

*Assimilation of TES  
observations for the analysis  
of processes governing the  
chemistry of the troposphere*

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# *Tropospheric models*

## ✧ AM2-Chem

- ✧ GCM developed at NOAA GFDL.
- ✧ B-grid model,  $2.0^\circ$  latitude x  $2.5^\circ$  longitude, 24 vertical levels (top level approx. 45km).
- ✧ Chemistry scheme based on MOZART-2 [Horowitz et al., 2003; Tie et al., 2004].
- ✧ Model dynamics constrained by Newtonian nudging to NCEP reanalyses.

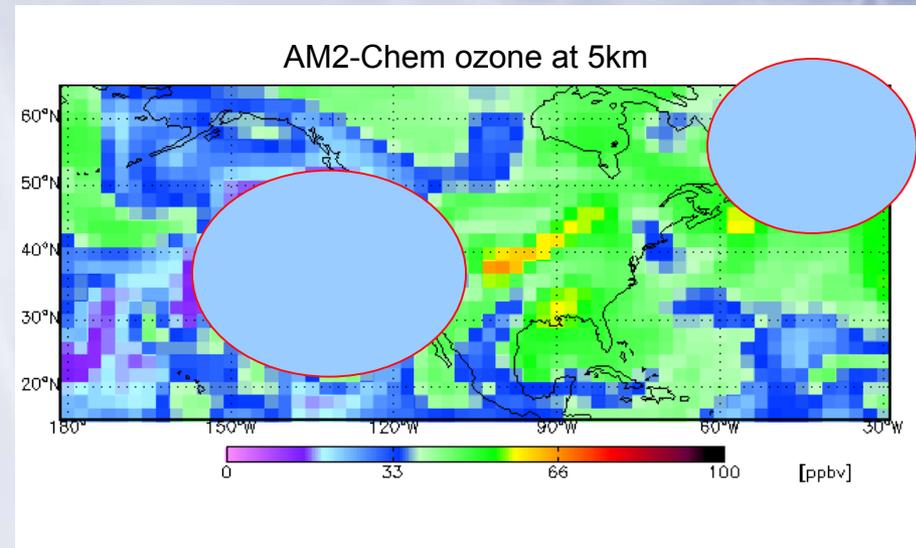
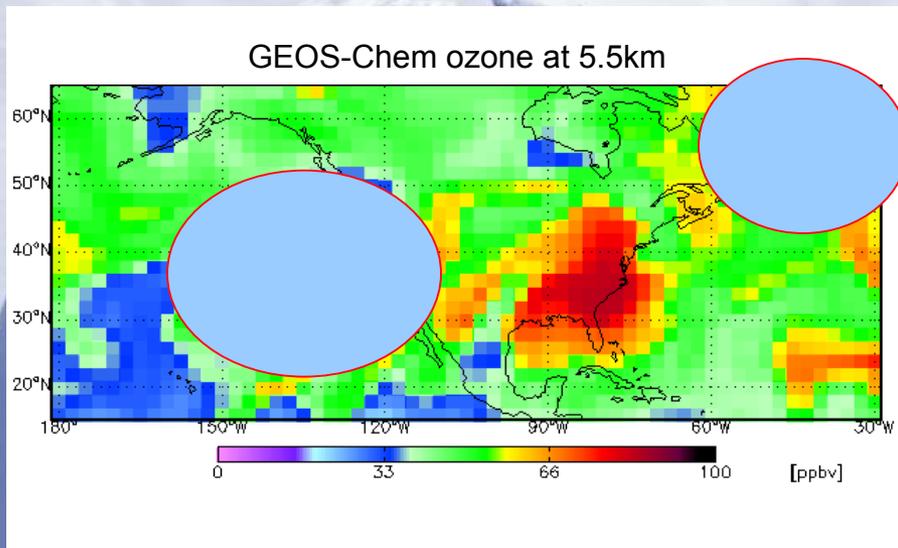
## ✧ GEOS-Chem

- ✧ CTM.
- ✧  $2.0^\circ$  latitude x  $2.5^\circ$  longitude, 30 vertical levels (top level approx. 80km).
- ✧ Model transport driven by GEOS-4 GMAO analyses.

Sequential Kalman filter approach to the assimilation of TES ozone and carbon monoxide profiles.

TES data assimilated for July 2005.

