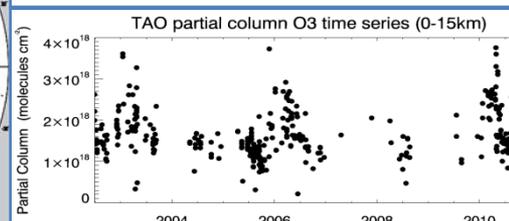
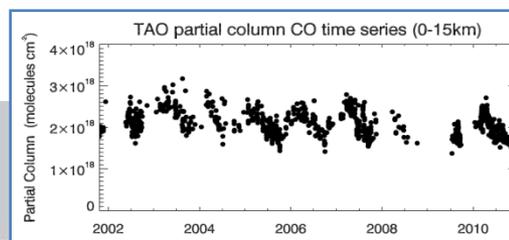
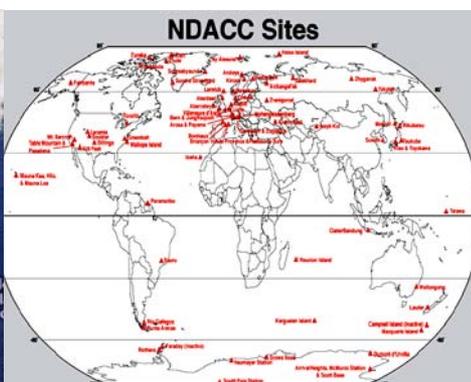
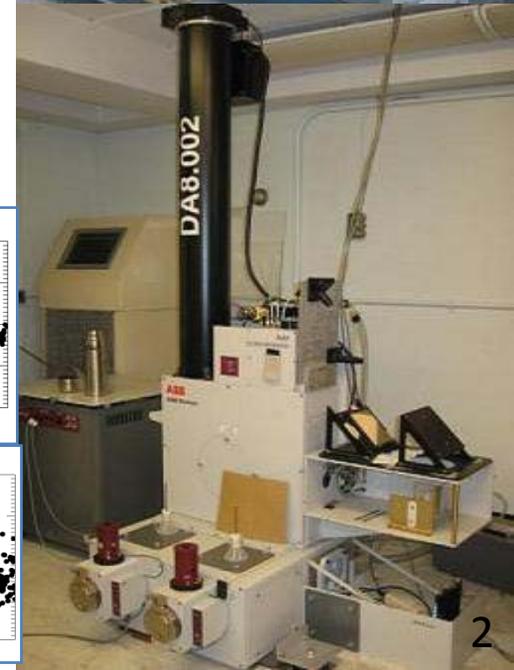
An aerial photograph of the Toronto skyline, featuring the CN Tower on the left and various skyscrapers in the background. The text is overlaid on the image.

Analysis of CO, O₃ and HCN using GEOS-Chem and ground-based FTIR observations from the Toronto Atmospheric Observatory

Cyndi Whaley, Dylan Jones, Thomas Walker, and Kimberly Strong
University of Toronto, Dept. of Physics
IGC5 2-5 May 2011

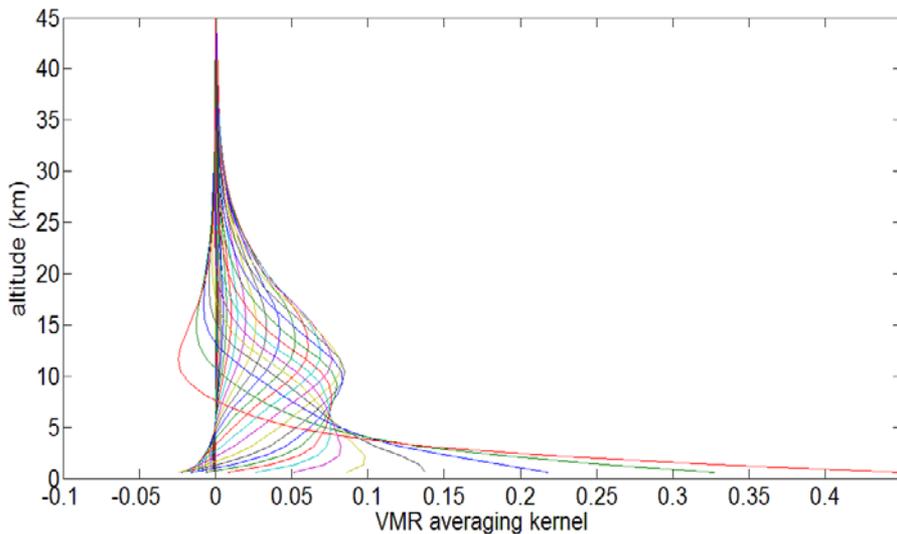
Toronto Atmospheric Observatory (TAO)

