Using transport diagnostics to understand chemistry climate model ozone simulations


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[1] We use observations of N$_2$O and mean age to identify realistic transport in models in order to explain their ozone predictions. The results are applied to 15 chemistry climate models (CCMs) participating in the 2010 World Meteorological Organization ozone assessment. Comparison of the observed and simulated N$_2$O, mean age and their compact correlation identifies models with fast or slow circulations and reveals details of model ascent and tropical isolation. This process–oriented diagnostic is more useful than mean age alone because it identifies models with compensating transport deficiencies that produce fortuitous agreement with mean age. The diagnosed model transport behavior is related to a model’s ability to produce realistic lower stratosphere (LS) O$_3$ profiles. Models with the greatest tropical transport problems compare poorly with O$_3$ observations. Models with the most realistic LS transport agree more closely with LS observations and each other.

We incorporate the results of the chemistry evaluations in the Stratospheric Processes and their Role in Climate (SPARC) CCMVal Report to explain the range of CCM predictions for the return-to-1980 dates for global (60°S–60°N) and Antarctic column ozone. Antarctic O$_3$ return dates are generally correlated with vortex Cl$_x$ levels, and vortex Cl$_y$ is generally correlated with the model’s circulation, although model Cl chemistry and conservation problems also have a significant effect on return date. In both regions, models with good LS transport and chemistry produce a smaller range of predictions for the return-to-1980 ozone values. This study suggests that the current range of predicted return dates is unnecessarily broad due to identifiable model deficiencies.


1. Introduction

[2] Chemistry climate models (CCMs) are the current state-of-the-art tools used to assess stratospheric ozone and make predictions of its future evolution [World Meteorological Organization (WMO), 2011, 2007]. Ozone distributions are controlled by transport, chemistry, and temperature (i.e., dynamics and radiation). In the stratosphere, the processes that control ozone are expressed by the ozone tendency equation,

\[
dO_3/dt = \text{Transport} \ + \ P - L(O_3) - L(NO_x) - L(Cl_x) - L(Br_x) - L(HO_x),
\]

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Table 1. Chemistry Climate Models Participating in CCMVal and WMO [2011]

<table>
<thead>
<tr>
<th>CCM Name</th>
<th>Atmospheric GCM</th>
<th>Reference</th>
<th>Horizontal Resolution of Advection, Levels, and Top</th>
</tr>
</thead>
<tbody>
<tr>
<td>AMTRAC3</td>
<td>AM3</td>
<td>Austin and Wilson  [2006]</td>
<td>~200 km, L48, 0.017 hPa</td>
</tr>
<tr>
<td>CAM3.5</td>
<td>CAM</td>
<td>Lamarque et al. [2008]</td>
<td>1.9° × 2.5°, L26, 3.5 hPa</td>
</tr>
<tr>
<td>CCSR NIES</td>
<td>CCSR/NIES AGCM 5.4g</td>
<td>Akiyoshi et al. [2009]</td>
<td>T42L34, 0.012 hPa</td>
</tr>
<tr>
<td>CAMM</td>
<td>AGCM3</td>
<td>Scinocca et al. [2008]</td>
<td>T31L71, 0.00081 hPa</td>
</tr>
<tr>
<td>CNRM-ACM</td>
<td>ARPEGE-Climat v4.6</td>
<td>Déqué [2007], Yessièdre et al. [2007]</td>
<td>T42L60, 0.07 hPa</td>
</tr>
<tr>
<td>E39ICA</td>
<td>ECHAM4</td>
<td>Stenke et al. [2008, 2009]</td>
<td>T30L39, 10 hPa</td>
</tr>
<tr>
<td>EMAC</td>
<td>ECHAM5</td>
<td>Jöckel et al. [2006]</td>
<td>T42L90, 0.01 hPa</td>
</tr>
<tr>
<td>GEOSSCM v2</td>
<td>GEO5</td>
<td>Pawson et al. [2008]</td>
<td>2° × 2.5°, L72, 0.015 hPa</td>
</tr>
<tr>
<td>LMDZepro</td>
<td>LMDz</td>
<td>Jourdain et al. [2008]</td>
<td>2.5° × 3.75°, L50, 0.07 hPa</td>
</tr>
<tr>
<td>MRI</td>
<td>MJ98</td>
<td>Shibata and Deshi [2008a, 2008b]</td>
<td>T42L68, 0.01 hPa</td>
</tr>
<tr>
<td>Niwa SOCOL</td>
<td>MAECHAM4</td>
<td>Schranner et al. [2008]</td>
<td>T30L39, 0.01 hPa</td>
</tr>
<tr>
<td>SOCOL</td>
<td>MAECHAM4</td>
<td>Schranner et al. [2008]</td>
<td>T30L39, 0.01 hPa</td>
</tr>
<tr>
<td>ULAQ</td>
<td>ULAQ-GCM</td>
<td>Pitari et al. [2002]</td>
<td>R6L26, 0.01 hPa</td>
</tr>
<tr>
<td>UMETRACb</td>
<td>HadAM3 L64</td>
<td>Austin and Butchart [2003]</td>
<td>2.5° × 3.75°, L64, 0.01 hPa</td>
</tr>
<tr>
<td>UMSLIMCAT</td>
<td>HadAM3 L64</td>
<td>Tian and Chipperfield [2005]</td>
<td>2.5° × 3.75°, L60, 0.01 hPa</td>
</tr>
<tr>
<td>UMUKA-METO</td>
<td>HadGEM-A</td>
<td>Morgenstern et al. [2009]</td>
<td>2.5° × 3.75°, L60, 84 km</td>
</tr>
<tr>
<td>UMUKA-UCA M</td>
<td>HadGEM-A</td>
<td>Morgenstern et al. [2009]</td>
<td>2.5° × 3.75°, L60, 84 km</td>
</tr>
<tr>
<td>WACCM</td>
<td>CAM</td>
<td>Garcia et al. [2007]</td>
<td>1.9° × 2.5°, L66, 6 × 10⁻⁶ hPa</td>
</tr>
</tbody>
</table>

a No mean age output submitted.

b No REF-B2 (future scenario) submitted.

