



Iodine in the Troposphere

*corresponding author:
ts551@york.ac.uk

Tomás Sherwen^{*a}, Mathew J. Evans^{a,b}, Lucy J. Carpenter^a
a: Department of Chemistry, University of York, Heslington, York, YO10 5DD, UK.
b: National Centre for Atmospheric Science, University of York, Heslington, York, YO10 5DD, UK

WHY IODINE?

Iodine is crucial for human health in diet; it perturbs the ozone budget & therefore impacts air-quality, crop yields and climate change. Reactive halogens (inc. iodine) catalytically destroy ozone through direct reaction followed by photolytic regeneration. In the troposphere this depletes ozone concentrations, decreasing the tropospheric long-wave absorption since ozone is a strong IR absorber. Of the halogens, the effect of bromine can now be considered by some global chemistry transport models (CTM) (Inc. GEOS-Chem), but the contribution of iodine cannot. The current understanding suggests that the combined impacts on ozone are greater than individual components [Read et al, 2008], & also with current uncertainties over sources [Sommariva et al, 2013; Carpenter et al, 2013] modelling can contribute to reconciling understanding & observations. Work is therefore being performed to introduce the current understanding of iodine chemistry into a established global CTM (GEOS-Chem), with the aim of developing the understanding of the role of iodine within the halogen-driven depletion of ozone & its broader impacts.

BACKGROUND/OBSERVATIONS

- Difficult to detect the source species directly for reasons including short lifetimes & low concentrations
- Oxidised species (e.g. IO) have been measured in the MBL with strong diurnal cycles (fig. 1) [Read et al, 2008]

