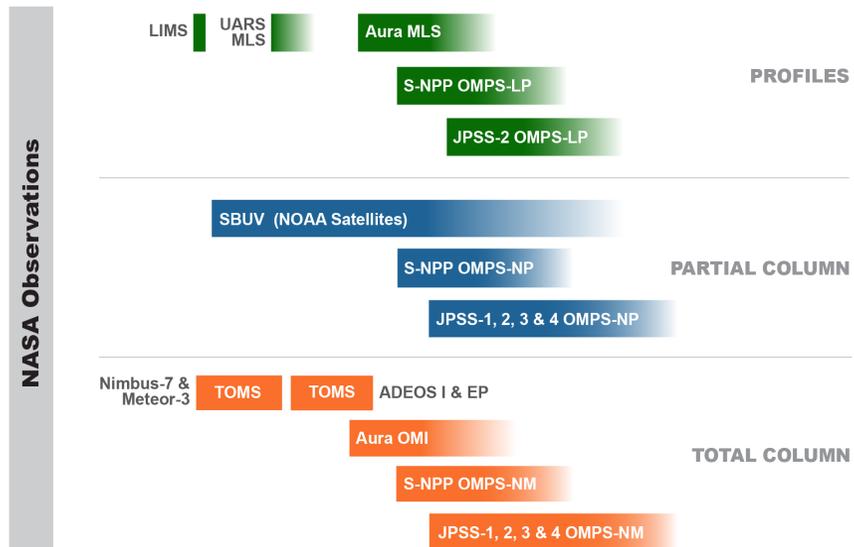


1960 2000 2040 2080



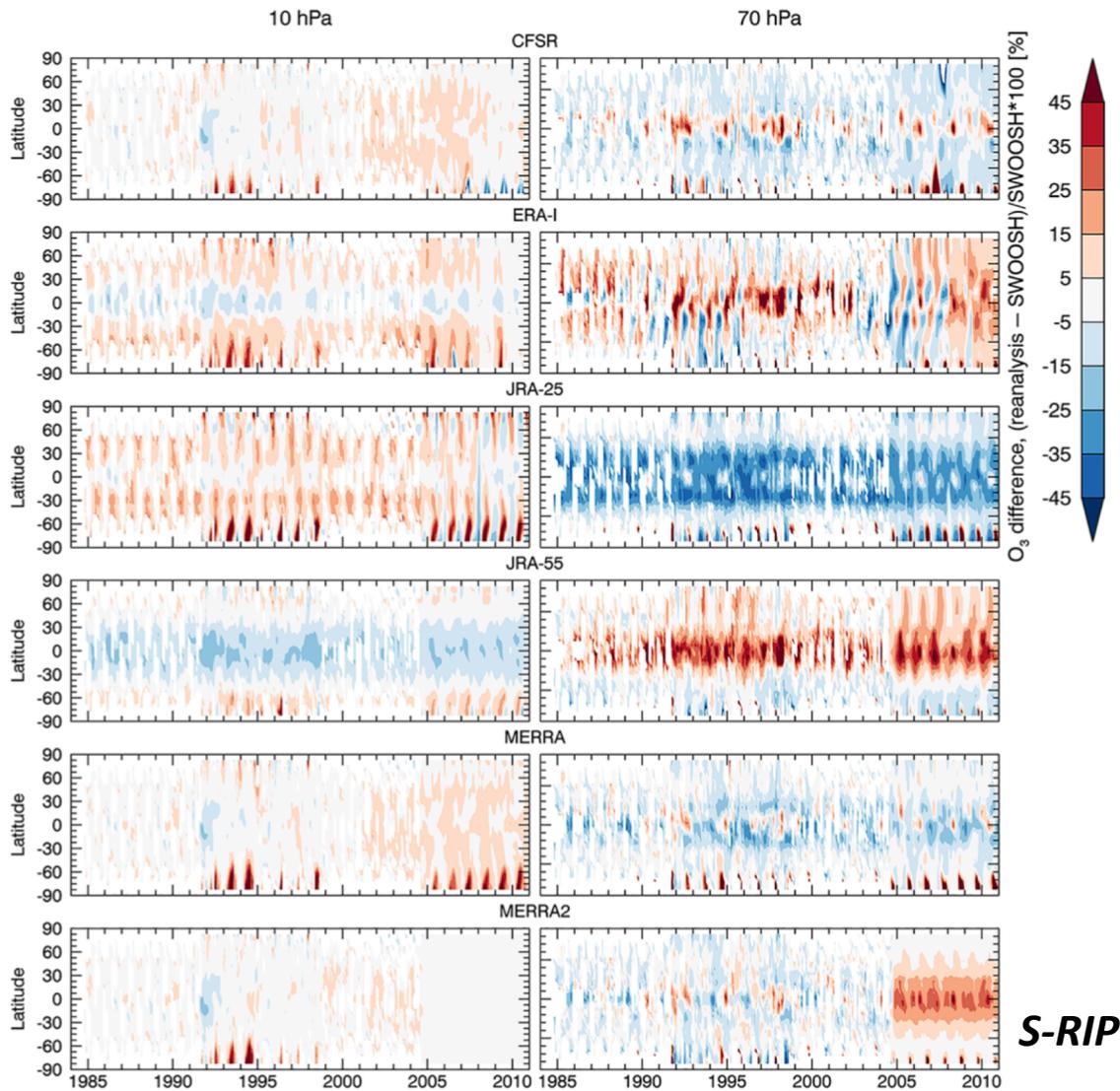
Tackling the challenges in the representation of trends and long-term variability in future stratospheric chemical reanalyses

K. Wargan & collaborators

Outline

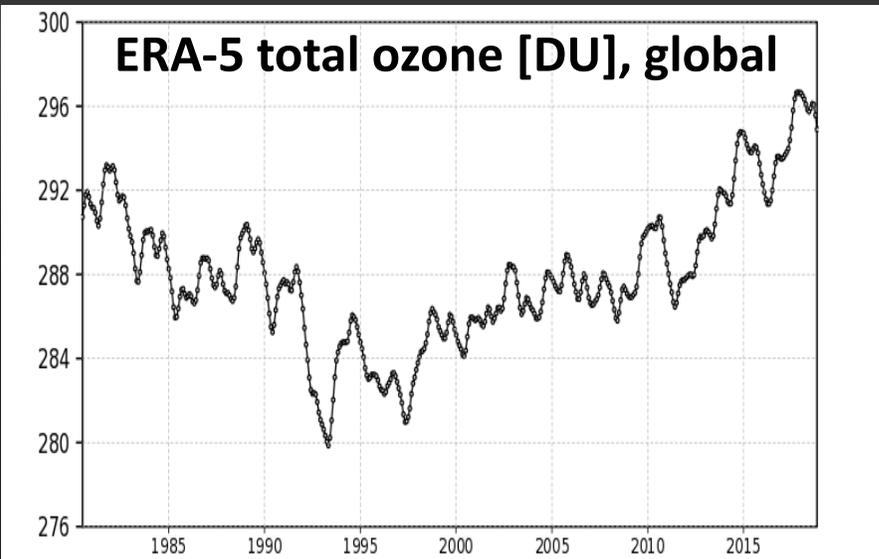
- Continuity of ozone in reanalyses for trend studies: addressing changing observing systems and representation of transport
- Stratospheric chemical reanalysis work at the GMAO
- Laundry list of thoughts and questions about the theme

SPARC-DAWG Workshop, 2019, Theme 1: chemical reanalysis



Discontinuities and drifts

Differences between six major reanalyses and the SWOOSH.
 Step changes in observations have consequences for studies of the long-term ozone variability

