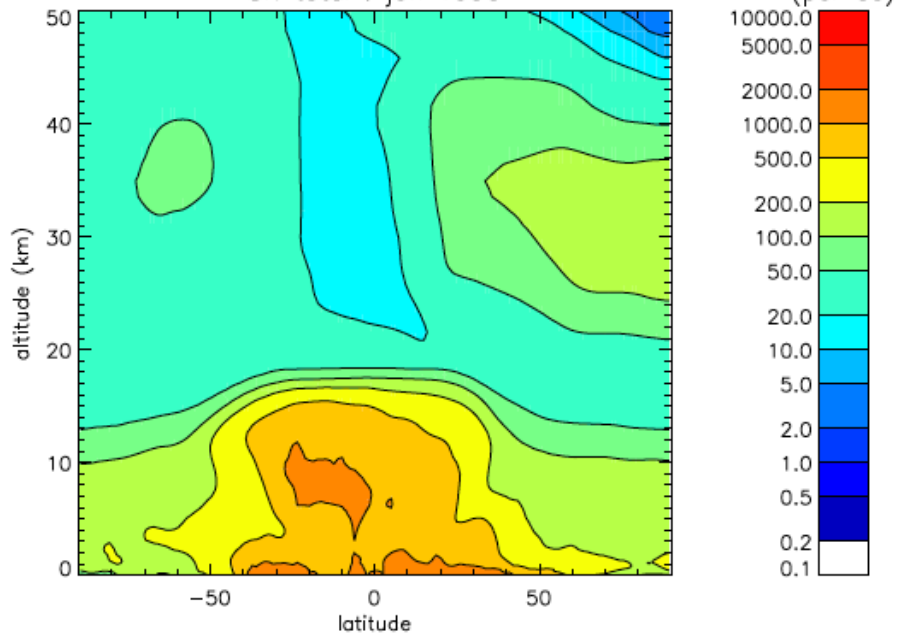
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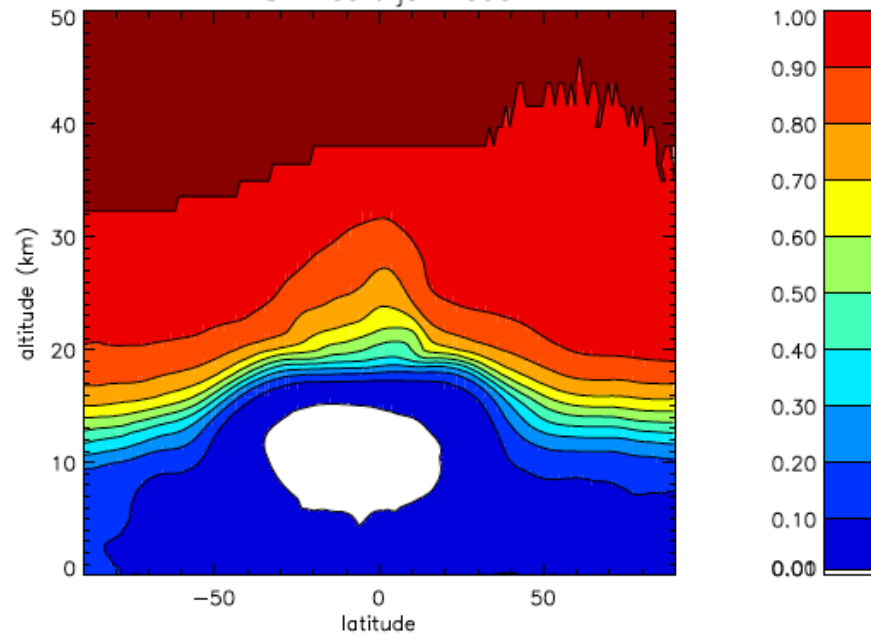
CN nonMSP: jan 1990



CN total : jan 1990

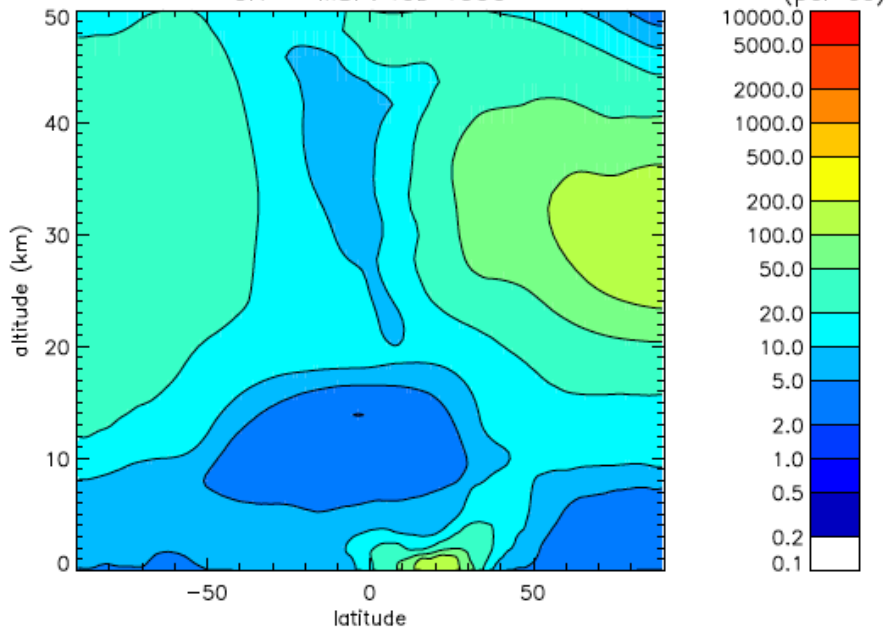


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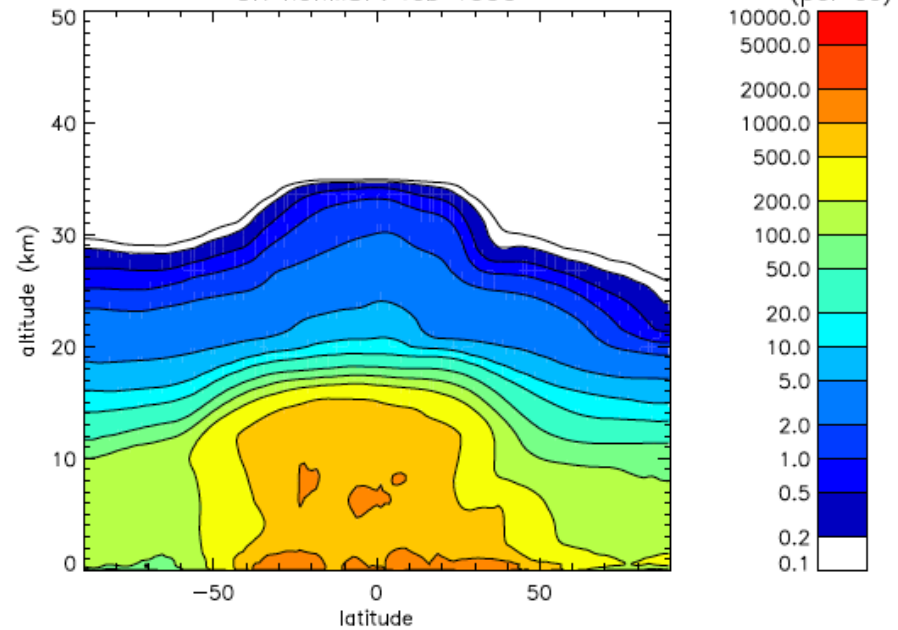




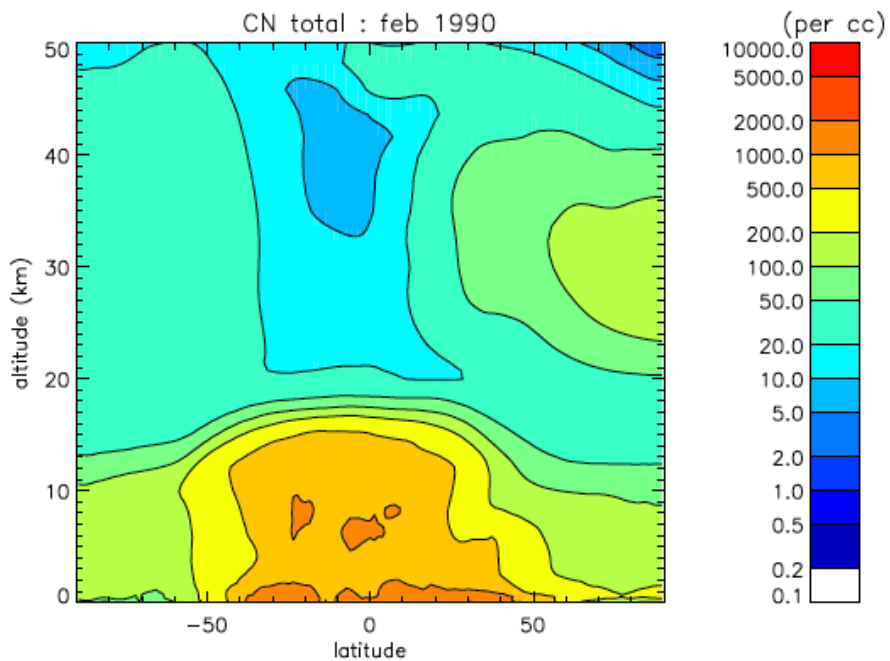
CN MSP: feb 1990



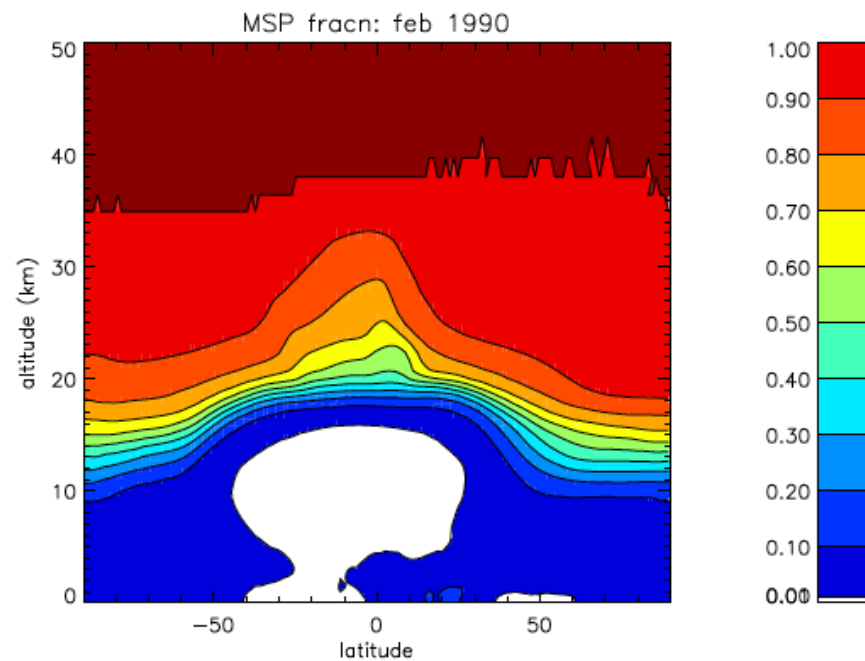
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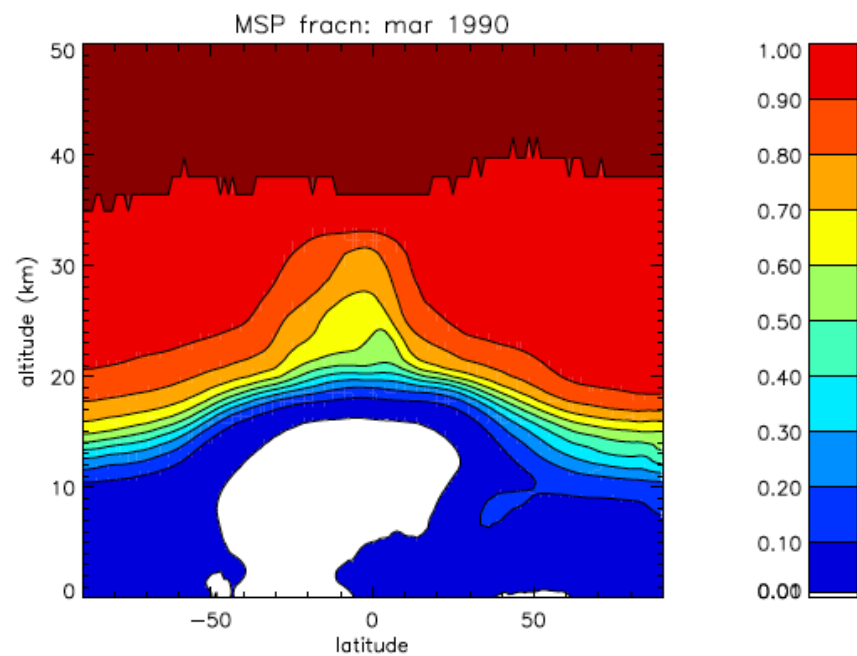
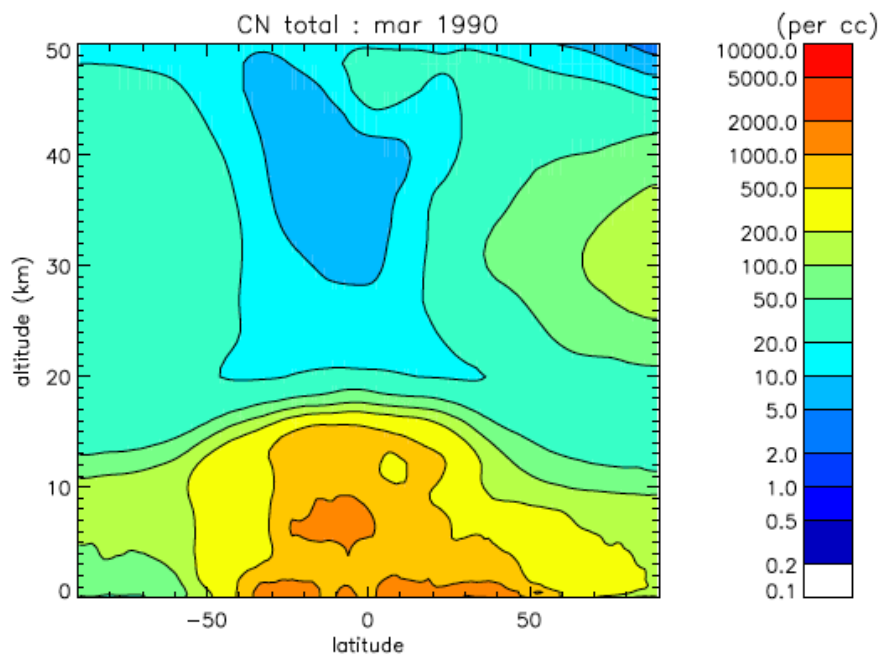
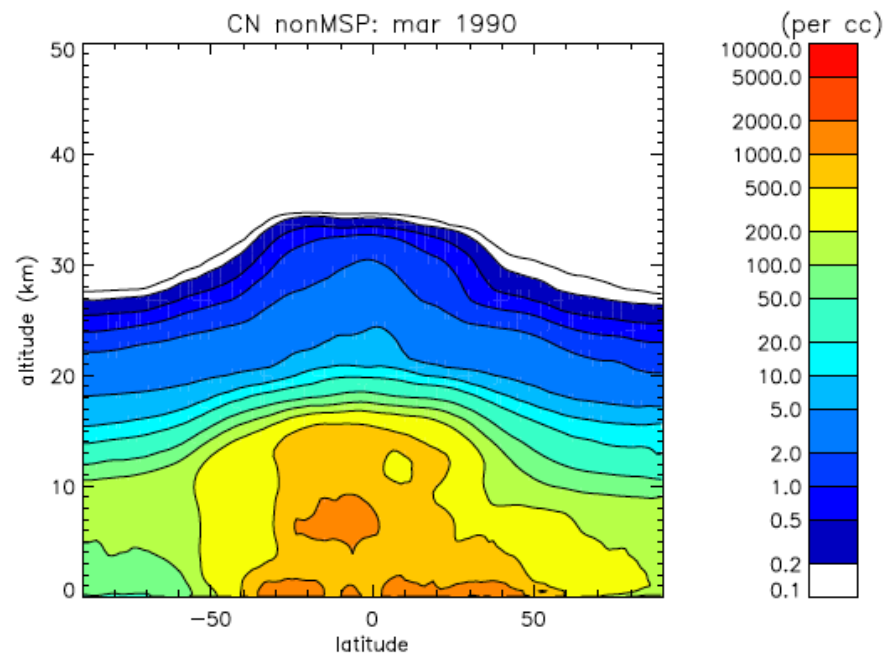
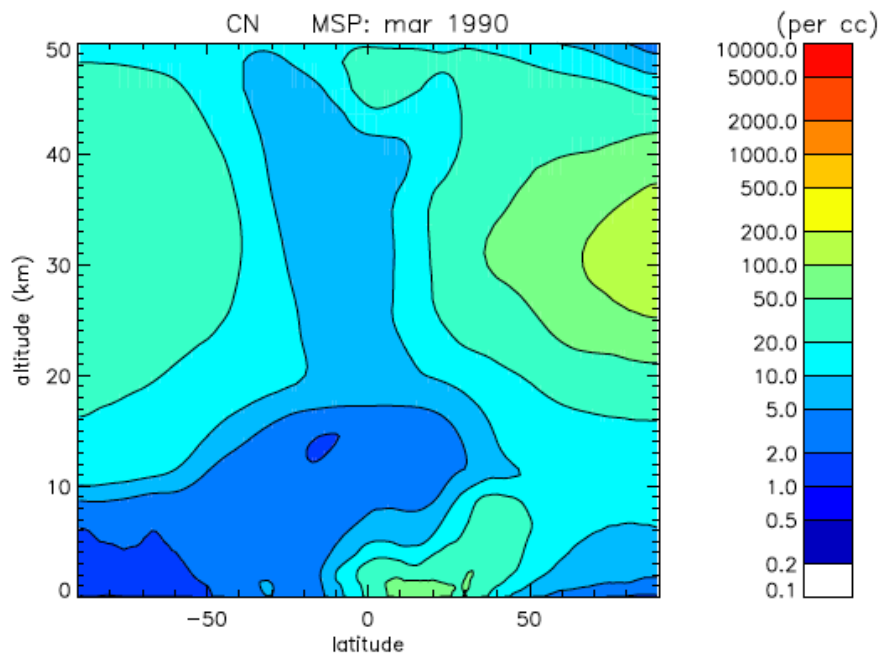