where P is O₃ photochemical production and the L terms are chemical loss processes due to different radical families [Stolarski and Douglass, 1985, and references therein]. In the lower stratosphere (LS), O₃ chemistry is slow and distributions are controlled primarily by transport. In the middle and upper stratosphere, photochemistry is fast but transport still plays an important role because it controls the distributions of long-lived families that produce radicals involved in O₃ loss processes [Perliski et al., 1989; Douglass et al., 2004]. Photochemistry and temperature control the steady state balance between radicals and their precursors, e.g., NOₓ/NO₃ and ClOₓ/Clₓ. Transport and chemistry in a model must both be physically realistic to produce a credible prediction of O₃. [3] The use of O₃ as a measure of realism in a simulation is fraught with problems. In some cases, compensating deficiencies in the processes affecting O₃ produce a realistic result. In other cases, an O₃ profile or column may be insensitive to some of the terms in the tendency equation. For example, Douglass et al. [1997] noted that while their model’s O₃ profiles agreed with observations their simulated long-lived tracer profiles did not. The tracer profiles were poor due to the dependence on horizontal transport, which was excessive in the model. The good agreement of the simulated ozone with observations indicated that the sum of O₃ loss processes was reasonable, but as the long-lived tracers were too high, the relative fractional losses from different cycles (i.e., NOₓ/NO₃, HOₓ, ClOₓ, and O₃) were probably incorrect [Douglass et al., 2004]. This presents a problem with regard to credibility of prediction. If the relative fractional losses are misrepresented in the present, the response of each term to the changing halogen and greenhouse gas (GHG) emissions and Brewer-Dobson circulation in the future will further change the relative size of each loss term, making it improbable that they will correctly simulate future O₃. [4] Douglass et al. [1999] proposed setting standards for model evaluation that were based on objective comparisons with observations, including quantitative scoring. This approach became feasible in the 1990s with the availability of multiyear, near-global stratospheric trace gas data sets such as those from the Upper Atmosphere Research Satellite (UARS) instruments. This approach is physically based and may identify areas where model improvement is needed. Objective, observationally based evaluations are also advantageous because they provide a way to quantitatively reassess a model after improvements have been made. [5] Eyring et al. [2006] and Waugh and Eyring [2008] adopted a physically based approach for evaluating CCM simulations by assessing the representation of processes that affect stratospheric ozone. They posit that diagnosing transport and dynamical processes in CCMs, rather than O₃, is a meaningful way to evaluate a model’s ability to make reliable projections of future composition. This approach was applied to the CCMs that participated in the WMO 2006 assessment [Eyring et al., 2007] to interpret model predictions of future O₃ levels. They found that the four CCMs showing the best agreement with a wide range of observations had a smaller variation in O₃ projections than did the full suite of 11 CCMs. The Stratospheric Processes and their Role in Climate (SPARC) CCMVal Report [SPARC CCMVal, 2010, hereinafter referred to as SCR] builds on this approach and represents the most comprehensive effort to date toward evaluating model processes by developing observationally based diagnostics for radiation, dynamics, transport, and chemistry. The 18 CCMs participating in the most recent WMO assessment [WMO, 2011] were evaluated in this report and are listed in Table 1. Details of the components of these models and the reference simulations performed is provided by Morgenstern et al. [2010]. [6] The transport evaluation in the SCR concluded that tropical ascent (i.e., circulation) and quasi-horizontal irreversible mixing of midlatitude air into the tropics (i.e., recirculation) were two fundamental processes that strongly affect the distributions of ozone and ozone-depleting substances. The SCR concluded that at least half of the par-
and is interpreted in terms of the models’ representation of tropical ascent and recirculation. Comparison of these results with O\textsubscript{3} profile data investigates the physical link between this transport diagnosis and model O\textsubscript{3} simulations in the tropical and midlatitude lower stratosphere. The N\textsubscript{2}O/mean age diagnostic complements the suite of transport diagnostics discussed in the SCR.

[9] The second part of the paper examines predictions of the return-to-1980 ozone columns for the 15 CCMs simulating the 21st century. Using the transport diagnostics presented here along with some of the SCR chemistry evaluations, we show that several transport and chemistry problems, by themselves, explain at least half of the range of model-predicted return dates for global (60°S–60°N) and October Antarctic ozone columns. We conclude that the current range of predicted return dates is unnecessarily large due to the modeling deficiencies identified here. While there are significant uncertainties in the return dates due to the unknown levels of future halocarbon and GHG emissions [Charlton-Perez et al., 2010], the use of transport and chemistry diagnostics to identify models with credible LS transport and photochemistry can reduce the uncertainty in return dates for a given scenario caused by unrealistic representation of some of the important processes controlling stratospheric ozone.

2. N\textsubscript{2}O and Mean Age Observations

[10] In this section we review the observations of N\textsubscript{2}O and mean age that are used to develop empirical constraints on lower stratospheric transport. A monthly N\textsubscript{2}O climatology derived from observations of the Atmospheric Chemistry Experiment (ACE) satellite instrument onboard SCISAT-1 (2004–2009) [A. Jones, personal communication, 2010] is used to calculate mean stratospheric N\textsubscript{2}O. The use of the 6 year ACE data mean reduces the effect of the quasi-biennial oscillation on LS N\textsubscript{2}O distributions. The latitudinal coverage of ACE measurements varies seasonally, and latitudes poleward of 70° are sampled infrequently. We therefore use an annual mean climatology restricted to 68°S–68°N, 150 hPa and above. Figure 1 shows zonal annual mean ACE N\textsubscript{2}O determined from the Jones climatology. The Aura Microwave Limb Sounder (MLS) also measured N\textsubscript{2}O during this time period [Livesey et al., 2007]. Where these data sets have spatial overlap in the LS, 68°S–68°N and 100–20 hPa, the annual mean differences are almost always less than 10%; for most of the LS they are less than 5%. In the ACE N\textsubscript{2}O validation study, Strong et al. [2008] report that the mean profile differences between ACE and MLS is ±5% from 1 to 100 hPa, with MLS showing low bias at pressures greater than 32 hPa.

[11] Figure 2 displays the mean age data used in this analysis. The tropical profiles come from three OMS balloon CO\textsubscript{2} profiles measured in February and November, 1997 [Andrews et al., 2001] and from two balloon CO\textsubscript{2} profiles measured in June, 2005 (A. Engel, personal communication, 2010). The average of the OMS and Engel profiles (red) and its 1σ uncertainty (yellow shading) are used in the analysis. The mean midlatitude mean age and uncertainty profiles were derived by Engel (pers. comm.) from CO\textsubscript{2} and SF\textsubscript{6} data from 27 balloon flights occurring mostly between May and October, 32°N–51°N, spanning
30 years [Engel et al., 2009]. Figure 2 (bottom) shows the 50 hPa (∼20 km) mean age as a function of latitude [Andrews et al., 2001]. It was derived from dozens of aircraft flights that sampled during most months, although there were no southern hemisphere flights in summer. These data sets represent our best estimate of the annual averaged mean age.

