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Quantifying uncertainty in projections of stratospheric ozone over the 21st century

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Abstract

Future stratospheric ozone concentrations will be determined both by changes in the concentration of ozone depleting substances (ODSs) and by changes in stratospheric and tropospheric climate, including those caused by changes in anthropogenic greenhouse gases (GHGs). Since future economic development pathways and resultant emissions of GHGs are uncertain, anthropogenic climate change could be a significant source of uncertainty for future projections of stratospheric ozone. In this pilot study, using an “ensemble of opportunity” of chemistry-climate model (CCM) simulations, the contribution of scenario uncertainty from different plausible emissions pathways for ODSs and GHGs to future ozone projections is quantified relative to the contribution from model uncertainty and internal variability of the chemistry-climate system. For both the global, annual mean ozone concentration and for ozone in specific geographical regions, differences between CCMs are the dominant source of uncertainty for the first two-thirds of the 21st century, up-to and after the time when ozone concentrations return to 1980 values. In the last third of the 21st century, dependent upon the set of greenhouse gas scenarios used, scenario uncertainty can be the dominant contributor. This result suggests that investment in chemistry-climate modelling is likely to continue to refine projections of stratospheric ozone and estimates of the return of stratospheric ozone concentrations to pre-1980 levels.

1 Introduction

Over the past four years the chemistry-climate modelling community has made progress in understanding and refining projections of stratospheric ozone over the 21st century. For the first time, a large number of Chemistry-Climate Models (CCMs) have been run using common ozone and climate forcings for the period 1960-2100 for the second phase of the Stratospheric Processes And their Role in Climate (SPARC) Chemistry-Climate Model Validation Project (CCMVal2). The set of model integrations provided by 18 modelling centres around the world has been extensively interrogated by a large number of authors and a comprehensive report detailing our current understanding of the CCMs was completed early in 2010. As part of CCMVal2, a new analysis of uncertainty in projections of total column ozone over the 21st century and dates of return of total column ozone has been produced (Chapter 9, SPARC CCMVal (2010)). This analysis, however, has been conducted for reference simulations of CCMs based on a common GHG forcing scenario. Many recent studies have pointed to the role that future changes in GHG concentrations might play in determining both global and regional ozone abundances over the 21st century (Shepherd (2008), Waugh et al. (2009), Jonsson et al. (2009), Oman et al. (2010a)). GHGs can affect ozone concentrations through the additional HOx and NOx production from increases in CH₄ and N₂O decomposition, through acceleration of catalytic ozone destruction cycles in the colder stratosphere and indirectly by influencing the strength of the Brewer-Dobson circulation and hence the meridional and vertical transport of ozone (Barnett et al. , 1974; Randeniya et al. , 2002; Rosenfield et al. , 2002; Haigh and Pyle , 1982; Chipperfield and Feng , 2003; Portman and Solomon , 2007; Ravishankara et al. , 2009; Butchart et al. , 2006)

Given the sensitivity of ozone amounts to GHG concentrations, there is an additional uncertainty in projections of ozone column amounts due to the uncertainty in the use of fossil fuels and other GHG producing compounds over the next 100 years. In the climate change community, this uncertainty is addressed by producing projections

of climate variables using a range of different possible GHG scenarios (SRES Naki-
cenovic et al. (2000)), consistent with economic models of global development. While
these scenarios do not necessarily span the full range of possible GHG emissions, they
do at least allow an estimate of the sensitivity of climate models to different economic
development paths.

In this study, additional simulations of the CCMs used for the CCMVal2 intercom-
parison forced with different GHG scenarios (and in some cases slightly different ODS
scenarios) are used to quantify the uncertainty in global and regional stratospheric
ozone projections associated with three factors: model uncertainty, scenario uncer-
tainty and internal variability. By separating the overall uncertainty of CCM projections
in this way, it is possible to identify regions in which:

- model uncertainty is dominant, such that projections could be improved by CCM
development leading to a narrowing of this uncertainty,
- scenario uncertainty is dominant, such that additional simulations of CCMs forced
with different GHG scenarios would enable a better understanding of future ozone
abundances, and
- internal variability is relatively large, such that there will be significant component
of the total uncertainty which is irreducible.

Identifying regions and time periods where these conditions are found will aid future
CCM ensemble design and inform policymakers about the potential to narrow uncer-
tainty in ozone projections.

2 Model Integrations

Model integrations used in this analysis form part of the CCMVal2 intercomparison
project. The core set of integrations used are so-called future reference simulations
(REF-B2) from the CCMVal2 activity. These simulations are described and assessed

in detail in the SPARC CCMVal report (SPARC CCMVal, 2010) and the participating CCMs are described in Morgenstern et al. (2010) and references therein. REF-B2 simulations use surface time series of halocarbons based on the adjusted A1 scenario from WMO (2007), and the long-lived GHG concentrations are taken from the SRES A1b scenario Nakicenovic et al. (2000). Sea surface temperatures (SSTs) and sea ice concentrations (SICs) are generally prescribed from coupled ocean model simulations, either from simulations of the same atmospheric GCM run with a coupled ocean, or from coupled ocean-atmosphere model simulations of a different AR4 GCM simulation under the same A1B GHG scenario. Only one of the 13 CCMs used here ran with a coupled ocean (CMAM).

To avoid artifacts from using different numbers of models as time progresses [Chapter 9 of SPARC CCMVal, 2010; Waugh and Eyring, (2008)], only the subset of 13 CCMVal REF-B2 simulations that provided output from 1960-2095 are used in this study.

In addition to these REF-B2 simulations, several other model integrations which primarily varied the GHG scenario are also analysed. Five different GHG scenarios are considered, with the models running each scenario listed in Table 1. Two scenarios were run with different SRES pathways (SRESA2 and SRESB1) but with an identical ODS scenario to the standard REF-B2 runs. Three additional scenarios were run by one model (CAM3.5), consistent with the new coordinated Climate Change Stabilization Experiments proposed for climate models (Meehl et al. , 2007). These new Representative Concentration Pathways (RCPs, Moss et al. (2008)) were generated by Integrated Assessment Models (IAMs) and were harmonized with the historical emissions from Lamarque et al. (2010) in both amplitude and geographical distribution. The RCP simulations that were performed with CAM3.5 are RCP 8.5 (Riahi et al., 2007), RCP 4.5 (Clarke et al., 2007), and RCP 2.6 (van Vuuren et al., 2007)). In addition to the large differences in GHG amount, the RCP scenarios also differ in the concentration of ODS amounts over the 21st century (although to a lesser degree than the differences in GHG amount).

Therefore the predominant influence on ozone amounts in the different scenarios considered here is that of changing GHG quantities. The following analysis will therefore attempt to quantify the impact of uncertainty in future GHG scenario on the evolution of stratospheric ozone. There also remains some uncertainty in future emissions of ODS amounts which is outlined in detail by Chapter 8 of WMO (2007). For example a hypothetical elimination of ODS emissions from both continued ODS production and halocarbon banks in 2006 would have accelerated the return of Effective Equivalent Stratospheric Chlorine (EESC) to pre-1980 levels by 15 years according to calculations in WMO (2007). These uncertainties are not addressed in this study, and should be considered as another source of potential scenario uncertainty that might be quantified in the future. Separation of the impacts of GHG and ODS changes on stratospheric ozone amounts for the CCMVal2 models is discussed in Eyring et al. (2010).

3 Methods for quantifying uncertainty

In two previous studies, Hawkins and Sutton (2009, 2010) quantified the uncertainty in global and regional near surface temperature and precipitation using a suite of model integrations from phase 3 of the Coupled Models Intercomparison Project (CMIP3). We adapt these methods to examine the sources of uncertainty in projections of stratospheric ozone for the 21st century, focussing on specific regions and the date of return of ozone to 1980 levels.

A detailed description of the methods used can be found in the appendix of Hawkins and Sutton (2009), but the approach is briefly outlined below for an ideal ensemble of CCM simulations. We also discuss some necessary changes to the methodology to accommodate issues specific to ozone projection.

3.1 Separating the sources of uncertainty

For an ideal ensemble of simulations (*which is not available for the CCMVal ozone projections*) the following approach to separating the sources of uncertainty in ozone projections could be followed.

5 It is assumed that the total variance in future ozone concentrations (T^2) as a function of time (t) can be separated into three components,

$$T(t)^2 = M(t)^2 + S(t)^2 + I^2,$$

where $M(t)^2$ represents uncertainty due to the use of different models, $S(t)^2$ represents the variance in projections associated with different future emissions scenarios and I^2 represents variance associated with internal variability in ozone concentrations,
10 which is assumed to be constant in time (this assumption is discussed further below).