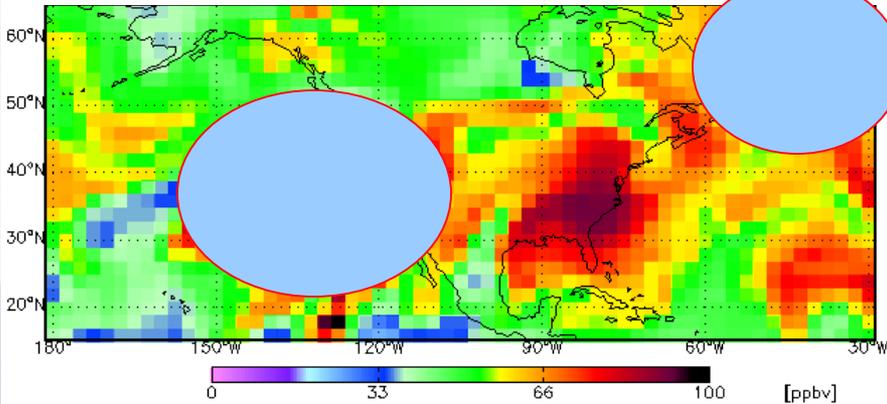
# *Tropospheric Ozone*



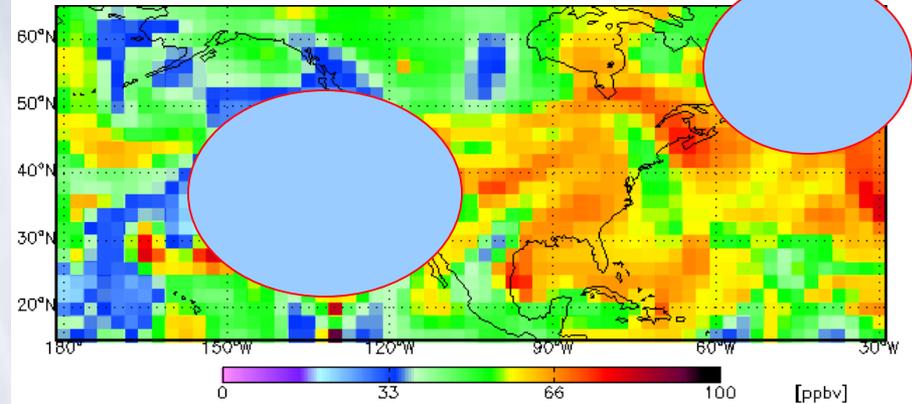
- ✧ Ozone field over North America at approx. 5km on 26th July 2005.
- ✧ Dynamical features correlate in both models.
- ✧ Ozone abundance is generally much greater in GEOS-Chem compared to AM2-Chem.
- ✧ Large discrepancy over SE USA due to more  $\text{NO}_x$  in GEOS-Chem, possibly due to lightning source in mid-troposphere.

