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The Brewer-Dobson circulation and total ozone from seasonal to decadal time scales

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Abstract

The effect of the winter Brewer-Dobson circulation (BDC) on the seasonal and decadal evolution of total ozone in both hemispheres is investigated using satellite total ozone data and outputs from two chemistry-climate models (CCM). Combining data from both hemispheres a linear relationship between the winter cumulative extratropical 100 hPa eddy heat flux and the ozone ratio with respect to fall ozone levels exists and is statistically significant for tropical as well as polar ozone. The high correlation at high latitudes persists well into the summer months until the onset of the next winter season. The anti-correlation of the cumulative eddy heat flux with tropical ozone ratios, however, breaks down in spring as the polar vortex erodes and changes to a weak positive correlation similar to that observed at high latitudes. The inter-annual variability and decadal evolution of ozone in each hemisphere in winter, spring, and summer

- are therefore driven by the cumulative effect of the previous winter's meridional circulation. This compact linear relationship is also found in two different CCMs (EMAC-¹⁵ FUB, DLR-E39C-A) indicating that current models realistically describe the variability
- FOB, DLR-E39C-A) indicating that current models realistically describe the variability in stratospheric circulation and its climate effect on total ozone. Both models show a positive trend in the winter mean eddy heat flux (and winter BDC strength) in both hemispheres until year 2050, however the inter-annual variability (peak-to-peak) is two to three times larger than the mean change between 1960 and 2050. It is, therefore, possible to detect a shift in this compact linear relationship related to past and future
- changes in the stratospheric halogen load. A similar shift is difficult to derive from observational data since the satellite era now spanning more than thirty years is still fairly short.

1 Introduction

²⁵ For many years the main focus in studies of long-term ozone changes was the impact on chemical ozone loss from halogen containing anthropogenic ozone depleting



substances (ODS) (WMO, 1999). Chemical mechanisms leading to chemical ozone losses, in particular heterogeneous ozone loss in the polar region (polar ozone loss), are now generally well understood (Solomon, 1999; WMO, 2003, 2007, 2011). More recently studies of dynamical aspects, related to stratospheric circulation and meteo-

- ⁵ rology, on ozone changes have become more important in particular with respect to the impact of climate change on ozone and vice versa. Ozone levels have started rising in accordance with the slowdown and reversal of stratospheric halogen increases, a consequence of the global phase-out of many ODS from the successful implementation of the Montreal Protocol and its amendments (WMO, 2007, 2011). The question arises how the climate and, in particular, associated changes in the stratospheric circulation,
- how the climate and, in particular, associated changes in the stratospheric cir will impact the speed of ozone recovery in coming decades.

The Brewer-Dobson circulation (BDC) generally describes the meridional transport of trace gases from the tropics to the poles (Brewer, 1949; Dobson, 1956). This circulation is now understood as a residual circulation since the general air flow remains zonal

- and planetary waves in the extratropics disturb this flow and cause a slow meridional drift (Haynes et al., 1991; Rosenlof and Holton, 1993; Newman et al., 2001; Plumb, 2002). The transport time of air in the stratosphere from the equator to the lowermost polar stratosphere is in the range of a few years (e.g., Waugh and Hall, 2002). The poleward transport of ozone is most effective in the winter hemisphere (e.g., Chipper-
- field and Jones, 1999). Several climate models and CCMs indicate that the strength of the winter BDC will increase with increasing greenhouse gas concentrations (GHG) and accelerate the expected ozone recovery (e.g., Rind et al., 1998, 2002; Butchart and Scaife, 2001; Schnadt et al., 2002; Sigmond et al., 2004; Eichelberger and Hartmann, 2005; Butchart et al., 2006, 2010; Jiang et al., 2007; Li et al., 2007; Eyring et al., 2007; Fomichev et al., 2007; Garcia and Randel, 2008).

The planetary wave activity driving the BDC is usually described by the convergence of the Eliassen-Palm (EP) flux F, i.e. $-\nabla \cdot F$. The convergence of the EP flux in the stratosphere is a measure for easterly momentum deposited to decelerate the westerly zonal flow in winter (Newman et al., 2001). Geostrophic balance then requires a small



meridional flow component that initiates the meridional or residual circulation (Andrews et al., 1987). The vertical component of the EP flux vector, F_z , which is proportional to the eddy heat flux, v'T', is a measure of the vertical propagation of planetary waves from the troposphere. Both EP flux convergence and the eddy heat flux are frequently used to describe variations in the BDC driving.