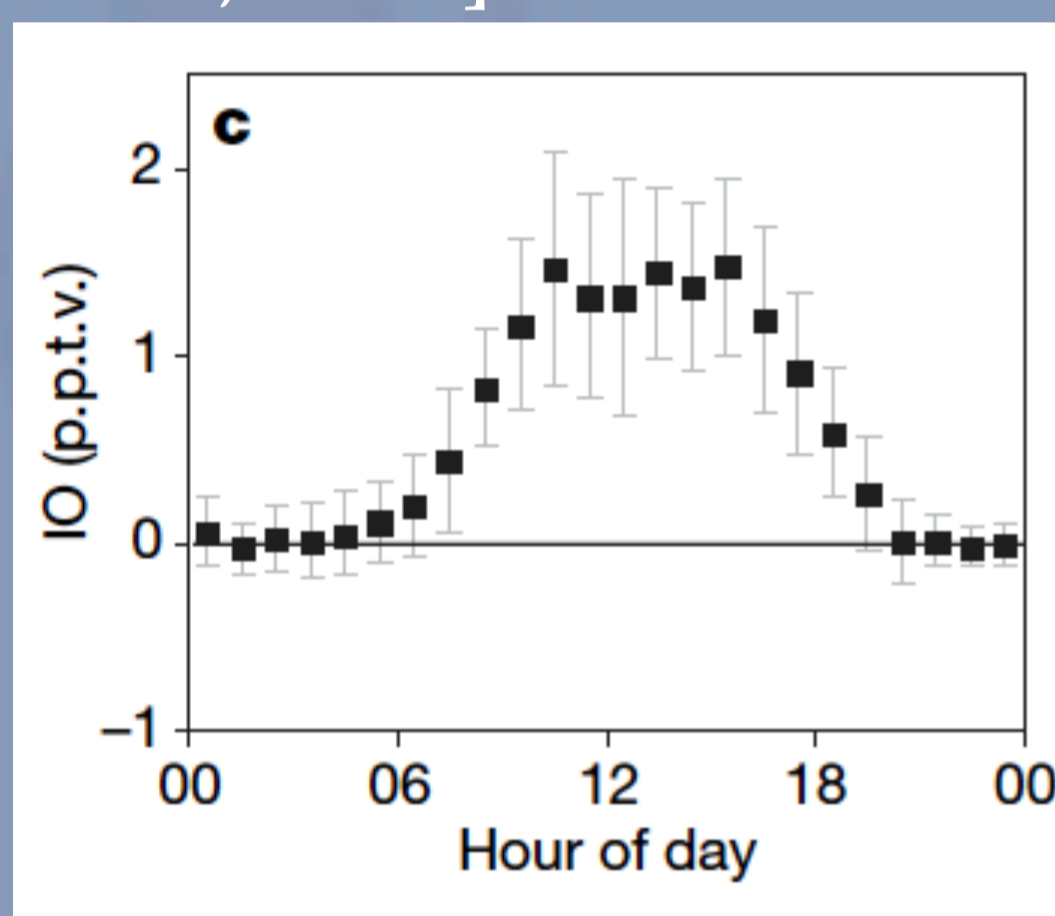