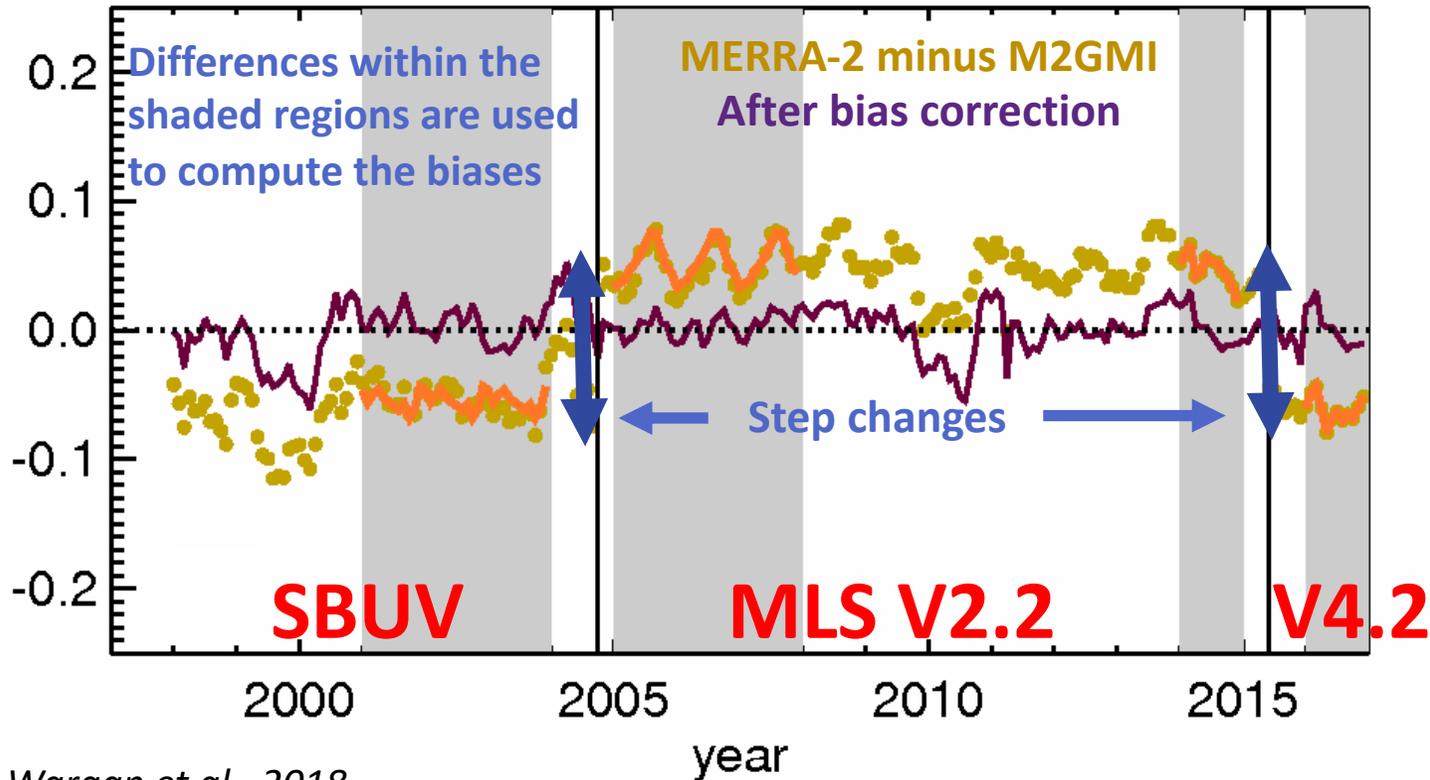


The uptick in the past decade not supported by observations

Correction of discontinuities

MERRA-2 – M2-GMI difference

70 hPa, tropics

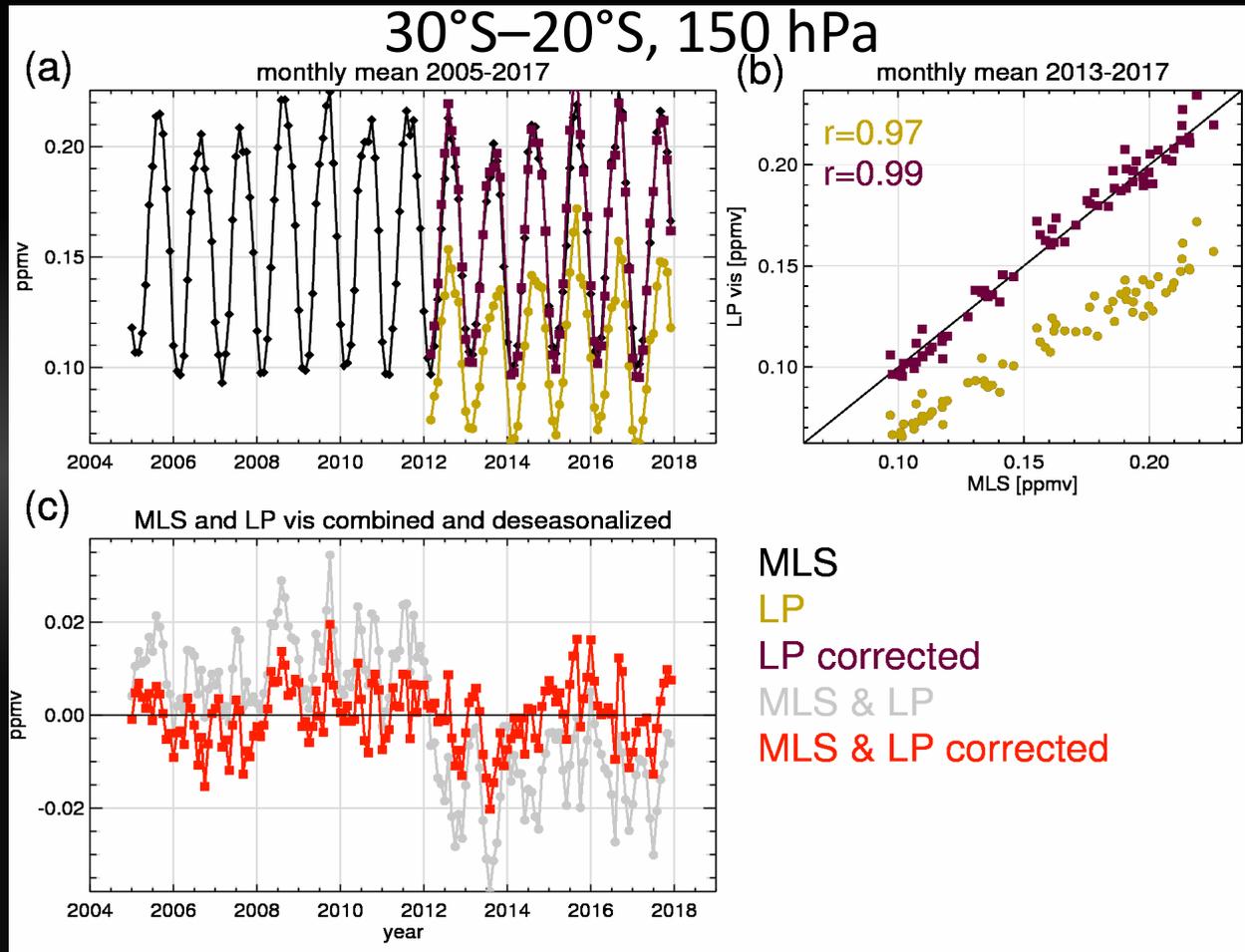


Wargan et al., 2018

In this approach we use a chemistry model simulation driven by assimilated meteorology (M2GMI) as a transfer standard to correct step changes in MERRA-2 ozone that arise from changes in the observing system.

It would be preferable to have a reanalysis output that is already discontinuity-free.

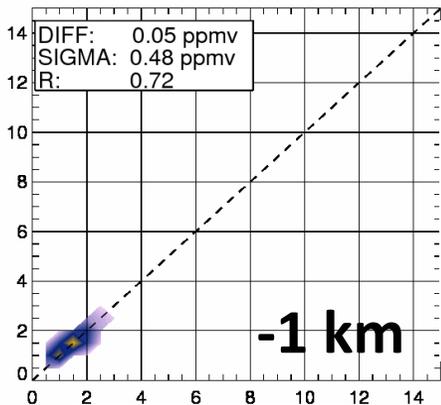
Homogenization of MLS and OMPS-LP data for reanalyses



- OMPS Limb Profiler on Suomi NPP (2012–present) and planned for JPSS-2, 3, & 4 well into the 2030s
- Switching from MLS to OMPS-LP (or to assimilating both) would lead to a discontinuity
- The discontinuity is eliminated by simple homogenization
- Assimilation of the homogenized data will eliminate the discontinuity but not the drift in OMPS-LP (not shown)

Sean suggested assimilating water vapor and ozone homogenized the way it was done in SWOOSH

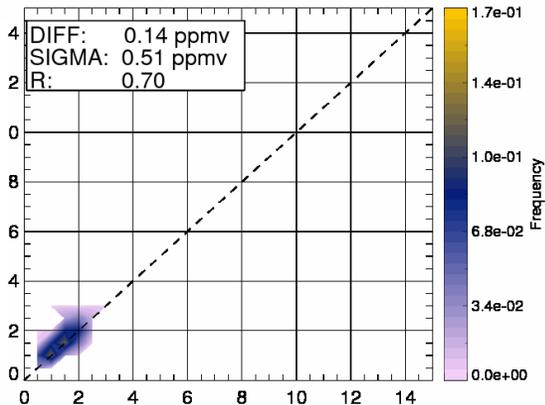
MLS analysis [mPa]



-1 km

sondes [mPa]

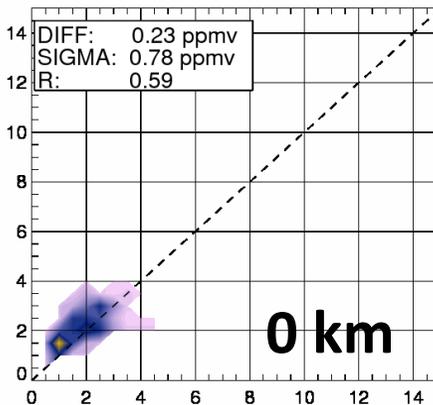
OMPS analysis



sondes [mPa]

254 sondes

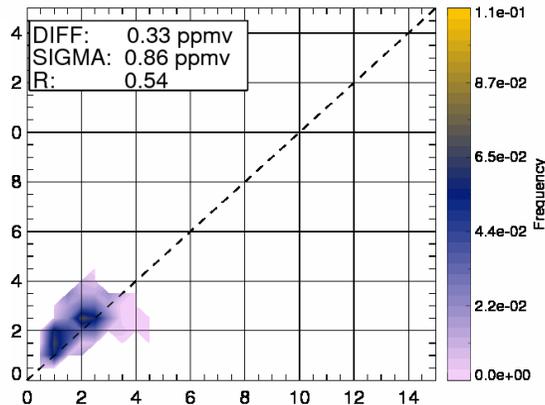
MLS analysis [mPa]



0 km

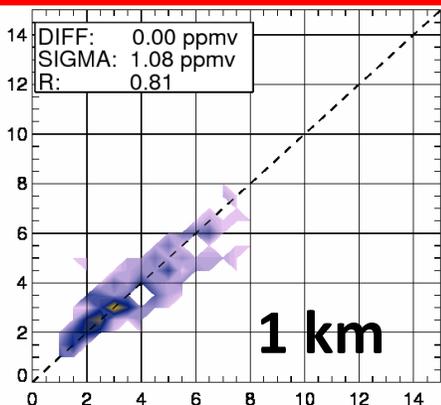
sondes [mPa]

OMPS analysis



sondes [mPa]

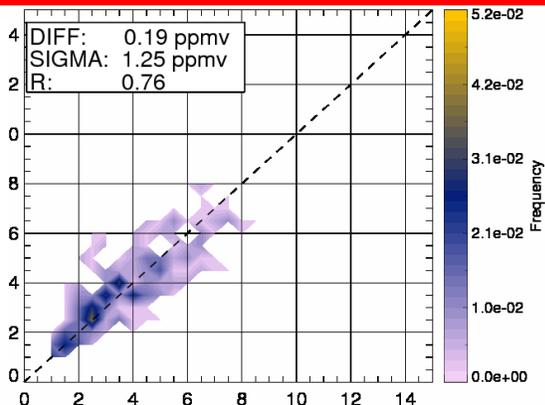
MLS analysis [mPa]



1 km

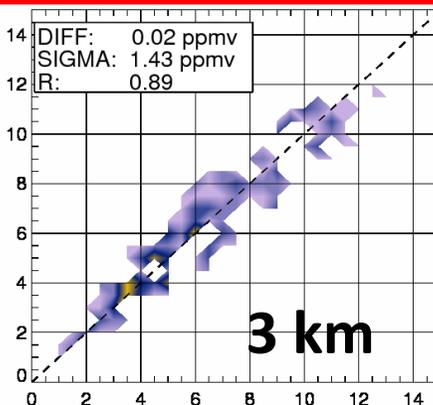
sondes [mPa]