To estimate the different components, a smooth curve is fitted to each model integration, and the variance of the residuals represents the contribution from internal variability. The multi-model mean of this quantity then represents an estimate of I^2 . In
15 Hawkins and Sutton (2009, 2010) a fourth order polynomial is used, but for the ozone projections, the smooth curve is created by using a Gaussian filter, with full width at half-maximum of 20 years, on the annual data. The variance of the ensemble around the multi-model mean is considered as the model uncertainty, and this is averaged over the scenarios to give an estimate of M^2 . Finally, the variance of the multi-model means
20 for each scenario is considered as the scenario uncertainty, S^2 .

Each model is assumed to be independent and equally realistic, i.e. no effort is made to weight the models by their ability to simulate historical changes in ozone. The analysis can be repeated for different seasonal means and regions.

In all cases, decadal mean ozone concentrations are considered, since ozone predictions from CCMs are typically used to examine and quantify the long-term changes
25 in ozone amounts rather than predict the amount of ozone present in any given year. This means that our estimates of internal variability (I^2) represent changes on decadal

or longer timescales, and do not consider (for example) the interannual variability in ozone which is large for some geographical regions and seasons.

Additionally, for ozone projections it is common to consider variations from 1980 total column amounts. To calculate the ozone offset, a smooth value for the 1980 total ozone column for the particular region and season considered is calculated by applying the same Gaussian filter to the raw total column ozone data. The procedure described above is then applied to anomalies from this smooth 1980 value in the total column ozone data.

3.2 Application to ozone projections

An ideal multi-model ensemble for this analysis would comprise integrations of a large number of different CCMs each forced by a large number of GHG scenarios. In the studies of Hawkins and Sutton, uncertainty in projections of surface temperature and precipitation are quantified using 15 coupled ocean-atmosphere GCMs which all ran simulations with three different GHG emissions scenarios. Unfortunately an ensemble of CCM simulations of similar size is not currently available (see Section 2). In order to make progress in quantifying the uncertainty in ozone projections it is necessary to make some approximations to the methodology described above. Since the simulations available comprise a large ensemble of model integrations with a single scenario and only a small ensemble of simulations from a subset of the models (13 CCMs for the REF-B2 ensemble and 2 CCMs for each alternative GHG scenario) with different scenarios, a logical first approximation is to restrict the calculation of model uncertainty and internal variability to the single scenario ensemble (labelled REF-B2 above and using SRESA1b GHG forcings). This step assumes that model uncertainty and internal variability in other scenarios is consistent with that in the REF-B2 simulations. This has the added benefit that estimates of model uncertainty should be consistent with the estimates previously made in the CCMVal2 report (Chapter 9, SPARC CCM-Val (2010)). Estimation of scenario uncertainty is more difficult, since it is important to avoid contaminating our estimate of scenario uncertainty with model uncertainty.

In this pilot study, two methods are used to estimate scenario uncertainty. The first method uses four simulations from three different CCMs (CCSRNIES, GEOSCCM and WACCM) which are forced with alternative scenarios from the SRES emissions sets. The second method uses runs from a single CCM (CAM3.5) which is forced with alternative scenarios from the RCP emissions set. This separation is motivated both by the slightly different nature of the simulations in the SRES and RCP sets and also by the fact that the SRES and RCP forcing sets are designed to separately estimate the potential spread in future economic and technological development over the 21st century, leading to different emissions of climate relevant gases. Note that the spread in emissions of carbon dioxide, methane and nitrous oxide are much larger in the RCP scenario set than for the SRES scenario set over the 21st century.

3.2.1 Method 1: Estimation of scenario uncertainty from single scenarios of multiple models

In an ideal ensemble of simulations, the scenario uncertainty would be estimated by first calculating the multi-model mean trend of an ensemble of CCM simulations forced with each chosen scenario, and then estimating the standard deviation of the scenario specific multi-model mean. Given the small number of scenario runs available, one way to make progress is to attempt to estimate the smooth evolution of a large multi-model ensemble based on the scenario runs available. These estimate of the multi-model ensemble mean for each scenario will be denoted as the *pseudo multi-model mean timeseries* for each scenario. Scenario uncertainty is then estimated by calculating the standard deviation of the multi-model mean for the reference scenario and the pseudo multi-model mean time series for the other GHG scenarios.

It is assumed that an estimate of the shift of the multi-model mean in each scenario can be made by first measuring the difference in the evolution of ozone in each model between its simulation in that scenario and in the REF-B2 scenario ($shf(t)_m$). For example,

$$shf(t)_{WACCM} = x(t)_{WACCM,SRESB1} - x(t)_{WACCM,REF-B2}$$

is the shift for the WACCM model. An estimate of the shift of the multi-model mean for each scenario can then be made by taking the mean of the model-specific shifts,

$$\overline{shf(t)} = \sum_m shf(t)_m.$$

The pseudo multi-model mean time series ($pmmt$) is then estimated from the REF-B2 multi-model smooth time series ($mmt(t)$) as (in this example for the SRESA2 scenario):

$$pmmt(t)_{SRESB1} = mmt(t) + \overline{shf(t)_{SRESB1}}$$

This procedure is equivalent to assuming that the difference between each model integration and the multi-model mean in each scenario is the same as the difference between that model and the multi-model mean in the REF-B2 run. It is also consistent with the assumption that the variance of the model ensemble remains the same in each scenario run. This is a simple assumption, but it has the advantage that it is easily reproducible and allows progress to be made in assessing the relative sizes of scenario and model uncertainty.

To test this procedure, it was applied to a set of six CCM simulations which assumed GHG concentrations remained fixed at their 1960 levels from 1960-2100 (note that these simulations are not used elsewhere in this study, but are extensively documented in Eyring et al. (2010)). This set of simulations will be referred to as the fGHG simulations). Fig. 1 shows the smooth trend in global mean total column ozone (relative to 1960 values) of the multi-model mean for the REF-B2 simulations of the six models (solid black line) and the multi-model mean for the fGHG simulations of the six models (dashed black line). The procedure for the estimation of the pseudo multi-model mean trend for the fGHG scenario based on a limited number of CCM simulations is illustrated by the grey, blue and yellow lines. In each case, the grey, blue or yellow line represents

an estimate of the fGHG multi-model mean calculated by adding $shf(t)$ calculated from a single CCM onto to the multi-model mean of the REF-B2 simulations. Each of the grey lines is an estimate of the dashed black line calculated by a simple average of the six fGHG simulations. Clearly, for a single CCM, the estimate of the multi-model

5 mean can be very different to the true multi-model mean. In the case of CCSRNIES (yellow line) and WACCM (blue line), which, along with GEOSCCM, ran SRES GHG sensitivity simulations used in the later analysis, both models somewhat overestimate the sensitivity of the global mean ozone column around the time of maximum ozone depletion, but by the end of century are close to the multi-model mean.

10 For the global mean and similar tests in other geographical regions (not shown) when the pseudo multi-model mean is calculated from two model integrations (red line in this example) the estimate is closer to the true multi-model mean. Of course, as more and more scenario simulations are added to the average shift calculation the pseudo multi-model mean timeseries will better approximate the true multi-model mean. The

15 example calculations here show that for two models, the pseudo multi-model mean is reasonably close to the true multi-model mean and generally approximates the main features of the true multi-model mean.

3.2.2 Method 2: Estimating scenario uncertainty from runs of a single CCM

An alternative method for estimating scenario uncertainty is to use multiple simula-

20 tions of a single CCM forced with a range of different scenarios. A set of simulations of this type is available for the RCP forcing scenarios from the CAM3.5 CCM. Since three simulations of this model with different RCP scenarios are available it is trivial to calculate the time evolving scenario uncertainty by computing the standard deviation of the ensemble of three CAM3.5 simulations. This estimate of scenario uncertainty

25 can be compared to the estimates derived from method 1, and together with the estimates of model uncertainty and internal variability from the REF ensemble to provide a breakdown of the uncertainty in ozone projection.

For the RCP scenario simulations of CAM3.5, there are significant differences in the

tropospheric ozone column between the runs, because tropospheric ozone pre-cursors vary between the different scenarios. Since the focus of this study is on uncertainties in projections of stratospheric ozone, when calculating scenario uncertainty from the RCP simulations of CAM3.5 only the ozone column above 200hPa is considered.

5 The major disadvantage of this approach is that it relies completely on an accurate simulation of the sensitivity of stratospheric ozone to each forcing scenario by the CAM3.5 model. If this model over or under estimates the sensitivity of stratospheric ozone to changes in for example GHG amounts then the estimate of scenario uncertainty will be similarly biased.