### 3. Relationship Between N₂O and Mean Age

Although mean age observations have less spatial coverage than satellite N₂O measurements, they do span a broad latitude and altitude range. Figure 3 shows ACE annual mean N₂O from the same latitudes and altitudes as the mean age observations shown in Figure 2. Mean age of less than 4.5 years and N₂O greater than 150 ppb form a compact and nearly linear relationship. Where mean age is >4.5 years, age and N₂O lose correlation. Hall et al. [1999] and Schoeberl et al. [2000] have explained this as a result of increasing N₂O loss (photochemical exposure) as air ages. The N₂O/mean age behavior is analogous to the compact correlation observed in the LS for N₂O and Cl [Plumb and Ko, 1992; Schaufler et al., 2003] that is a consequence of slope equilibrium. Slope equilibrium, as originally explained by Mahlman et al. [1986], creates a compact relationship between two long-lived constituents due to the mixing on quasi-horizontal surfaces being much faster than time scales for chemical loss or transport through the surfaces. For mean ages ≤4.5 years, tropical (red) and extratropical (blue and green) points show the same compact relationship, suggesting that for the extrapolar lower stratosphere where N₂O > 150 ppb (i.e., where N₂O and mean age are in slope equilibrium), mean age can be estimated from N₂O.

[13] All collocated mean age and N₂O > 150 ppb data were fit to a line. The fitted line is shown in Figure 3 in blue, where the yellow shading represents the approximate 1σ uncertainty range for the observations, ~5% uncertainty for N₂O and ~25% for mean age. Figure 4 was created using the fit to the observed compact N₂O/mean age correlation to estimate mean age for values of N₂O over the range 160–315 ppb. The heavy yellow line marks the 160 ppb N₂O contour, indicating the upper limit of the LS domain in which the compact correlation is expected to occur. Note that this relationship cannot be determined in polar regions due to the latitude range of the N₂O climatology (68°S–68°N).

[14] Nearly 20 years ago, Plumb and Ko [1992] suggested that the compact correlation observed between long-lived tracers could be applied to models to evaluate their balance between the large-scale circulation and horizontal mixing.
In section 4 we apply the observationally derived compact correlation between N\textsubscript{2}O and mean age to evaluate CCM transport.

4. Interpretation of Mean Age and N\textsubscript{2}O in Chemistry Climate Models

[15] We now examine the correlation between N\textsubscript{2}O and mean age in the CCMs that participated in the 2010 WMO ozone assessment to assess their LS transport characteristics. Table 1 lists the CCMs analyzed here; details of the participating models and additional references can be found in the SCR and is provided by Morgenstern et al. [2010]. This analysis uses zonal monthly mean output of N\textsubscript{2}O and mean age from the last decade of the REF-B1 simulation of the recent past (1960–2006) to compute annual means. This period was chosen because it reflects the present-day circulation, as do the observations; we average over 10 years to reduce any bias caused by the presence of a QBO in a model. Three of the participating CCMs did not provide mean age output and thus could not be evaluated here (CCSRNIES, E39CA, and EMAC).

[16] Figure 5 compares the observed global N\textsubscript{2}O/mean age relationship with 15 CCMs for the LS (150–50 hPa, 65°S–65°N, and up to 30 hPa from 40°S–40°N); the observations where the compact correlation is observed and the fitted line and uncertainties from Figure 3 are shown in black with yellow shading. The interpretation may be fairly simple when the model points (red) fall on a curve that is flatter or steeper than the observations (black). For example, most of the first and second rows of Figure 5 (AMTRAC3, CNRM-ACM, Niwa-SOCOL, SOCOL, and UMETRAC) show maximum mean age less than observed while showing an N\textsubscript{2}O range about the same as observed. This relationship suggests a fast circulation that may have good horizontal mixing. The fast circulation quickly transports N\textsubscript{2}O upward in the stratosphere where it is photo-chemically destroyed, resulting in a low mean age for a given value of N\textsubscript{2}O. The fifth row of Figure 5 (LMDZ and the UMUKCA models) shows model curves that are steeper than observed and a maximum mean age that is 1–3 years older than observed. For LMDZ, where the range of N\textsubscript{2}O values is about the same as observed, the high mean age suggests a slow circulation. The UMUKCA models have both high mean age and low N\textsubscript{2}O. This behavior may be the result of a slow circulation, but may also be caused by too much recirculation (mixing) between the tropics and midlatitudes. UMSLIMCAT and the models in the third and fourth rows of Figure 5 (CAM3.5, CMAM, MRI, WACCM, ULAQ, and GEOS) have curves that agree well with the observed slope and the range of N\textsubscript{2}O and mean ages.

[17] While all models show a compact and nearly linear relationship throughout this LS domain, CNRM-ACM has a much flatter slope than the observations. Figure 3 showed a large change in slope for points above 30 hPa in the mid-latitudes. The nearly flat slope found in CNRM-ACM may indicate that parcels found near the top of this domain (30 hPa) have already spent significant time at high altitudes where N\textsubscript{2}O is destroyed (10 hPa or higher). N\textsubscript{2}O loss rates below 30 hPa are very small, resulting in an N\textsubscript{2}O lifetime \(\gg 1\) year in the domain considered here.

[18] The processes controlling the slope and range of the compact correlation, tropical ascent and mixing, can be better assessed by examining N\textsubscript{2}O and mean age in the tropics. The fourth column of Figures 6a–6c shows the tropical subset of N\textsubscript{2}O/mean age points from Figure 5 (10°S–10°N, 100–30 hPa). The first, second, and third columns of Figures 6a–6c show three different profile comparisons with observations that diagnose the effects of (1) ascent and mixing together (tropical mean age), (2) tropical ascent rate (horizontal mean age gradient), and (3) horizontal mixing (tropical N\textsubscript{2}O). The quantity in each panel is the difference between the observed and simulated value; thus a profile centered on zero indicates exact agreement with the observations. Tropical mean age (first column of Figures 6a–6c) increases with height as a function of both the ascent rate and the horizontal mixing strength. The agreement of model and observed tropical mean age only shows that the combined effects of ascent and mixing produce a realistic mean age in the model. The second and third columns of Figures 6a–6c identify how ascent and horizontal mixing individually contribute to the overall tropical transport seen in the first column. These comparisons may reveal whether a problem lies with circulation or mixing, or both. The mixing evaluated here is cumulative (i.e., total mixing that has occurred during ascent) and relative to the model’s ascent rate; this differs from the analysis in the SCR that diagnosed mixing as a function of pressure. The horizontal mean age gradient is calculated as the difference between the mean ages averaged over 35°N–50°N and 10°S–10°N. For most models, their tropical residual mean vertical velocity, \(w^*\), agrees well with the age gradient derived quantity (SCR, chapter 5) except where noted.