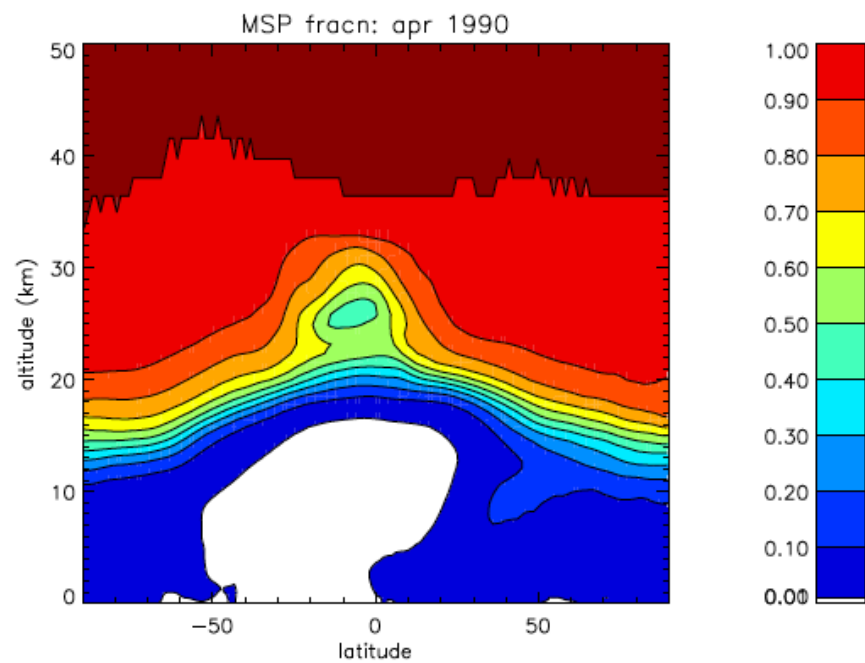
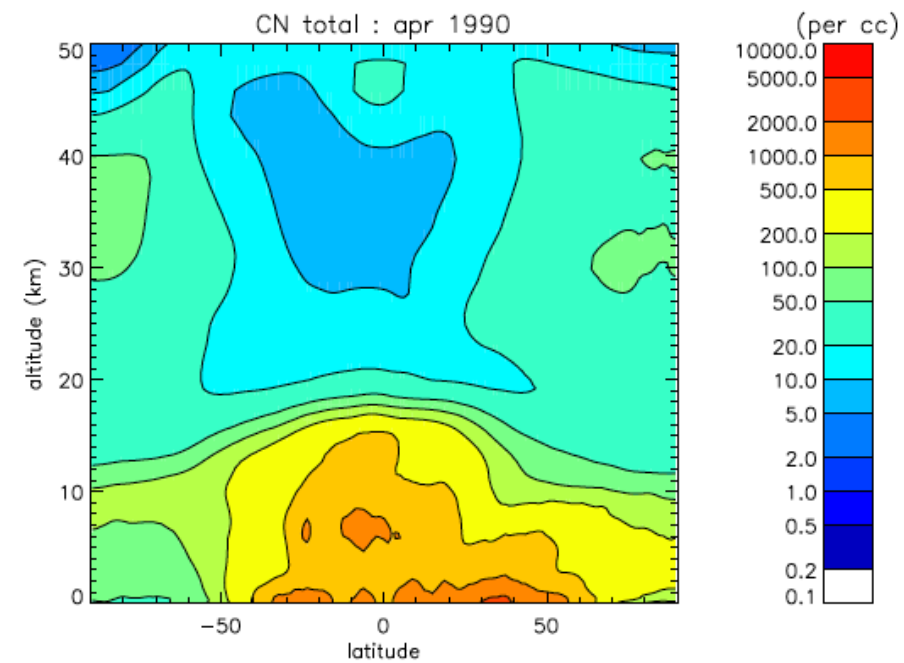
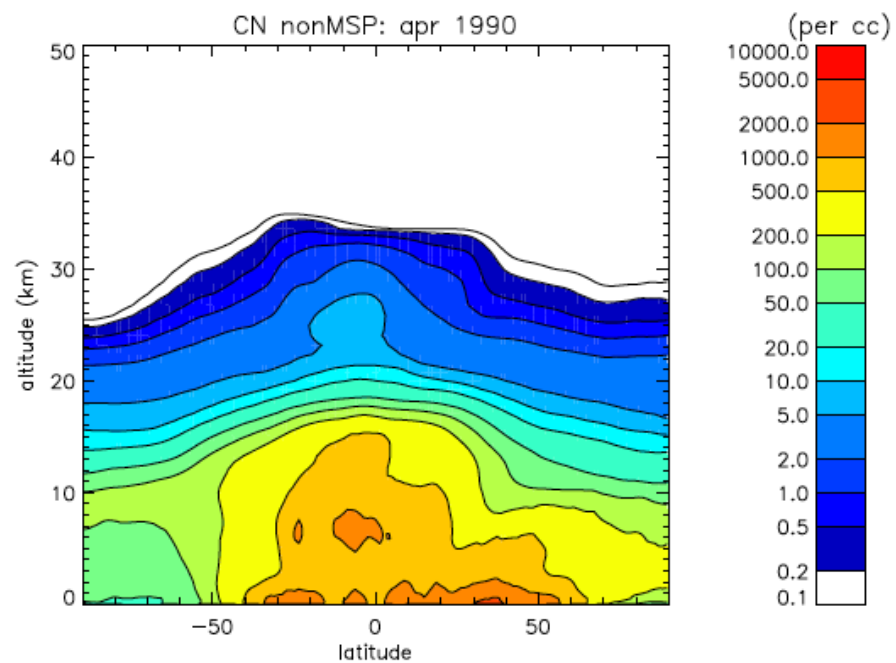
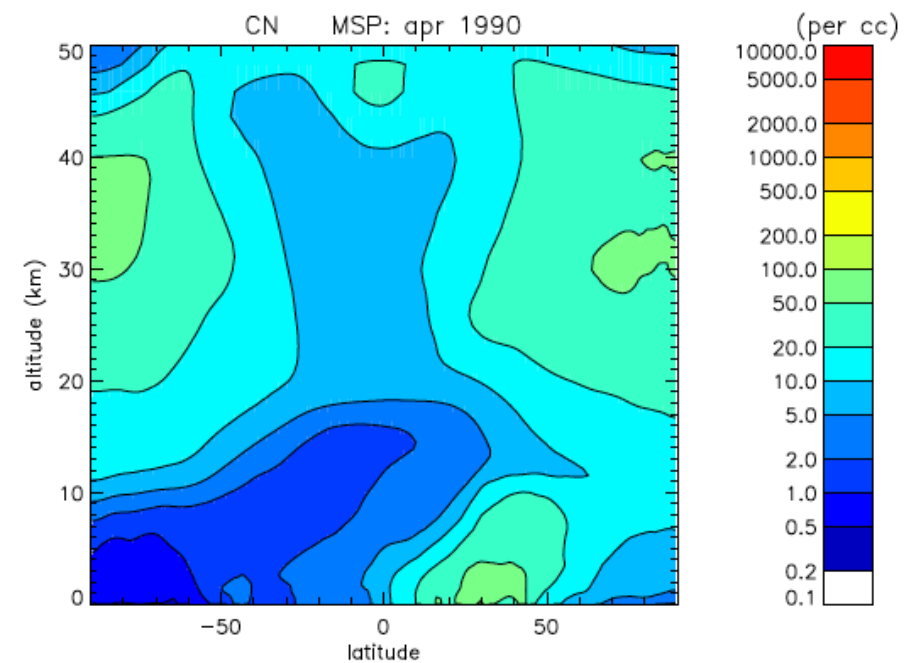
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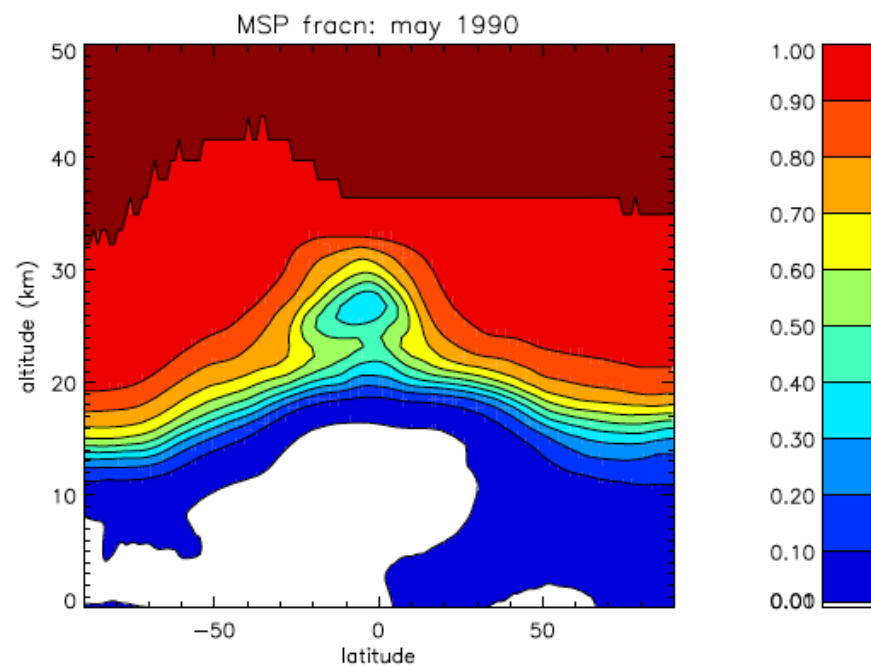
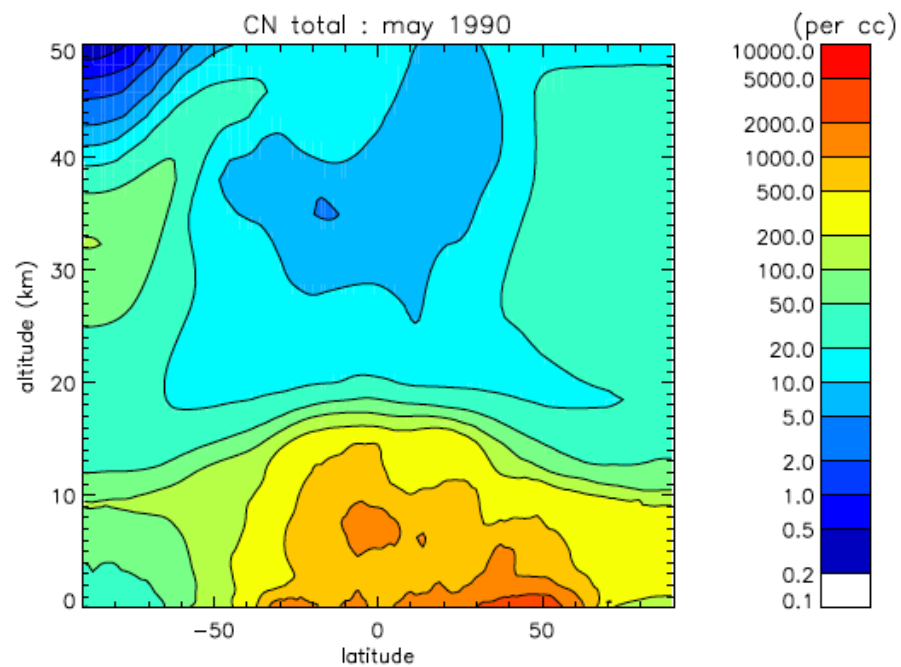
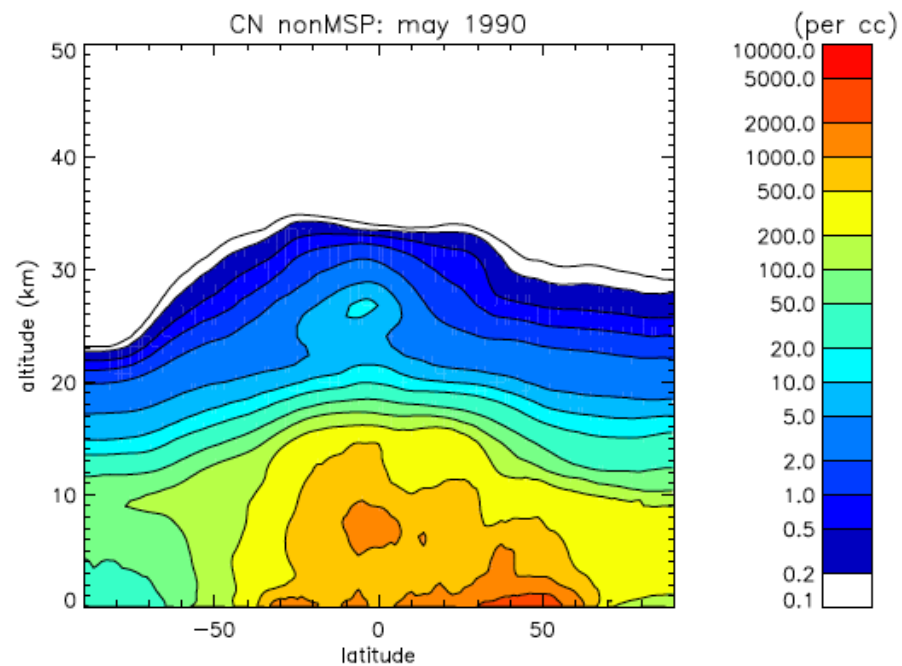
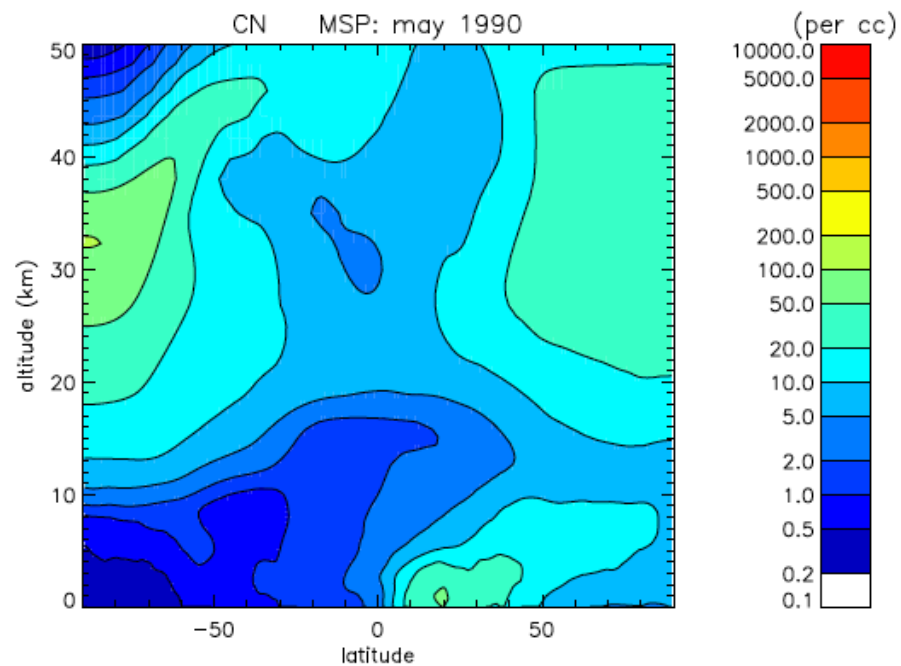