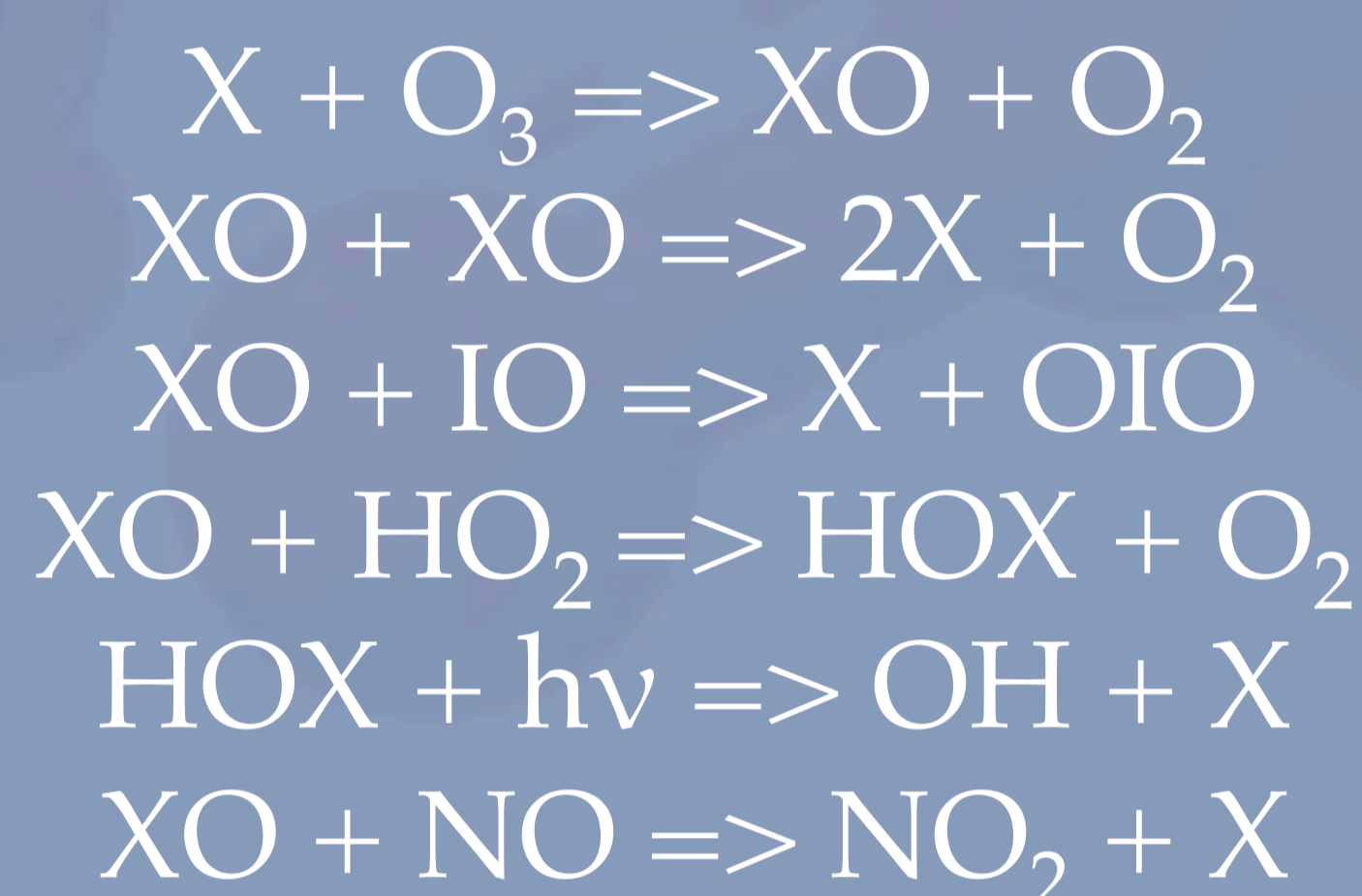