[19] In the limit of low vertical diffusion, the horizontal age gradient profile (second column in Figures 6a–6c) is an empirical measure of ascent rate because it reflects the transit time through Brewer-Dobson cell [Neu and Plumb, 1999; SCR]. It is independent of horizontal mixing across
the subtropics because tropical and midlatitude mean ages will be affected equally by the mixing. (Large vertical diffusion acts to decrease mean age everywhere, which biases the ascent rate high.) When the horizontal age gradient is less (greater) than observed, the ascent rate is too fast (slow).

The comparison of modeled and observed tropical N₂O profiles (third column of Figures 6a–6c) is a measure of the cumulative degree of horizontal mixing into the tropics. Below 30 hPa tropical N₂O has essentially no loss, so ascent alone has no effect on its mixing ratio. The decreasing

Figure 5. N₂O and mean age relationship in 15 CCMs (red) from the same domain as the compact correlation. Observations are shown in black. The yellow shaded area represents the uncertainty of the fitted line (from Figure 3). Model curves that fall off faster (slower) than observed indicate a fast (slow) circulation. Models results are sorted by fastest circulations at the top, slowest circulations at the bottom. Results are from the REF-B1 (1960–2006).
Figure 6a. Diagnostic plots evaluating tropical ascent and tropical-midlatitude (horizontal) mixing in CCMs. Results are from the REF-B1 simulation (1960–2006). The first, second, and third columns show the difference between model and observed profiles of tropical mean age (years), the horizontal mean age gradient (years), and tropical N\textsubscript{2}O (ppb), respectively. The yellow shading indicates the $\pm 1\sigma$ uncertainty in the observations. The fourth column compares the simulated (red) and observed (black) mean age/N\textsubscript{2}O relationship in the tropical LS, 100–30 hPa. A dashed red line in the second and third columns indicates an issue that complicates the interpretation of that diagnostic; see text. These CCMs have the youngest ages of the 15 CCMs.
mixing ratio with height is due only to the cumulative effects of horizontal mixing as the air ascends. Model \( N_2O \) that is higher (lower) than observed can be interpreted as too little (much) mixing of older air into the tropics. The models’ tropical \( N_2O \) profiles are normalized to the ACE data at 100 hPa and the yellow shading indicates the range of uncertainty in the LS based on validation against several satellite instruments [Strong et al., 2008]. If midlatitude

Figure 6b. Same as Figure 6a but for the 5 CCMs with the most realistic tropical circulation and mixing.
Figure 6c. Same as Figure 6a but for the 5 CCMs with slow circulations or excessive horizontal mixing. Note the dashed line in the third column for ULAQ, METO, and UCAM.
N$_2$O is biased high or low in a model, this diagnostic will have to be considered in light of the bias. For example, if the observed-model N$_2$O profile were good but the model’s midlatitude N$_2$O was biased low, this would imply that mixing was actually too weak. Most models’ midlatitude profiles agreed reasonably well with observations; those with bias are indicated by a dashed red line in the third column of Figures 6a–6c.

[20] All models in Figure 6a are too young in the tropics, particularly above 70 hPa (first column). The second column in Figure 6a shows that CNRM-ACM, Niwa-SOCOL, SOCOL, and AMTRAC3 have fast ascent rates with about the right amount of horizontal mixing relative to their ascent rates, although SOCOL borders on too much mixing near 30 hPa in spite of young tropical mean age there. All of these models have excellent agreement with ACE N$_2$O profiles at 40°N. Note that if ascent rates decreased in these models without any change in horizontal mixing, their resulting N$_2$O profiles would indicate too much mixing. CNRM-ACM has the youngest mean age and fastest ascent of any of the CCMs. CNRM-ACM, Niwa-SOCOL, and SOCOL have residual mean velocities that agree with the observationally derived $w^*$ of Schoeberl et al. [2008] but not with the age-gradient-derived ascent rate. The disagreement could be caused by excess vertical diffusion in these models that leads to a high bias for the age-derived ascent rate. UMTRAC has young tropical mean age yet appears to have a good ascent rate and good horizontal mixing. Its midlatitude N$_2$O is not biased. The $w^*$ was not available for UMTRAC or AMTRAC3, so the cause for UMTRAC’s young mean age is unknown. The SCR was also unable to reconcile young mean age in UMTRAC with its tropical transport diagnostics.

[21] The models in Figure 6b show the best agreement overall with all tropical diagnostic quantities, indicating they have the best ascent rates and horizontal mixing of the CCMs. CMAM has slightly rapid ascent near 50 hPa, but otherwise shows very good tropical transport behavior. UMSLIMCAT has variable ascent rates, a bit slow below 50 hPa and a bit fast above, but the overall tropical transport is good. WACCM shows borderline fast ascent and strong mixing at 50 hPa and above, with no bias in its midlatitude N$_2$O. These effects combine to give WACCM a good tropical mean age profile. GEOCCM ascent is a little slow below 50 hPa but otherwise agrees very closely with the diagnostics. All models in Figure 6b have very good agreement with ACE N$_2$O profiles at 40°N except for CAM3.5.

[22] The CAM3.5 model in the third row of Figure 6b has an age gradient that indicates fast ascent while tropical mean age and mixing appear correct. This is explained in part by the high bias in CAM3.5 midlatitude N$_2$O. While the mixing diagnostic (third column of Figure 6b) suggests mixing is about right (dashed line), it must actually be too strong because its midlatitude N$_2$O is too high: more mixing must occur in order to sufficiently lower the tropical mixing ratio. Thus, tropical mean age appears correct because the fast ascent, which lowers mean age, is compensated by excessive mixing, which increases age. However, this model has $w^*$ that agrees closely with observations. The inconsistency between the age gradient diagnostic and $w^*$ may be a result of the CAM3.5 low lid, giving it a shorter Brewer-Dobson circulation path and hence a faster age-derived transit time. The CAM3.5 $w^*$ is the same as WACCM and it uses the same atmospheric GCM, but WACCM has a much higher lid than CAM3.5 ($5 \times 10^{-6}$ hPa versus 2.2 hPa).