10 **4 Results**

4.1 Global mean ozone

To introduce the concept of our quantification of uncertainty we first show a time series of the global and annual mean, total column ozone (Fig. 2). The modeled evolution of ozone for both the past and projected future is shown (in grey), calculated by computing the ensemble mean of the REF and SRES simulations and is identical in both panels. 15 Estimates of the model uncertainty and internal variability are drawn from the REF-B2 simulations, and are identical in the two panels. Shaded regions show the one standard deviation uncertainty associated with each of the three uncertainty components. The two panels use the two alternative methods of estimating scenario uncertainty using the SRES scenario simulations (top panel) and the RCP scenario simulations (bottom 20 panel).

Considering first the top panel, it is clear that when using the SRES scenario simulations, total uncertainty is similar throughout the 21st century, with a small reduction around the 2040s as ozone returns to its 1980 values. The relative contributions of 25 model uncertainty (blue) and scenario uncertainty (green), however, change markedly over the course of the 21st century. For short lead times, close to peak ozone depletion,

model uncertainty is large, and is likely related to the variation in stratospheric inorganic chlorine (Cl_y) (Eyring et al., 2007) and related variations in chemistry, dynamics and transport between the models. By the end of the 21st century, when Cl_y concentrations have returned to pre-1980 levels, model uncertainty is significantly smaller.

5 It is worth noting in passing that while running initial condition ensembles of ozone projections would help to better determine the mean smooth trend of ozone, the uncertainty in the projections related to internal atmospheric variability would remain. As described in the methods section, internal variability is estimated from the model integrations themselves and is assumed to be constant throughout the period of the runs. Examination of the time-evolving residuals for both the global ozone and other
10 geographical regions revealed that in general there are no significant changes in the internal variability estimated from the 13 models in the REF-B2 ensemble. For global mean values, there are variations of around 0.1 DU in the one standard deviation estimate of internal variability, with a small trend toward smaller internal variability over the course of the 21st century. For the polar regions, there is slightly more variation in the estimate of internal variability (as expected given the smaller averaging area and shorter averaging period) with larger variations over Antarctica of around 2-3 DU and a peak in internal variability near to the period of maximum ozone depletion. We consider these variations to be small enough that an assumption of constant internal variability
15 is a valid first approximation for the analysis conducted in this study. It is also important to note that since internal variability is estimated from the models themselves, it could be larger than the estimate shown here if the models under-represent tropospheric and stratospheric internal variability, but given the limited observational record (black dots in Fig. 2) it is difficult to assess the magnitude of the internal variability. Additionally, if projections of annual mean global mean ozone were considered, (rather than
20 projections of decadal mean global mean ozone) internal variability would be a larger contributor to the total uncertainty in ozone projections (increasing by around a factor of 2).

From the middle of the 21st century, scenario uncertainty begins to play a bigger role

in determining the total uncertainty, as expected, since the input GHG concentrations only start to significantly diverge in the second half of the 21st century. By the end of the 21st century, when GHG concentrations in the various scenarios have significantly diverged, their impacts on stratospheric temperatures, and hence on the rates of gas-phase ozone loss cycles and changes in stratospheric circulation can be clearly seen by the growth of the scenario uncertainty component. Despite this growth in scenario uncertainty in the SRES simulations, model uncertainty remains the largest contributor to the total uncertainty in global ozone throughout the 21st century.

In contrast to the SRES scenario simulations, scenario uncertainty is a larger component of the total uncertainty in the RCP scenario simulations. Given the larger spread in GHG amounts in the RCP simulation this would have been expected a priori. However, scenario uncertainty from the RCP scenarios begins to dominate uncertainty in global mean ozone only by the later third of the 21st century, and not before global mean ozone has returned to its pre-1980 values. The enhanced scenario uncertainty also means that the total uncertainty in projections of total ozone is larger, but only by a factor of around 20% by the end of the 21st century when compared to the total uncertainty computed using SRES scenario uncertainty.

One benefit of dividing ozone projection uncertainty into model, scenario and internal variability components is that it gives an indication of the benefit in narrowing projection uncertainty that can be made by improving CCMs and where this improvement is likely to be most beneficial. In discussing these improvements, it is assumed that by improving CCMs, model uncertainty will be reduced, whilst scenario uncertainty will remain constant, since scenario uncertainty is primarily related to future global development paths. Under this assumption, results for both the SRES and RCP scenario cases suggest that it should be possible to narrow uncertainty in ozone projections throughout the 21st century by reducing model uncertainty. Projections of the increase in global column ozone amounts above 1980 values by the end of the 21st century are more dependent upon the concentrations of GHGs at this time. In general, scenarios with larger emissions of GHGs (SRESA2, RCP8.5) tend to produce larger amounts of

ozone at the end of the 21st century.

4.2 Uncertainty in ozone projections in different geographical regions

A similar quantification of uncertainty can be performed for total column ozone projections for various geographical regions and seasons to highlight the different behaviour of ozone and the importance of different physical processes in determining future ozone behaviour. In this analysis we consider similar regions to WMO assessments of future ozone listed in Table 2.

Uncertainty in the three different components, along with the total uncertainty is displayed in units of one standard deviation in Fig. 3. Broadly, the same behaviour as identified in the global ozone concentrations is seen, with total uncertainty (black lines) roughly constant throughout the 21st century when SRES simulations are used to assign scenario uncertainty. In general, there is also good agreement in both the estimated magnitude and temporal evolution of scenario uncertainty between the two independent estimates derived from the SRES (solid green lines) and RCP (dashed green lines) scenarios. Note that scenario uncertainty for both the SRES and RCP estimates is close to but not equal to zero at initial time, at this time in both the SRES and RCP scenarios there is almost no difference in GHG concentrations, suggesting that the small offset here is related to uncertainties in the estimation of the $S(t)^2$ term.

In most regions, when SRES scenarios are considered, model uncertainty remains the dominant term in the uncertainty budget throughout the 21st century. When scenario uncertainty is calculated from the RCP scenarios, in some regions (Global mean, SH mid-latitudes) total uncertainty undergoes a transition from model dominated uncertainty to scenario dominated uncertainty at some time in the last third of the century. However, there are significant differences to this behaviour in different regions.

In the tropics, model uncertainty remains a large contributor to the uncertainty throughout the 21st century and grows slightly from 2000 to 2100. SRES and RCP scenario uncertainty remains small throughout the 21st century. The evolution of tropical column ozone is determined by the balance between increases in upper stratospheric

ozone due to stratospheric cooling and decreases in lower stratospheric ozone due to increases in the strength of the Brewer-Dobson circulation. Comparison of vertically resolved ozone changes (not shown) indicate that a robust feature in the GHG sensitivity simulations is that, in the tropical upper stratosphere, the lower GHG scenarios generally result in lower ozone (due to smaller GHG-induced ozone increases), whereas in the lower stratosphere the lower GHG scenarios result in higher ozone (due to less tropical upwelling that advects ozone-poor air from the troposphere). However, in the upper stratosphere, factors other than GHG induced cooling play a role in the ozone decreases, so that e.g. the ozone evolution in the SRES A2 and SRES A1b simulations are very similar (Oman et al. , 2010a).

All CCMs also consistently predict an increase of tropical upwelling with increasing GHGs, but the significant variation in Brewer-Dobson circulation strength that was found amongst the CCMVal2 models (Chapter 4, SPARC CCMVal (2010), see also Fig. 7 of (Eyring et al., 2010)) plays a significant role in determining the large model uncertainty and its growth with time (since models with larger tropical upwelling mass flux also tend to exhibit larger trends in mass flux over the 21st century (Chapter 4, SPARC CCMVal (2010))). Consequently, uncertainty in projections of tropical ozone is likely to significantly benefit from improvements in CCMs transport schemes as well as improvements in the underlying dynamics of the General Circulation Models (GCMs).

Over the polar regions, while internal variability plays a larger role than in the other geographical regions, model uncertainty remains the dominant term in the total uncertainty for most of the 21st century. The role of internal variability is most prominent over the Arctic, where it is larger than scenario uncertainty till around 2050 for both scenario uncertainty estimates.

Model uncertainty over the Antarctic reduces markedly over the course of the 21st century, suggesting that most of this uncertainty is related to the representation of heterogeneous chemical ozone depletion, as expected in the SH. As ODS concentrations are reduced, and ozone concentrations recover to their pre-1980 values, CCMs are in greater agreement. However, even by the end of the 21st century, model uncer-

tainty remains as large or larger than SRES scenario uncertainty over the Antarctic. In contrast, over the Arctic model uncertainty increases throughout the 21st century. One hypothesis for the growth of model uncertainty over the Arctic is that because the response to GHG changes has a wide spread between models, model uncertainty increases linearly with time as the spread in the changes becomes larger (but the response to GHG changes evolves consistently within each model).