Figure 1: Averaged diurnal profiles for a typical open ocean site, with 1σ errors. (Reproduced from: Read et al, 2008)

- Iodine also interacts with NO_x & HO_x cycles. A simplification of these reactions is shown below:



IODINE MODULE

The current knowledge has been compiled & written into a module for GEOS-Chem:

- Chemistry: Gas-Phase & Aerosol
Reactions schemes have been compiled from existing knowledge inc.: [Sommariva et al, 2012; Perchtl et al, 2006; Vogt et al, 1999]
- Photolysis
Cross-sections from JPL [Sander et al, 2011, 2006] & prepared for Fast-J
- Dry & Wet deposition
Relevant species incorporated into existing GEOS-Chem approach [Wesley, 1989]

EMISSIONS

- The majority of halogen emissions are oceanic in source
- A preliminary scheme has been imposed with set emission fluxes from oceanic gridboxes
- The scheme is a zonal extrapolation of observations by latitude, based on Jones et al, 2010
- Average values applied where no data is held
- Resultant observed concentrations are of expected order (fig. 2)

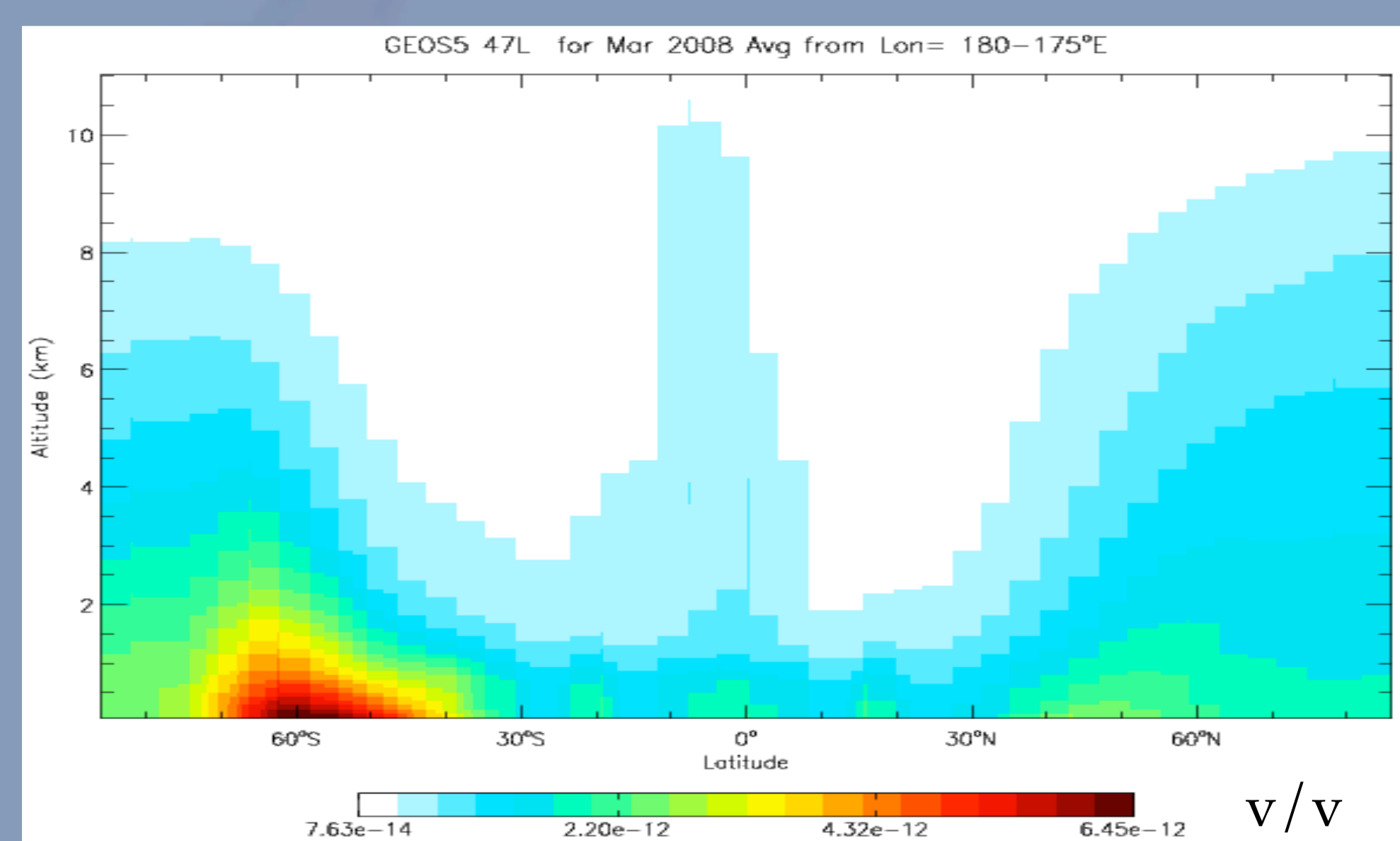


Figure 2: Averaged latitude profile of CH₃I with altitude for model with set CH₃I & CH₂I₂ fluxes run for lower 10km of atmosphere