OMPS analysis



sondes [mPa]

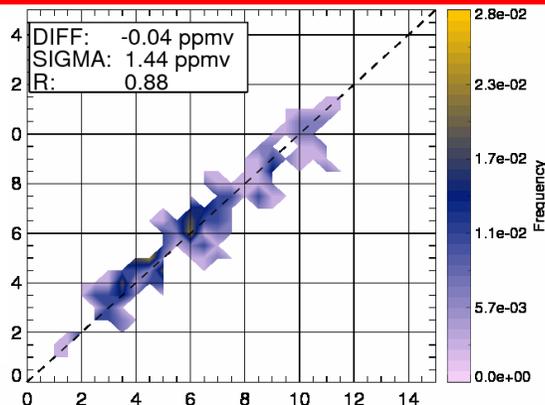
MLS analysis [mPa]



3 km

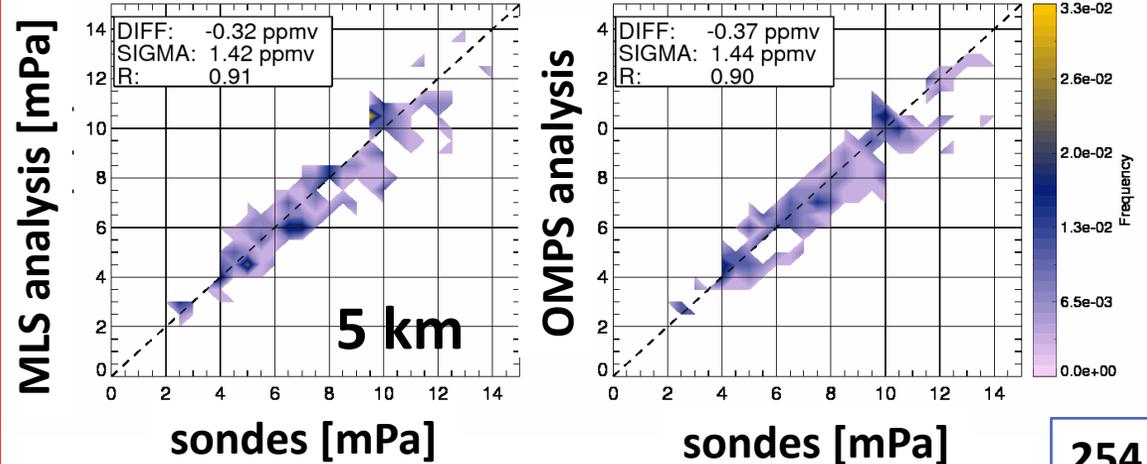
sondes [mPa]

OMPS analysis

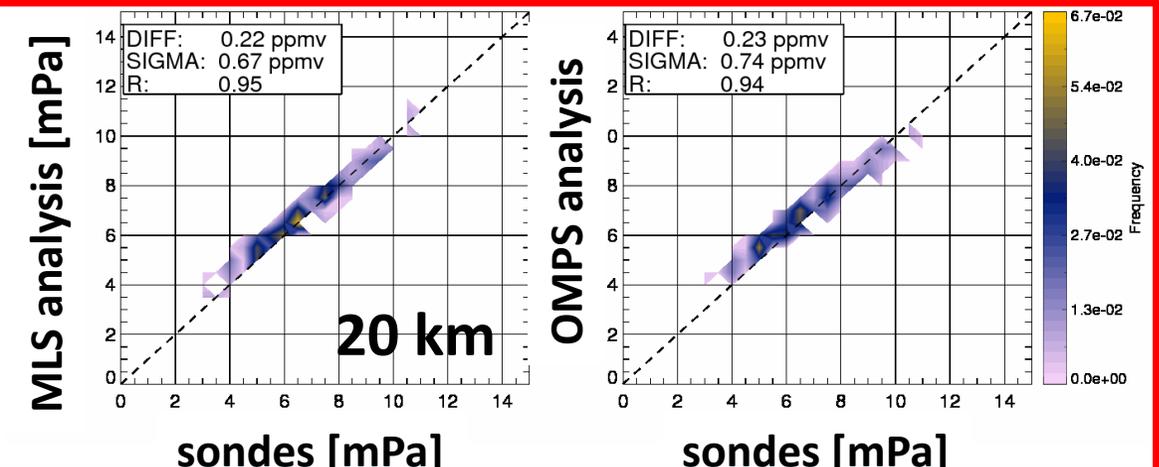
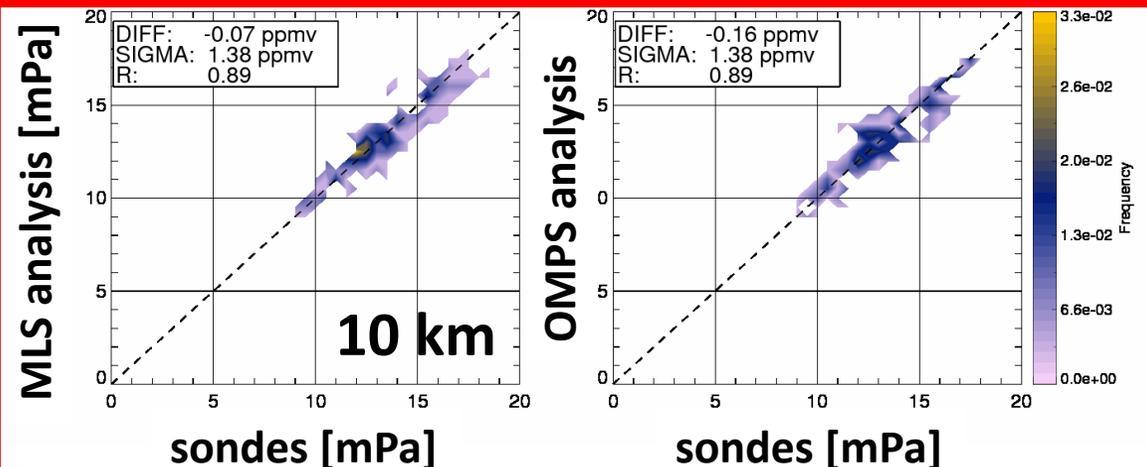
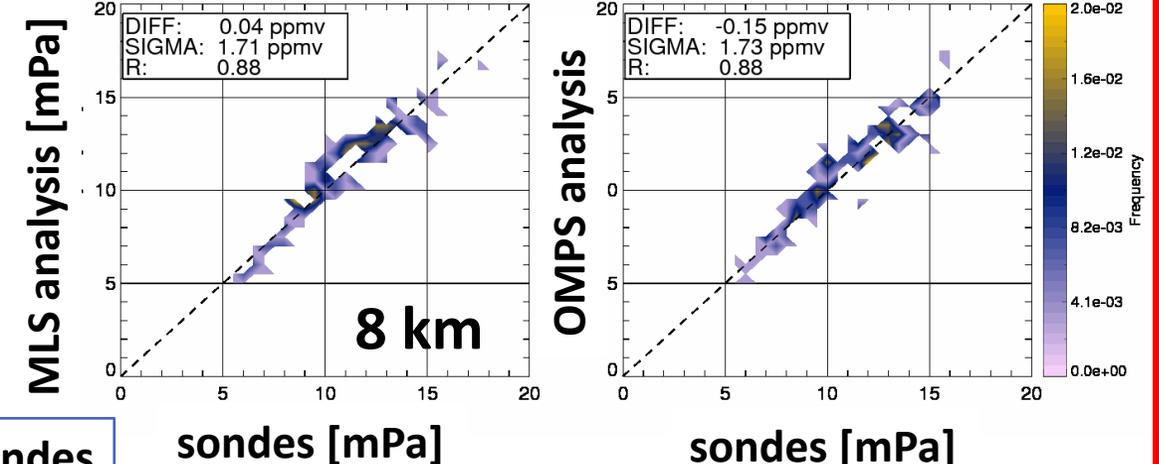


sondes [mPa]