While scenario uncertainty for both SRES and RCP scenarios grows over the 21st century for both polar cap regions there is a larger difference between the estimation of this uncertainty from the SRES and RCP scenarios over the Arctic than over the Antarctic. For the Antarctic, scenario uncertainty is broadly similar for the SRES and RCP scenario sets, and has similar size to model and internal variability from around 2050. For the Arctic, RCP scenario uncertainty grows much more strongly than SRES scenario uncertainty over the 21st century, but is smaller than model uncertainty in both cases. For regions like the Arctic where model uncertainty is large, scenario uncertainty estimates likely have larger error bars because they rely on runs of a small number of CCMs which may have different responses to GHG changes. Of the models used to estimate scenario uncertainty, Arctic ozone in the last third of the 21st century in the REF-B2 simulations is close to the multi-model mean for CAM3.5 and WACCM, larger than the multi-model mean in GEOSCCM and smaller than the multi-model mean in CCSRNIES (Fig. 9.11, Chapter 9, SPARC CCMVal (2010)).

Over the polar caps, these results suggests that there is very different potential benefits to improving CCMs. Over the Arctic, reducing model uncertainty could have a significant impact on the total uncertainty throughout the 21st century, since this term is large and increases substantially from 2000 to 2100. Over the Antarctic, the reduction in model uncertainty and the growth of scenario uncertainty means that the impact of a reduction in model uncertainty would be largest for the first two-thirds of the 21st century. The Antarctic also has a minimum in total uncertainty around 2050, similar to that seen in projections of global mean and regional surface temperature (Hawkins and Sutton, 2009).

4.3 Signal-to-noise ratio of ozone projections

As in Hawkins and Sutton (2009, 2010), the uncertainty estimates shown above can be used to determine other important properties of the ozone projections, such as the signal-to-noise ratio (S/N),

$$S/N = \frac{\text{multi-model mean ozone change}}{\text{total uncertainty}}.$$

5 Where S/N exceeds one, there is a robust change in ozone predicted by the multi-model ensemble. For near-surface temperature, Hawkins and Sutton (2009) used S/N to quantify periods in which the potential information content of projections was greatest. In the case of stratospheric ozone projections, we consider the change in total column ozone from 2000, the year chosen as the start of our projection period and close to the minimum in ozone column amounts. Where S/N is greater than one, this suggests that the model ensemble predicts a detectable change from 2000 values. Fig. 4 shows estimates of S/N for three of the geographical regions described above, with SRES scenario uncertainty (solid), RCP scenario uncertainty (dotted) and without scenario uncertainty (dashed) included. The S/N is estimated with and without scenario uncertainty to allow a more direct comparison with many previous studies and assessments of future ozone amounts which have tended to focus on a single GHG scenario.

15 It is clear from Fig. 4 that, in all three regions, including either SRES or RCP scenario uncertainty significantly degrades the ability of the model ensemble to predict changes in column ozone amounts, as expected. The date at which projections in all regions have S/N greater than one, and hence predict a statistically robust change in column ozone, is delayed by around 5 years in the global mean and 10 years for the Antarctic and tropical cases. In the global mean case, if the RCP scenario uncertainty is used instead of the SRES scenario uncertainty, S/N remains approximately constant after 2050, as scenario uncertainty becomes larger than model uncertainty. When scenario uncertainty is not considered, the signal-to-noise ratio continues to grow, indicating

increased confidence in projections. This is an alternative way of showing that CCM improvements would likely benefit projections of column ozone by the largest amount in the early part of the 21st century, up to the date of ozone return to 1980 values.

Over the Antarctic, although the signal of ozone increases from 2000 values is larger than the global mean signal, the signal to noise ratio is smaller because of the greater total uncertainty in all three components. The difference between the estimates of S/N using SRES and RCP scenario uncertainty estimates are very small.

In the tropical region, the evolution of S/N is somewhat different, with a noticeable peak in mid-century (where the difference between current and future column amounts is greatest) before a decline in the second half of the century. The difference between S/N with and without scenario uncertainty is less pronounced for the tropics, with only a small reduction in S/N when either SRES or RCP scenario uncertainty is included. It should also be noted, however, that the size of S/N in the tropics is smaller than for either the global mean or the Antarctic.

4.4 Uncertainty in ozone return date

Finally, we consider how different sources of uncertainty influence the calculation of the return date for total-column ozone to its 1980 value. To do this, a different calculation procedure is required.

It is assumed that the calculation of return date will be post-hoc (i.e. based on the transition of a *smooth* estimate of the long-term trend in ozone across the zero line). It is therefore simple to estimate the uncertainty in return date due to model uncertainty (by calculating the variance of return dates across the model ensemble) and due to scenario uncertainty (by calculating the variance of return dates across the REF-B2 and pseudo multi-model mean estimates for each scenario using method 1 and across the RCP ensemble using method 2). To calculate the contribution of internal variability to uncertainty in return date, a Monte-Carlo approach is adopted. For each model, the residuals from the smooth model trend are randomly permuted and added back to the model trend to give an alternative evolution of column ozone. By fitting a smooth

trend to this new synthetic data and estimating the return date, and then repeating this procedure 10,000 times, an estimate of the uncertainty in return date due to internal variability in each model can be made. The mean of the internal variability uncertainty for the 13 models in the REF-B2 ensemble is used to estimate the overall uncertainty due to internal variability. Note that this estimate may be slightly smaller than the real contribution of internal variability to the uncertainty in return date, since we neglect to preserve the decadal variability present in the noise term when we permute the residuals using this method.

Fractional standard deviation in return date for Arctic, Antarctic and global ozone is shown in Fig. 5 as bars which add up to 1 (or 100% of the total standard deviation). As in previous calculations, it is clear that a substantial part of the uncertainty in return date is associated with model uncertainty. This conclusion is not dependent upon the source of the scenario uncertainty estimate, although scenario uncertainty in return date is greater for the SRES simulations over the Antarctic and greater for the RCP simulations over the Arctic. Over these three key averaging regions, the maximum contribution to the fractional standard deviation in ozone return date is around 45% for the Antarctic using SRES scenario uncertainty and the minimum contribution is around 10% for the Antarctic using RCP scenario uncertainty estimates. There is therefore strong potential to improve estimates of the return date for ozone in these regions by continued CCM development.

5 Conclusions

In this study, an attempt to quantify the uncertainty in future projections of total ozone column amounts has been made. In particular, a method introduced and tested for the CMIP3 models has been adapted and developed for models run as part of the Chemistry-Climate Model Validation activity (CCMVal2). Projection uncertainty has been divided into three components representing model uncertainty, scenario uncertainty and uncertainty due to internal variability of the system. Scenario uncertainty

has been estimated using two different methods which used scenario simulations from both SRES and RCP future emission scenarios.

When considering scenario uncertainty in the SRES scenarios, total uncertainty in ozone projections is roughly constant throughout the 21st century and is dominated by model uncertainty. Although scenario uncertainty in the SRES scenario simulations grows throughout the 21st century, for most geographical regions and seasons it remains smaller than model uncertainty.

When considering scenario uncertainty in the RCP scenarios, total uncertainty grows slightly throughout the 21st century, due to a larger growth in scenario uncertainty than for the SRES simulations. There is an important transition between model uncertainty and scenario uncertainty for some regions (notably the global mean ozone and ozone in the SH mid-latitudes) with scenario uncertainty dominating projections by the last third of the 21st century. This has important consequences for the ability of models to predict significant changes to ozone column amounts, which is degraded by scenario uncertainty during this period.

In general, the consistency of the two independent estimates of scenario uncertainty for most diagnostics lends confidence in these estimates. However, the application of the scenario uncertainty should proceed with some caution. The GHG scenarios used by the models are designed to provide an estimate of the potential range of future emissions based on vastly different global economic and technological development. The scenarios are not equally likely, do not take into account all possible futures and should not be taken to indicate an upper bound on the range of potential future emissions. The newer RCP scenarios have a larger spread in emissions of GHGs compared to the SRES scenarios used, partly due to the inclusion of the very low GHG emission scenario (RCP2.6) which represents a future scenario with peak radiative forcing around 2050. This scenario does not have an obvious equivalent in the SRES scenarios considered.

An additional source of uncertainty in projections of ozone which we did not consider in this study are the mean state biases between CCMs. In all cases, CCM time series

of ozone were corrected to their 1980 values as is customary in the CCM community. As shown in, for example, Chapter 9 of the CCMVal report, in many regions models have large basic state biases in the total ozone column. If these uncertainties were taken into account they would likely further increase the relative contribution of model uncertainty to the total uncertainty estimates we have derived.