PRELIMINARY DATA

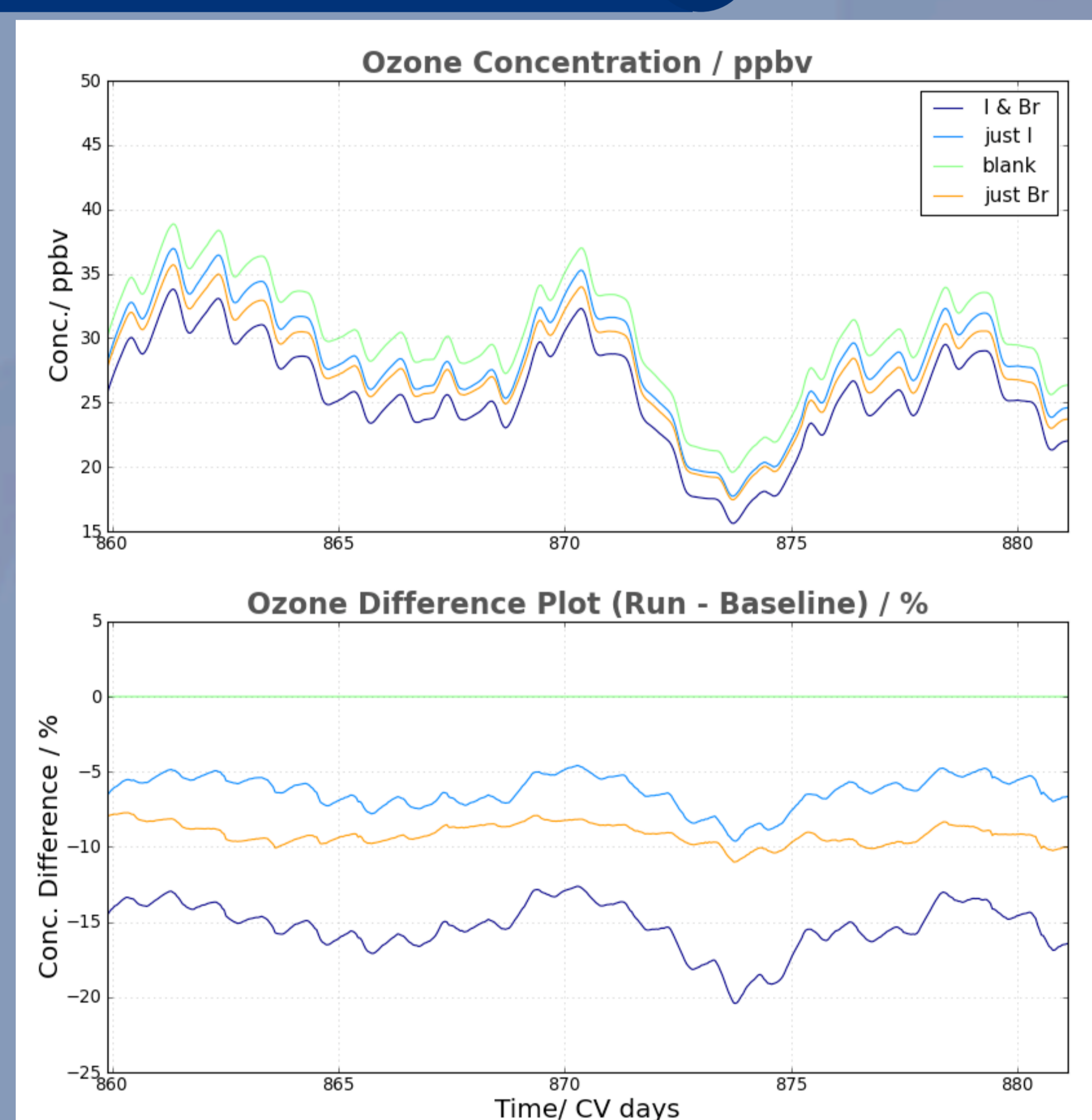


Figure 4: Ozone depletion modelled at a typical open ocean site (Cape Verde) over a 3 week period in May 2008: baseline, green; just iodine, blue; just bromine, light blue; both, dark blue

- The model shows a spatial distribution limited to oceanic surface, as expected (fig. 3)
- Oxidised iodine (IO) shows total ocean coverage
- Antarctic maximum shown in other models has been replicated

- Depletion of ozone is clearly observable with the model (fig. 4)
- The joint impact of iodine & bromine on troposphere ozone is greater than either individual component
- The magnitude of ozone depletion for both species is of the order report by literature [Read, 2008]
- The values of IO being replicated by the model are still lower than those observed