- **Location:** 43.66° N, 79.40° W, 174 m
- **Operational since 2002 → 9 years of data**
- **Primary instrument:** Bomem DA8 FTIR with solar tracker
- TAO is an NDACC (Network for Detection of Atmospheric Composition Change) site
- We use the Optimal Estimation Method for retrievals
 - Implemented with SFIT2 v3.94
 - Using HITRAN 2008 spectral line database
 - Employing harmonized retrieval parameters recommended by the NDACC Infrared Working Group
- GEOS-Chem profiles are smoothed with TAO averaging kernels before comparisons are made.



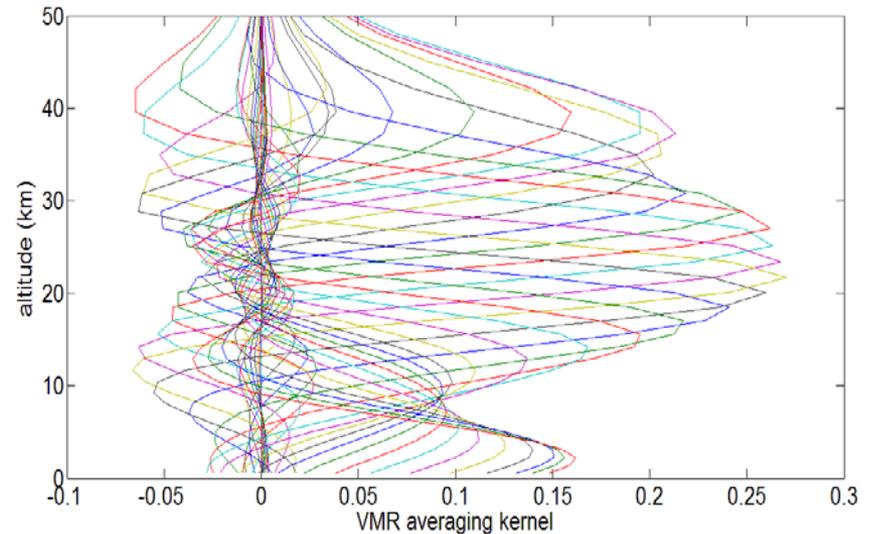
TAO Tropospheric Sensitivity: Typical Averaging Kernels