The quantification of uncertainty derived in our study also provides some insight into the ability of the Chemistry-Climate modelling community to refine projections of future ozone amounts by the continued development and improvement of CCMs. The results of our analysis suggest that this work would be beneficial for projections of ozone amounts in the first half of the 21st century in most regions and particularly in the determination of dates of return of ozone amounts to pre-1980 values (a key metric for ozone defined by the 2006 WMO/UNEP ozone assessment). In future, similar methods could be used to quantify the uncertainty in projections of other parameters of the stratospheric climate system in order to attribute the scenario uncertainty to different physical processes. It is also important to note, however, that the relationship between spread and skill for geophysical models is far from trivial. While it is reasonable to wish to reduce overall uncertainty in projections of future ozone, it does not necessarily follow that narrowing the uncertainty in these projections will also improve their skill.

The quantification of uncertainty in ozone projections in this study is far from ideal and is sensitive to the assumptions and models used, particularly in constructing estimates of scenario uncertainty. It is hoped that the results will provide guidance to the chemistry-climate community on the benefits that running large multi-scenario ensembles of CCMs would provide. It is clear from our initial analysis that the benefits in fully quantifying and exploring the projection uncertainty from running multi-scenario ensemble simulations vary substantially between different geographical regions, seasons and time-periods. We re-iterate that an ideal ensemble of simulations for the quantification of uncertainty introduced here would require a large ensemble of models to all run at least three different GHG scenarios.

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References

- Akiyoshi, H., L. B. Zhou, Y. Yamashita, K. Sakamoto, M. Yoshiki, T. Nagashima, M. Takahashi, J. Kurokawa, M. Takigawa, and T. Imamura, (2009) A CCM simulation of the breakup of the Antarctic polar vortex in the years 1980-2004 under the CCMVal scenarios, *J. Geophys. Res.*, 114, D03103, doi:10.1029/2007JD009261.
- Austin, J., and R.J. Wilson, (2009) Sensitivity of polar ozone to sea surface temperatures and chemistry, *J. Geophys. Res.*, submitted.
- Barnett J. J., J. T. Houghton and J. A. Pyle (1974) The temperature dependence of the ozone concentration near the stratopause, *Quart. J. Roy. Met. Soc.*, 101, 245-257
- Butchart N., A. A. Scaife, M. Bourqui, J. de Grandpré, S. H. E. Hare, J. Kettleborough, U. Langematz, E. Manzini, F. Sassi, K. Shibata, D. Shindell, M. Sigmond (2006) Simulations of anthropogenic change in the strength of the Brewer-Dobson circulation, *Climate Dynamics*, 27, 727-741, doi:10.1007/s00382-006-0162-4
- Chipperfield, M. P., and W. Feng, Comment on: Stratospheric Ozone Depletion at northern mid-latitudes in the 21st century: The importance of future concentrations of greenhouse gases nitrous oxide and methane (2003) *Geophys. Res. Lett.*, 30, 1389, doi: 10.1029/2002GL016353.
- Clarke, L., J. Edmonds, H. Jacoby, H. Pitcher, J. Reilly, and R. Richels, Scenarios of Greenhouse Gas Emissions and Atmospheric Concentrations. (2007) Sub-report 2.1A of Synthesis and Assessment Product 2.1 by the U.S. Climate Change Science Program and the

- Subcommittee on Global Change Research. Department of Energy, Office of Biological & Environmental Research, Washington, 7 DC., USA, 154 pp.
- de Grandpré, J., S. R. Beagley, V. I. Fomichev, E. Griffioen, J. C. McConnell, A. S. Medvedev and T. G. Shepherd, (2000) Ozone climatology using interactive chemistry: Results from the Canadian Middle Atmosphere Model, *J. Geophys. Res.*, 105, 26475-26491.
- 5 Déqué, M., Frequency of precipitation and temperature extremes over France in an anthropogenic scenario: model results and statistical correction according to observed values. (2007) *Global and Planetary Change*, 57, 16-26.
- Egorova, T., E. Rozanov, V. Zubov, E. Manzini, W. Schmutz, and T. Peter, (2005) Chemistry-climate model SOCOL: a validation of the present-day climatology, *Atmos. Chem. Phys.*, 5, 1557-1576.
- 10 Eyring, V., N. Butchart, D. W. Waugh, H. Akiyoshi, J. Austin, S. Bekki, G. E. Bodeker, B. A. Boville, C. Brhl, M. P. Chipperfield, E. Cordero, M. Dameris, M. Deushi, V. E. Fioletov, S. M. Frith, R. R. Garcia, A. Gettelman, M. A. Giorgetta, V. Grewe, L. Jourdain, D. E. Kinnison, E. Mancini, E. Manzini, M. Marchand, D. R. Marsh, T. Nagashima, P. A. Newman, J. E. Nielsen, S. Pawson, G. Pitari, D. A. Plummer, E. Rozanov, M. Schraner, T. G. Shepherd, K. Shibata, R. S. Stolarski, H. Struthers, W. Tian, and M. Yoshiki, (2006) Assessment of temperature, trace species and ozone in chemistry-climate model simulations of the recent past, *J. Geophys. Res.*, 111, D22308, doi:10.1029/2006JD007327.
- 15 Eyring, V., D. W. Waugh, G. E. Bodeker, E. Cordero, H. Akiyoshi, J. Austin, S. R. Beagley, B. Boville, P. Braesicke, C. Brhl, N. Butchart, M. P. Chipperfield, M. Dameris, R. Deckert, M. Deushi, S. M. Frith, R. R. Garcia, A. Gettelman, M. Giorgetta, D. E. Kinnison, E. Mancini, E. Manzini, D. R. Marsh, S. Matthes, T. Nagashima, P. A. Newman, J. E. Nielsen, S. Pawson, G. Pitari, D. A. Plummer, E. Rozanov, M. Schraner, J. F. Scinocca, K. Semeniuk, T. G. Shepherd, K. Shibata, B. Steil, R. Stolarski, W. Tian, and M. Yoshiki, (2007), Multimodel projections of stratospheric ozone in the 21st century, *J. Geophys. Res.*, 112, D16303, doi:10.1029/2006JD008332.
- 20 Eyring, V., I. Cionni, G. E. Bodeker, A. J. Charlton-Perez, D. E. Kinnison, J. Scinocca, D. W. Waugh, H. Akiyoshi, S. Bekki, M. P. Chipperfield, M. Dameris, S. Dhomse, S. M. Frith, H. Garny, A. Gettelman, A. Kubin, U. Langematz, E. Mancini, M. Marchand, T. Nakamura, L. D. Oman, S. Pawson, G. Pitari, D. Plummer, E. Rozanov, T. G. Shepherd, K. Shibata, W. Tian, P. Braesike, S. C. Hardiman, J.F. Lamarque, O. Morgenstern, J. A. Pyle, and Y. Yamashita (2010) Multi-model assessment of ozone return dates and ozone recovery in CCMVal2 mod-
- 25
- 30

- els, *Atmos. Chem. Phys.*, submitted.
- Garcia R. R., D. R. Marsh, D. E. Kinnison, B. A. Boville, and F. Sassi, (2007) Simulation of secular trends in the middle atmosphere, 1950-2003, *J. Geophys. Res.*, 112, D09301, doi:10.1029/2006JD007485.
- 5 Haigh, J. D., and J. A. Pyle (1982), Ozone perturbation experiments in a two-dimensional circulation model. *Q. J. R. Meteorol. Soc.*, 108, 551-574.
- Hawkins E and Sutton R., (2009), 'The potential to narrow uncertainty in regional climate predictions', *Bull. Am. Met. Soc.*, 90, p1095, doi: 10.1175/2009BAMS2607.1
- Hawkins E and Sutton R., (2010), 'The potential to narrow uncertainty in projections of regional precipitation change', *Clim. Dyn.*, submitted
- 10 Jonsson, A. I., V. I. Fomichev and T. G. Shepherd (2009) The effect of nonlinearity in CO2 heating rates on the attribution of stratospheric ozone and temperature changes, *Atmos. Chem. Phys.*, 9, 8447-8452
- Jourdain, L., S. Bekki, F. Lott, and F. Lefvre, (2008) The coupled chemistry-climate model LMDz-REPROBUS: description and evaluation of a transient simulation of the period 1980-1999, *Annales Geophysicae*, 26, 6, 1391-1413.
- 15 Lamarque J.-F., D. E. Kinnison, P. G. Hess, F. M. Vitt, (2008) Simulated lower stratospheric trends between 1970 and 2005: Identifying the role of climate and composition changes, *J. Geophys. Res.*, 113, D12301, doi:10.1029/2007JD009277.
- 20 Lamarque, J.-F., T. C. Bond, V. Eyring, C. Granier, A. Heil, Z. Klimont, D. Lee, C. Liousse, A. Mieville, B. Owen, M. G. Schultz, D. Shindell, S. J. Smith, E. Stehfest, J. Van Aardenne, O. R. Cooper, M. Kainuma, N. Mahowald, J. R. McConnell, V. Naik, K. Riahi and D. P. van Vuuren, Historical (1850-2000) gridded anthropogenic and biomass burning emissions of reactive gases and aerosols: methodology and application, (2010), *Atmos. Chem. Phys.*, submitted.
- 25 Meehl, G. A. and Hibbard, K. A. A Strategy for Climate Change Stabilization Experiments with AOGCMs and ESMs (2007) WCRP Informal Report No. 3/2007, ICPO Publication No. 112, IGBP Report No. 57, World Climate Research Programme, Geneva
- Moss T., M. Babiker, S. Brinkman, E. Calvo, T. Carter, J. Edmonds, I. Elgizouli, S. Emori, L. Erda, K. Hibbard, R. Jones, M. Kainuma, J. Kelleher, J.-F. Lamarque, M. Manning, B. Matthews, G. Meehl, L. Meyer, J. Mitchell, N. Nakic'enovic', B. O'Neill, T. Pichs, K. Riahi, S. Rose, P. Runci, R. Stouffer, D. van Vuuren, J. Weyant, T. Wilbanks, J. P. van Ypersele, and M. Zurek, (2008), *Towards New Scenarios for Analysis of Emissions, Climate Change, Impacts, and Response Strategies.*, Intergovernmental Panel on Climate Change, Geneva, 132 pp.,