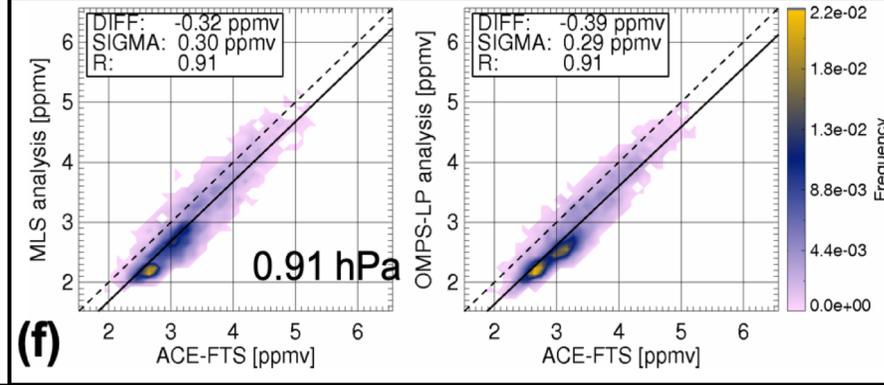
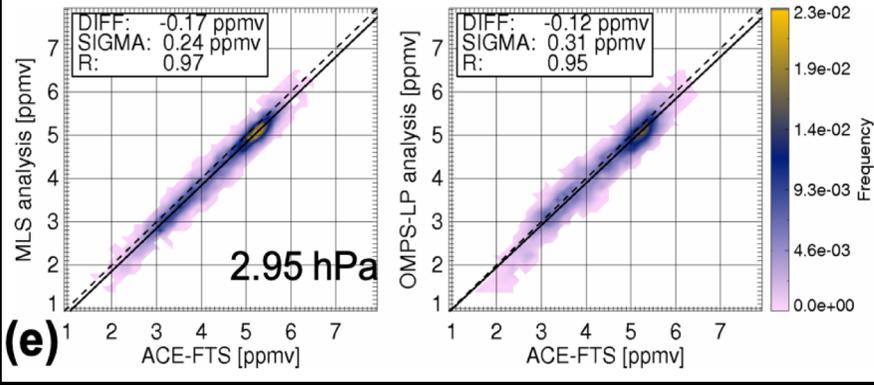
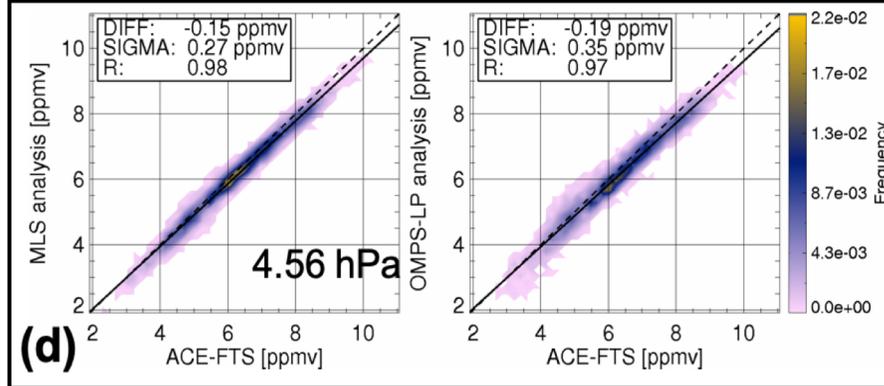
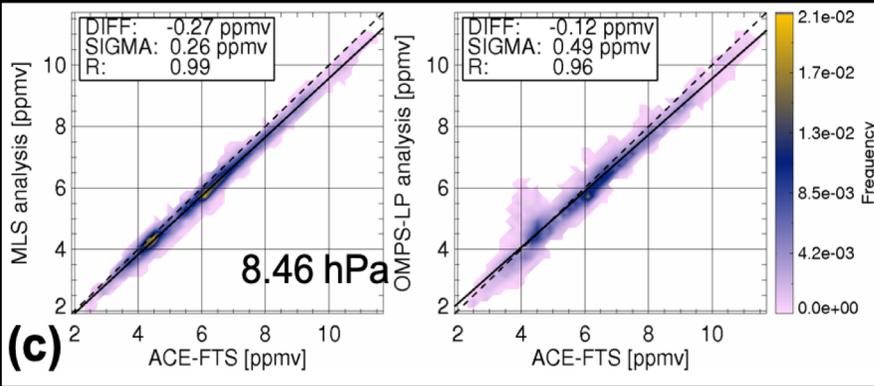
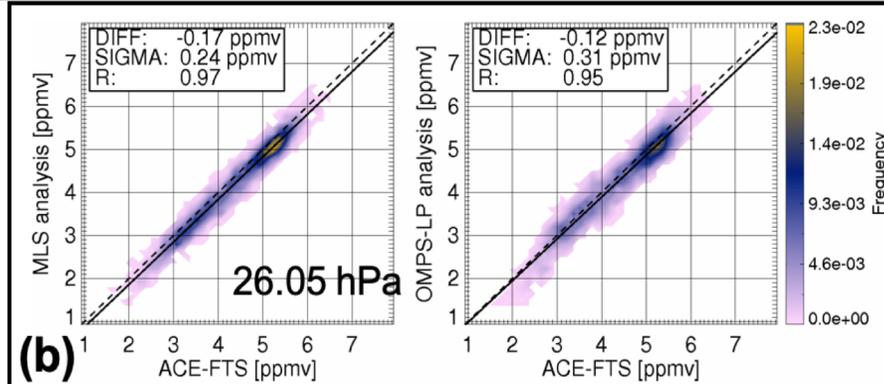
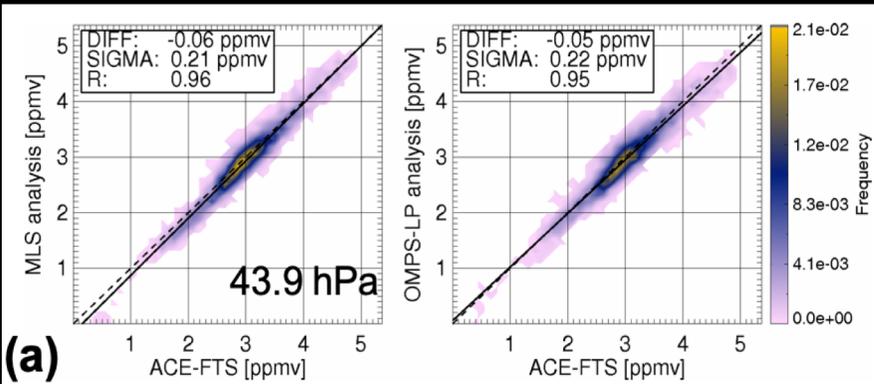
Joint probability distributions of global ozonesonde data and the two analyses at different levels relative to the tropopause. **MLS and OMPS-LP analyses exhibit very similar characteristics**



254 sondes



Joint probability distributions of global ozonesonde data and the two analyses at different levels relative to the tropopause. **MLS and OMPS-LP analyses exhibit very similar characteristics**



Joint probability distributions of global ACE-FTS ozone and the two analyses at selected pressure levels. **MLS and OMPS-LP analyses exhibit very similar characteristics**

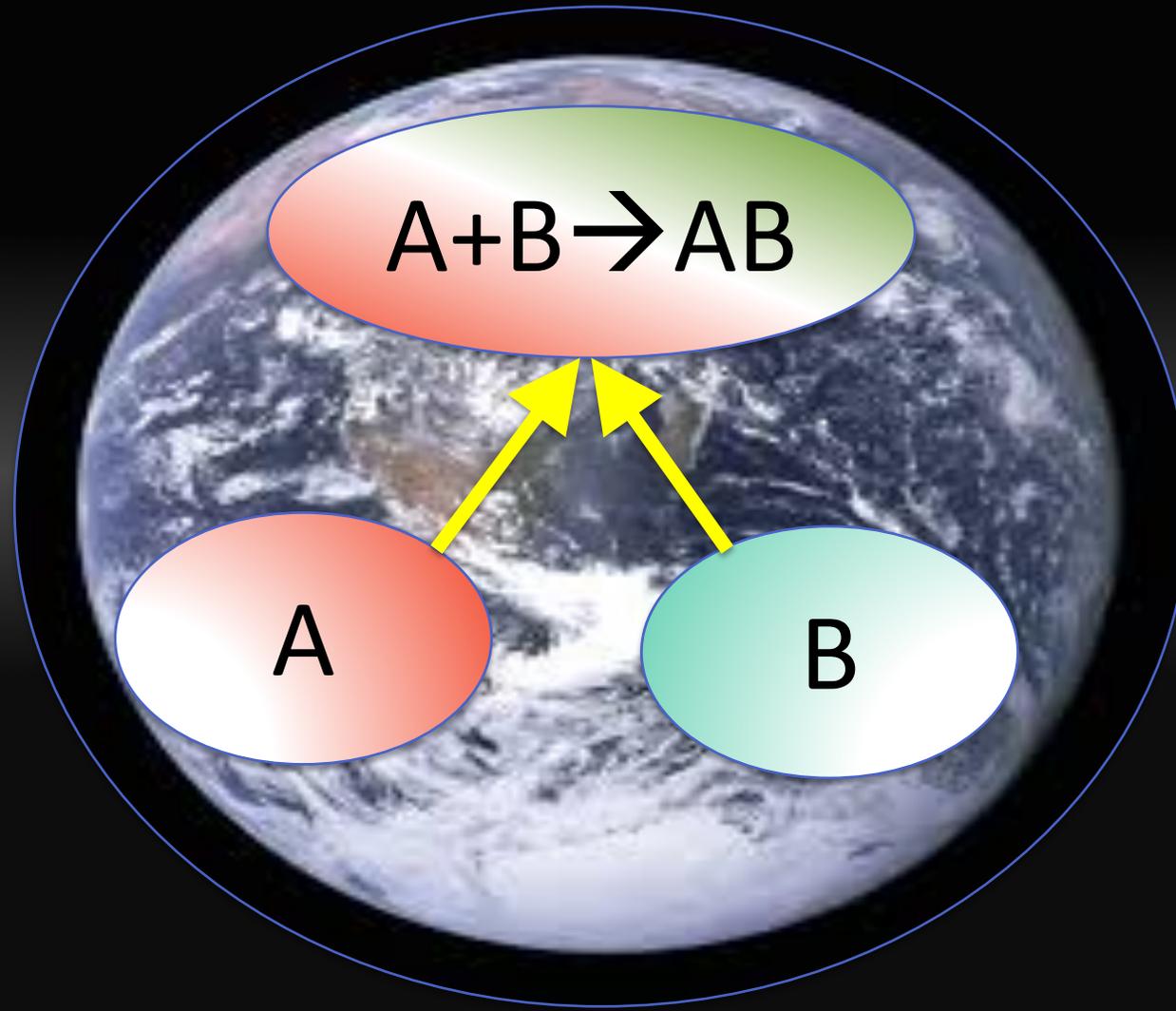
...and the analyses perform very well, within estimated ACE-FTS/MLS uncertainties