- Good sensitivity to the lower tropospheric signal



CO

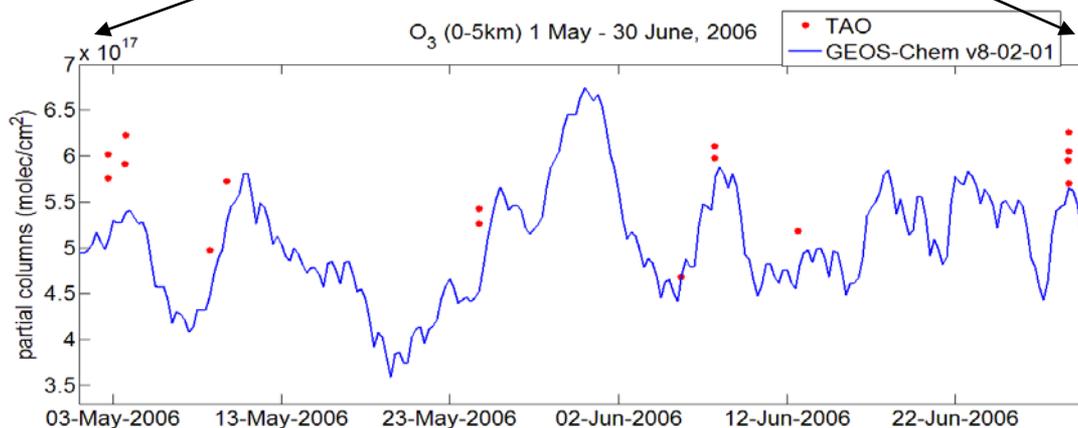
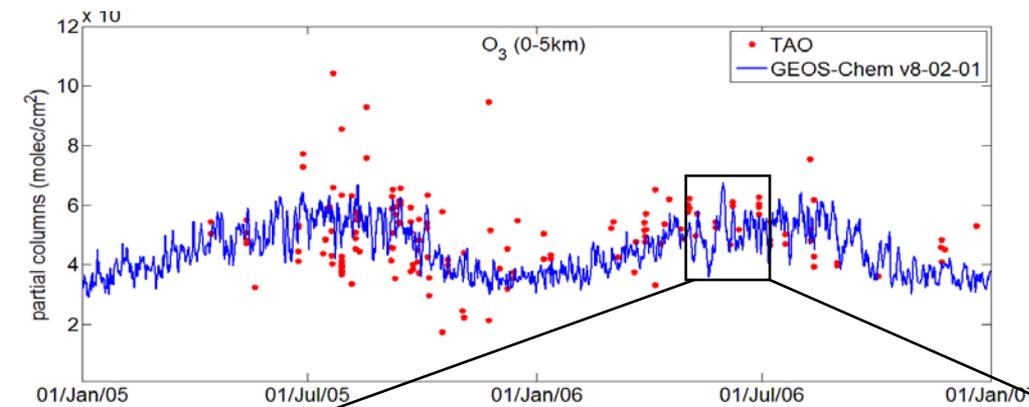
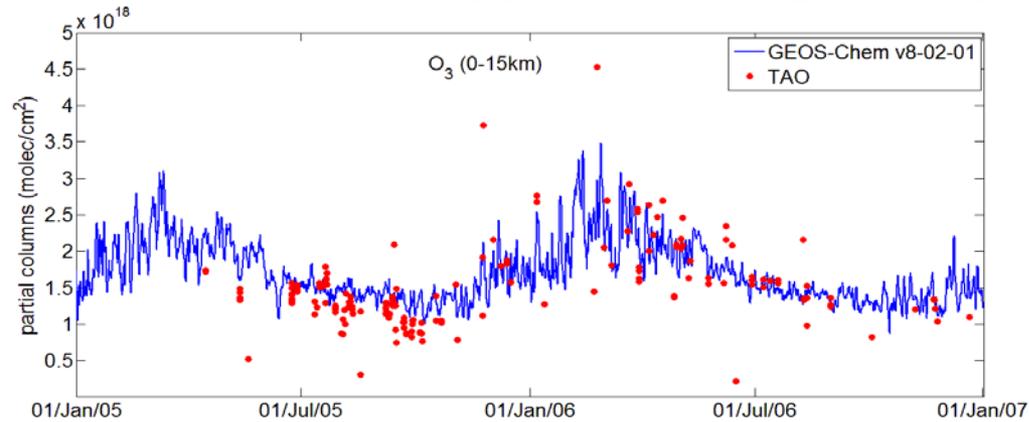
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0-15km
DOFS=1 for
0-2.3km
(10-09-2009)