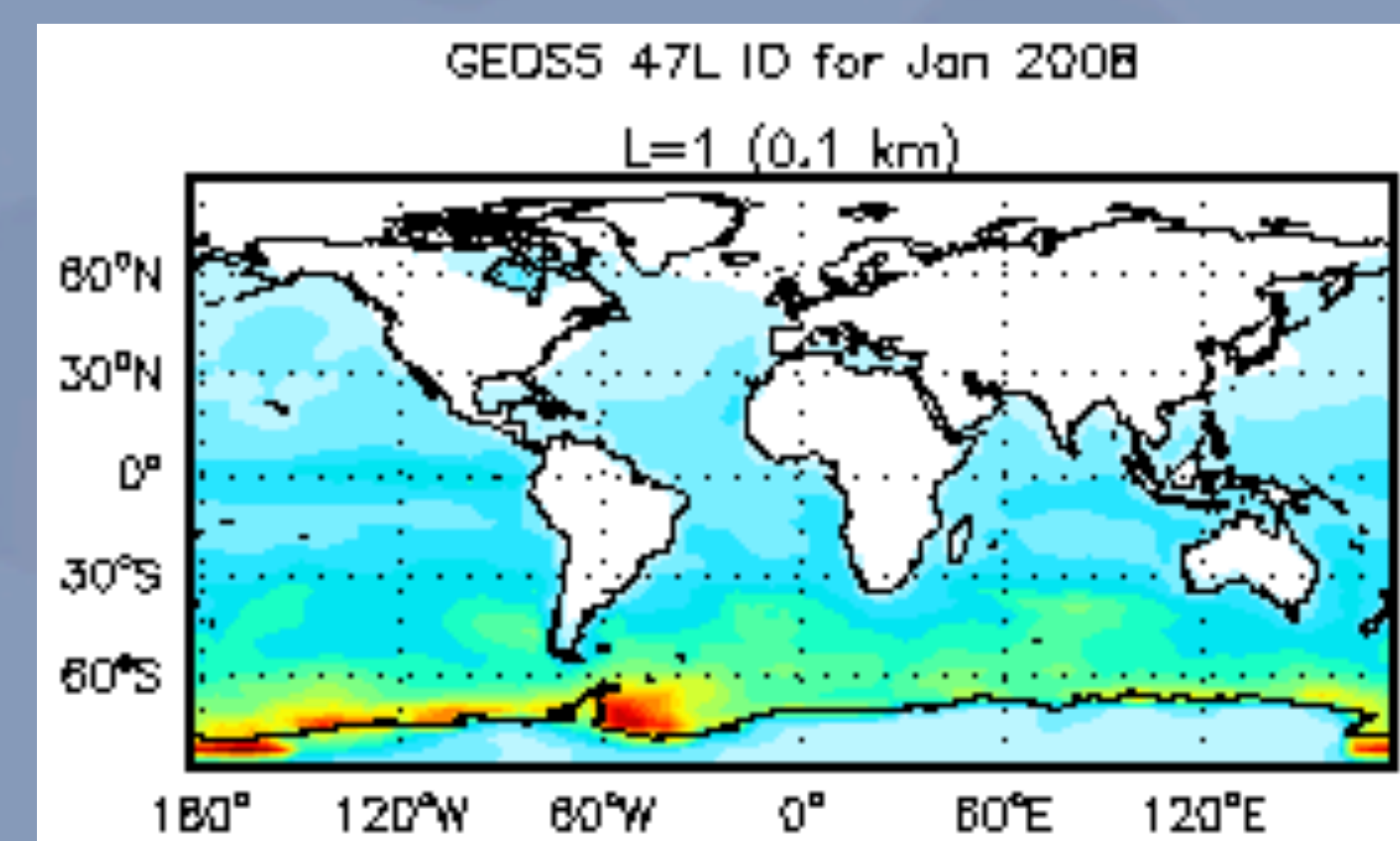


Figure 3: IO distribution plot averaged over altitude fields

CONCLUSIONS & FUTHER WORK

- The ozone depleting capability of iodine has been modelled within GEOS-Chem successfully
- Bromine & iodine combined have a greater effect than either in isolation
- This preliminary model data requires further sensitivity testing of deposition factors, photolytic rates, refining representation of chemistry and crucially, further development of the current approximated emissions scheme
- Experimental work to provide improved & further parameterisation of iodine sources species form the ocean surface
- Investigating evidence for hypothesised additional iodine sources on a global scale
- Quantifying implications on ozone budgets & mercury lifetimes

ACKNOWLEDGEMENTS

TS gratefully acknowledges NERC for PhD Studentship funding, support in studies from the University of York, and student & academic members of the Atmospheric Chemistry Group for continued assistance.