[23] The models in Figure 6c show a mixture of tropical transport problems. Of these models, only LMDZ and MRI have good agreement with ACE N$_2$O profiles at 40°N. MRI produces a good tropical mean age profile at all levels but the ascent and mixing diagnostics indicate this is fortuitous. MRI ascent and mixing look good from 100 to 70 hPa, but at 50 hPa and above mixing is too strong while ascent is too fast. The results for ULAQ are similar above 50 hPa, where fast ascent combines with too much mixing to produce good tropical mean age. Below 50 hPa, ULAQ has slow ascent, yet apparently good mean age and horizontal mixing. ULAQ has low midlatitude N$_2$O from 70 to 100 hPa, so the good performance on the mixing diagnostic actually means that there is too little mixing. At 30 hPa the ULAQ midlatitude N$_2$O is biased high, thus mixing is even stronger than indicated by this diagnostic. The ULAQ $w^*$ is considerably slower than the age gradient ascent rate from 100 to 30 hPa, suggesting that its age derived ascent rate may be affected by vertical diffusion. LMDZ has very slow ascent from 100 to 70 hPa. By 50 hPa, the ascent rate appears correct and mixing is good at all levels. The UMUKCA models show very slow ascent, especially below 50 hPa. However, both models have very low midlatitude N$_2$O. When low N$_2$O mixes into the tropics, less mixing is required to produce the ‘correct’ tropical N$_2$O. We can only conclude that slow ascent contributes to much older than observed age; the mixing diagnostic is not useful for these models. The results presented here are generally consistent with the transport conclusions in the SCR.

5. Effects of Transport and Chemistry on CCM Ozone Simulations

5.1. Transport Effects on O$_3$ in the Lower Stratosphere

[24] In this section we examine models’ LS O$_3$ profiles to see if their agreement with observations can be related to transport characteristics. We include all models whose REF-B1 (1960–2006) transport could be evaluated or that ran the REF-B2 (1960–2100) scenario. Oman et al. [2010] report that among 14 CCMs, differences in the lower stratospheric O$_3$ are responsible for most of the column differences in the evolution of O$_3$ in the 21st century. Because O$_3$ is strongly influenced by gas phase chemistry and temperature above 30 hPa and by heterogeneous chemistry in the polar LS, the best place to look for a link between ozone and transport is the extrapolar LS.

[25] Figure 7 shows the difference between annual mean O$_3$ profiles from Aura MLS and 16 CCMs for the SH and NH midlatitudes and the tropics. Although the CCSRNIES model did not submit the necessary mean age output for the transport diagnostic, its O$_3$ profiles are included here. The model annual means are a 10 year average calculated from the last 10 years of the REF-B1 run, usually 1997–2006. An annual zonal mean was calculated from 4 years (2005–2008) of Aura MLS v2.2 O$_3$ [Livesey et al., 2007]; precision error is negligible because of the large number of profiles in the average. The uncertainty of MLS O$_3$ as a function of pressure ranges from up to 30% in the lowermost stratosphere to 5–8% at 50 hPa and above. The shaded area about the zero
line indicates MLS systematic uncertainties. No model agrees with MLS within the uncertainties at all locations shown. The five models with the best representation of circulation and mixing, identified by their tropical \(\text{N}_2\text{O}/\text{mean age}\) relationship in Figure 6b, are plotted in red. (CAM3.5 is shown as dashed red, indicating lower confidence in its transport.) Overall, these models show closer agreement with MLS \(\text{O}_3\) than most models, and the spread among them, except for CAM3.5, is small. The five models shown in blue or green have the greatest LS transport problems, either very fast ascent (CNRM-ACM, light blue), very slow ascent (UMUKCA-METO and UMUKCA-UCAM, dark blue), or combined ascent and mixing problems (MRI and ULAQ, green dashed). Their performances are consistent with their LS transport diagnoses. Below the \(\text{O}_3\) maximum, slow ascent and excessive mixing each act to increase mean age and hence \(\text{O}_3\): the UMUKCA models, MRI, and ULAQ all have higher than observed \(\text{O}_3\). (For...
ULAQ this is only true in the tropics). Fast ascent produces younger mean age and lower ozone, and CNRM-ACM consistently has too low O₃.

[26] The six models shown in black (AMTRAC3, CCSRNIES, LMDZ, Niwa-SOCOL, SOCOL, and UMETRAC) have identifiable transport deficiencies yet they often compare as well with observations from 100 to 50 hPa as do the models with realistic transport. (The SCR transport evaluation characterized CCSRNIES as having a slightly fast ascent rate.) These models have less serious transport deficiencies and smaller disagreements with the observed O₃ than the blue and green line models discussed above. The center panel shows that three of the six models shown in black have much higher than observed tropical O₃ near 30 hPa (CCSRNIES, Niwa-SOCOL, and SOCOL), suggesting a possible chemistry or temperature problem. Figure 7 demonstrates that the LS transport diagnostics are able to physically link poor LS O₃ profiles with transport behavior.

Figure 8. Annual mean 60°S–60°N column O₃ anomalies with respect to 1980 for 15 CCMs. Results from the 15 CCMs that integrated the REF-B2 simulation (1960–2100) are shown in each plot. Observations from the Merged Ozone Data Set [Stolarski and Frith, 2006] are shown with black asterisks. (top) The 5 CCMs (red) with the most realistic transport (CAM3.5, CMAM, GEOSCCM, UMSLIMCAT, and WACCM). (middle) The 3 CCMs in blue have been diagnosed with slow ascent (LMDZ, UMUKCAM, and UMIUKA-UA). The CCM in orange is diagnosed with fast ascent and too much mixing (MRI). The CCM with a dashed blue/orange line has slow LS ascent and too much mixing (ULAQ). (bottom) The 5 CCMs (green) with a fast circulation (AMTRAC3, CCSRNIES, CNRM-ACM, Niwa-SOCOL, and SOCOL).
only in cases where the tropical transport is quite poor. The ability of models with identifiable problems to reasonably simulate \(O_3\) profiles serves as a reminder that \(O_3\) itself is not a good diagnostic quantity.

5.2. Return-to-1980 Column Ozone 60°S–60°N

[27] Roughly half of the ozone column resides in the LS, but to what degree can transport diagnostics explain the entire model column or the return-to-1980 date? Figure 8 shows the annual global mean (60°S–60°N) of the column \(O_3\) anomaly (with respect to 1980) from the REF-B2 (1960–2010) simulation for 15 CCMs and observed annual mean column \(O_3\) anomaly from the Merged Ozone Data Set [Stolarski and Frith, 2006]. Most CCMs do not explicitly calculate tropospheric ozone, but because comparisons are made with column anomalies this should not be a source of bias. Differences in approaches to tropospheric \(O_3\) treatment, as well as differences between in the implementation of the REF-B2 forcings, such as SST projections, will have some impact on the models’ \(O_3\) predictions; details of these differences are provided by Morgenstern et al. [2010]. E39CA, EMAC, and UMETRAC did not submit a REF-B2 simulation and are not included in the remainder of the discussion.