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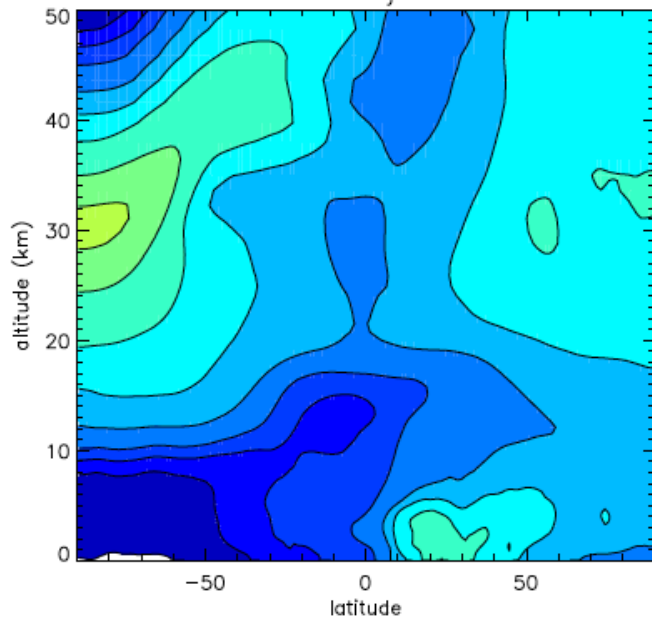




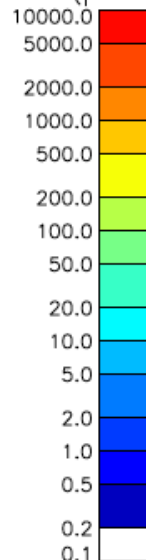




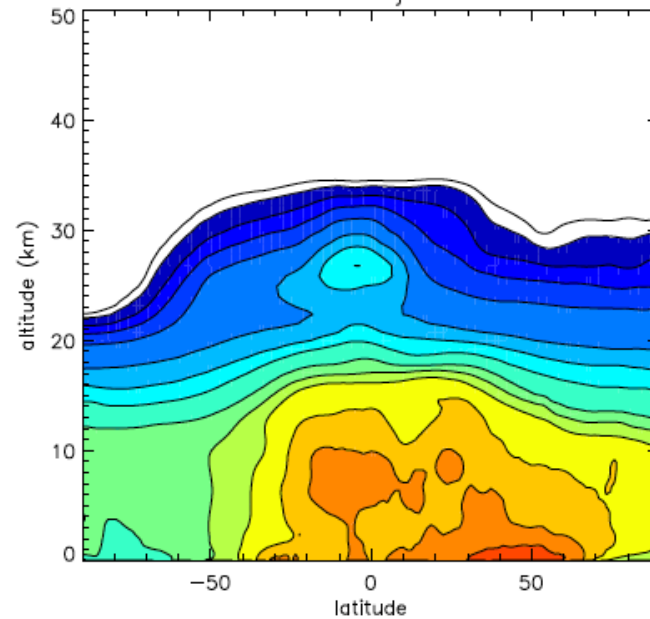
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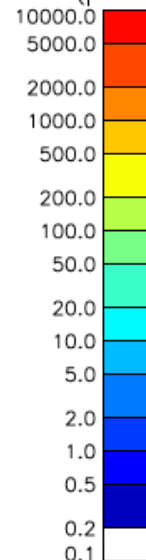
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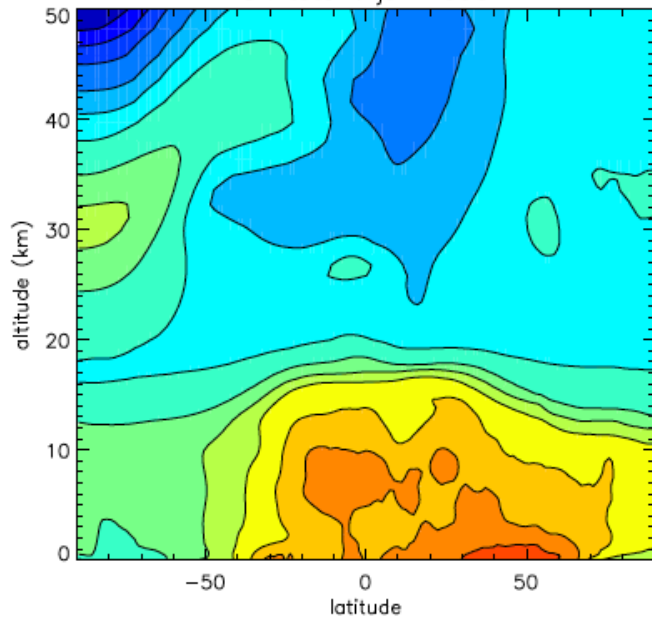
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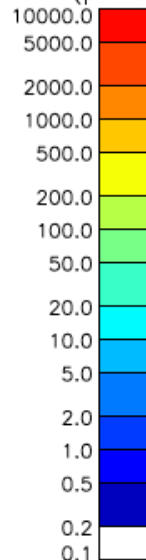
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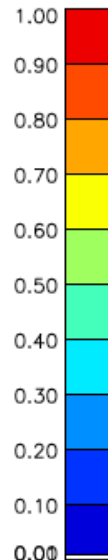
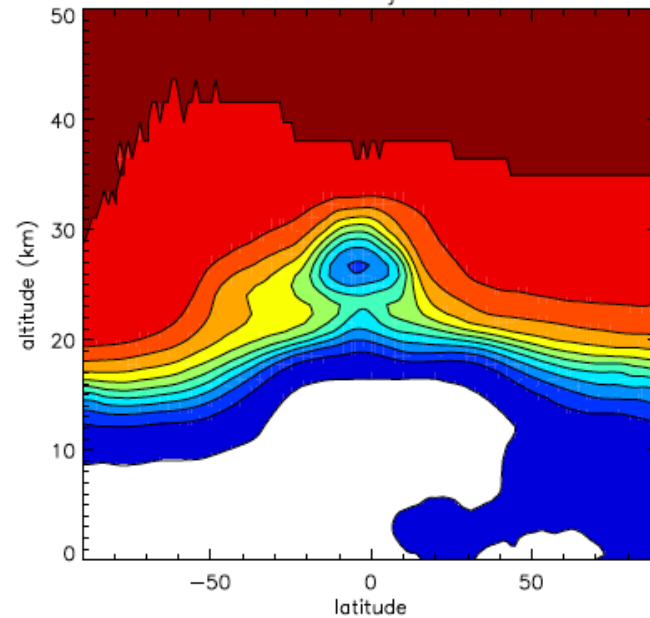
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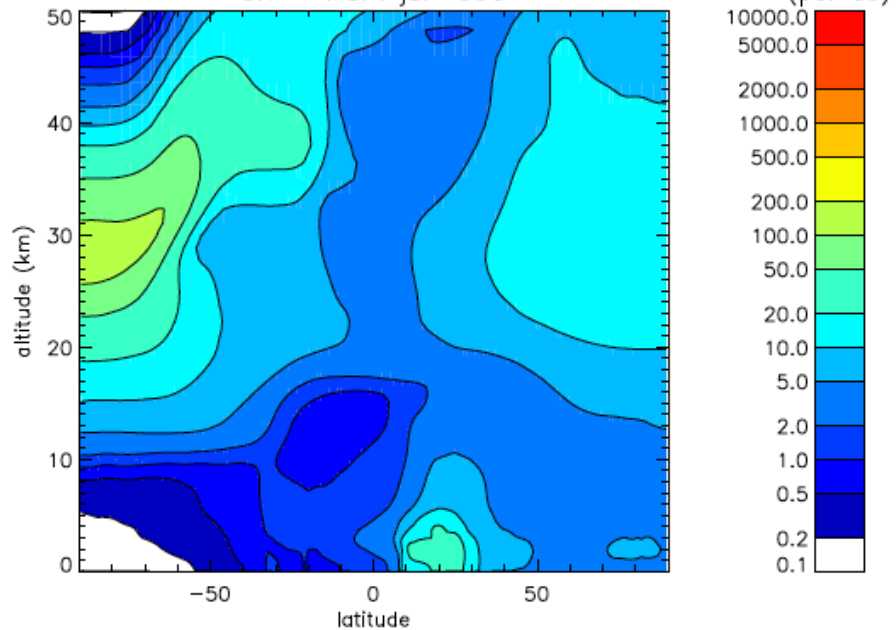