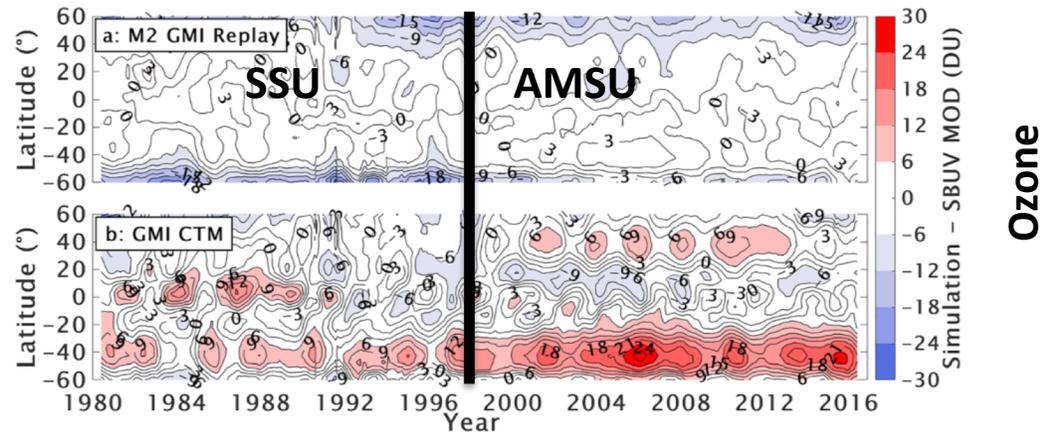


Points made so far

- **Achieving continuity of reanalysis ozone**

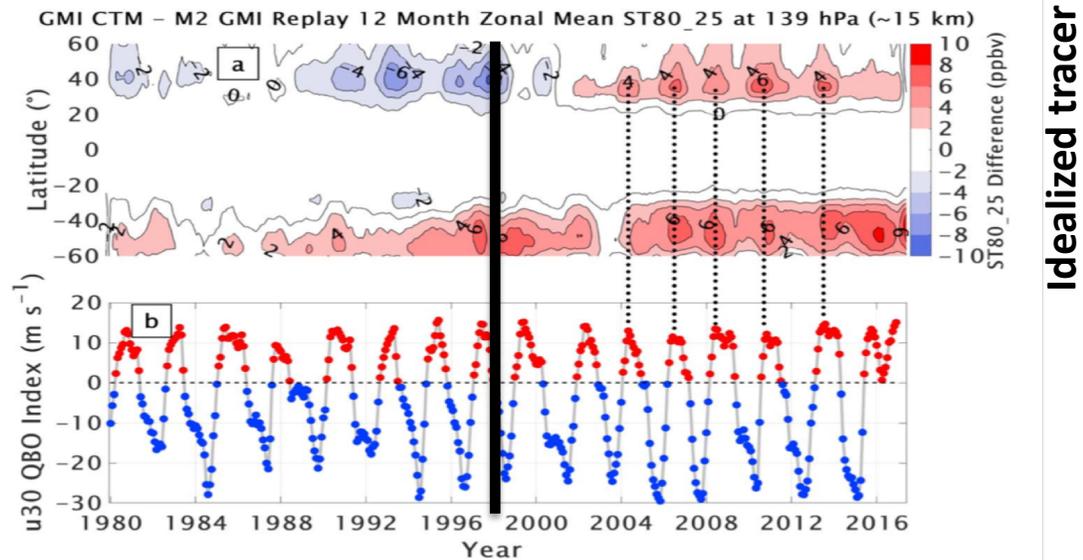
Transport





Ozone

Figure 6. Smoothed 12-month zonal means of differences between simulated and SBUV MOD total O_3 columns. (a) M2 GMI Replay. (b) GMI CTM. The black dashed line on each panel indicates the year 1998.



Idealized tracer

Two ways of doing specified dynamics:

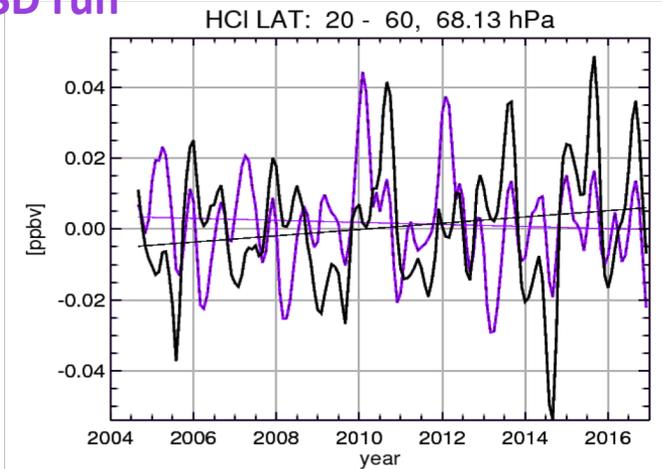
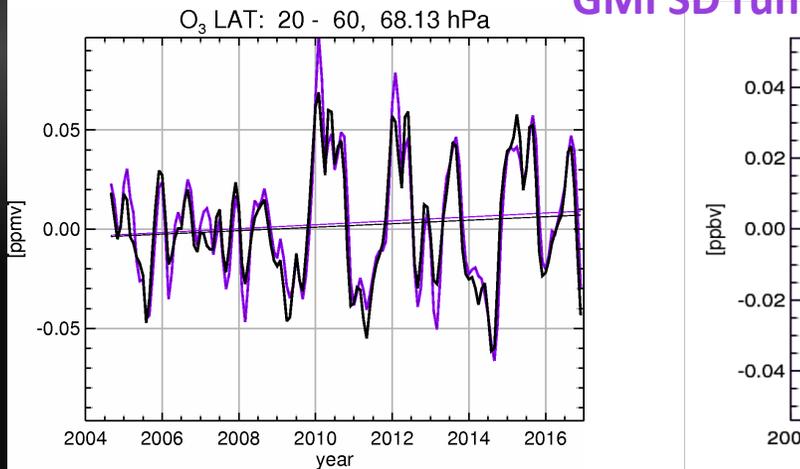
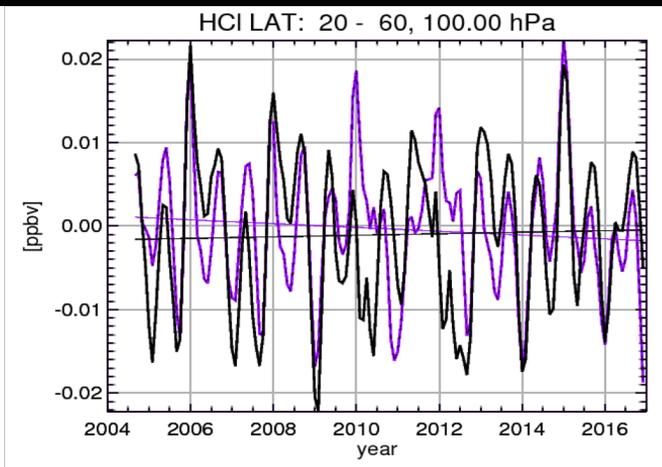
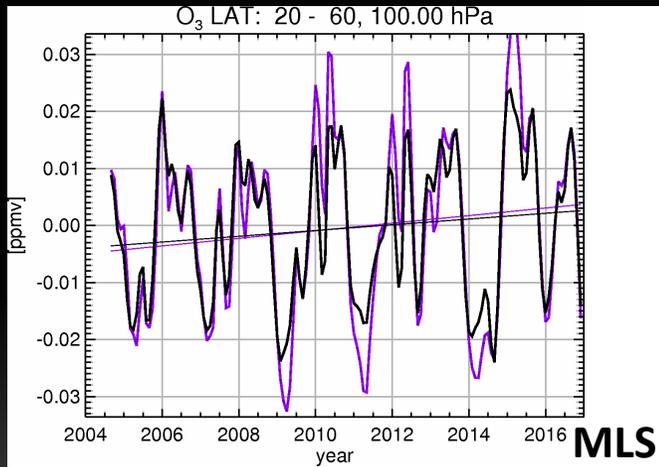
- CTM
- 'Replay'

Both driven by MERRA-2, both use versions of the GMI chemistry model; CTM has somewhat updated chemistry

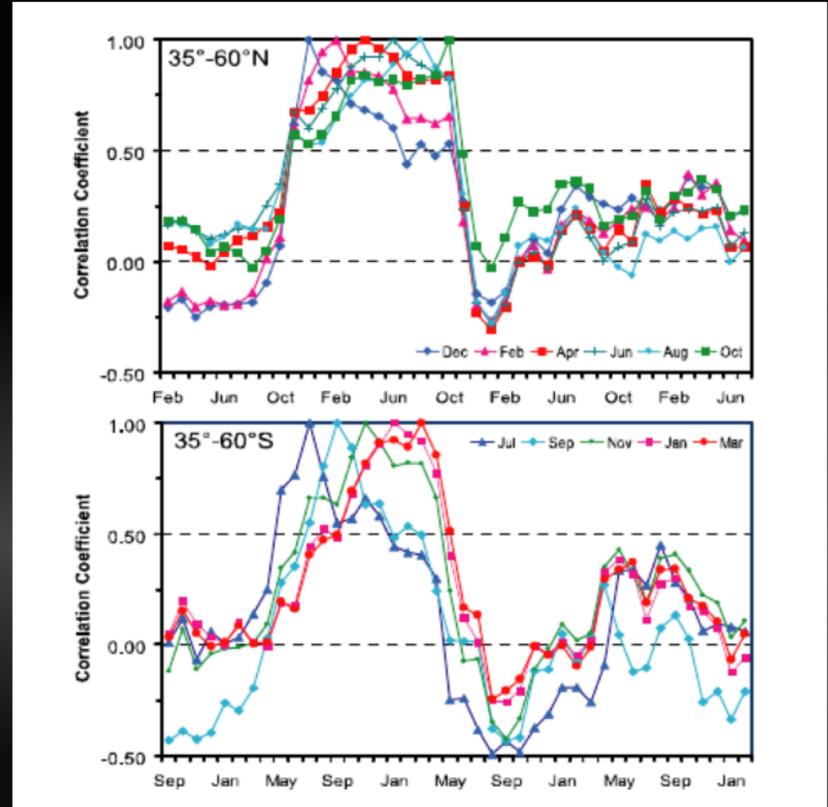
Very different response to the 1998/1999 observing system change (TOVS/ATOVS transition), apparently related to QBO-induced transport.