available at: <http://www.ames.ucar.edu/docs/>.

- Morgenstern, O., P. Braesicke, M. M. Hurwitz, F. M. O'Connor, A. C. Bushell, C. E. Johnson, and J. A. Pyle, (2008) The World Avoided by the Montreal Protocol, *Geophys. Res. Lett.*, 35, L16811, doi:10.1029/2008GL034590.
- 5 Morgenstern, O., et al., (2009) Evaluation of the new UKCA climate-composition model. Part 1: The stratosphere. *Geosci. Model Dev.*, 1, 43-57.
- Morgenstern, O., M. A. Giorgetta, K. Shibata, V. Eyring, D.W. Waugh, T. G. Shepherd, H. Akiyoshi, J. Austin, A. J. G. Baumgaertner, S. Bekki, P. Braesicke, C. Brhl, M. P. Chipperfield, D. Cugnet, M. Dameris, S. Dhomse, S. M. Frith, H. Garny, A. Gettelman, S. C. Hardiman, 10 M. I. Hegglin, D. E. Kinnison, J.-F. Lamarque, E. Mancini, E. Manzini, M. Marchand, M. Michou, T. Nakamura, J. E. Nielsen, G. Pitari, D. A. Plummer, E. Rozanov, J. F. Scinocca, D. Smale, S. Strahan, H. Teysedre, M. Toohey, W. Tian, and Y. Yamashita, (2010) Review of present-generation stratospheric chemistry-climate models and associated external forcings, *J. Geophys. Res.*, submitted.
- 15 Nakicenovic N., J. Alcamo, G. Davis, B. de Vries, J. Fenhann, S. Gaffin, K. Gregory, A. Grbler, T. Y. Jung, T. Kram, E. Lebre La Rovere, L. Michaelis, S. Mori, T. Morita, W. Pepper, H. Pitcher, L. Price, K. Riahi, A. Roehrl, H.-H. Rogner, A. Sankovski, M. Schlesinger, P. Shukla, S. Smith, R. Swart, S. van Rooijen, N. Victor, Z. Dadi (2000) IPCC Special Report on Emissions Scenarios
- 20 Oman L., D. W. Waugh, S.R. Kawa, R. S. Stolarski, A. R. Douglass, P. A. Newman (2010) Mechanisms and feedbacks causing changes in upper stratospheric ozone in the 21st century, *J. Geophys. Res.*, in press
- Oman, L. D., D. Plummer, D. W. Waugh, J. Austin, J. Scinocca, A. R. Douglass, R. J. Salawitch, H. Akiyoshi, S. Bekki, P. Braesicke, N. Butchart, M. Chipperfield, S. Dhomse, V. Eyring, S. Frith, R. R. Garcia, A. Gettelman, S. C. Hardiman, D. Kinnison, J. F. Lamarque, E. Mancini, M. Marchand, M. Michou, O. Morgenstern, S. Pawson, G. Pitari, T. Peter, J. Pyle, E. Rozanov, T.G. Shepherd, K. Shibata, R. Stolarski, H. Teysedre, and W. Tian, (2010) Multi-model assessment of the factors driving the ozone evolution over the 21st century, *J. Geophys. Res.*, submitted.
- 25 30 Pawson, S., R. S. Stolarski, A. R. Douglass, P. A. Newman, J. E. Nielsen, S. M. Frith, and M. L. Gupta, (2008) Goddard Earth Observing System chemistry-climate model simulations of stratospheric ozone-temperature coupling between 1950 and 2005, *J. Geophys. Res.*, 113, D12103, doi:10.1029/2007JD009511.

- Pitari G., E. Mancini, V. Rizi and D. T. Shindell: Impact of Future Climate and Emission Changes on Stratospheric Aerosols and Ozone, (2002) *J. Atmos. Sci.*, 59.
- Portmann, R. W., and S. Solomon, Indirect radiative forcing of the ozone layer during the 21st century (2007) *Geophys Res. Lett.*, 34, L02813, doi:10.1029/2006GL028252.
- 5 Randeniya, L. K., P. F. Vohralik, and I. C. Plumb (2002) Stratospheric ozone depletion at northern mid latitudes in the 21st century: The importance of future concentrations of greenhouse gases nitrous oxide and methane, *Geophys. Res. Lett.*, 29 (4), 1051, doi: 10.1029/2001GL014295
- 10 Ravishankara, A. R., J. S. Daniel, and R. W. Portmann, Nitrous Oxide (N₂O): The Dominant Ozone Depleting Substance Emitted in the 21st Century (2009) *Science*, 326, 123 682-125.
- Riahi, K., A., Gruebler, and N. Nakicenovic, (2007) Scenarios of long-term socio-economic and environmental development under climate stabilization, *Technological Forecasting and Social Change*, 74, 7, 887-935.
- 15 Rosenfield, J.E., A.R. Douglass, and D.B. Considine (2002) The impact of increasing carbon dioxide on ozone recovery, *J. Geophys. Res.*, 107 (D6), 4049, doi: 10.1029/2001JD000824.
- Schraner, M., E. Rozanov, C. Schnadt-Poberaj, P. Kenzelmann, A. Fischer, V. Zubov, B. P. Luo, C. Hoyle, T. Egorova, S. Fueglistaler, S. Brnnimann, W. Schmutz and T. Peter, (2008) Chemistry-climate model SOCOL: version 2.0 with improved transport and chemistry/ micro-physics schemes, submitted to ACP, February.
- 20 Scinocca, J. F., N. A. McFarlane, M. Lazare, J. Li and D. Plummer, (2008) Technical Note: The CCCma third generation AGCM and its extension into the middle atmosphere, *Atmos. Chem. Phys.*, 8, 7055-7074.
- Shepherd T.G. (2008) Dynamics, stratospheric ozone, and climate change. *Atmos. Ocean*, 46, 117-138.
- 25 Shibata, K. and M. Deushi, (2008a) Long-term variations and trends in the simulation of the middle atmosphere 1980-2004 by the chemistry-climate model of the Meteorological Research Institute, *Annales Geophysicae*, 26, 1299-1326
- Shibata, K. and M. Deushi, (2008b) Simulation of the stratospheric circulation and ozone during the recent past (1980-2004) with the MRI chemistry-climate model, CGER's Supercomputer Monograph Report Vol.13, National Institute for Environmental Studies, Japan, 154 pp.
- 30 SPARC CCMVal, SPARC CCMVal Report on the Evaluation of Chemistry-Climate Models, V. Eyring, T. G. Shepherd, D. W. Waugh (Eds.), (2010), SPARC Report No. 5, WCRP-X, WMO/TD-No. X, <http://www.atmos.physics.utoronto.ca/SPARC>.