Fusco and Salby (1999) and Salby and Callaghan (2004a) showed that inter-annual variation of the eddy heat flux well correlates with year-to-year changes of extratropical column ozone and at the same time anti-correlates with ozone changes in the tropics, documenting the year-to-year variation in hemispheric ozone transport. Part

- of the long-term trend in extratropical total ozone in the Northern Hemisphere can be explained by changes in the winter BDC (Fusco and Salby, 1999; Randel et al., 2002; Dhomse et al., 2006). The magnitude of this contribution to the overall ozone trend strongly depends on the choice and details of dynamical proxy terms used in the trend regression (Steinbrecht et al., 1998; Hood and Soukharev, 2005; Reinsel et al., 2005;
- Dhomse et al., 2006; Wohltmann et al., 2007). Various proxies have been used in the 15 trend models individually and in combination to describe dynamical ozone changes, like the potential vorticity or tropopause height, teleconnection patterns, e.g. Arctic Oscillation (AO), equivalent latitude proxy, and eddy heat flux. They are, however, not independent of each other as is discussed for the AO and eddy heat flux as an exam-

ple in the following. 20

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Figure 1 shows the evolution of middle latitude December to February (DJF) total ozone in the Northern Hemisphere starting in the 1960s. Also shown are the DJF Arctic oscillation in the middle panel (which is closely related to the North-Atlantic oscillation (NAO) defined by the surface pressure difference between Azores/Lisbon and Iceland)

and the stratospheric halogen load (inverse scale for clarity) as represented by the 25 equivalent effective stratospheric chlorine (EESC) (e.g., Newman et al., 2007) in the bottom panel.

The winter AO index represents changes in stratospheric dynamics in recent decades and they correlate with ozone changes (see smoothed lines in figure and



Appenzeller et al., 2001). The long-term changes in the AO index indicate possible long-term changes in the BDC (Fusco and Salby, 1999; Randel et al., 2002). Winters with extreme positive (negative) NAM values (northern annular mode, a new term for the Arctic oscillation, see Thompson et al., 2003), correspond to years with small (high)

- winter eddy heat fluxes and strong (weak) polar night jets (Hartmann et al., 2000). An extreme negative AO index combined with an easterly QBO (quasi-biennial oscillation) phase lead to very high total ozone throughout the Northern Hemisphere (NH) in 2010 (Steinbrecht et al., 2011). The study by Kiesewetter et al. (2010a) showed that for winters in a high (low) annular mode state, negative (positive) upper stratospheric ozone
- anomalies in fall result in low (high) column ozone anomalies in polar spring. Low ozone in winter/spring is also expected from low winter planetary wave activity (weak BDC) and low polar temperatures during the winter season (Weber et al., 2003; Kawa et al., 2005; Sinnhuber et al., 2006).
- Figure 1 seems to indicate that long-term ozone changes are correlated to both
 ¹⁵ changes in stratospheric halogen and atmospheric dynamics. This close coupling between atmospheric dynamics and chemistry is also evident on inter-annual time scales as shown by the compact linear relationship between the cumulative hemispheric winter eddy heat flux and spring-to-fall polar total ozone ratio combining data from both hemispheres as reported by Weber et al. (2003). They also showed a corresponding
 ²⁰ anti-correlation with OCIO columns from GOME observations, an indication of chlorine activation inside the polar vortex, that prove year-to-year variability in polar ozone loss are driven by coupled variations in atmospheric dynamics and polar chemistry (Weber et al., 2003; Tegtmeier et al., 2008).

The goal of this paper is to elucidate further on the compact linear relationship between winter eddy heat flux and the ozone build-up as reported in our earlier study (Weber et al., 2003) by using in addition to observations results from two CCMs. First, CCMs are compared to current observations and secondly the CCM data are used to look how such a dynamics-chemistry coupling may change in the future in connection with climate change and ozone recovery. Section 2 briefly describes the total ozone



data sets and the two CCMs (DLR-E39C-A and EMAC-FUB) used in this study. Section 3 summarises results on the similarity of changes in the winter eddy heat flux, stratospheric temperature, and observed spring ozone in both hemispheres since the start of the satellite era at the end of the 1970s. The update on the compact linear relationship between the winter BDC and spring-to-fall ozone ratio using more years of data from the GOME, SCIAMACHY, and GOME2 merged data set (GSG data) is presented in Sect. 4. The ozone anomaly as observed in spring persists throughout the summer months until begin of next fall and its connection to the previous winter's BDC is described and discussed in Sect. 5. Model results are presented in Sect. 6.