# Impact of TES data

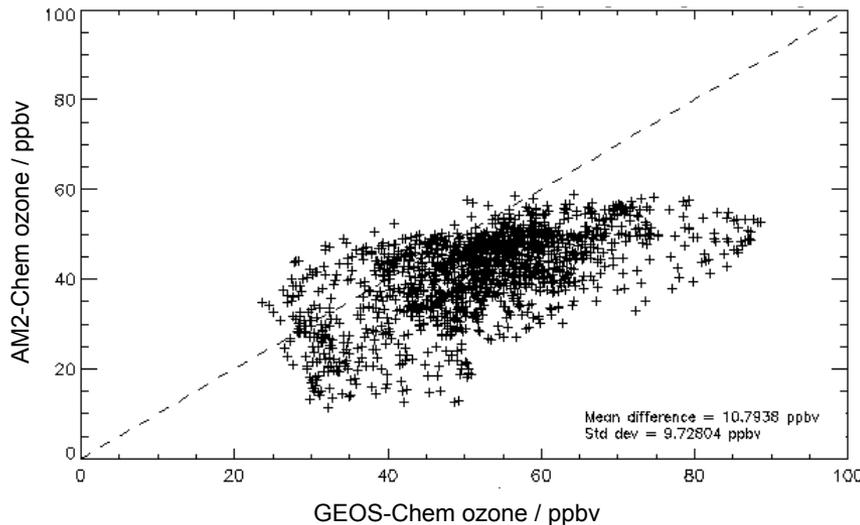
GEOS-Chem ozone analysis at 5.5km



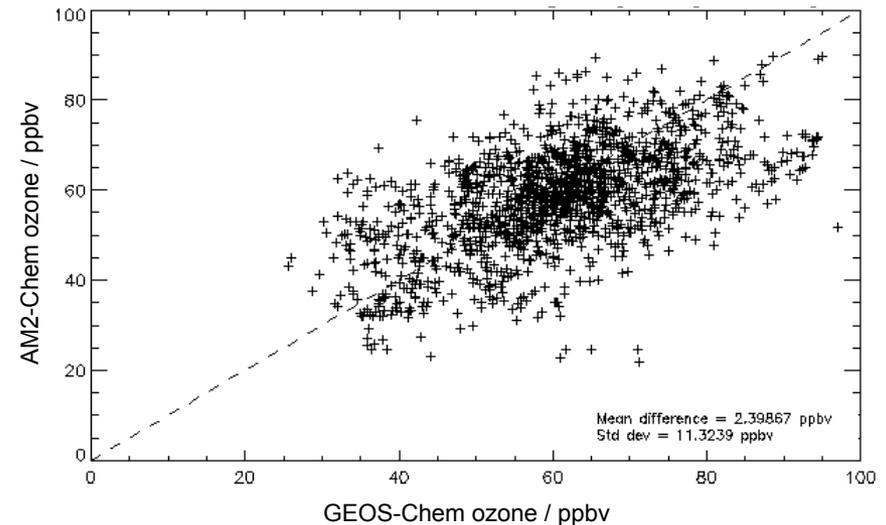
AM2-Chem ozone analysis at 5km



Scatter of model ozone fields at 5km



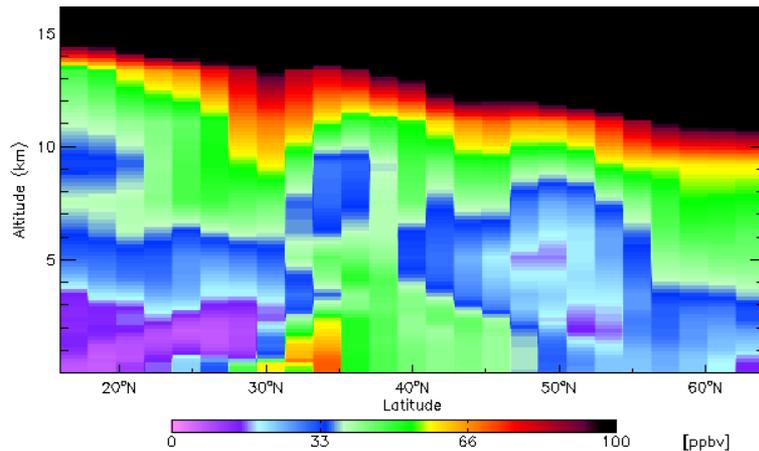
Scatter of model ozone analysis fields at 5km



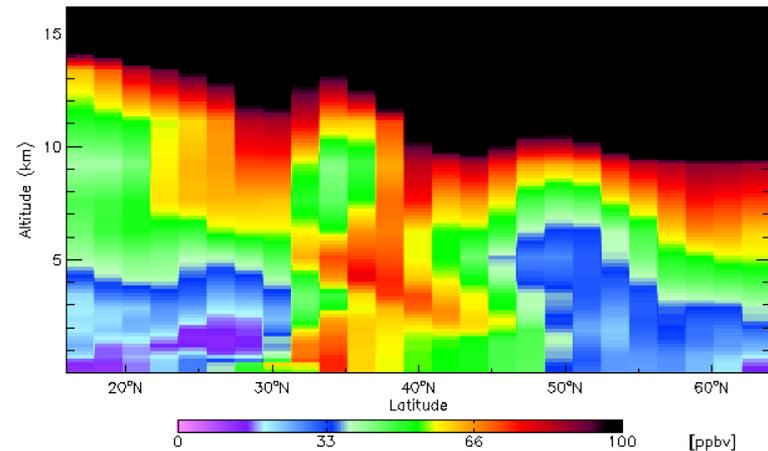
Model ozone analyses “converge” with a reduction in the mean difference from 10.7 ppbv to 2.4 ppbv.