- 5 Teyssède H., M. Michou, H. L. Clark, B. Josse, F. Karcher, D. Olivié, V.-H. Peuch, D. Saint-Martin, D. Cariolle, J.-L. Attié, P. Nédélec, P. Ricaud, V. Thouret, R. J. van der A, A. Volz-Thomas, and F. Chéroux, (2007) A new tropospheric and stratospheric Chemistry and Transport Model MOCAGE-Climat for multi-year studies: evaluation of the present-day climatology and sensitivity to surface processes, *Atmos. Chem. Phys.*,
- 10 Tian, W, and M. P. Chipperfield, (2005) A new coupled chemistry-climate model for the stratosphere: the importance of coupling for future O3-climate predictions, *QJRMS*, 131, pp281-303.
- 15 Tian, W., M. P. Chipperfield, L. J. Gray, and J. M. Zawodny, (2006) Quasi-biennial oscillation and tracer distributions in a coupled chemistry-climate model, *J. Geophys. Res.*, 111, D20301, doi:10.1029/2005JD006871.
- van Vuuren, D., M. den Elzen, P. Lucas, B. Eickhout, B. Strengers, B. van Ruijven, S. Wonink, R. van Houdt., (2007) Stabilizing greenhouse gas concentrations at low levels: an assessment of reduction strategies and costs. *Climatic Change*, doi:10.1007/s/10584-006-9172-9.
- 20 Waugh, D. W., and V. Eyring, Quantitative performance metrics for stratospheric-resolving chemistry-climate models, (2008) *Atmos. Chem. Phys.*, 8, 5699-5713.
- Waugh, D. W., L. Oman, S. R. Kawa, R. S. Stolarski, S. Pawson, A. R. Douglass, P. A. Newman, and J. E. Nielsen (2009), Impacts of climate change on stratospheric ozone recovery, *Geophys. Res. Lett.*, 36, L03805, doi:10.1029/2008GL036223.
- World Meteorological Organization (WMO)/United Nations Environment Programme (UNEP), (2007) *Scientific Assessment of Ozone Depletion: 2006*, World Meteorological Organization, Global Ozone Research and Monitoring Project, Report No. 50, Geneva, Switzerland.

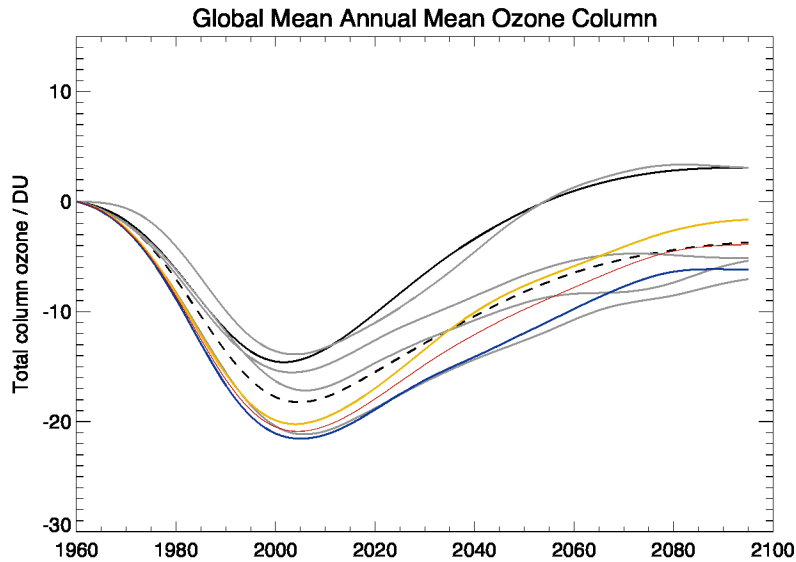


Fig. 1. Multi-model mean smooth trend in ozone for six models (CCSRNIES, CMAM, MRI, SOCOL, ULAQ and WACCM) for the standard REF scenario (black solid line) and for a scenario with fixed greenhouse gas concentrations at 1960 levels (black dashed line). Grey lines show estimate of the multi-model mean trend in the fixed GHG scenario, calculated by adding the shift between the REF and fixed GHG scenario runs and adding this shift to the multi-model mean trend from the REF scenario. Estimates produced using the CCSRNIES (yellow) and WACCM (blue) models are highlighted in colour. The estimate of the multi-model mean calculated by averaging the shift in the CCSRNIES and WACCM models is shown in the red line.

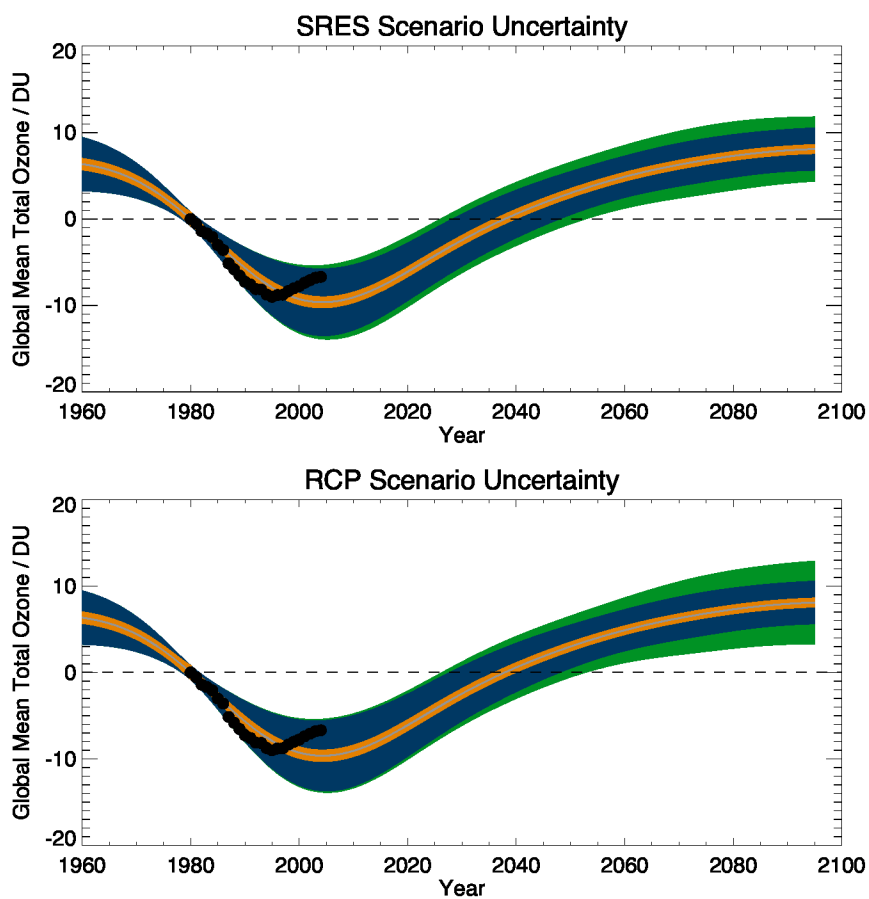
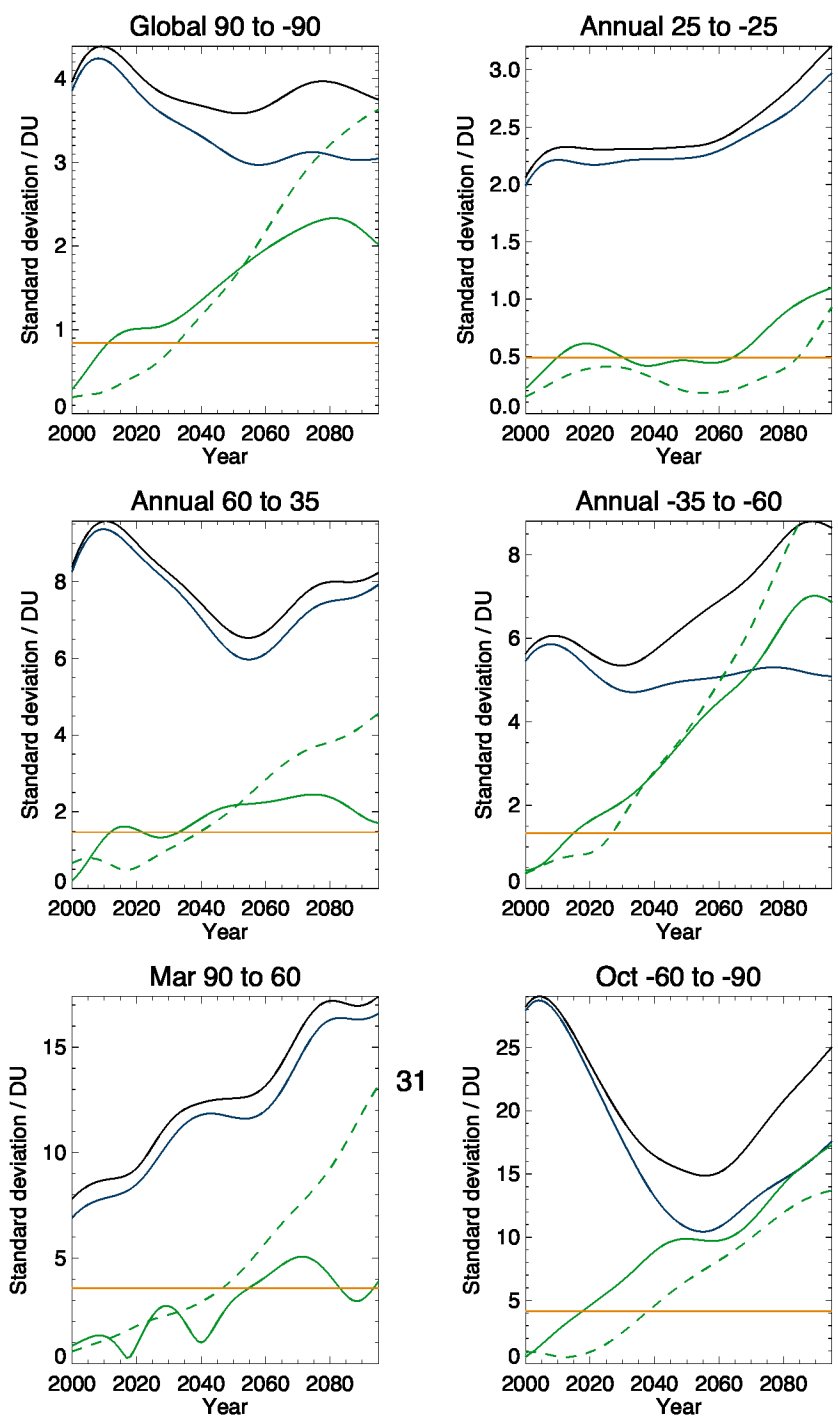