It matters how specified dynamics is done

Ozone QBO fit HCl



autumn ozone "resetting"



Fioletov & Shepherd 2003

Nothing like this for HCl.
There is no one such thing as transport

Good QBO fit for ozone

Bad for HCl



Points made so far

- **Achieving continuity of reanalysis ozone**
- **Importance of transport**

A GEOS chemical reanalysis of the stratosphere: work in progress

- 4-year project funded by MAP (Modeling, Analysis, and Prediction)
- A significant extension of NASA GMAO's GEOS Data Assimilation System to include assimilation of several stratospheric constituents beyond ozone
- Currently assimilating: water vapor, HNO₃, HCl from **MLS**
- Planning: N₂O and potentially ClO
- **Goal:** produce an MLS mission-long reanalysis of the stratosphere for chemistry, composition and transport studies. Note, this is similar to the BASCOE Reanalysis of Aura MLS v2 (BRAM2)
 - **Perform a high-resolution multiyear scientific analysis of polar processing during winter and spring in both hemispheres**
 - **Assess the predictability of polar stratospheric ozone and water vapor (WV) fields on short to seasonal time scales**
 - **Investigate the lower stratospheric WV budgets and WV-ozone interactions in the middle latitudes**

Data assimilation system

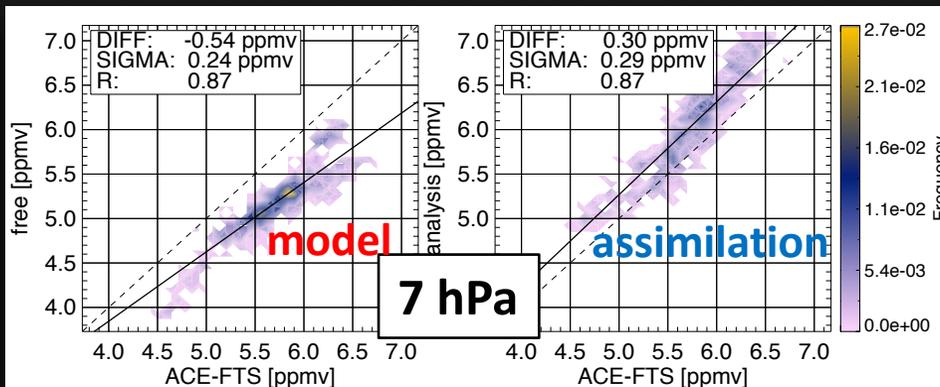
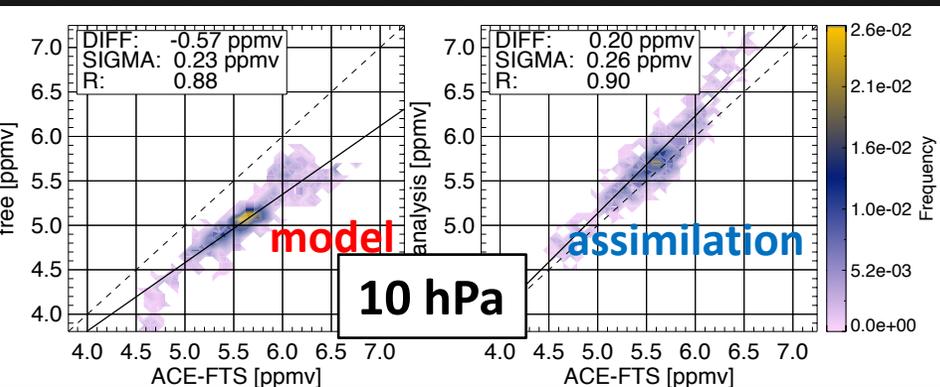
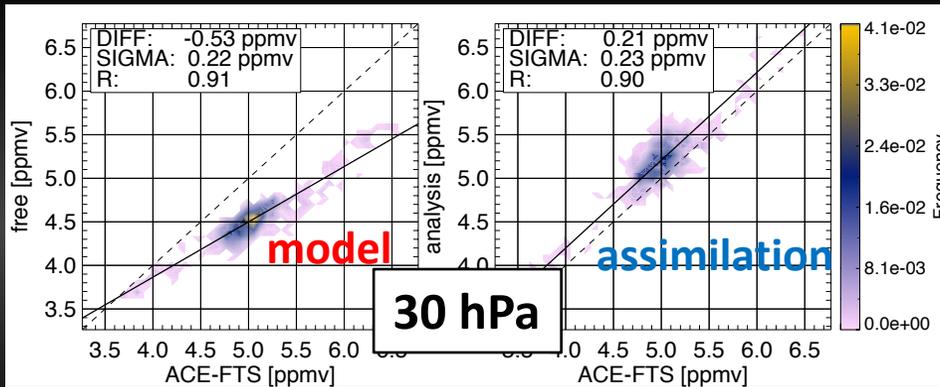
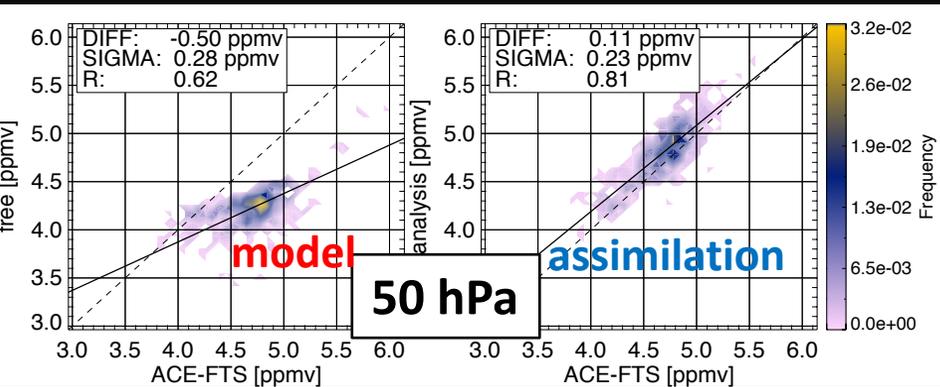
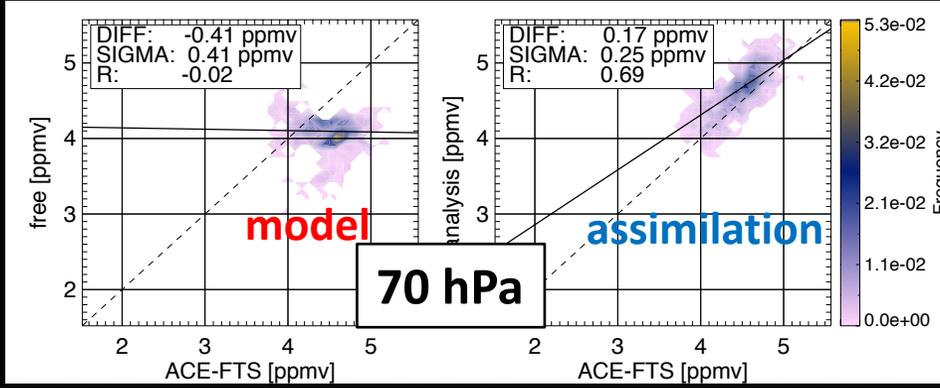
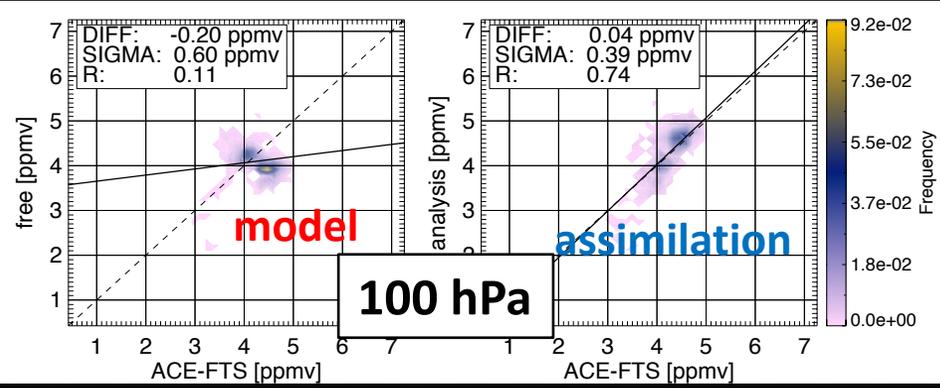
- This work uses a version of the GEOS general circulation model with a **stratospheric chemistry model** driven by MERRA-2 meteorology; GMAO analyses to date have used a simple parameterized chemistry scheme
- **The chemistry model, StratChem:**
 - 51 transported and 17 derived species
 - 149 gas-phase and 39 photolysis reactions
 - Reaction rates follow the recommendations in *JPL 2015*
 - Includes a PSC scheme and heterogeneous reactions
- Currently assimilating ozone, water vapor, HNO₃, and HCl data from MLS and total ozone from OMI

Water vapor

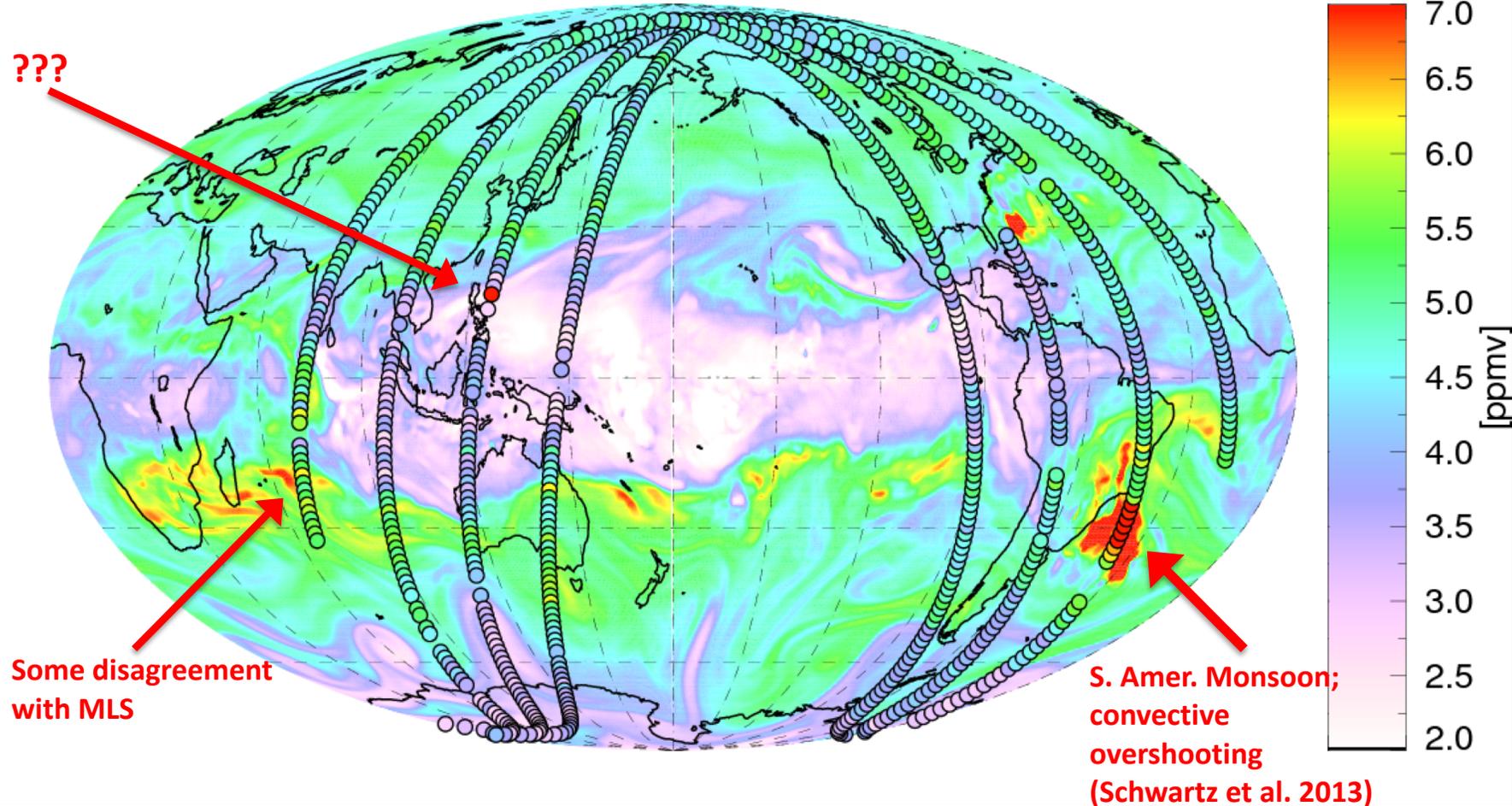
Joint probability distributions: ACE-FTS vs. free run and ACE-FTS vs. assimilation.