O₃

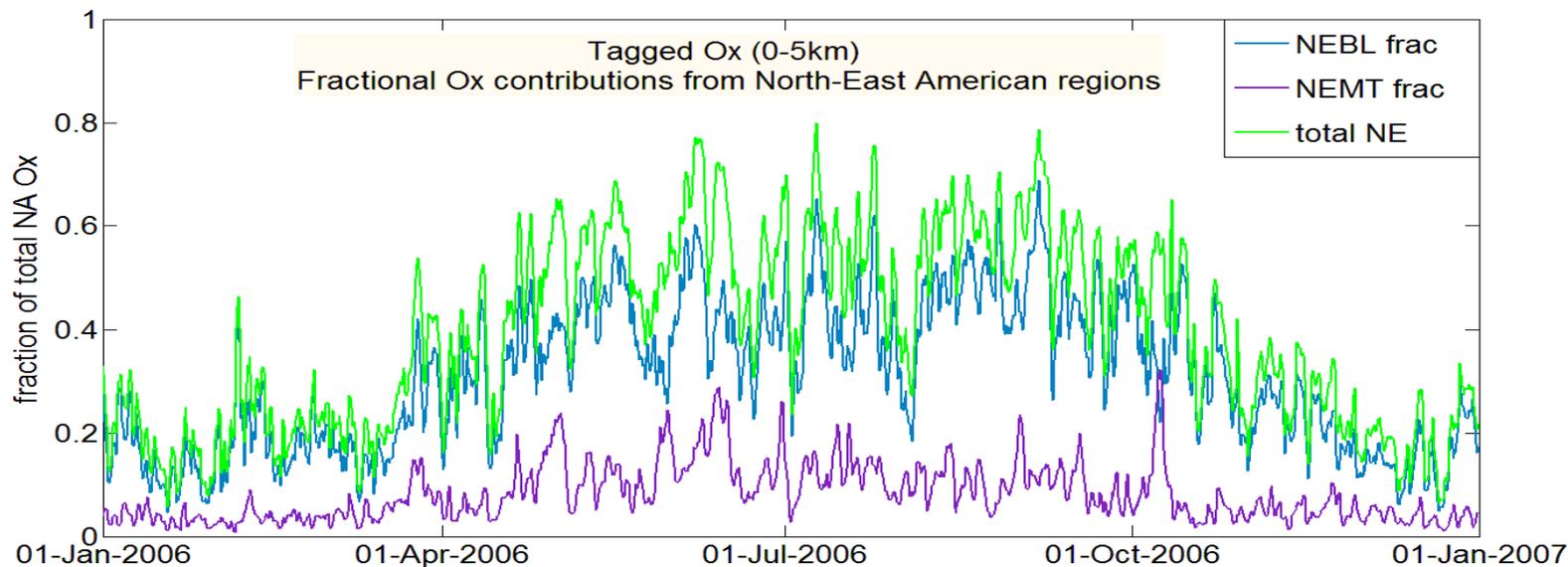
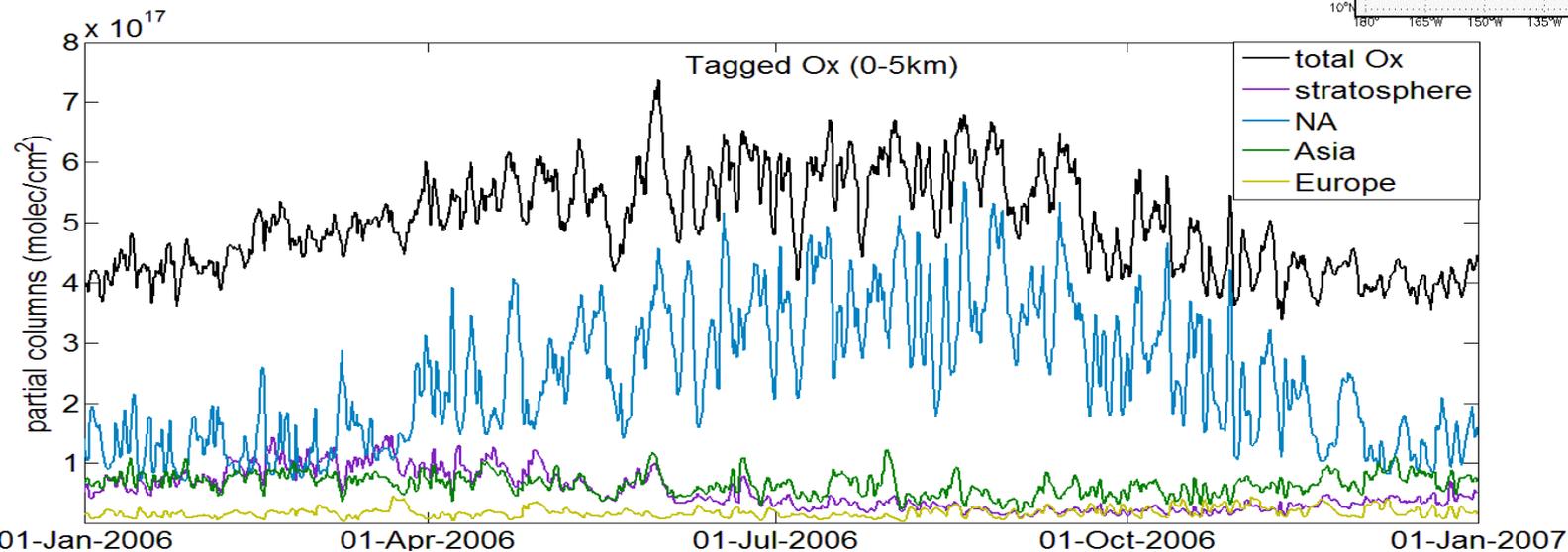
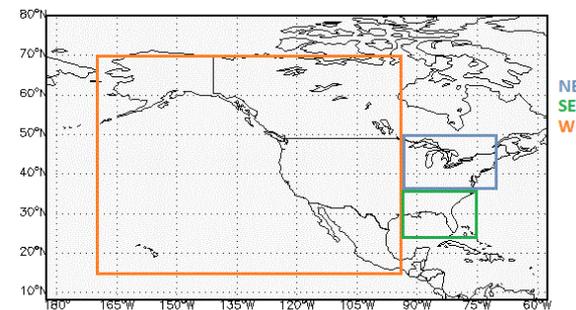
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0-15km
DOFS=1 for
0-7 km
(11-05-2007)