Fig. 2. Changes in annual-mean, global-mean, total-column ozone amounts relative to 1980 values. Black dots indicate observed changes from the NIWA combined and patched ozone column dataset. The grey solid line shows the multi-model, multi-scenario mean. Coloured wedges indicate the cumulative uncertainty in this projection associated with one standard-deviation estimates of internal variability (orange), model uncertainty (blue) and scenario uncertainty (green). The top panel shows calculations of scenario uncertainty using four model simulations run with SRES scenarios. The bottom panel shows calculations of scenario uncertainty using three integrations of CAM3.5 run with RCP scenarios.



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Fig. 3. Uncertainty in the predicted total-column ozone amounts, relative to 1980 values, estimated for different geographic regions. Uncertainty is expressed as the one standard deviation estimate for interannual variability (orange), model uncertainty (blue) and scenario uncertainty (green) and the total (black). Solid green lines show calculations of scenario uncertainty using four model integrations run with SRES scenarios. Dashed green lines show calculations of scenario uncertainty using three integrations of CAM3.5 run with RCP scenarios.

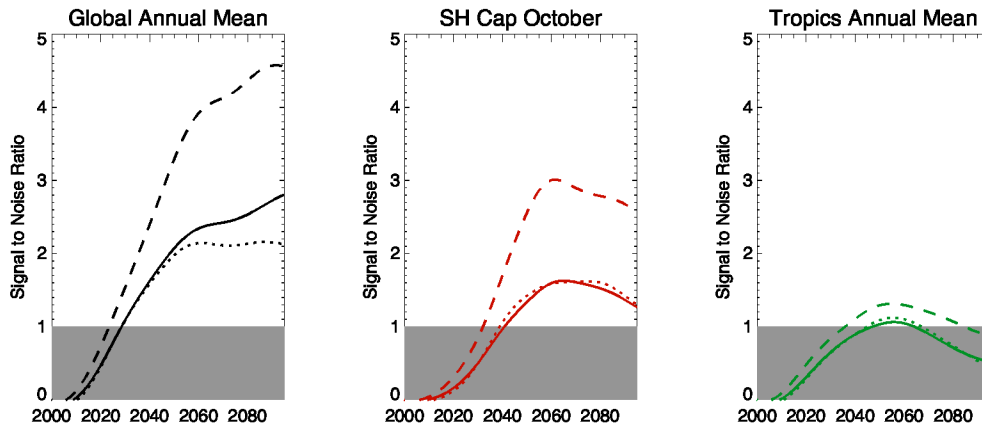


Fig. 4. Estimated signal-to-noise ratio (S/N, see text for details) of the change in total-ozone column amounts relative to the multi-model mean amount in 2000. Solid lines show estimates which include model uncertainty, internal variability and scenario uncertainty from four model integrations run with SRES scenarios. Dotted lines show estimates which include model uncertainty, internal variability and scenario uncertainty from three model integrations of CAM3.5 run with RCP scenarios. Dashed lines show estimates which exclude scenario uncertainty. Different coloured lines indicate averages over different geographical regions and seasons. Black lines show the global and annual average, red lines show the springtime SH polar cap (60-90°S, October) and the green lines show the tropical region (25°N-25°S), annual average).

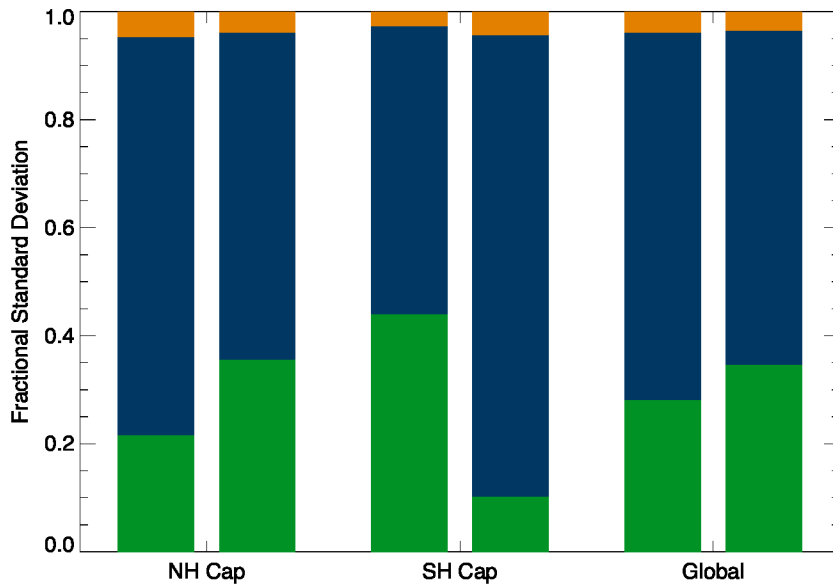


Fig. 5. Estimates of the fractional uncertainty in the date of return to 1980 values of total-column ozone amounts due to different sources of uncertainty, for three different geographical regions and seasons. The coloured areas indicate the fractional contribution of that type of uncertainty to overall uncertainty in the date of ozone return date (orange - internal variability, blue - model uncertainty, green - scenario uncertainty). Bars are shown for the northern polar cap during spring (60-90°N, March) for the southern polar cap during spring (60-90°S, October) and for the global and annual. Left-side bars show calculations of scenario uncertainty using four model integrations run with SRES scenarios. Right-side bars shows calculations of scenario uncertainty using three integrations of CAM3.5 run with RCP scenarios.

Table 1. Simulations used in the analysis

Scenario Name	GHG and ODS forcing	Models	Reference
REF-B2	SRESA1B	AMTRAC3 CAM3.5 CCSRNIES CNRM-ACM CMAM GEOSCCM LMDZrepro MRI MRI SOCOL ULAQ UMSLIMCAT UMUKCA-UCAM WACCM	Austin and Wilson (2009) Lamarque et al. (2008) Akiyoshi et al. (2009) Déqué et al. (2007) Teyssède et al. (2007) Scinocca et al. (2008) de Grandpré et al. (2000) Pawson et al. (2008) Jourdain et al. (2008) Shibata and Desushi (2008a) Shibata and Deushi (2008b) Schraner et al. (2008) Egorova et al. (2005) Pitari et al. (2002) Eyring et al. (2006) Eyring et al. (2007) Tian and Chipperfield, (2005) Tian et al. (2006) Morgenstern et al. (2008) Morgenstern et al. (2009) Garcia et al. (2007)
SCN-GHG-SRESA2	SRESA2	CCSRNIES GEOSCCM	Akiyoshi et al. (2009) Pawson et al. (2008)
SCN-GHG-SRESB1	SRESB1	CCSRNIES WACCM	Akiyoshi et al. (2009) Garcia et al. (2007)
SCN-GHG-RCP2.6	RCP2.6	CAM3.5	Lamarque et al. (2008)
SCN-GHG-RCP4.5	RCP4.5	CAM3.5	Lamarque et al. (2008)
SCN-GHG-RCP8.5	RCP8.5	CAM3.5	Lamarque et al. (2008)

Table 2. Geographical regions considered in the analysis

Region	Area	Season
Global Mean	90°S-90°N	Annual
Tropics	25°S-25°N	Annual
NH Mid-latitudes	35°N-60°N	Annual
SH Mid-latitudes	60°S-35°S	Annual
NH Polar Cap (Arctic)	60°N-90°N	March
SH Polar Cap (Antarctic)	90°S-60°S	October