Large positive impact of the assimilation on all statistics at pressures > 50 hPa

Assimilation improves the mean at all levels compared to the model



Analysis and MLS water vapor at 100 hPa, 23 December 2015 21UTC



Good fit to the assimilated data, although there are some outliers. This is the first time stratospheric water vapor has been successfully assimilated in GEOS-DAS.

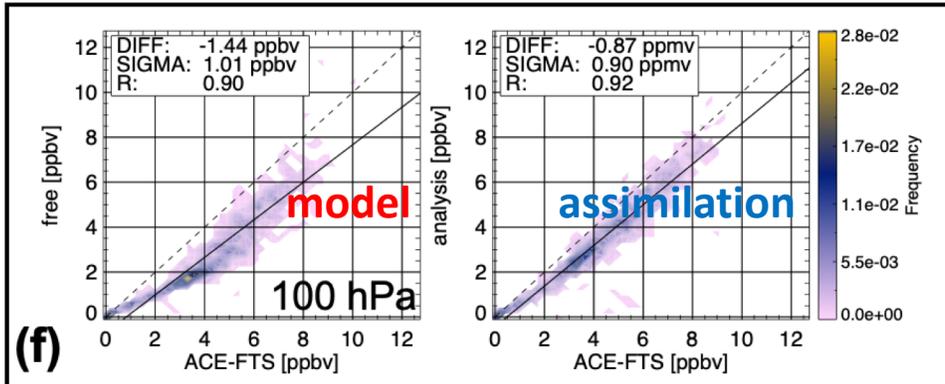
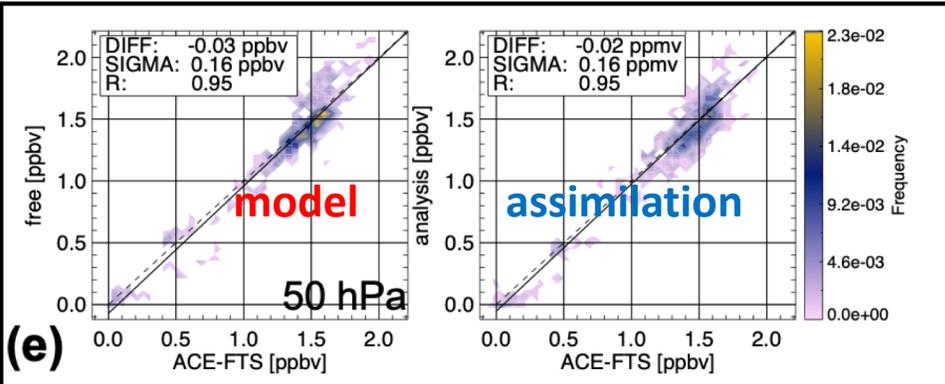
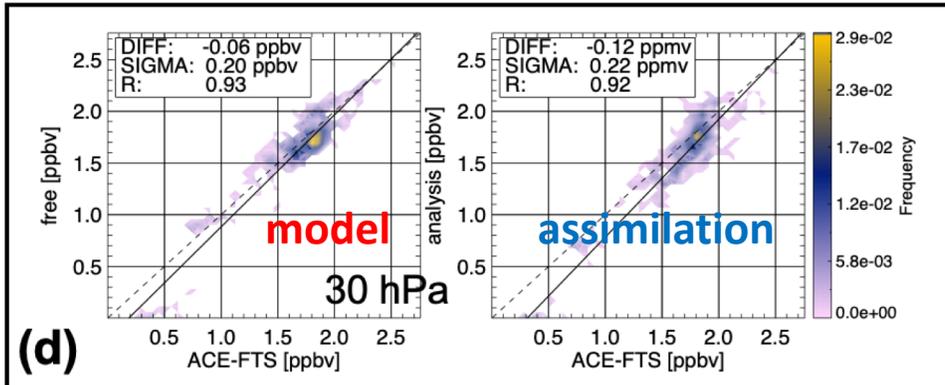
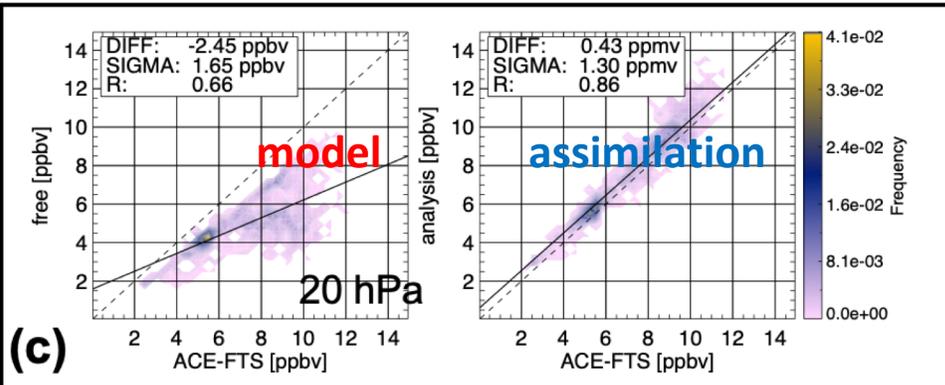
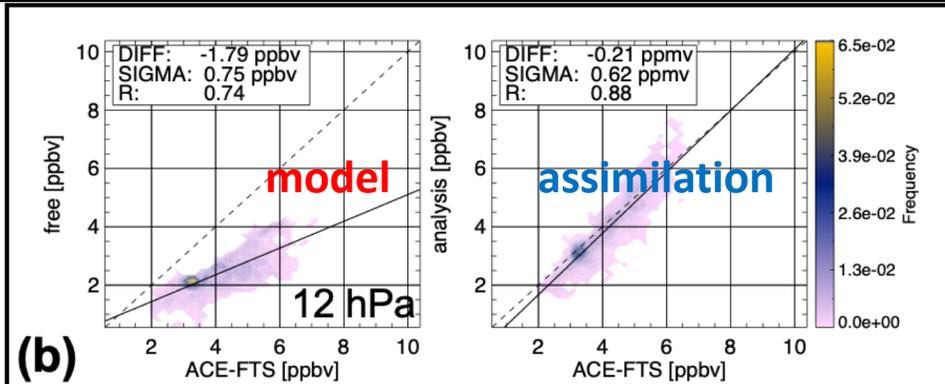
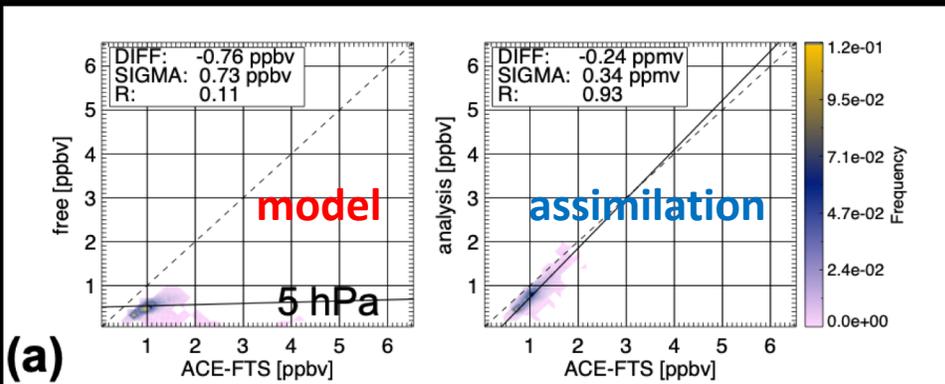
MLS observations are color-coded by mixing ratio