# Stratosphere-Troposphere Exchange

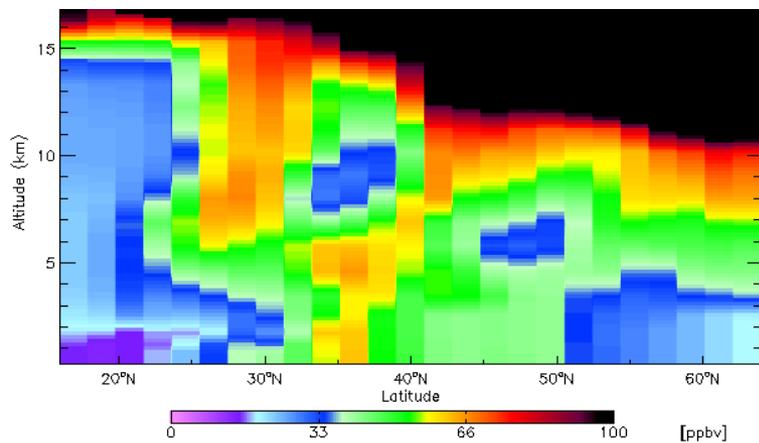
AM2-Chem ozone



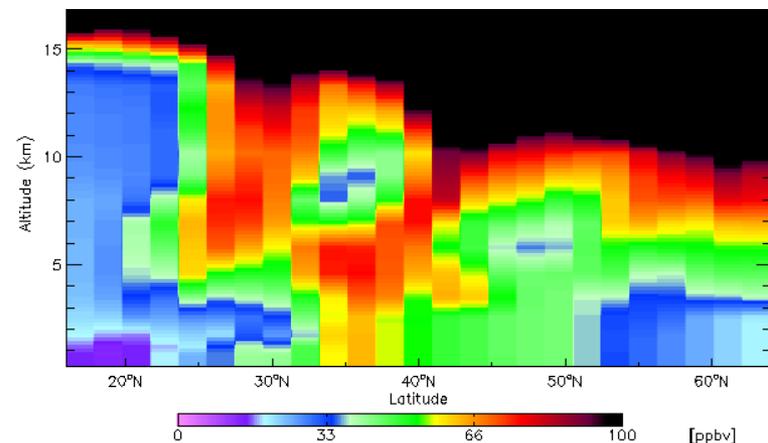
AM2-Chem ozone analysis



GEOS-Chem ozone



GEOS-Chem ozone analysis



Altitude-latitude cross-sections at 120°W, 21:00 GMT.

Stratospheric intrusion at 40N more pronounced in both models following assimilation.

# Partitioning of $NO_x$

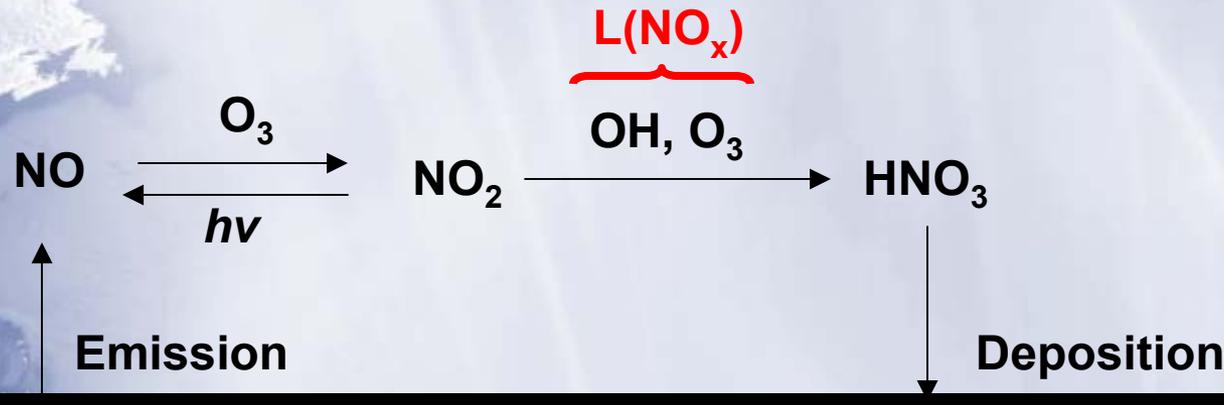
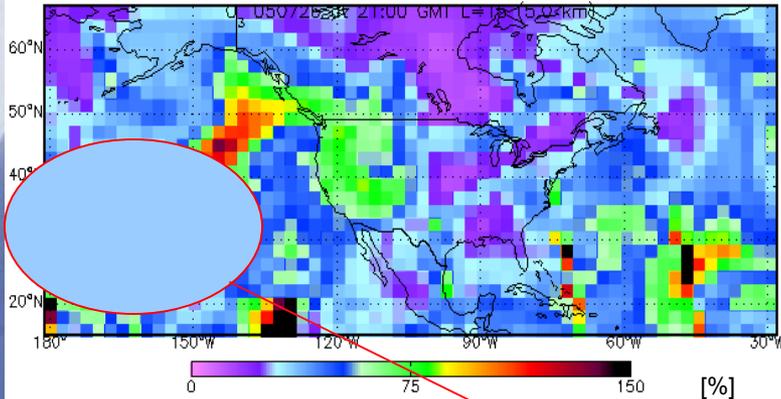