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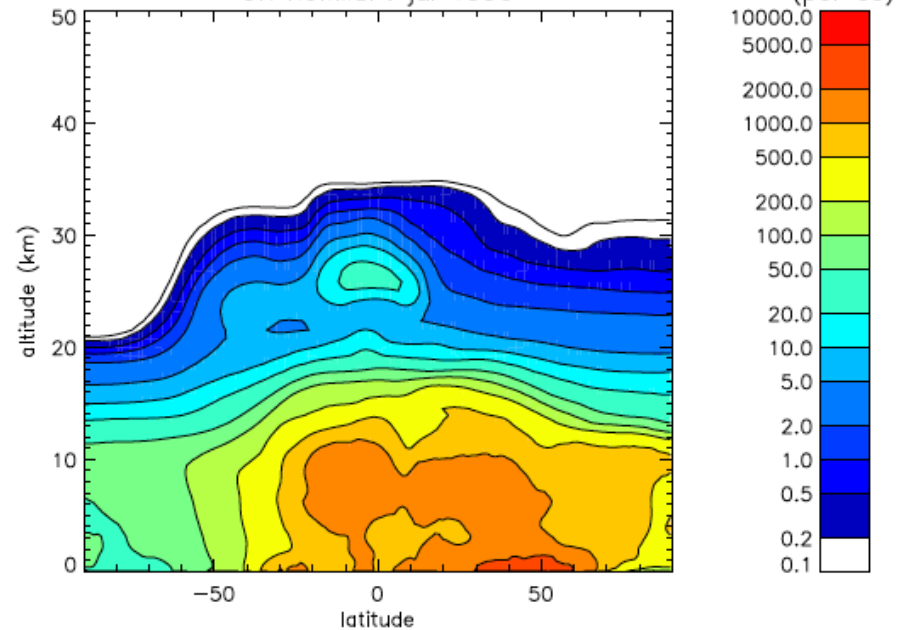