TAO and GEOS-Chem Ozone



- The O_3 simulation agrees well with TAO
- Different partial columns show different seasonal cycle \rightarrow lower tropospheric columns capture summertime O_3 production

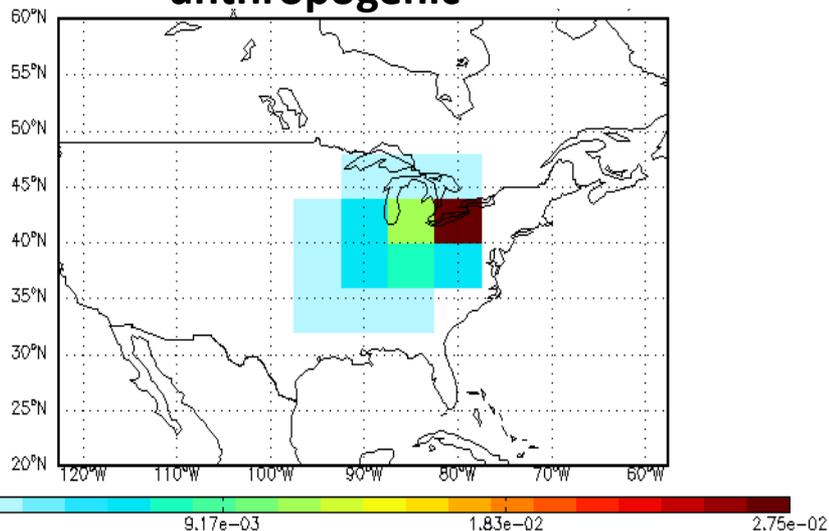
GEOS-Chem Tagged Ox Sources



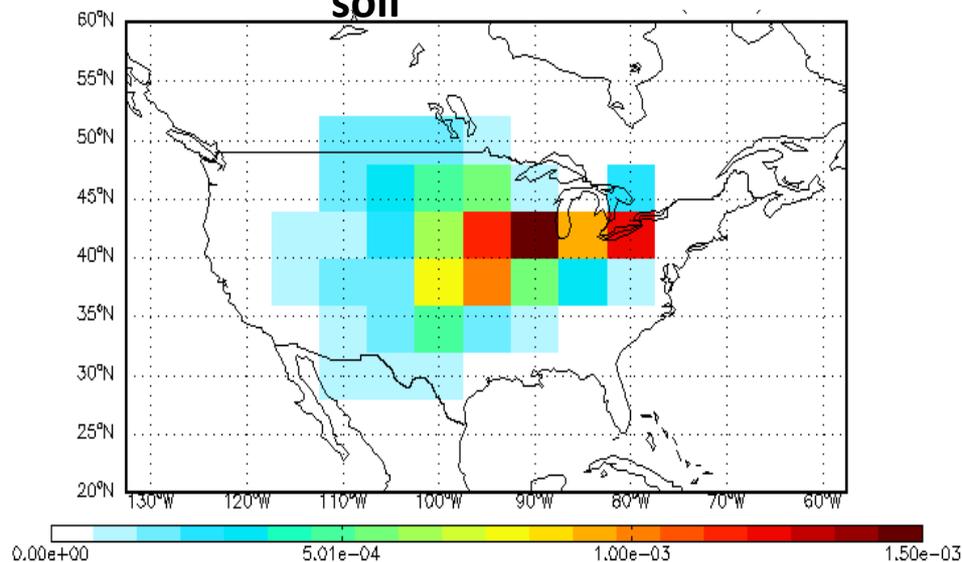
Adjoint Sensitivity Maps of O₃ Above Toronto to NO_x Emissions in July 2006

- Capturing mostly the local anthropogenic emissions
- Also sensitive to soil emissions from American agriculture, and biomass burning

