Ozone Temperature Correlations in the Upper Stratosphere as a Measure of Chlorine Content Richard S. Stolarski and Anne R. Douglass



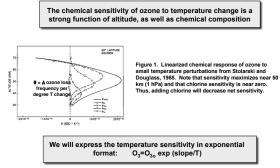
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A little history:

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1975 Barnett et al first suggest that chemical time constant can be determined from ozonetemperature correlations

1985-6 Douglass, Rood show that ozone temperature correlations are also affected by dynamical perturbations that affect both ozone and temperature. They also pointed out that time delays of the ozone response can impact correlations. They looked for meteorologically stable conditions as optimum for detecting chemical sensitivities and suggested that changing chlorine would impact the sensitivity.



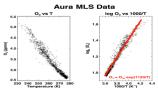


Figure 2: Ozone vs temperature at 1 hPa using data from the MLS experiment on Aura. Ozone shows a small curvature as a function of temperature that is removed by plotting the log of ozone vs 1000/T. The slope of this straight line is the temperature coefficient.

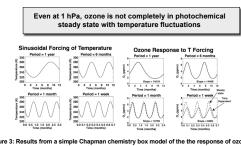


Figure 3: Results from a simple Chapman chemistry box model of the the response of ozone to sinusoidal temperature variations. For a one-year period (seasonal cycle), the time-dependent result matches the steady-state response and the exponential temperature coefficient is the same for both (1451K). For a 6-month period, the results are nearly the same, but for a one-week period the temperature coefficient is 446K. In the atmosphere, temporal and spatial variations will consist of a mixture of time scales that will confuse the deduction of photochemical temperature sensitivities.

MLS data on Aura illustrate the seasonal cycle of ozone and temperature at 1 hPa for mid latitudes

MLS Aura Data at 1 hPA (2004-2007)

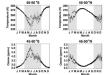
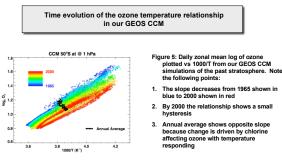


Figure 4:MLS data for ozone and temperature for northem and southern mid latitudes illustrate a seasonal cycle that follows the solar zenith angle over much of the year with ozone anti-correlated to temperature. In the winter, planetary wave disturbances add noise to this picture and increase the mean temperatures above the continuation of the summer cycle.

GEOS CCM

(Goddard Earth Observing System Chemistry Climate Model) GEOS 4 Atmospheric GCM combined with Goddard Stratospheric CTM Coupled radiatively through ozone, carbon dioxide, methane, nitrous oxide, and CFCs Run at 2x2.5 latitude-longitude resolution with 55 vertical levels Pawson et al., J. Geophys. Res., 113, D12103, doi:10.1029/2007JD009511, 2008



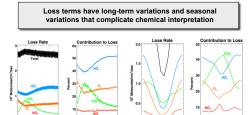


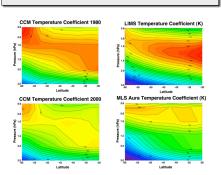
Figure 6: Loss rates at 1 hPa 50°S from GEOS CCM simulations. Note that as chlorine contribution increases, HO₂ and O₂ contributions decrease. Right panels show the mean seasonal variation for each catalytic loss cycle. Again, chlorine varies in the opposite sense from HO₄ and O₂ with the phase slightly different.

2000 2025 2050

1975 2000 2025 2050 2075 210

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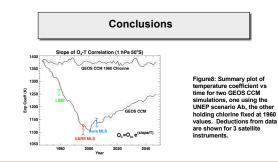
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Temperature coefficient exhibits strong altitude and

latitude dependence as shown in CCM and data

Figure 7: Cross sections of the temperature coefficient vs latitude and pressure altitude for 1980 and 2000 (averaged over 5 years) from GEOS CCM simulations in left panels. Right panels show temperature coefficient derived from LIMS data for 1978-79 (7 months) and MLS Aura data for 2004-2007. At lower altitudes, the coefficient is strongly affected by dynamical influences and there are differences with the model.



Ozone temperature correlations can be used to see the impact of chlorine on upper stratospheric ozone

The method has many difficulties:

- Sensitivity is a function of altitude, latitude, and season
 Deduced sensitivity depends on time scale of variations
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 Transport affects both ozone and temperature

It is critical to use measurements of ozone an temperature that are contemporaneous and have good vertical resolution