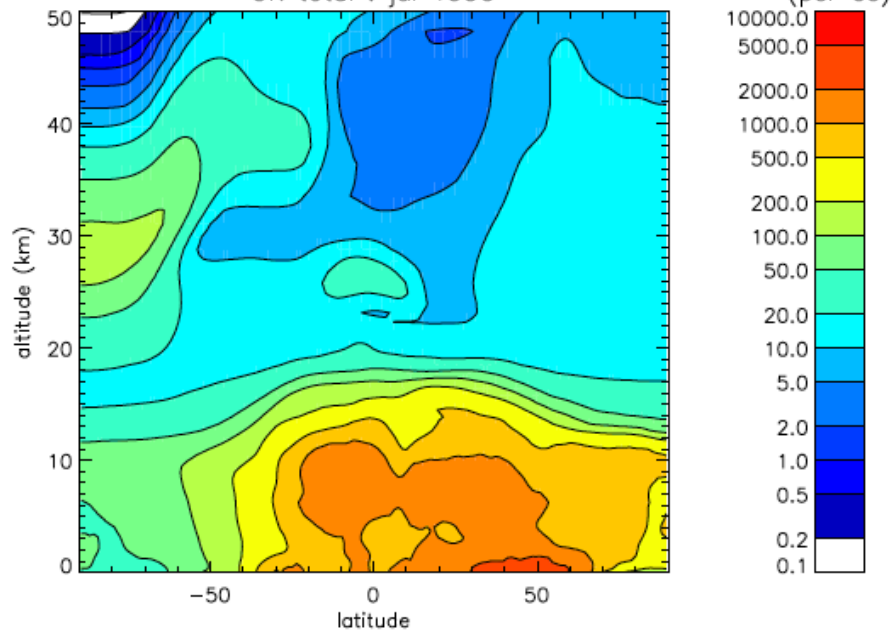
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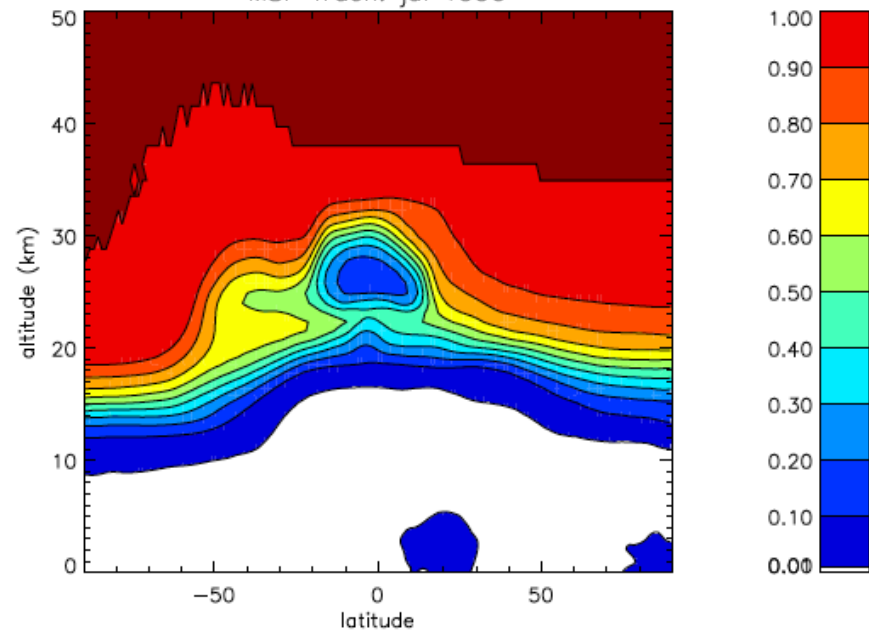
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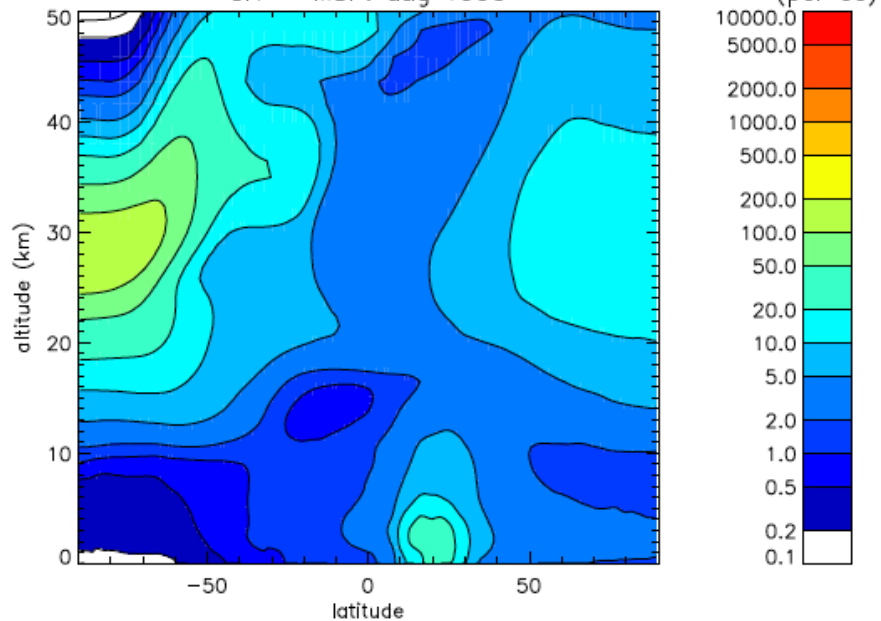
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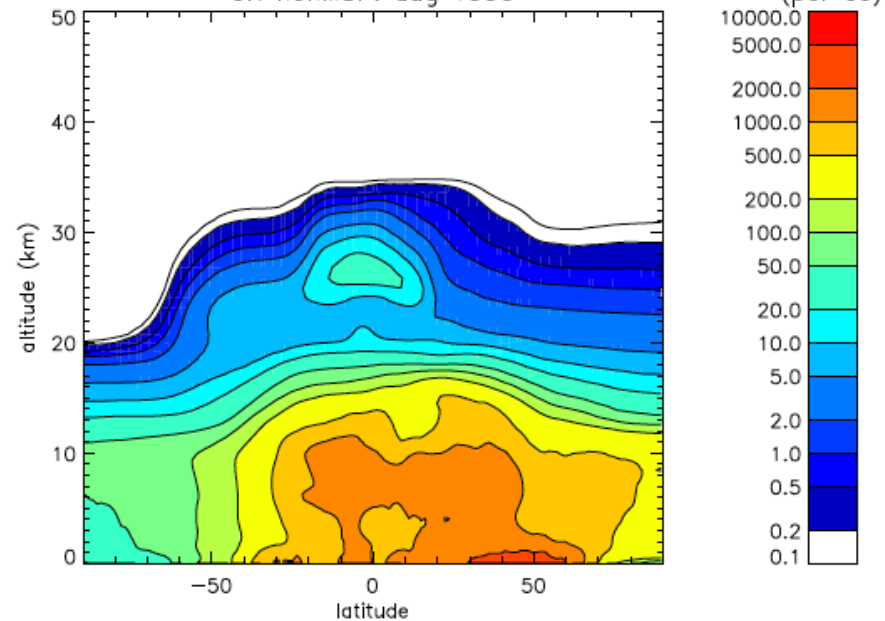
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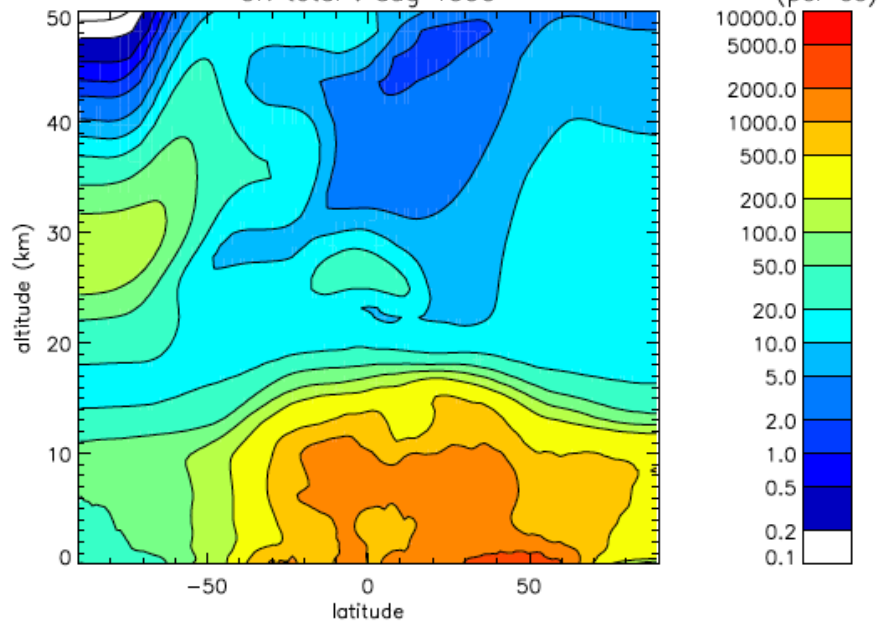
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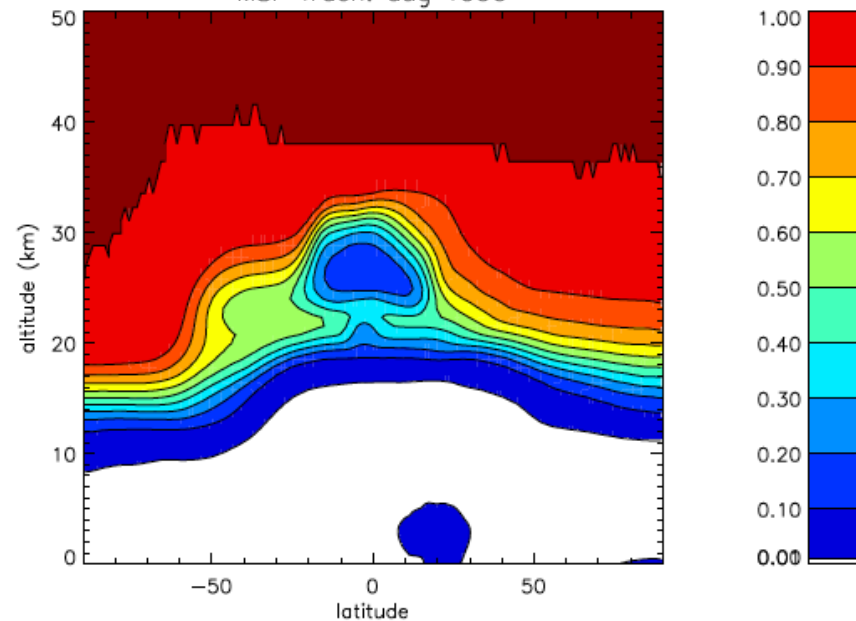
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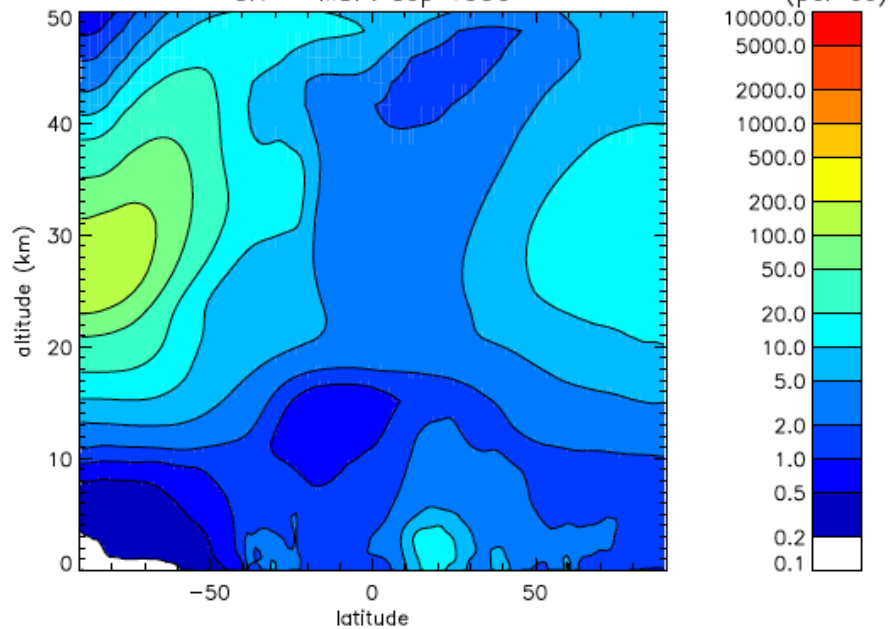
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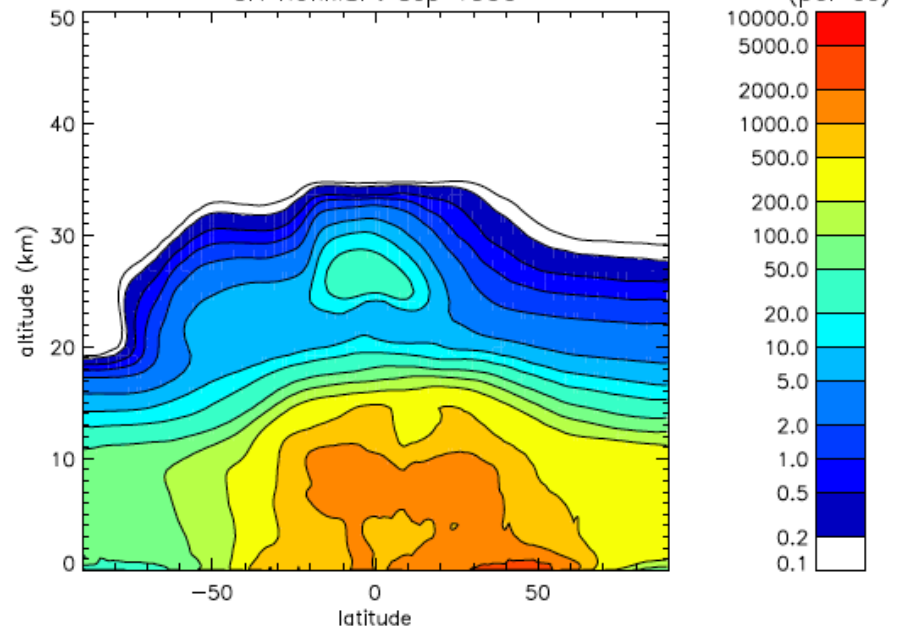
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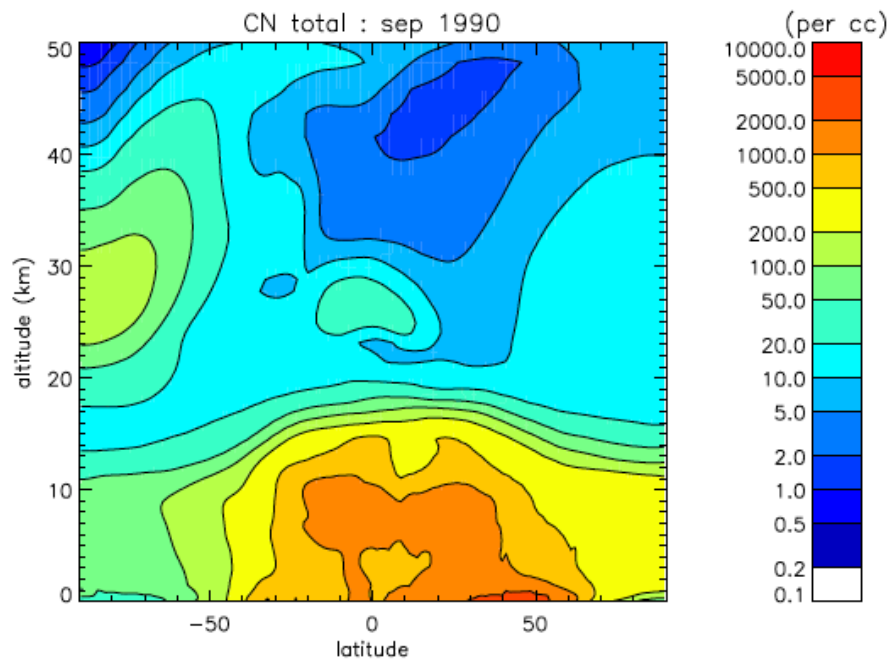
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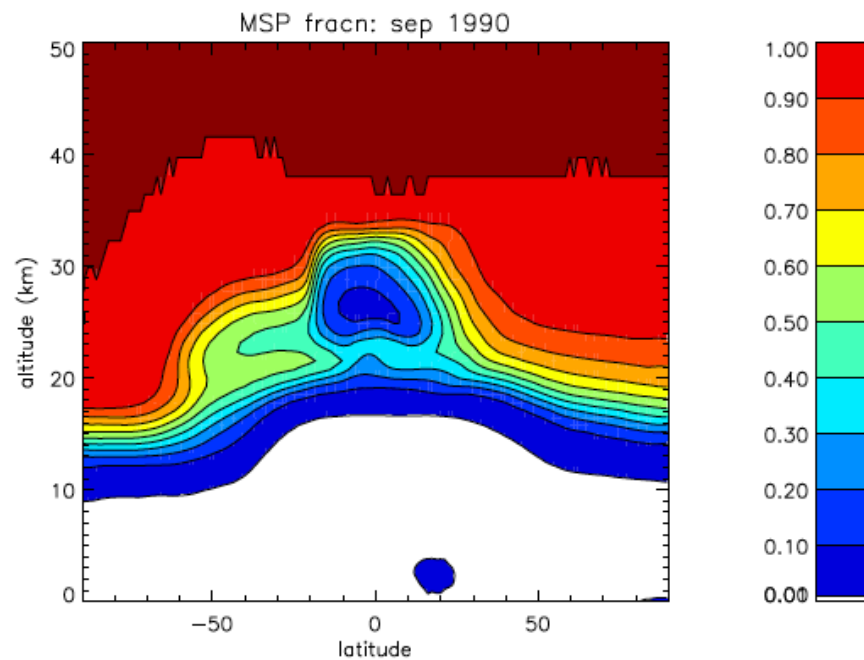
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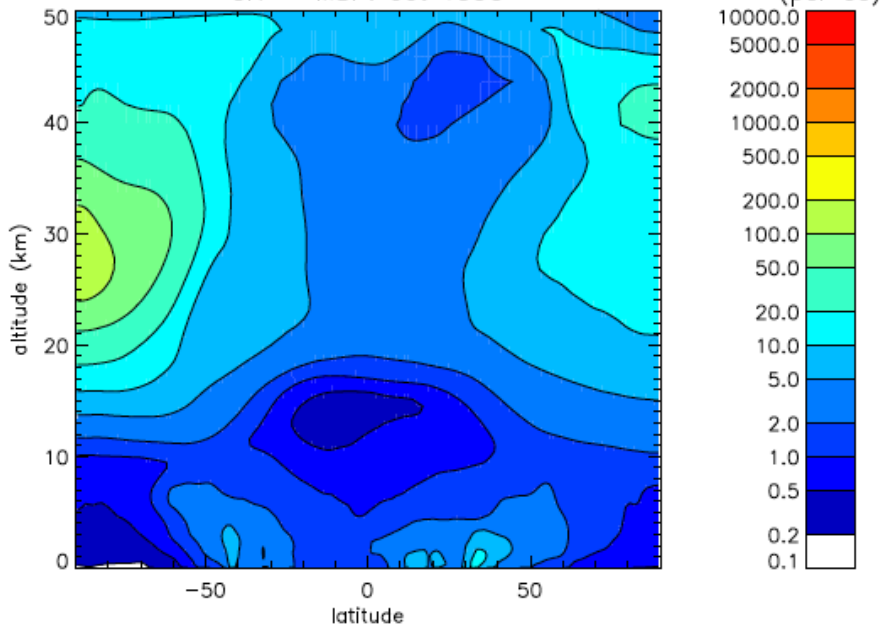
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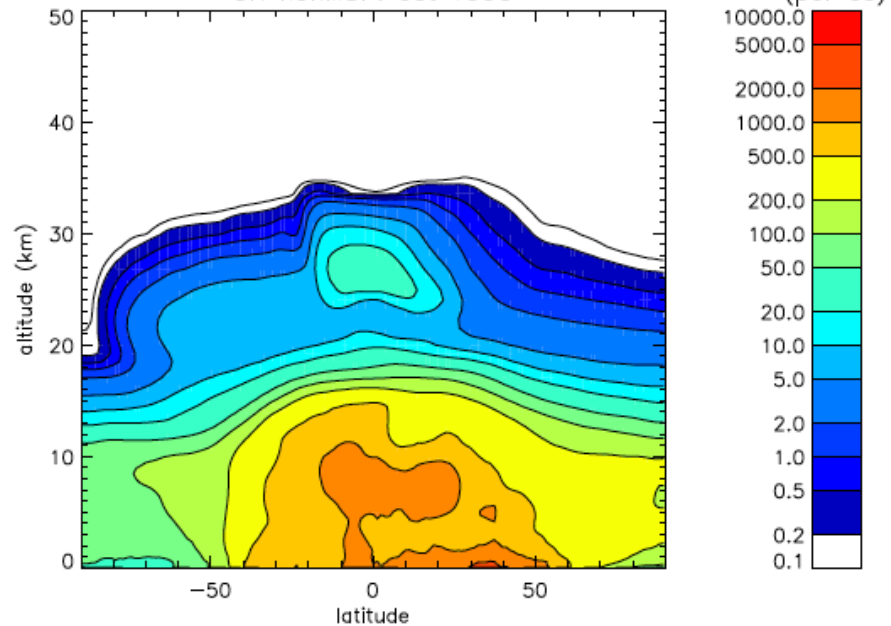
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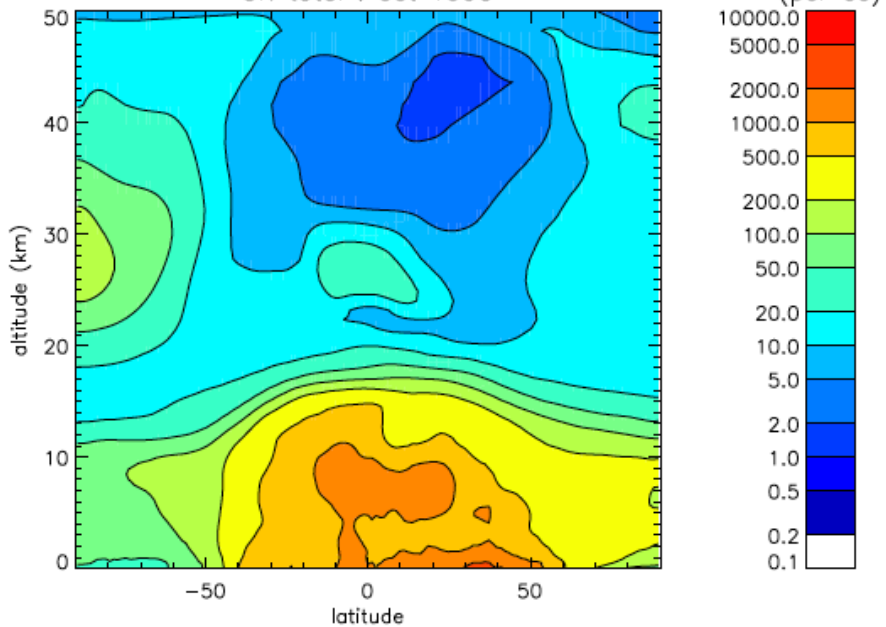
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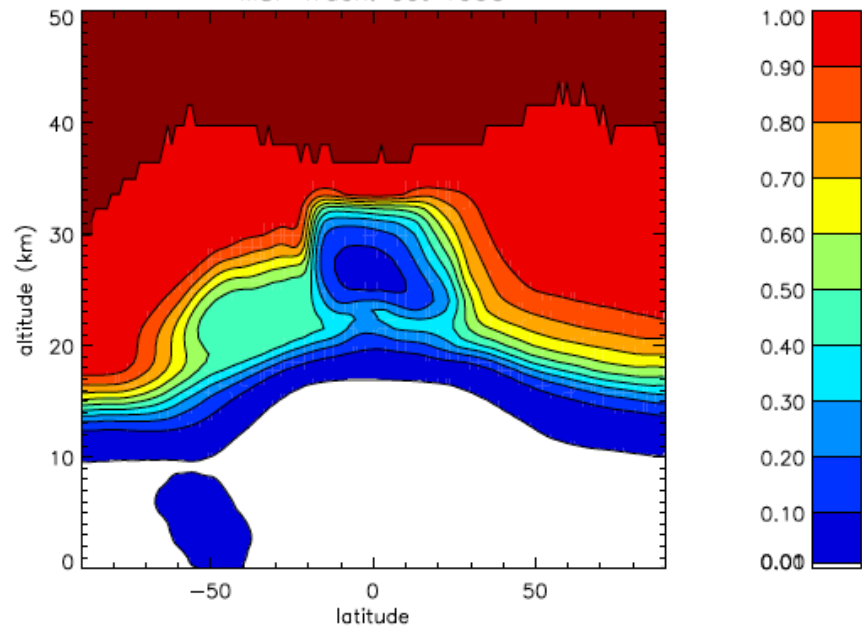
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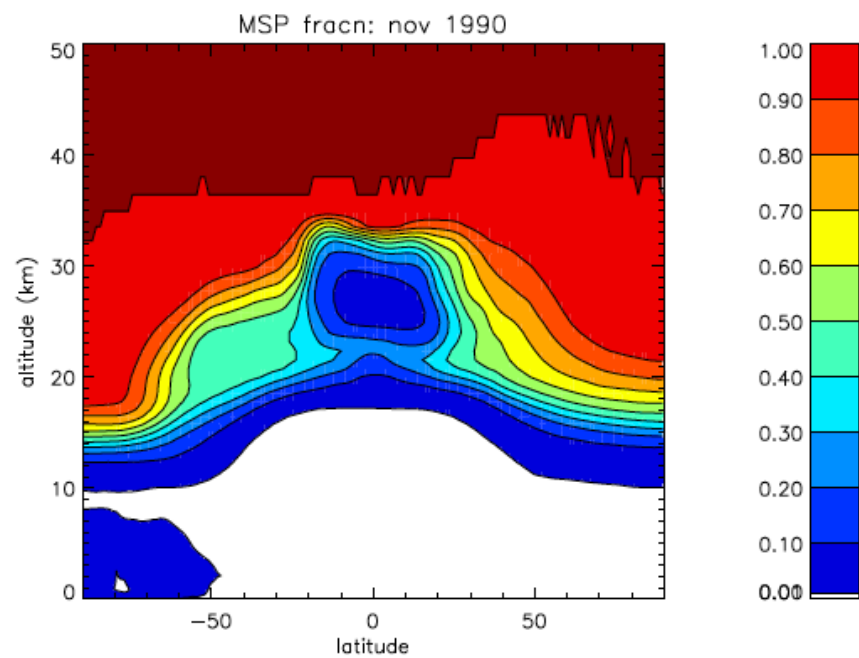
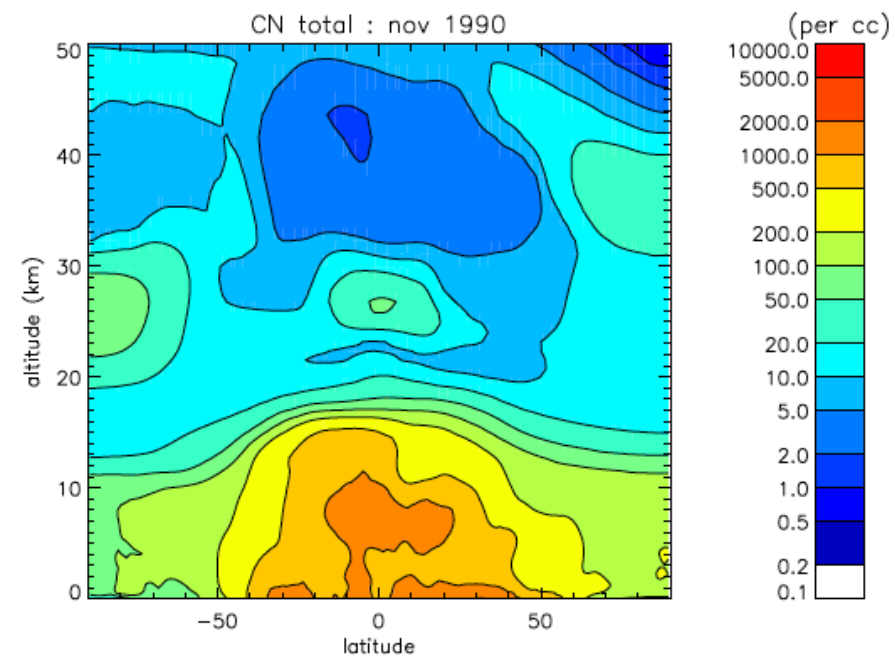
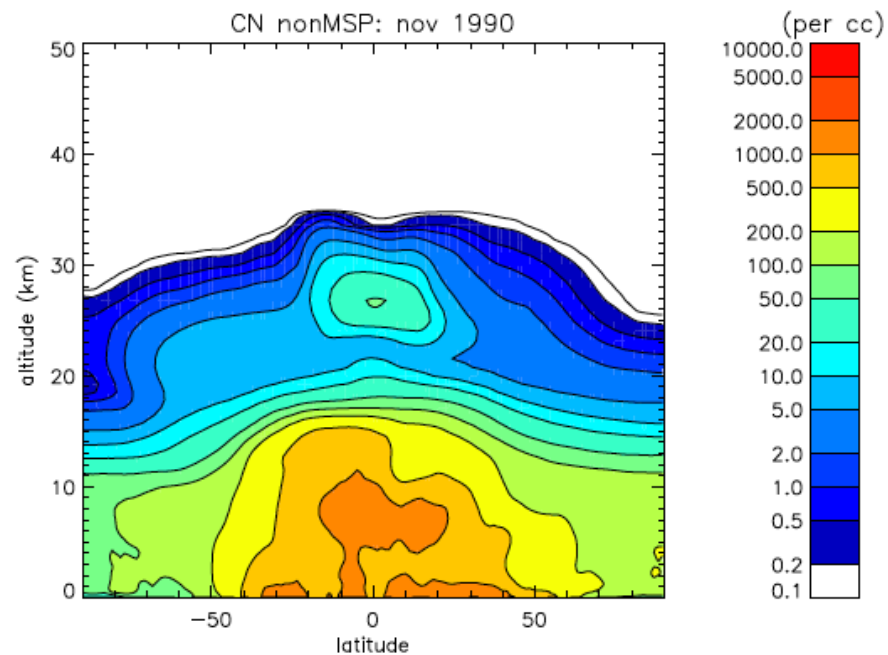
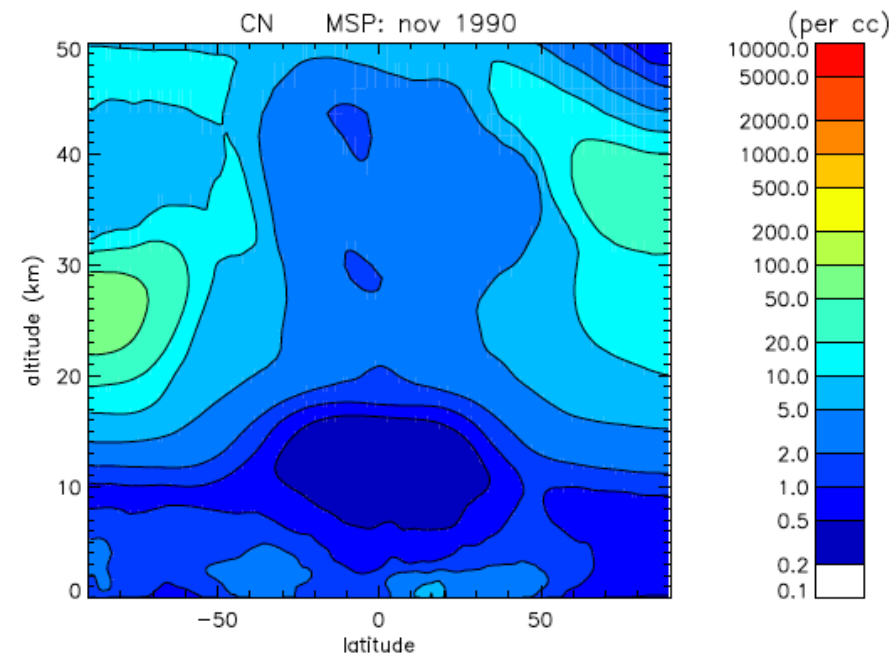