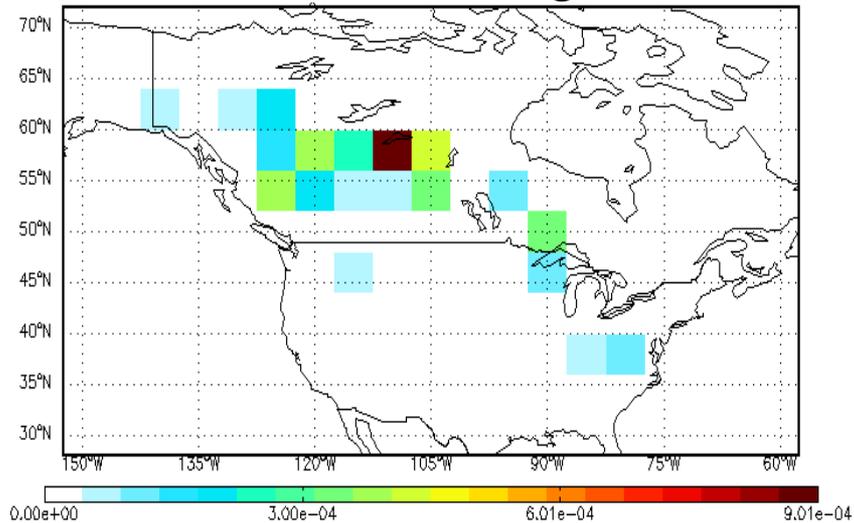
anthropogenic



soil

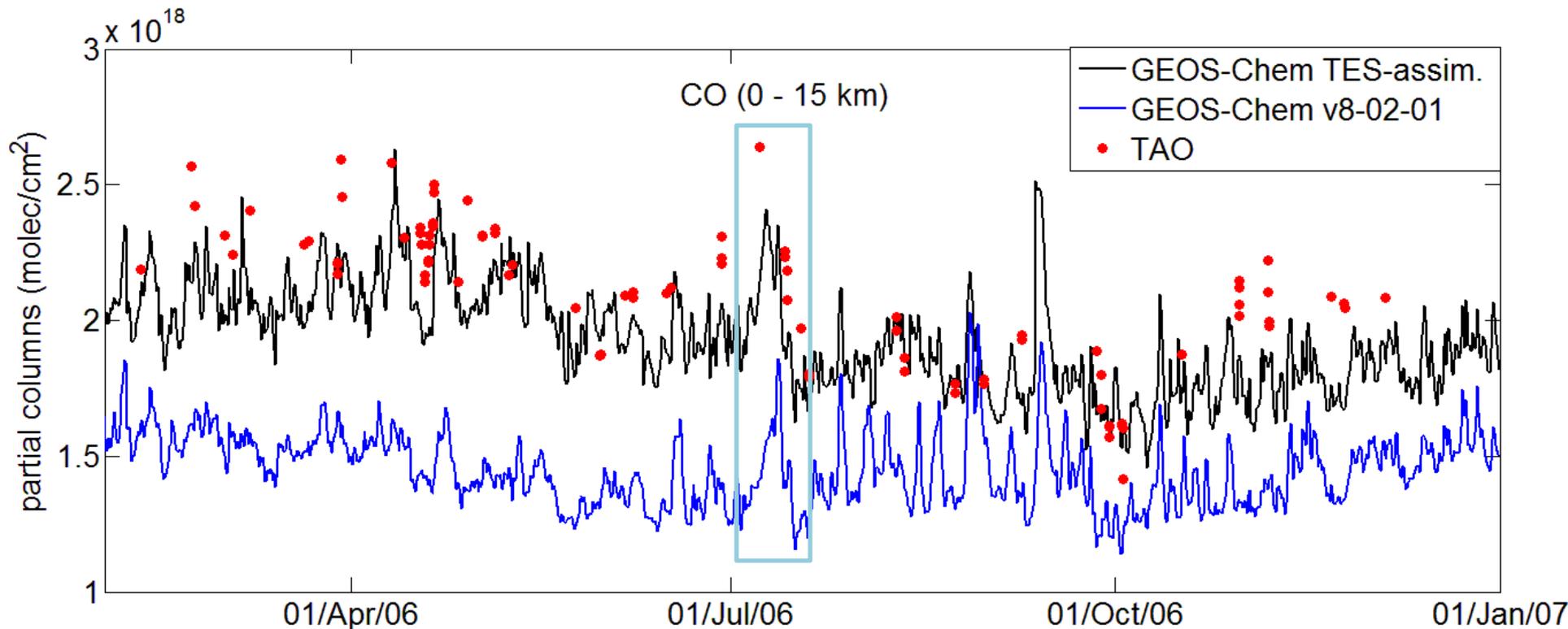


biomass burning



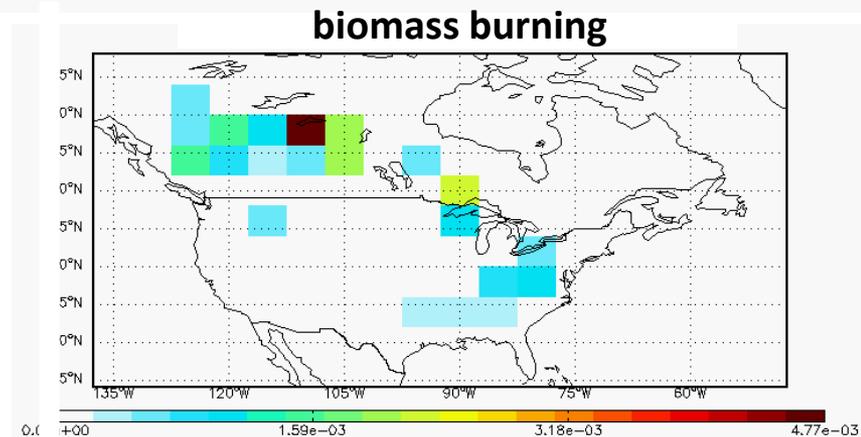
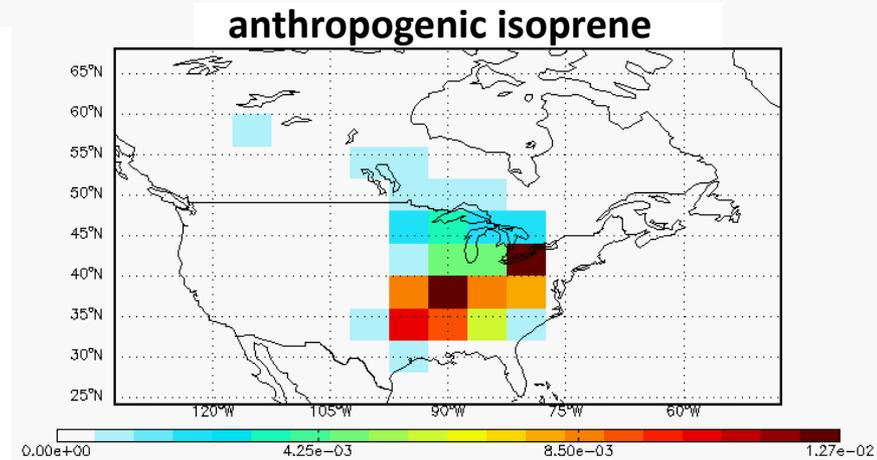
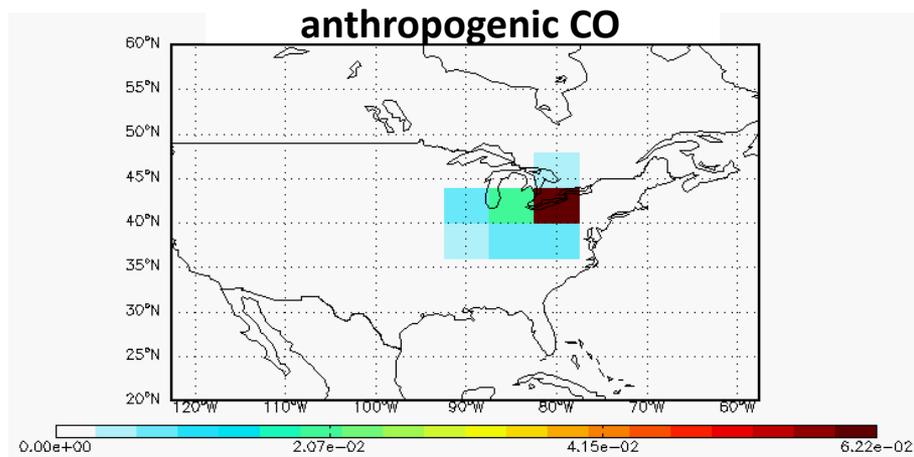
TAO and GEOS-Chem CO

- Despite the good agreement in O_3 , GEOS-Chem CO does not agree well with TAO
 - Kopacz et al. (2009) suggested that GEOS-Chem had a low CO bias due to an underestimate of fossil fuel emissions in Northern mid-latitudes
 - When TES observations are assimilated into the model, there is better agreement with TAO → clearly a problem with the emission inventories in the model



Adjoint Sensitivity Maps of CO Above Toronto to Emissions in July 2006

- Adjoint and tagged runs provide maps of CO from VOCs, fossil fuels and biomass burning
- Relative importance of various sources differs for CO and O₃



Summary and Conclusion



- TAO currently has 9 years of tropospheric trace gas measurements, revealing seasonal and short-term variabilities
- Short-term variations in CO and O₃ have been analysed with GEOS-Chem and sources of pollution affecting air quality in Toronto have been identified
 - Local anthropogenic emissions are a dominant source for O₃, and anthropogenic *and* biogenic sources are important for CO
- Future Work:
 - Expand GEOS-Chem analysis to include more species (C₂H₆, HCN, CH₂O), adjoint for CO, and higher resolution (nested grid) simulations
 - Quantify long-term trends as the TAO dataset increases in time
 - Preliminary investigation of HCN (not shown) reveals that the model is biased low. Work needed to assess the model situation.