[28] Figure 8 groups the return-to-1980 column \(O_3\) predictions for the 15 CCMs by the results of the transport diagnosis in section 4. In Figure 8 (top) the CCMs with the best transport characteristics (in red: CMAM, GEOSCCM, UMSLIMCAT, WACCM, and CAM3.5) produce a narrow range of predicted return dates (2026–2040) compared to the full range of predicted dates (2004–2060). Figure 8 (middle) shows the four models that demonstrated slow ascent below 50 hPa (in blue: LMDZ, ULAQ, UMUKCA-METO, and UMUKCA-UCAM). They too show a narrow range of return dates, 2023–2035, about 4 years earlier than the models with the best LS transport. The difference in the mean return dates for the two sets of models results is not statistically significant. Two of the slow models (UMUKCA) showed considerable high bias with respect to MLS \(O_3\) in the LS (Figure 7), yet the return dates are not significantly different from the models with good transport and little or no bias. While transport affects LS \(O_3\) distributions, the return date for ozone columns depends on model \(O_3\) sensitivities, especially to temperature and Cl [Stolarski and Douglass, 1985; Oman et al., 2010]. The models with strong tropical recirculation (MRI and ULAQ) are shown in orange in Figure 8 (middle).

[29] Figure 8 (bottom) shows that the five CCMs diagnosed with fast transport (in green: AMTRAC3, CCSRNIES, CNRM-ACM, Niwa-SOCOL, and SOCOL) span the full range of predicted return dates (2004–2060). It is interesting that two models with very similar transport diagnostics, AMTRAC3 and Niwa-SOCOL (e.g., Figure 6a), that also showed reasonable agreement with present-day \(O_3\) profiles produce widely different return dates, 2060 and 2004, respectively. While these CCMs have similar transport diagnostics, clearly they must have a different balance of processes controlling \(O_3\).

5.3. Role of Chemistry

[30] The role of chemistry must be considered in order to understand differences in CCM \(O_3\) predictions. Oman et al. [2010] have evaluated the contributions of halogens (\(Cl,Br\)), \(NO_x\), and temperature to tropical (25°S–25°N) \(O_3\) profiles changes over the 21st century for most of the CCMs evaluated in this study. AMTRAC3 stands out from the other models by having a much greater \(NO_x\) contribution to \(O_3\) loss and a much smaller halogen contribution to \(O_3\) increase in the tropical upper stratosphere during the 21st century. AMTRAC3 has just over half the abundance of halogens compared to most other models. While most models show a LS \(NO_x\) decrease and upper stratospheric increase over the 21st century, AMTRAC3 shows \(NO_x\) increases in both the lower and upper stratosphere, leading to more \(O_3\) loss (relative to other models) and a slower recovery. Low halogens and high \(NO_x\) levels in the AMTRAC3 tropics, compared to the other CCMs, explain why AMTRAC3’s predicted return date is much later than other CCMs.

[31] Niwa-SOCOL and SOCOL have the earliest 60°S–60°N return-to-1980 dates of the 15 CCMs, 2004 and 2020. The Niwa-SOCOL REF-B2 simulation was not evaluated by Oman et al. [2010], however they did evaluate SOCOL, which has the same chemical mechanism and transport scheme as Niwa-SOCOL [Morgenstern et al., 2010]. Oman et al. [2010] show that the halogen contribution to SOCOL’s tropical \(O_3\) increase during the 21st century is much larger than AMTRAC3’s and most of the CCMs. Of all the CCMs, SOCOL shows nearly the highest level of halogens in the year 2000 in the tropical stratosphere between 20 and 1 hPa. Levels of ~4 ppb are simulated, which is in excess of the Cl mixing ratio boundary conditions. While Oman et al. [2010] show that SOCOL has a tropical \(O_3\) sensitivity to changing halogens that is very similar to other models, SOCOL has a much larger abundance of Cl, and photochemistry that produces too much ClO for a given Cl (chapter 6, SCR). At the same time, SOCOL has a low \(NO_x/NO_y\) ratio and thus a smaller sensitivity to \(NO_y\) (chapter 6, SCR). In general, increasing \(NO_x\) during the 21st century causes additional \(O_3\) loss and hence slows down \(O_3\) recovery. SOCOL’s and implicitly Niwa-SOCOL’s early return dates may be a consequence of a larger fraction of their middle and upper stratospheric \(O_3\) losses being due to halogens while the relatively small fraction of \(NO_x\) loss does not retard the recovery.

[32] Chapter 6 of the SCR evaluated chemistry in CCMs by looking at radicals and radical precursors, and photolysis rates. The remainder of this section summarizes some of the chemistry issues identified in the SCR that affect \(O_3\) simulations. In the SCR, a photochemical steady state (PSS) model was used to calculate radicals using each model’s radical precursor and temperature profiles as input. If the PSS model results match the CCM’s radicals, then the CCM is deemed to have the same (i.e., correct) chemical mechanism as the PSS. Niwa-SOCOL did not participate in the PSS comparisons, but it uses the same photolysis and chemistry schemes as SOCOL. Model precursors (e.g., Cl, \(NO_x\), \(O_3\)) were evaluated based on their correlation with \(N_2O\).

[33] CCSRNIES, CNRM-ACM, MRI, and SOCOL had the most disagreements with observations for the precursors, particularly Cl, and MRI and SOCOL had poor agreement with several radicals (e.g., O³D, HO³, \(NO_x\), and ClO). MRI greatly overestimates ClO/Cl, due to a missing loss reaction.
for ClO. AMTRAC3 has minor disagreements with the ClO/Cl\textsubscript{y} ratio in the middle and upper stratosphere. CCSRNIES, Niwa-SOCOL, SOCOL, and UMUKCA-METO have total chlorine in the upper stratosphere that exceeds total Cl\textsubscript{y} emitted at the surface, indicating a lack of conservation of Cl\textsubscript{y}. (The UMUKCA-METO model has excessive Cl\textsubscript{y} due to mistreatment of tropospheric HCl removal.) Simulated tropospheric Cl\textsubscript{y} was much higher than expected for CCSRNIES, CNRM-ACM, MRI, SOCOL, ULAQ, and UMUKCA-METO. In these models, the excess Cl\textsubscript{y} was found to extend into the lowermost stratosphere. This presents a larger problem for CCSRNIES, CNRM-ACM, SOCOL, and ULAQ because they also have LS ClO/Cl\textsubscript{y} that is too high, increasing these models’ potential for ozone loss for a given level of Cl\textsubscript{y}. HO\textsubscript{y} is quite low and NO\textsubscript{y}/NO\textsubscript{y} is quite high in the LS in UMUKCA-METO; UMUKCA-UCAM did not participate in the PSS comparison but uses the same chemistry scheme as UMUKCA-METO and its results are expected to be similar. The PSS comparison identified no major issues with radicals or precursors for CMAM, EMAC, GEOSCCM, LMDZ, UMSLIMCAT, or WACCM.