¹⁰ The implications on future changes in the ozone – BDC interaction are also discussed in this section followed by conclusions from this study.

2 Data

2.1 Observations

The GOME/SCIAMACHY/GOME2 total ozone merged data set, also called in short the GSG data set, has been compiled from observations made by GOME (Coldewey-Egbers et al., 2005; Weber et al., 2005), SCIAMACHY (Bracher et al., 2005), and GOME2 (Weber et al., 2007) instruments and starts in 1995. In the GSG data set the SCIAMACHY (2002–present) and GOME2 (2007–present) data have been adjusted to the GOME data record (1995–present) by determining a mean scaling fac-

- tor (GOME2 and SCIAMACHY) and trend (SCIAMACHY only) in the monthly mean zonal mean ratios. Using the criterion of maximum global sampling, the GSG data set is then composed of GOME until June 2003, SCIAMACHY (2003–2006), and GOME2 after 2006. More details can be found at and data are available from http://www.iup.uni-bremen.de/gome/wfdoas/.
- ²⁵ We also use the TOMS/SBUV merged total ozone data sets (http://acdb-ext.gsfc. nasa.gov/Data_services/merged/), which combines TOMS, SBUV(/2), and OMI total



ozone data by removing biases and drifts between data sets in overlapping periods (Stolarski and Frith, 2006). These data encompass the time period from 1978 to present and the latest version is MOD V8. Since this data at the moment of writing only extends to end of 2009, OMI data (2004–2010) are also shown based on the TOMS V8 retrieval, also called OMTO3 (OMI-ATBD (2002), http://disc.sci.gsfc.nasa.gov/Aura/data-holdings/OMI/omto3_v003.shtml).

2.2 Chemistry-climate models

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The output from two CCMs is used to compare with observations and to discuss past and future changes in the relationship between the BDC and ozone. A major differ ence between the CCM EMAC-FUB and DLR-E39C-A is that the first is a high-top model with altitude levels up to the mesosphere (0.01 hPa) and the latter a low top model (10 hPa) with better vertical resolution in the UTLS (upper troposphere lower stratosphere). Both models have taken part in the SPARC CCMval activities (Eyring et al., 2005, 2007; SPARC-CCMVal, 2010). The DLR-E39C-A model is based on the general circulation model ECHAM4, coupled with the chemistry module CHEM and contains 39 layers extending to 10 hPa (Stenke et al., 2009). The EMAC-FUB CCM consists of the ECHAM5-MESSy CCM (Jöckel et al., 2006) and includes the interactive chemistry module MECCA (Sander et al., 2005) as well as an improved short-wave radiation parameterisation (Nissen et al., 2007). It was run with a horizontal resolution of T42 (2.8 × 2.8) and 39 levels with the top at 0.01 hPa.

In this study model runs from 1960 until 2050 with prescribed GHG concentrations from the A1B scenario (IPCC, 2000) and surface mixing ratios of ozone depleting substances (ODS) based on the halogen A1 scenario from WMO (2007) are used. Both models include volcanic aerosols and assumed background stratospheric aerosols af-

ter year 2006. Sea surface temperature and sea ice coverage are prescribed in the CCMs according to climate model projections. The solar activity cycles as well as the QBO are considered in both CCMs to include their impact on ozone-climate interaction. Details of the respectively chosen boundary conditions can be found in Eyring et al. (2008) and Morgenstern et al. (2010).



3 Coupling of chemistry and dynamics of total ozone

A key parameter for coupling atmospheric dynamics and chemistry is the stratospheric temperature (Newman et al., 2001; Salby and Callaghan, 2002). At lower temperatures the gas-phase catalytic cycles destroying ozone in the upper stratosphere slow down (Jonsson et al., 2004). This would mean that in a future changing climate with 5 a stratosphere getting colder due to enhanced infrared GHG emission (e.g., Shindell et al., 1999), a super ozone recovery (a recovery to higher ozone levels than in the 1960s) could be expected (WMO, 2007). In the lower stratosphere, where the largest fraction of the total ozone column resides, temperature changes are primarily driven by atmospheric dynamics. An enhancement in planetary wave activity results in instan-10 taneous adiabatic compression in the polar region, in extreme cases in major stratospheric warmings, and corresponding adiabatic expansion in the tropics and cooling, driving temperatures in each case away from radiation equilibrium (Andrews et al., 1987; Kodera, 2006; Salby, 2008). The relaxation back to equilibrium temperatures then drives the residual or diabatic circulation with slow ascent in the tropics and descent in the polar region (Fels, 1982; Andrews et al., 1987).