HNO₃

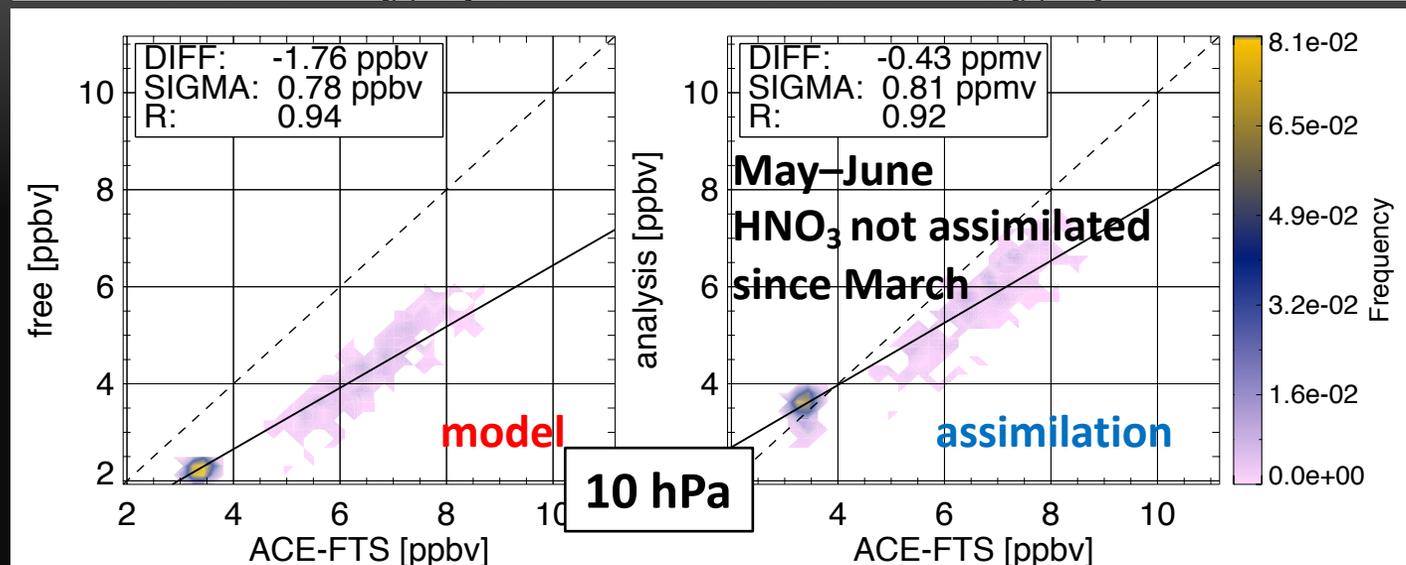
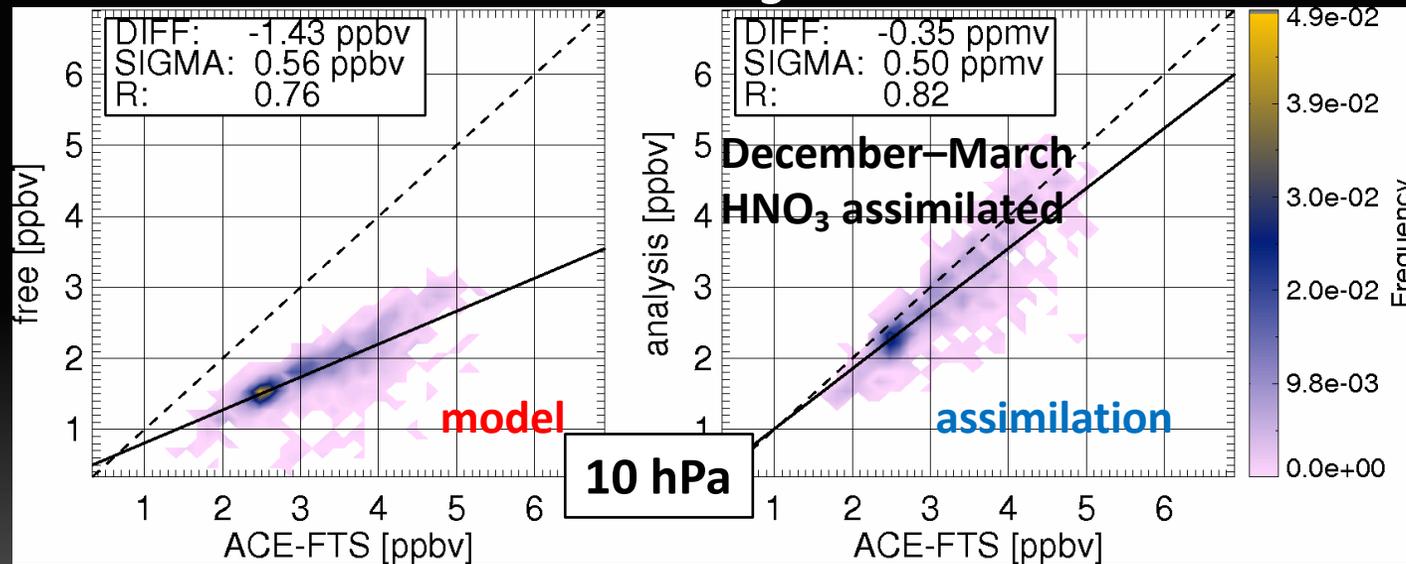
Joint probability distributions: ACE-FTS vs. free run and ACE-FTS vs. assimilation.

Assimilation improves difference standard deviations and correlations in the upper and middle stratosphere.

It improves the mean difference almost everywhere.



HNO₃



An example of feedback through chemistry

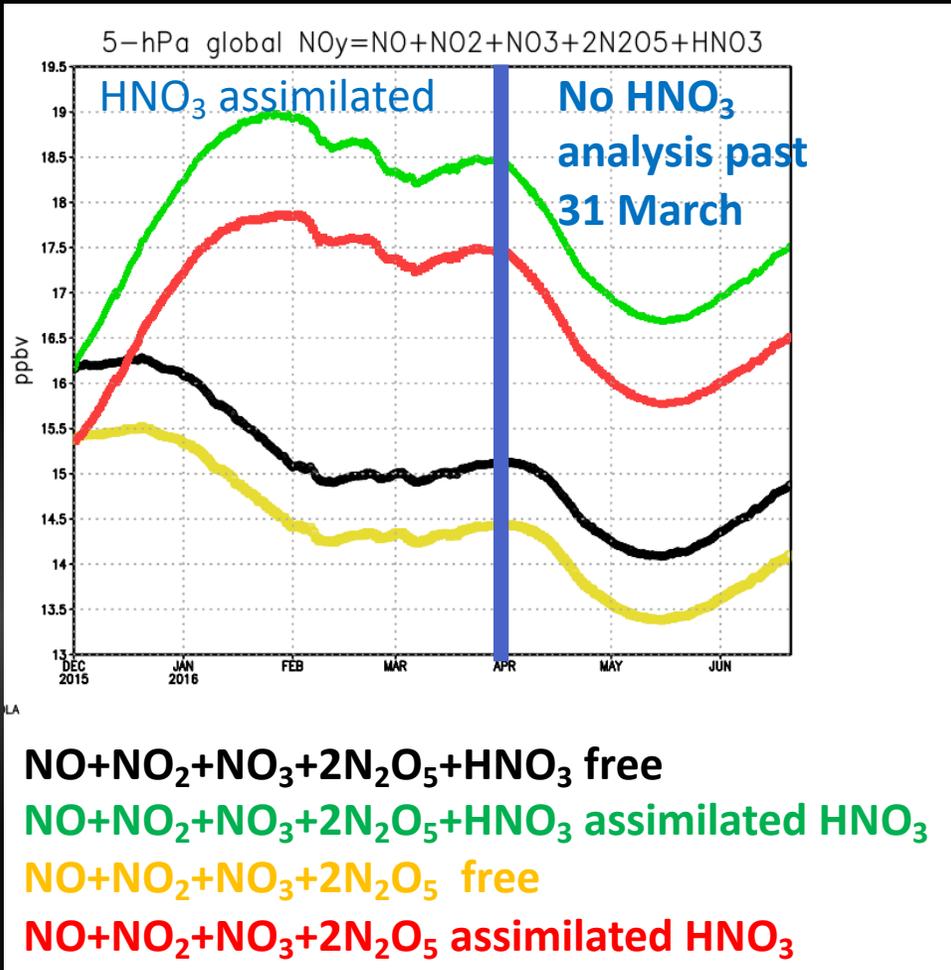
Assimilation has lasting effects on the mean HNO₃ several months after the system ceased to see the data. This is very long compared to its chemical lifetime (~1 day).

Predictability

Assimilation of HNO₃ alters the NO_y budget.

$$\begin{aligned} \text{NO}_y &= \\ &= \text{NO} + \text{NO}_2 + \text{NO}_3 + \\ &+ 2\text{N}_2\text{O}_5 + \text{HNO}_3 + \text{ClONO}_2 \end{aligned}$$

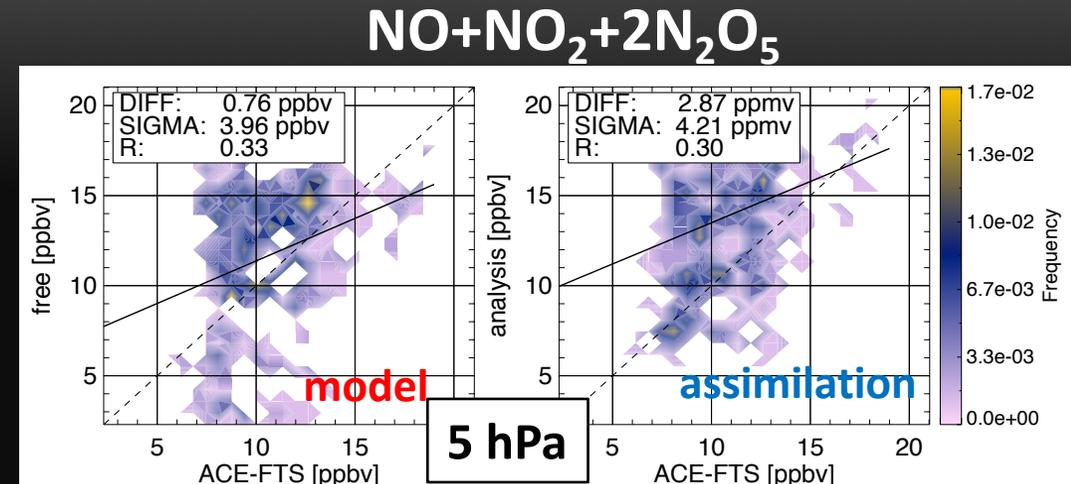
So, is this good news?



HNO_3 assimilation affects total NO_y and nitrogen partitioning.

This reduces HNO_3 bias but does it improve the other nitrogen species? Not in StratChem

NO , NO_2 , N_2O_5 are poorly represented in StratChem but also in GMI. How about other models?



Points made so far

- **Achieving continuity of reanalysis ozone**
- **Importance of transport**
- **Interconnectivity, predictability, and feedbacks in chemical DA; impact on non-assimilated species**

Challenges, questions

- MLS and MIPAS provide observations of many key constituents for chemical reanalyses of the stratosphere, 2003 (to be generous) to present. What happens when MLS is gone? Connection with Theme 3.
- More fundamentally: what is the minimal set of observed stratospheric constituents?
- What is the best strategy for eliminating systematic differences between data sources?
- What do we do about drifts (MLS WV and N₂O, OMPS-LP ozone)
- Focus on transport:
 - Do we assimilate constituent data into a SD simulation? What's the right way to do specified dynamics to achieve fidelity of transport?
 - Do we assimilate constituents within a full data assimilation system (along with meteorology)? Is that computationally feasible?
 - In either case, how do we address discontinuities in the analyzed meteorology?
- What is the impact of constituent assimilation on non-assimilated species/families? How well are the latter represented? Implications for predictability

Potential discussion points

- What science questions can stratospheric chemical reanalyses address?
 - A comprehensive assessment of what a reanalysis can and cannot do would be useful
 - How can we effectively communicate all this to other researchers?