Figure adapted from Jacob [1999]

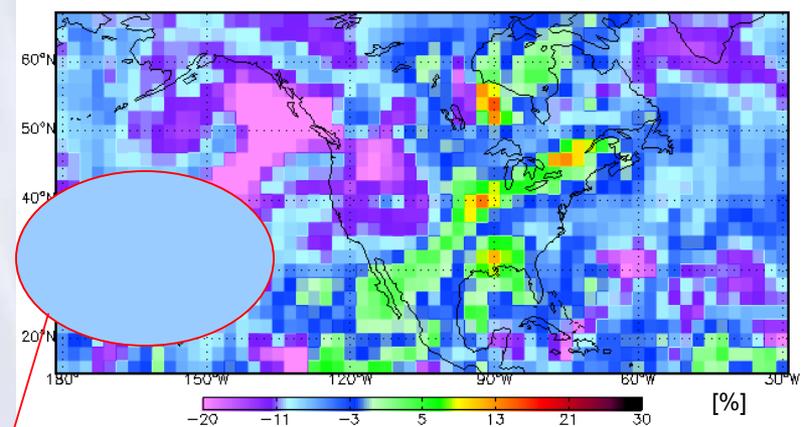
- ✧ The  $NO_x$  cycle in the troposphere implies that the balance of  $NO:NO_2$  is sensitive to the ozone abundance.
- ✧ Assimilation of TES data significantly increases the ozone tracer field in the models.
- ✧ Increased ozone in the model will decrease the  $NO_x$  lifetime.

# Impact of assimilation on $\text{NO}_x$ chemistry

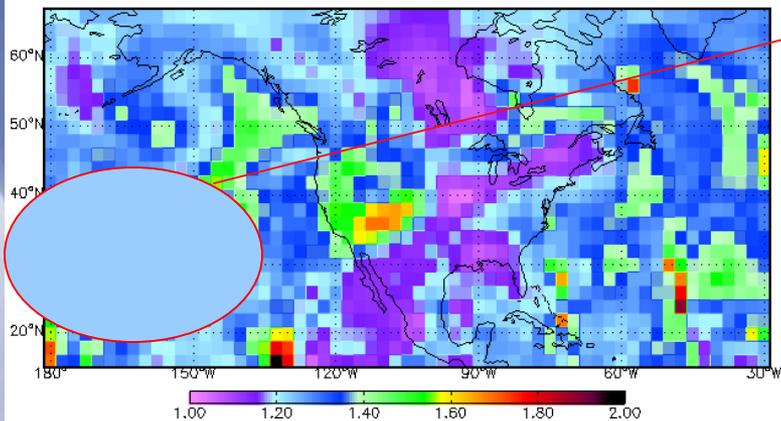
Percent change in AM2-Chem  $\text{O}_3$



Percent change in AM2-Chem  $\text{NO}_x$



$[\text{NO}:\text{NO}_2]_{\text{noassim}}/[\text{NO}:\text{NO}_2]_{\text{assim}}$



- ✧ Increases in ozone lead to a shift in the partitioning of  $\text{NO}:\text{NO}_2$  and an increase in  $\text{NO}_2$  reflected by high values in the ratio of  $[\text{NO}:\text{NO}_2]_{\text{noassim}}/[\text{NO}:\text{NO}_2]_{\text{assim}}$
- ✧  $\text{NO}_2$  increase leads to a decrease in the lifetime of  $\text{NO}_x$  and subsequent decrease in  $\text{NO}_x$  abundance.

$z = 5\text{km}$ , Time = 21:00 GMT

# *Conclusions and outlook*

## ✧ Tropospheric ozone

- ✧ The AM2-Chem and GEOS-Chem models show significant differences in tropospheric ozone abundances. Assimilation of TES data brings the ozone abundances closer together in both models.
- ✧ Dynamical features (e.g. stratospheric intrusion) are better represented in the ozone distribution in both models.

## ✧ NO<sub>x</sub> chemistry

- ✧ Large changes in NO<sub>x</sub> chemistry correspond to changes in the ozone distribution.
- ✧ Assimilation of NO<sub>x</sub> observations (OMI) will further constrain model chemistry.
- ✧ Chemical data assimilation has the potential to provide a powerful tool for understanding the processes controlling chemistry in the troposphere.