34 The SCR photochemical intercomparison (‘Photo-Comp’) evaluated the accuracy of J values; nine CCMs participated. AMTRAC3, CCSRNIES, and Niwa-SOCOL showed the greatest inaccuracies, while UMSLIMCAT, WACCM, GEOSCCM, and LMDZ had highly accurate J values.

35 Models with both transport and chemistry problems will have different responses to the changing forcings of the 21st century, i.e., increasing SSTs and the concomitant circulations changes, and changing GHG and halocarbon emissions. All the fast circulation models in Figure 8 (bottom), as well as most of the models in Figure 8 (middle), have transport and Cl chemistry problems. Together they span a 56 year range of return dates. MRI has too much tropical recirculation and significant ClO/Cl\textsubscript{y} problems including a missing loss process for ClO. Niwa-SOCOL has a fast circulation, lack of conservation of Cl\textsubscript{y}, and inaccuracies in its J values. CCSRNIES, CNRM-ACM and SOCOL have fast circulations, excessive ClO/Cl\textsubscript{y} in the LS, and disagreements with radical precursors; CCSRNIES, Niwa-SOCOL, and SOCOL also lack Cl\textsubscript{y} conservation. AMTRAC3 has a fast circulation and mixed accuracy for J values. Problems were found in its ClO/Cl\textsubscript{y} and Cl\textsubscript{y}/N\textsubscript{2}O relationships, which are probably due to the parameterization used to calculate Cl\textsubscript{y}. ULAQ and UMUKCA-METO have circulation and/or mixing problems and a problem with high Cl\textsubscript{y} in the lowermost stratosphere.

36 Of the four models with the best LS transport (CMAM, GEOSCCM, UMSLIMCAT, and WACCM), none had any major chemistry problems. CAM3.5 has fairly good transport and minor problems with ClO/Cl\textsubscript{y} and NO\textsubscript{y}/NO\textsubscript{y}. All five of these CCMs, except for UMSLIMCAT, agree closely with column O\textsubscript{3} observations (discussed in section 6). It is worth noting that the LMDZ model performed very well in the photolysis, radical, and precursor evaluations and its only identified transport deficiency is slow ascent in the tropics below 50 hPa. Oman et al. [2010] examined partial column O\textsubscript{3} recovery in the tropics in these CCMs and found very small changes to the 500–20 hPa columns over the 21st century; the greatest O\textsubscript{3} recovery in all CCMs occurs outside the tropics. Because much of midlatitude O\textsubscript{3} is transported from the production region in the tropical middle stratosphere, tropical circulation problems below 50 hPa may have only a minor effect on the 60°S–60°N recovery. The LMDZ return date is within the range predicted by the five models with the best LS transport.

5.4. Return to 1980 Column O\textsubscript{3} in the Antarctic

37 The N\textsubscript{2}O/mean age transport diagnostic cannot be directly applied to polar regions because most high latitude N\textsubscript{2}O and mean age values there are outside the range of the compact correlation. However, the results of the transport analysis are indirectly relevant because polar descent is linked to tropical ascent through the circulation. In fact, CCMs that performed best on tropical ascent evaluations also performed best on polar descent (SCR, chapter 5). In this section we look at the relationship between vortex Cl\textsubscript{y}, Cl chemistry, and column O\textsubscript{3} return dates. Vortex Cl\textsubscript{y} is a more sensitive diagnostic of transport than mean age because Cl\textsubscript{y} is both time scale and pathway dependent [Waugh et al., 2007]. For example, two models may have the same mean age in the Antarctic vortex, but if one model’s circulation takes parcels to higher altitudes where more CFCs are photolysed, that model will have higher vortex Cl\textsubscript{y}. Differences in modeled Antarctic Cl\textsubscript{y} have been shown to be related to differences in the return-to-1980 O\textsubscript{3} values [Eyring et al., 2006, 2007]. Activated Cl, i.e., ClO, is required for O\textsubscript{3} loss, so a realistic response to Cl requires both a good simulation of Cl\textsubscript{y} (transport) and good photochemistry (ClO/Cl\textsubscript{y}). Other factors are important too, such as polar temperatures, PSC distributions, and the presence of a mixing barrier at the edge of the Antarctic vortex.

38 Figure 9 compares each model’s Antarctic return date with its simulated October 2005 mean Cl\textsubscript{y} at 80°S. The models shown in red have no significant problem with Cl chemistry or conservation. For this set of 8 models there is a clear, positive correlation between Cl\textsubscript{y} in the LS vortex and the return-to-1980 date. This is relationship is generally true for the full set of CCMs not just the ones having correct Cl chemistry, although there is more scatter.

39 Each of the models in black was identified in the SCR as having high Cl\textsubscript{y} and/or high ClO/Cl\textsubscript{y} in the upper troposphere/lower stratosphere. A high ClO/Cl\textsubscript{y} allows a model to have a later return date in spite of low Cl\textsubscript{y} because more ClO is produced for a given Cl\textsubscript{y} level. In the case of MRI, the cause of the high ratio is likely the missing ClO loss reaction. UMUKCA-METO has a known problem in the removal of tropospheric HCl that results in excessive stratospheric Cl\textsubscript{y} (by about 0.5 ppb) but its ClO/Cl\textsubscript{y} in the LS is not high. Understanding this model’s return date is more complex because it also has high NO\textsubscript{y}/NO\textsubscript{y}, low HO\textsubscript{y}, and high vortex temperatures. The other five models in black have high ClO/Cl\textsubscript{y} in the UT/LS but the cause was not determined in the SCR (chapter 6). This high ratio alone may explain why Niwa-SOCOL, SOCOL, and ULAQ have a later return date than would be expected for their low Cl\textsubscript{y} levels. However, CCSRNIES and CNRM-ACM also have a high ratio but do fall on the fitted line, so the ClO/Cl\textsubscript{y} problem cannot be the only factor. CNRM-ACM has very young mean age in the Antarctic and incorrect fractional release for CFC-11 and CFC-12 (SCR, chapter 5); this explains why high Cl\textsubscript{y} is found in the CNRM-ACM LS.
vortex in spite of the transport problems, but the cause for the excessive fractional release is unknown. Fractional release could not be assessed for CCSRNIES, but overall CCSRNIES performed below the multimodel average for precursors and radicals.