The coupling of winter polar stratospheric temperature variations from year to year with corresponding changes in the winter mean eddy heat flux and NH polar ozone in spring is shown in Fig. 2. Atmospheric dynamics drives the inter-annual variability

- of polar stratospheric temperatures and ozone transport into the polar region (represented by the winter eddy heat flux) during winter. Another important aspect is that temperature controls polar ozone loss due to heterogeneous chemistry requiring polar stratospheric clouds to be formed (usually below 195 K). The inter-annual variation of chlorine activation with variations in the winter eddy heat flux was shown from OCIO
- ²⁵ data from GOME (Weber et al., 2003) that are indicative for chlorine activation and subsequent catalytic ozone losses inside the polar vortex (Solomon, 1999; Wagner et al., 2001; Richter et al., 2005). Figure 3 shows the corresponding figure for the SH. The correlation between temperature, winter planetary wave activity, and spring ozone in



each hemisphere is particularly good for the late period after the middle 1990s and somewhat lower before.

The nearly linear decrease in SH polar spring ozone in the 1980s is not accompanied with corresponding changes in planetary wave activity and can be therefore ⁵ more directly associated with upward trends in the halogen loading (see bottom of Fig. 1). The impact of the stratospheric aerosols from the Mt. Pinatubo major eruption in 1991 apparently also reduces this correlation. Following the Pinatubo event an increase in the winter mean eddy heat flux is evident in each hemisphere, but without corresponding increases in polar ozone. Sulfate aerosols contribute to ozone deple-10 tion via heterogeneous reactions that needs not to be limited to the polar vortex (Hofmann et al., 1994). However studies have shown that polar stratospheric temperatures where reduced in subsequent NH winters as well as tropical lowermost stratospheric temperature increased (Dhomse et al., 2006, Fig. 15) a typical signature of a reduced Brewer-Dobson circulation. Several other studies indicate that the stratospheric circu-15 lation was reduced and winter AO enhanced following the Pinatubo eruption (see for

instance Stenchikov et al. (2004) and Fig. 1 in this paper)

4 Linear correlation between BDC and ozone build-up

The compact linear relationship between spring-to-fall polar ozone ratio and the winter mean eddy heat flux calculated for each hemisphere, respectively, is shown in Fig. 4. This figure combines data from the Southern and Northern Hemisphere and is an

- ²⁰ This figure combines data from the Southern and Northern Hemisphere and is an update from Weber et al. (2003) and WMO (2007, Chap. 4) and includes more recent ozone data from SCIAMACHY and GOME2 and covers the period 1995 to 2009. The triangles show the SH data and circles the NH data. The SH winter eddy heat flux is negative but plotted here in absolute (positive) values. By combining data from both
- hemispheres the broad range of variation in the winter BDC strength from minimum values in the SH up to maximum values typical for NH winters and associated changes in the winter ozone build-up is visualised.



The spring ozone distribution is shown for selected winter seasons at the top of this figure demonstrating the gradual changes from ozone hole conditions typically in the SH to ozone rich distributions seen only in the NH. The BDC is known to be weaker in the SH, where polar ozone loss outweighs the net transport into high latitudes, thus