CN total : oct 1990

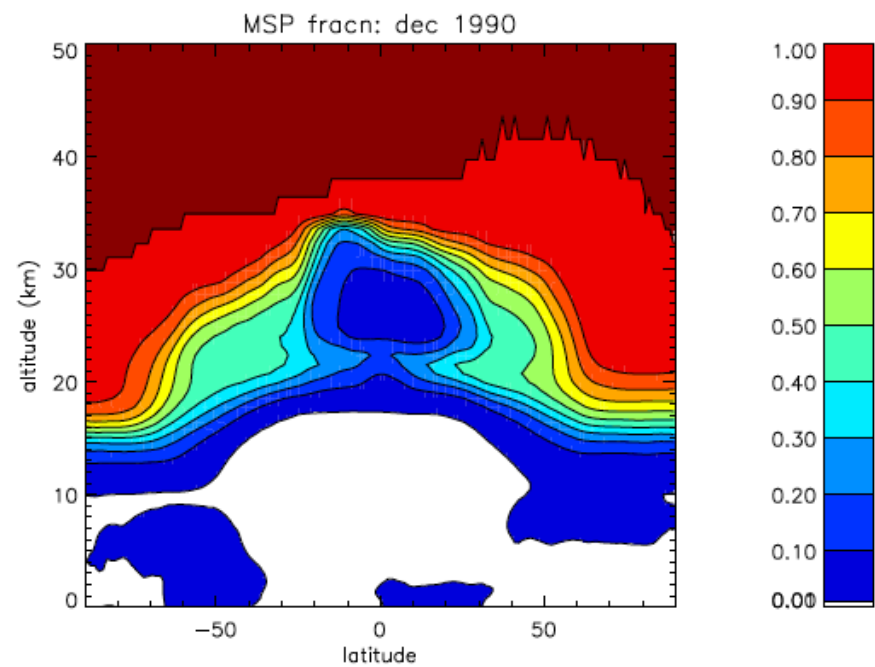
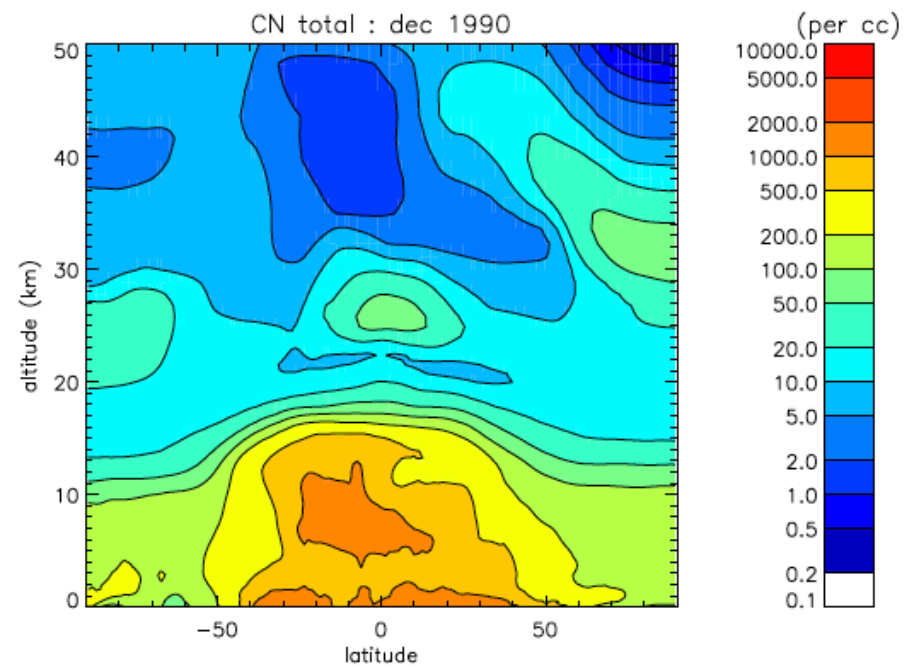
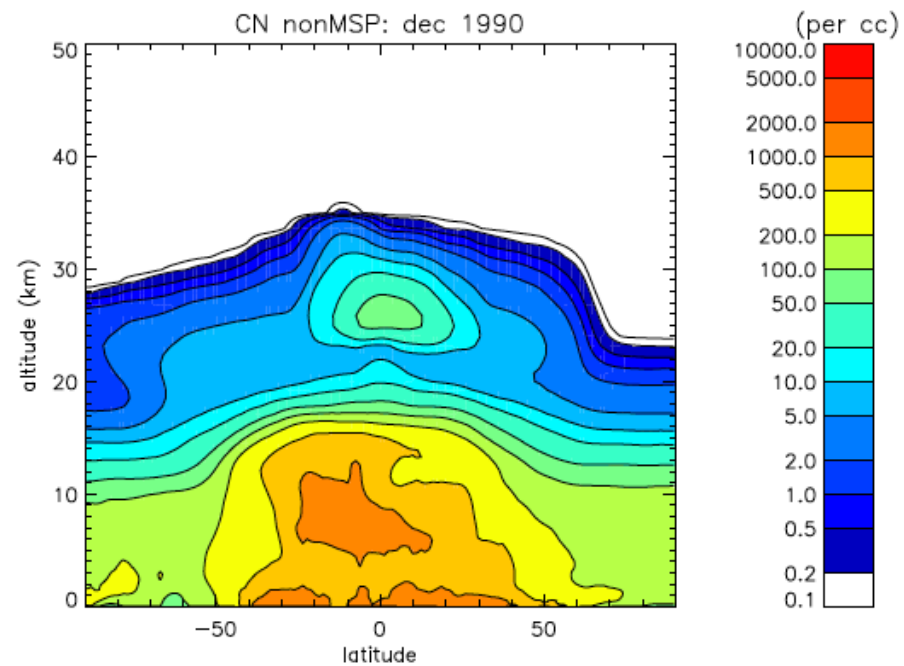
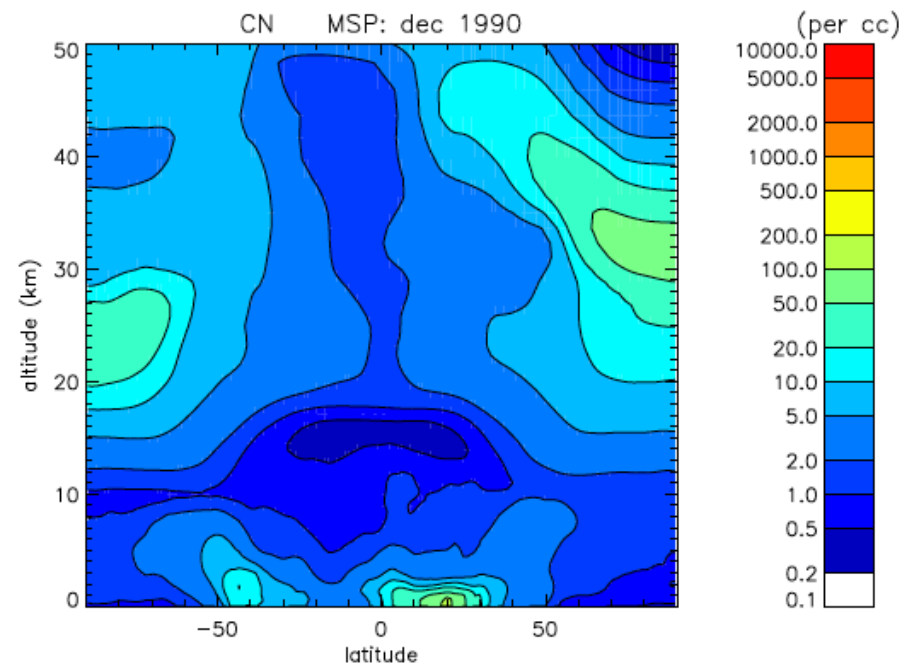