[40] Antarctic O₃ recovery is not strictly correlated with vortex Clᵧ (i.e., transport) because the amount of Clᵧ that a model converts into active Cl is dependent on so many processes. The amount of vortex Clᵧ in these CCMs is roughly correlated with their transport circulation: many fast models have low Clᵧ (e.g., CCSRNIES, Niwa-SOCOL, SOCOL) while those with slower circulations have high Clᵧ (e.g., UMUKCA-METO, UMUKCA-UCAM, ULAQ). However, model problems with Cl photochemistry and conservation prohibit a strong correlation with circulation strength from standing out in Figure 9.

[41] For models that do not have significant chemistry problems, we find a smaller range of predicted Antarctic return dates. Figure 9 shows a range of 39 years (2028–2067) for Antarctic column O₃ return-to-1980 dates for the 15 CCMs. When we consider only the models that do not have Cl chemistry problems (8 models), we find a range of 30 years (2037–2067). Of these 8 models, 3 have been diagnosed with transport problems (UMUKCA-UCAM, LMDZ, and AMTRAC3). The five models that demonstrated the most realistic LS transport and chemistry (CAM3.5, CMAM, GEOSCCM, UMSLIMCAT, and WACCM) produce a 19 year range of return dates, 2037–2056.

6. Discussion and Conclusions: Understanding Model Predictions

[42] Figure 10 shows 60°S–60°N column O₃ and column anomalies for 15 CCMs. In Figure 10 (top) the models whose column O₃ agrees to within 5% of observations (~15 DU) are plotted in blue. Figure 10 (middle) shows column anomalies, indicating when each model returns to its 1980 60°S–60°N mean value. The 10 models that agree best with global mean observations (blue) span the full 54 year range of return dates (2004–2058). No reduction in the range of predictions is found by selecting models based on their ability to reproduce ozone observations, although a reduction in range for a randomly selected group of 10 results would be expected 57% of the time. Figure 10 (bottom) shows the same column anomalies where the models with the best LS transport, as determined by this analysis, are plotted in red; the range of predicted recovery dates is 14 years (2026–2040). This reduction is unlikely to result from the reduced sample size (5 CCMs instead of 15). The random selection of 5 results gives a range of 14 years or less only 7% of the time. The 10 models shown in black have identifiable problems with transport, such as fast or slow ascent, or inappropriate tropical isolation. Seven of those 10 models also have a problem with their ClO/Clᵧ, and three of those also lack Clᵧ conservation.

[43] Column ozone is influenced by so many processes that a simulated column may be close to the observed value due to compensating effects from multiple problems. For this reason, column O₃ should not be used to gauge model credibility. Similarly, CCMs with good LS transport and chemistry will not necessarily produce good agreement with observed columns for several reasons, including lack of tropospheric chemistry, problems with upper stratospheric temperatures, and representation of heterogeneous chemical processes.

[44] The overall goal of this paper is to understand ozone predictions in CCMs. The initial approach was to evaluate model transport, then determine if transport problems had a
noticeable impact on simulated LS $O_3$. We found that large transport discrepancies (e.g., very fast or slow circulations, or strong mixing) did affect $O_3$ profiles in the lower stratosphere, while smaller transport problems did not. Later, using some of the SCR chemistry evaluations and results from Oman et al. [2010], we showed how $O_3$ simulations are strongly affected by chemistry problems, particularly those involving the Cl family.

[45] Ozone recovery in CCMs is a result of decreasing CFCs that are forced by the mixing ratio boundary conditions used in the scenario. Chemistry and transport evaluations of CCMs provide the means to explain much of the variation in return-to-1980 dates. Antarctic recovery is a function of reactive Cl. If a model’s photochemistry is correct, then realistic Cl$_x$ can be produced if LS Cl$_y$ is also realistic, and the Cl$_y$ levels are to a large extent controlled by transport. If there are problems with ClO/Cl$_y$, then the relationship between transport and Cl$_y$, and hence return date, will be less clear. For the Antarctic and the 60°S–60°N average, the models with good chemistry and good LS

Figure 10. (top) Annual mean 60°S–60°N column $O_3$ in 15 CCMs from 1970 to the present. Results are from the REF-B2 simulation (1960–2100). Observations are shown with black asterisks in each plot. The models in blue show agreement within 5% of the observations; models in black do not agree within 5% of the observations. (middle) The same CCM output and observations, but plotted as anomalies with respect to 1980, from 1960 to 2080. The dashed vertical lines show the earliest and latest predicted return dates for the models with the best agreement with observations (blue), which is the same range of all 15 CCMS (2004–2058). (bottom) The same CCM output but models with the best LS transport are shown in red. The dashed vertical lines show the earliest and latest predicted return dates for the models in red, 2026–2040.
transport, as identified by close agreement with observations of N₂O and mean age, show a range of return dates that is much less than the range of the full set of 15 CCMs. In the Antarctic the range is reduced by half, from 39 to 19 years, and for the 60°S–60°N mean, the range is reduced by 75%, from 56 to 14 years. Realistic chemistry may have a greater impact than realistic transport on O₃ simulations, but this can’t be assessed for this set of CCMs because none was found to have excellent chemistry but poor transport. However, given the importance of Cl₂ to Antarctic recovery and the sensitivity of Cl₂ to transport, we conclude that both good chemistry and good transport are required for credible predictions.

[46] While there is much uncertainty in future ozone levels due to the uncertainties in future emissions of ozone-depleting substances and greenhouse gases [e.g., Charlton-Perez et al., 2010], the current range of predicted return dates is unnecessarily large due to identifiable model transport and chemistry deficiencies. It is remarkable, and perhaps an encouraging sign of progress in chemistry climate modeling, that a group of models with different dynamical cores, transport schemes, chemical solvers, and spatial resolutions can produce very similar lower stratospheric O₃ and column O₃ return dates. The explanation for the similar behavior in these models is suggested by the diagnostics presented here: these models have credible representation of important physical and chemical processes that affect the distribution of ozone and other trace constituents involved in O₃ chemistry. Having models with credible physical processes that produce similar O₃ predictions increases our confidence in our current understanding of the essential chemical and dynamical processes controlling ozone.

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References


Garcia, R. R., R. J. Wilson (2006), Ensemble simulations of the decline and recovery dates. The explanation for the similar behavior in these models is suggested by the diagnostics presented here: these models have credible representation of important physical and chemical processes that affect the distribution of ozone and other trace constituents involved in O₃ chemistry. Having models with credible physical processes that produce similar O₃ predictions increases our confidence in our current understanding of the essential chemical and dynamical processes controlling ozone.