- ⁵ providing ozone ratios below one. One exception is the perturbed Antarctic ozone hole anomaly with the first major stratospheric warming observed in the SH (e.g., Weber et al., 2003; Sinnhuber et al., 2003; Richter et al., 2005; von Savigny et al., 2005) where the spring-to-fall polar ozone ratio is well above one. Polar ozone losses in the NH are more sporadic and ozone transport is generally more effective.
- Normally the winter build up of ozone is strongest after late fall (November) as discussed in Fioletov and Shepherd (2003) and Salby and Callaghan (2004b), but here we use September in the NH and March in the SH, respectively, as starting point for the winter build-up. They are the months in the seasonal cycle where total ozone reaches its minimum (Weber et al., 2003; Fioletov and Shepherd, 2003).
- Severe ozone losses in March were observed during the cold Arctic winters in the 1990s (1995/96, 1996/97, and 1999/2000) (Chap. 4 in WMO, 2007). Both the cold Arctic winters in the 1990s and the Antarctic ozone hole anomaly of 2002 represent in terms of the linear relationship shown in Fig. 4 an intermediate case between typical Antarctic and Arctic winters. The Antarctic winter 2006 was characterized by the
- ²⁰ largest ozone hole area (area covered by total ozone below 220 DU) observed in the GOME/SCIAMACHY era and was a result of an extremely weak winter BDC fostering very low polar stratospheric temperatures and, therefore, is located at the left lower end of the linear relationship shown in Fig. 4. Nevertheless, all special events, the Antarctic ozone hole in 2006, the ozone hole anomaly in 2002, and the cold Arctic winters are well accurated by the linear relationship shown in Fig. 4.
- ²⁵ well covered by the linear relationship observed in Fig. 4.

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5 Seasonal persistence of total ozone variability

In several studies it was shown that the inter-annual variations seen in polar total ozone anti-correlate with tropical ozone changes as a result of the overturning BDC (Fusco and Salby, 1999; Salby and Callaghan, 2004b). This is also evident from the change

- ⁵ of the positive correlation of the winter accumulated eddy heat flux with polar ozone ratios into an anticorrelation in the tropics as is shown in Fig. 5. Here, the ozone ratios of February over September (NH) and August over March (SH) are displayed for several zonal bands ranging from the tropics to the polar region. This corresponds to a five month winter build up of ozone starting from September (NH) and March (SH),
- ¹⁰ respectively. The correlation coefficients are very high and close to 0.9 and -0.9 in the polar and tropical region, respectively. The turnaround latitude where the correlation changes sign is within the latitude band between 40° and 50°, which can be interpreted as the zonal band where upwelling at low latitudes changes to downwelling in the extratropics as branches of the residual circulation.
- Ozone anomalies at the end of winter persist throughout the summer months in the extratropics until fall, when photochemical equilibrium is reached and the residual circulation starts building up ozone for the next winter (Fioletov and Shepherd, 2003). The summertime changes in ozone are mainly governed by NO_x chemistry in the absence of dynamic activity (Brühl et al., 1998; Brühl and Crutzen, 2000). As a consequence summertime trends are closely tied to winter trends (Fioletov and Shepherd, 2003, 2005).

In Fig. 6 the correlation coefficients of the linear relationship between ozone ratios and the average eddy heat flux for various zonal bands are plotted as a function of time from fall to fall next year. North of 45° latitude the correlation is positive and negative at lower latitudes throughout the winter and early spring similar to that shown in Fig. 5 for a specific winter month. The positive correlation remains high for polar latitudes until next fall. In late spring and early summer the region of positive correlation extends from the polar region to the subtropics. A possible explanation is the break-up of the



polar vortex and the removal of the polar transport barrier that mixes or dilutes polar air into middle latitudes and to the subtropics (e.g., Knudsen and Grooß, 2000; Ajtić et al., 2004; Fioletov and Shepherd, 2005). The anticorrelation observed at tropical latitudes goes to zero by early summer. The most likely explanation is the absence of dynamic activity in the summer hemisphere. Since variations in the tropics are lower than at high latitudes, the photochemical equilibrium is probably reached much faster than in the extratropics.

6 Model results

- The analysis of the BDC impact on total ozone variability on time scales from seasonal to inter-annual described so far provides a valuable diagnostics for CCMs. The question arises how well do models compare with observations and what can models tell us about past and future changes. In Figs. 7 and 8 the seasonal correlation maps are shown for both CCMs, DLR-E39C-A and EMAC-FUB, respectively. The model data are shown for years 1985 to 2010 to be comparable to observations shown in Fig. 6.
- The correlation maps with positive values in polar latitudes and negative in the tropics are qualitatively very similar to observations. The transition to summer conditions as characterized by the expansion of the positive correlation into middle latitudes also agrees with observations. The overall patterns in the winter/summer correlation appear realistic and are consistent with observations.
- Differences between models and observations are found in the transition zone from positive to negative correlations during winter. Observations indicate 45° latitude (Fig. 6), while it lies equatorward of 40° in both models. Figures 7 and 8 also show the changes with time in the transition zone. The dotted and dashed lines show the correlation of the 30°-40° zonal bands for the periods 1960-1985 (past) and 2010 2050 (future), respectively. In the DLR model it appears that the transition zone is nearly unaltered, only in the past it was slightly shifted to lower latitudes as evident from the higher and positive correlation. The FUB model shows a different behavior. In