MSP fracn: oct 1990

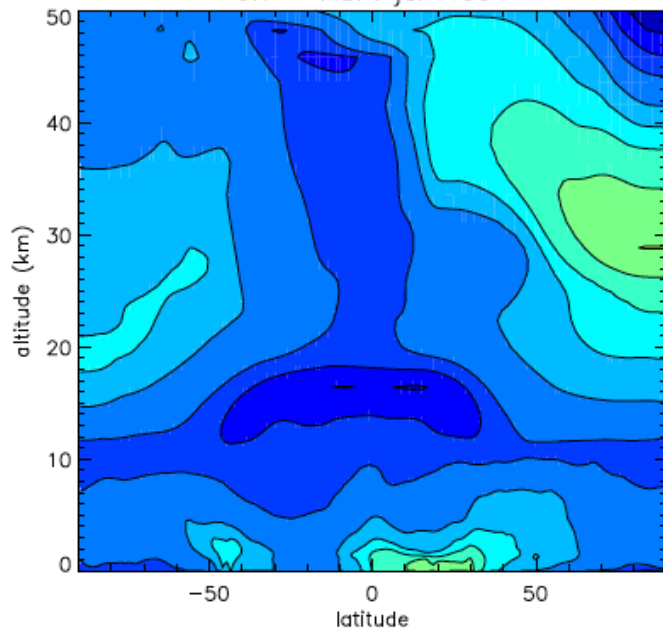




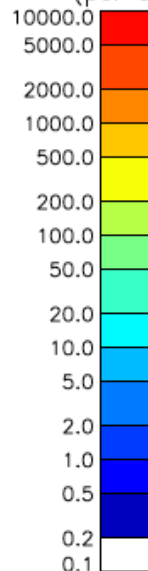




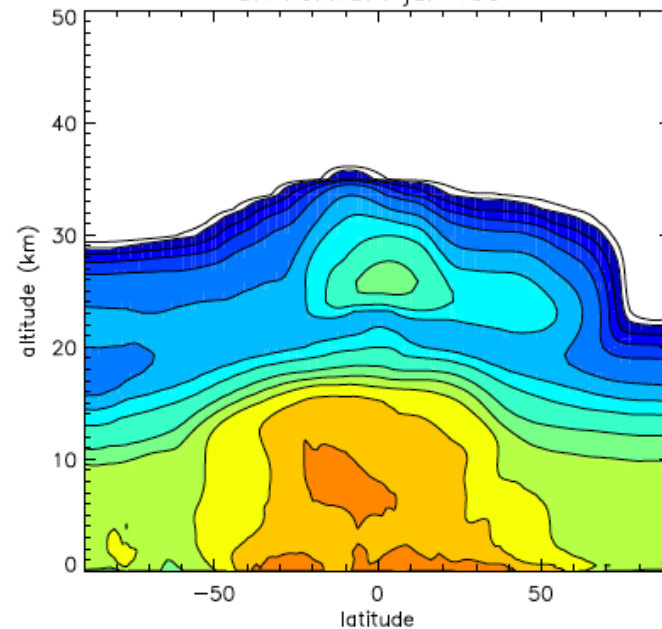
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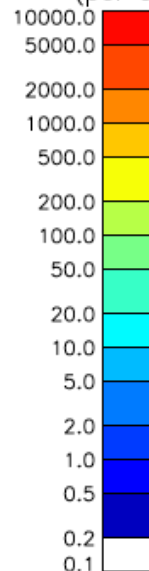
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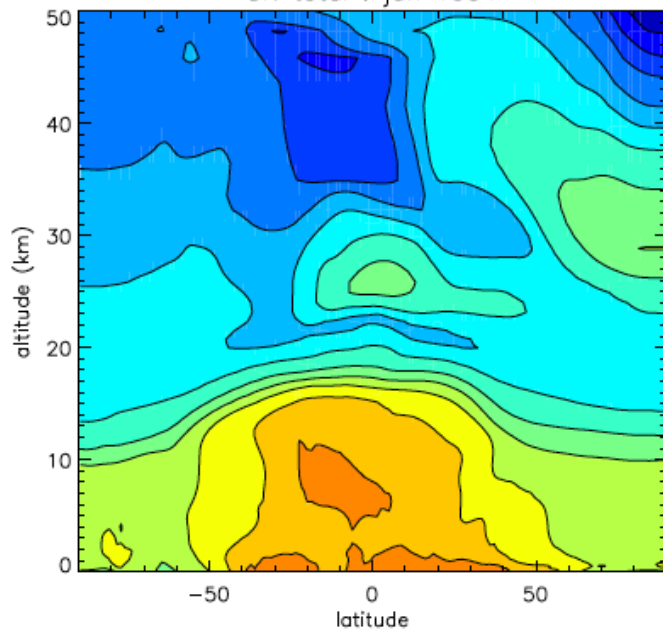
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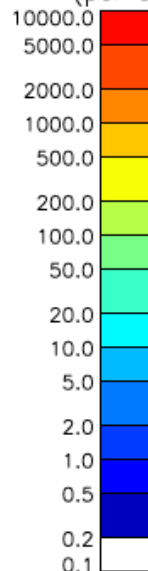
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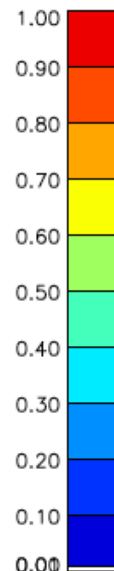
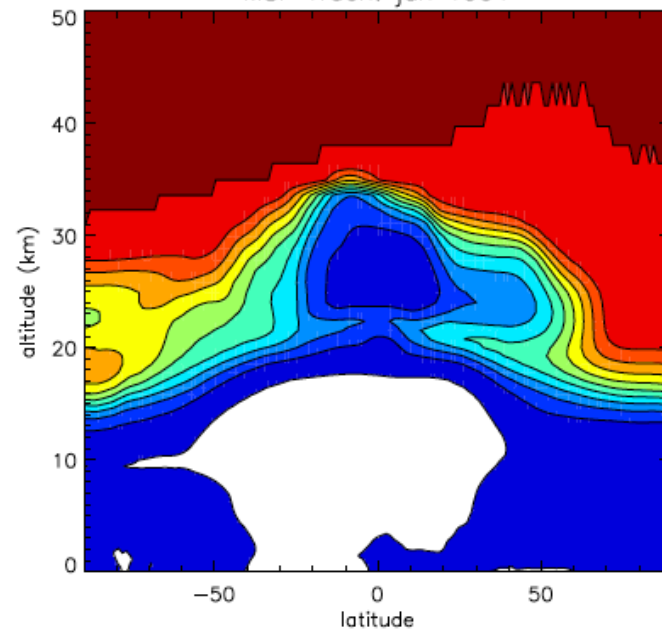
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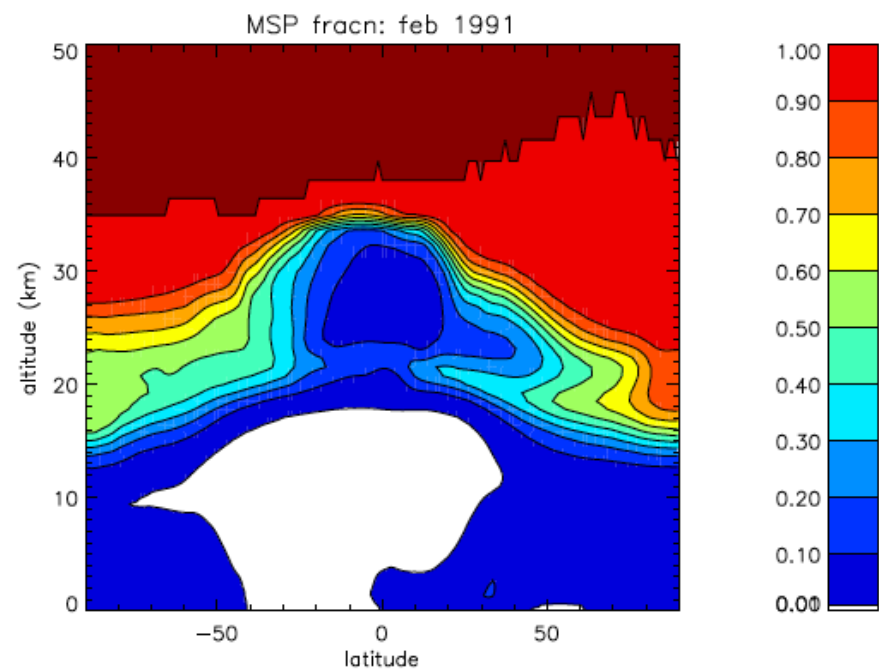
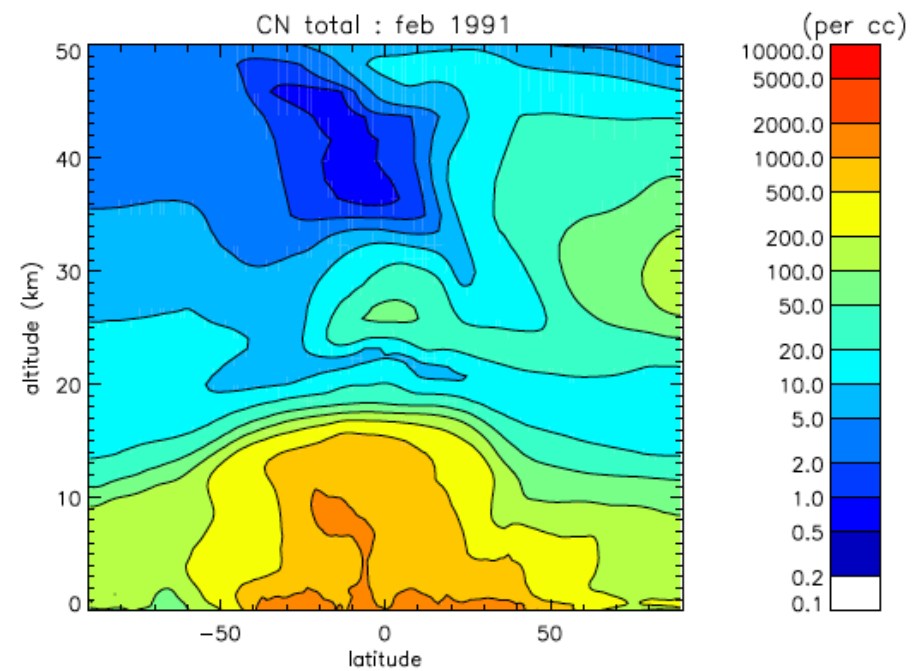
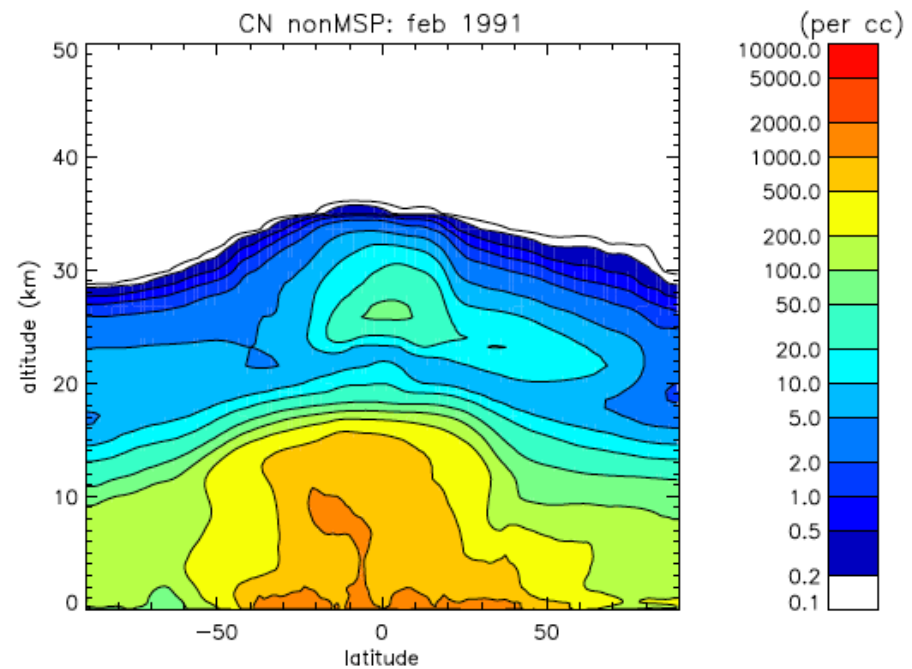
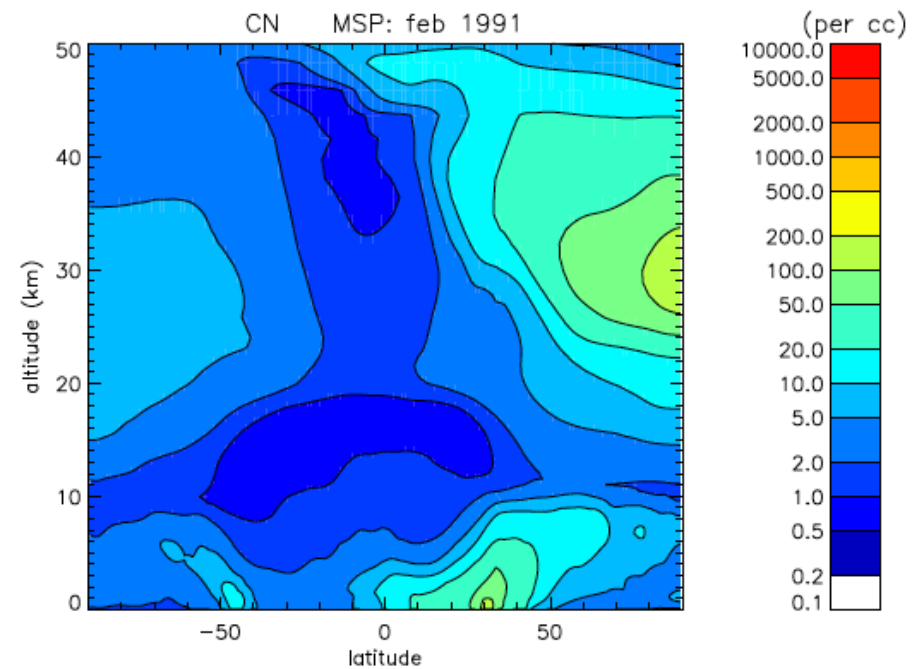