both past and future decades the transition is closer to about 35° latitude and slightly shifted to higher latitudes in the period 1985–2010. The rough position of the modeled transition near 35° latitude is close to the latitude separating tropical upwelling at lower latitudes from downwelling at higher latitudes shown in other model data (Austin and Li, 2006).

During the summer months the correlation of polar ozone (and the anti-correlation of tropical ozone) with the average eddy heat flux from last winter starts to weaken about a month earlier in the DLR model compared to the FUB model and observations.

Figures 9 and 10 show the relationship of the spring-to-fall polar total ozone ratio
(September over March in the SH and March over September ratio in the NH) for the DLR and FUB models, respectively. Also shown are the GSG data set from Fig. 4. All model data from 1960–2050 are shown. Different colors are representing different time periods, grey for the past (1960–1985), black the present (1985–2010), and blue the future period (2010–2049). The corresponding regression line (dashed) for the three different periods is also shown. Since the ozone values in the DLR model show a consistent bias of about 50 DU with respect to observations, an offset of –50 DU was applied to all DLR model data.

In general, the correlation in the relationship between polar ozone ratio and winter eddy heat flux is remarkably high and reproduces very well the observations. However,

- there are some differences with respect to observations. The spring-to-fall polar ozone ratios in the SH are in most cases above one, while observations are always below one except for the anomalous Antarctic ozone hole in 2002 indicating that ozone transport outweighs SH polar ozone loss in the models as opposed to observations. In both hemispheres the ozone ratios are in most cases higher than observations. A possible
- explanation for this could be that either modeled polar ozone losses are too low or that the polar vortex is more permeable due to numerical diffusion, meaning more mixing with middle latitude air occurs in the model (Austin et al., 2010).

One interesting aspect from the model studies is the evolution of this compact linear relationship on decadal time scales from the past into the future. The ozone ratios



are smallest during the present period (1985–2010) compared to the past and future in both models. The long-term changes in the linear fits in Figs. 9 and 10 coincides with changes in the halogen loading (see Fig. 1 and Newman et al., 2007). During the period 1985–2010 the halogen loading reached a maximum and ozone losses in

- ⁵ spring were maximum leading to lower spring-to-fall ozone ratios under identical meteorological/dynamical conditions of the polar winter stratosphere. Due to the phase-out of ozone depleting substances (WMO, 2007, 2011), the stratospheric halogen starts a slow decrease so that an increase in future ozone ratio and thus an upward shift in the linear fit curves as shown in Figs. 9 and 10 is expected.
- The upward shift of the regression line is smaller for the EMAC-FUB model than for DLR-E39C-A. This indicates that the ozone recovery is slower in the FUB model. However, both models are among the CCMs with faster ozone recovery (Austin et al., 2010; SPARC-CCMVal, 2010).

First signs of ozone recovery related to the turnaround in the halogen loading were
observed in the upper stratosphere (Newchurch et al., 2003; Steinbrecht et al., 2009).
However in the lower stratosphere, where the dominant contribution to the total ozone column comes from, the fingerprint of recovery is less clear due to the large influence from atmospheric dynamics (see Fig. 1). The largest contribution to the strong increase in NH total ozone in the extratropics since the middle 1990s is due to changes in the stratospheric circulation (Dhomse et al., 2006; Wohltmann et al., 2007; Harris et al., 2008). Nevertheless, the upward shift in the linear regression in Figs. 9 and 10 confirms that ozone will steadily increase with sinking stratospheric halogen loading despite the large variability seen from year-to-year.

The variability in the modelled winter eddy heat fluxes (Figs. 9 and 10) is quite large in each hemisphere (spread across the *x*-direction of the plots) and is mostly in the range what is seen in the observations. The variability in the NH is somewhat smaller for the FUB model, when compared to the DLR model and observations. Both FUB and DLR model have individual NH winters with extremely low winter eddy heat fluxes and spring-to-fall ozone ratios that are similar to Antarctic values. Such cases have not



been seen in observations during the satellite era (since 1978). On the other hand, none of the models was able to produce SH conditions similar to the Antarctic ozone hole anomaly observed in 2002.

No significant changes with time in the ranges of the winter eddy heat flux are evident in the models so that the largest changes with time in the compact linear relationship appear be due to changes in halogens and associated ozone chemistry. In Fig. 11 the time evolution of the CCM winter eddy heat flux in each hemisphere (averaged from September to March in the NH and March to September in the SH) is displayed. Also shown are the decadal means of the same quantities (blue for SH and red for NH) and the linear fit to the decadal means. A slow increase in the strength of the winter BDC

- and winter eddy heat flux from 1960 until 2050 is evident, which is $\sim +0.5 \text{ Km s}^{-1}$ over the ninety year time span. The year-to-year variations are on the order of $1-2 \text{ Km s}^{-1}$ which strongly masks the long-term trend. For a few decades the trends may even become reversed, e.g. 1990–2010 in the NH (DLR model) and 2000–2020 in the SH (FUB model), such that polar ozone losses can be intensified for one to two decades
- despite the slow decline in the stratospheric halogen loading.

In Fig. 11 the decadal means from NCEP reanalysis during the satellite period (1980–2010) are also shown. Over the three decades of the satellite era the increase in the observed winter eddy heat flux is a factor of two to three larger than the long-

term change seen in the CCMs. From the variability evident in the model data it is most likely that the current observed trend may not be maintained over a longer time period into the future. Also, it is clear that the 30-yr satellite data record is still comparatively short for a complete disentangling of long-term changes in ozone columns due to atmospheric dynamics and halogen changes.

25 **7** Summary and conclusion

In this paper we investigated the impact of the BDC on total ozone from seasonal to decadal time scales using satellite observations and two CCMs. From the long-term data record it is clear that long-term changes in total ozone in the extratropics,



representative for lower stratospheric ozone, have contributions from changes in the upper atmospheric circulation (atmospheric dynamics) and in ozone chemistry (polar ozone losses and gas-phase losses) due to the halogen load. The chemical trends are tied to changes in meteorology and atmospheric dynamics as shown in Fig. 1.

- This is also evident on shorter time scales when looking at inter-annual variations in stratospheric temperatures (as a proxy for chemistry), extratropcial ozone, and BDC strength, here expressed by the winter eddy heat flux in each hemisphere (Figs. 2 and 3).
- Combining data from both hemispheres a compact linear relationship exists between
 the winter eddy heat flux and the spring-to-fall ozone ratio as shown in Fig. 4. This figure is an update of Fig. 3 in Weber et al. (2003) with more years of data added and now showing even a higher correlation coefficient (*r* = 0.992). This compact linear relationship marks as one extreme the Antarctic winters with a weak BDC, colder polar lower stratospheric temperatures, the ozone hole, and the other extreme the Arctic
 winters with higher stratospheric temperatures, larger ozone transport, and reduced polar ozone losses. The anomalous SH ozone hole (Sinnhuber et al., 2003; Richter et al., 2005; von Savigny et al., 2005) with unusually enhanced ozone transport into polar latitudes and the cold Arctic winters in the 1990s with reduced ozone transport and larger polar ozone losses fit into this remarkable relationship. The tropical ozone zo ratios in contrast are anti-correlated to the winter-eddy heat flux as shown in Fig. 5.

The correlation between the eddy heat flux averaged from fall to the respective month and the corresponding ozone ratios remain high throughout spring until the end of the summer (Fig. 6). The positive correlation as seen in the extratropics during late spring and summer also extends into the tropical region after the polar vortex breaks up. This ²⁵ is consistent with results from Fioletov and Shepherd (2003, 2005) who investigated the seasonal autocorrelation of total ozone and its impact on long-term trends. The transition zone between negative correlation (tropics) and positive correlation (extratropics) during winter is at about 45° latitude. This is consistent with the fact that the sign of stratospheric temperature departure from radiative equilibrium reverses at the



same latitude, meaning it is negative equatorward (tropical cooling and upwelling) and positive poleward of this latitude as noted by Ueyama and Wallace (2010).