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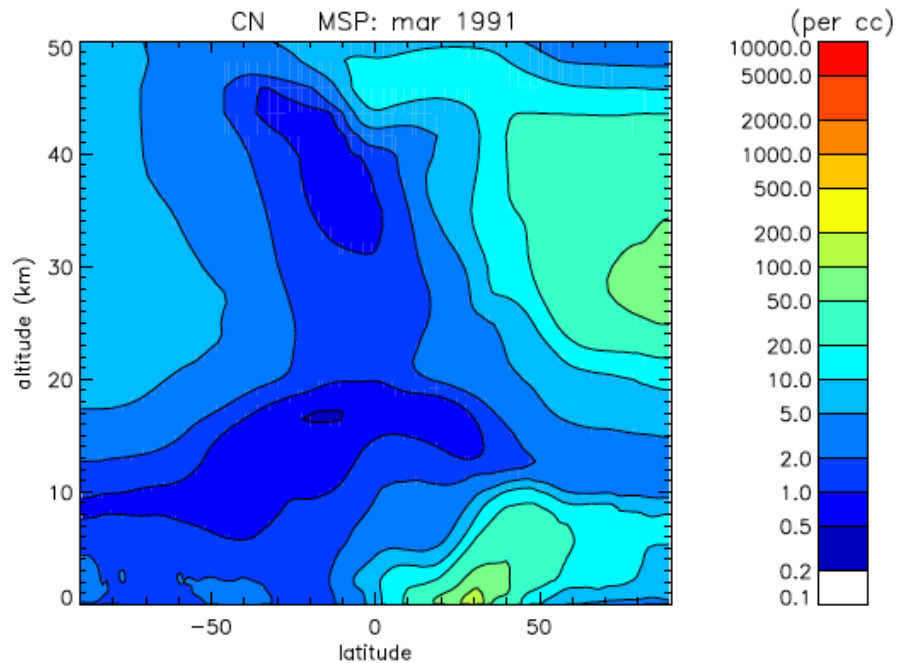


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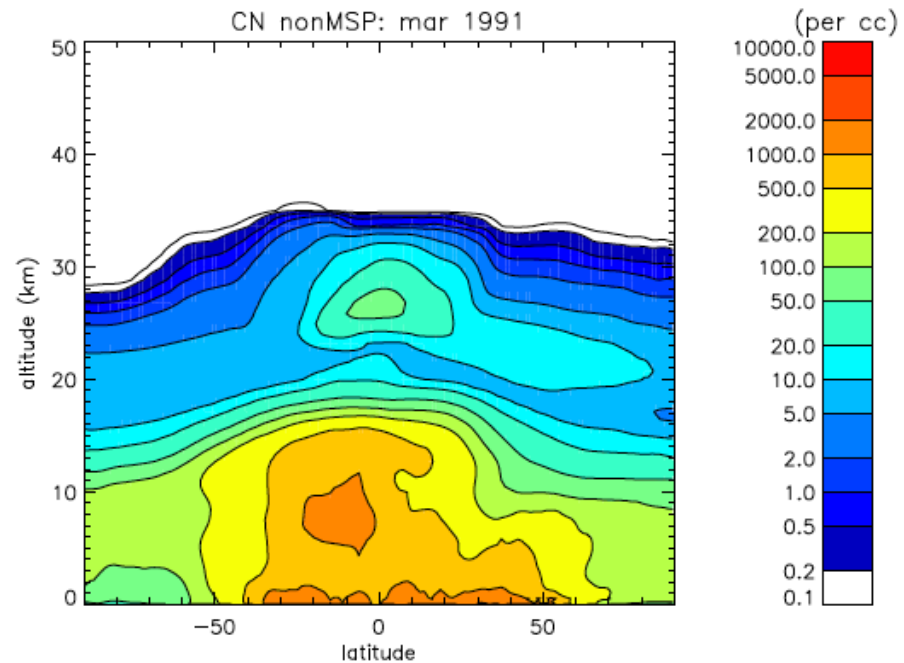




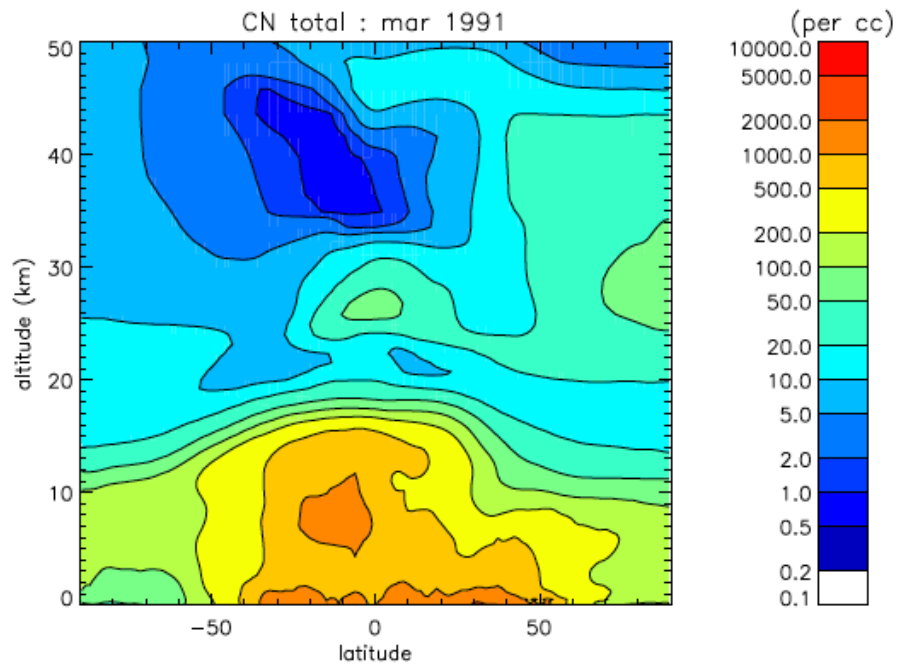
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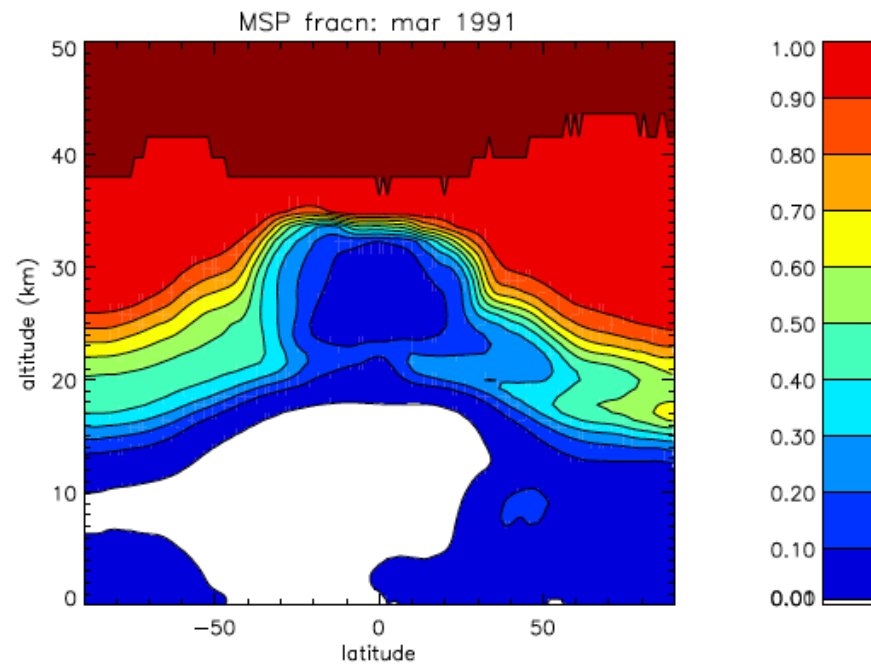
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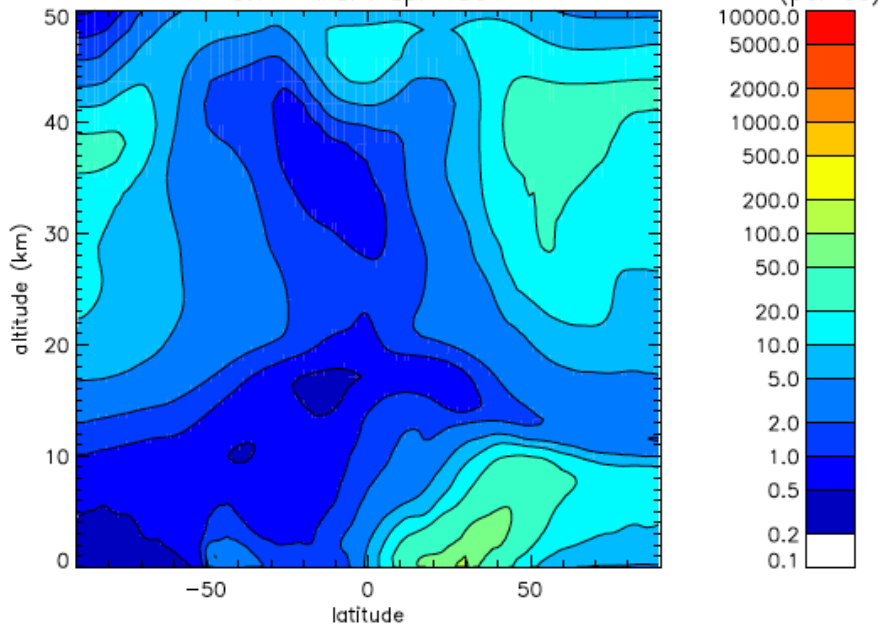
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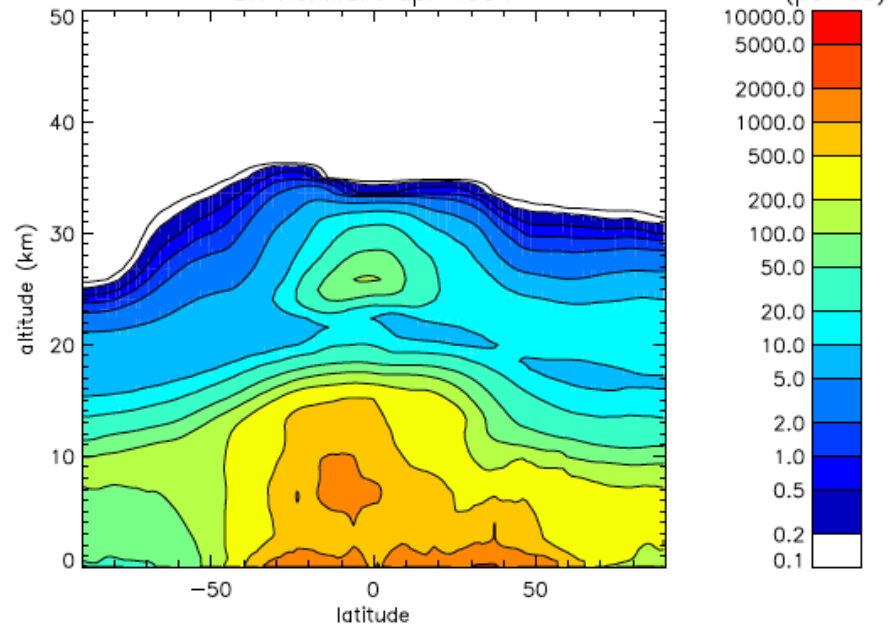
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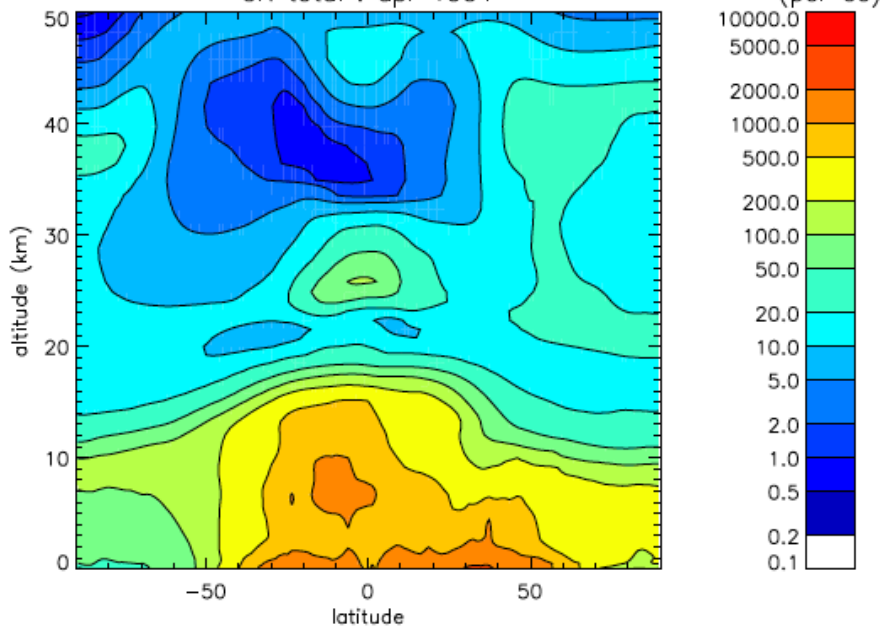
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CN nonMSP: apr 1991



CN total : apr 1991



MSP fracn: apr 1991