The seasonal correlation pattern as shown for observational data is also very well reproduced by two chemistry-climate models, the DLR-E39C-A and the EMAC-FUB

- (Figs. 7 and 8), that were run from year 1960 to 2050. One notable difference with respect to observations is that the transition zone between positive correlations in the extratropics and negative correlations in the tropics is somewhat shifted towards lower latitudes near 35° latitudes. A compact linear relationship between polar spring-to-fall ozone ratio and the modelled winter eddy heat flux is seen in both models, clearly is a set of the transition of the transition of the transition of the transition.
- ¹⁰ indicating that the overall BDC pattern on total ozone is well reproduced by the models (Figs. 9 and 10). However, the spring-to-fall ozone ratios are generally higher than the observed ones. The linear regression of this relationship shifts due to the reversal of the stratospheric halogen trends meaning that under near identical meteorological conditions and dynamic state of the atmosphere the recovery of total ozone in the future
- is evident. A similar shift in the linear regression is difficult to derive from observations since the satellite data record now spanning more than 30 yr appears to be still too short.

Both models show a positive trend in the winter eddy heat flux indicating that in a changing climate the BDC may intensify in coming decades, a feature which is ob-²⁰ served by many other CCMs (see Sect. 1). Nevertheless the year-to-year variability in the winter BDC strength is a factor of three to four higher than the decadal mean change from 1960 to 2050. For one or two decades as e.g. 1990–2010 in the NH (DLR model) and 2000–2020 in the SH (FUB model) a negative trend in the winter eddy heat flux may occur that could lead to more severe polar ozone loss despite the slow decline ²⁵ in the stratospheric halogen load.

When attributing processes to the current total ozone trends, it is difficult to clearly separate chemical and dynamical contributions (Kiesewetter et al., 2010b), which is understandable from the fact that both processes are closely coupled as shown in this study.



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Fig. 1. Top: JFM total ozone area-weighted between 45° N and 60° N as measured from ground (Fioletov et al. (2002), WOUDC web site, ftp://ftp.tor.ec.gc.ca/Projects-Campaigns/ZonalMeans/gb_1964-2010_za.txt) and satellites (SBUV/TOMS/OMI merged data, MOD V8, in red, GOME/SCIAMACHY/GOME2 merged data, GSG, in blue, and OMI, OMTO3, in light brown). The thick black line is the three point triangular smooth of the ground data. Middle panel: DJF Arctic oscillation index (blue values correspond to positive AO values) obtained from http://www.cpc.ncep.noaa.gov/products/precip/CWlink/daily_ao_index/ao.shtml. Thick black line is the three point triangular smooth of the AO index. Bottom panel: Equivalent effective stratospheric chlorine (EESC) drawn with a reversed *y*-axis.





Fig. 2. Top: March total ozone area-weighted between 50° N and 90° N as measured from satellites (solid line: SBUV/TOMS/OMI merged data, dashed line: SCIAMACHY/GOME data). Middle panel: NH area weighted polar cap temperature (70° N–90° N) from different meteorological analyses. Bottom panel: extra-tropical eddy heat flux at 100 hPa averaged from September to March and area weight averaged between 45° N and 75° N derived from different meteorological analyses.





Fig. 3. Top: October total ozone area-weighted between 50° S and 90° S as measured from satellites (solid line: SBUV/TOMS/OMI merged data, dashed line: SCIAMACHY/GOME data). Middle panel: SH area weighted polar cap temperature (70° S–90° S) from different meteorological analyses. Bottom panel: extra-tropical eddy heat flux at 100 hPa averaged from March to September and area weight averaged between 45° S and 75° S from different meteorological analysis.





Fig. 4. Spring-to-fall ratio of observed polar cap total ozone (> 50°) as a function of the absolute extratropical winter mean eddy heat flux (September to March and March to September in the respective hemispheres) derived from ECMWF ERA-Interim data separately in the respective hemisphere. Data from the Southern Hemisphere are the triangles (September over March ozone ratios) and from the Northern Hemisphere are solid circles (March over September ratios).





Fig. 5. Winter-to-fall ratio of observed total ozone for various zonal bands from the polar region to the tropics as a function of the mean absolute extratropical cumulative eddy heat flux between September and February in the NH and between March and August in the SH from ERA-Interim. Data from the Southern Hemisphere are the triangles (August over March ozone ratios) and from the Northern Hemisphere are solid circles (February over September ratios). Colors correspond to individual SH and NH winters as indicated in the bottom of Fig. 4.

















Fig. 8. Same as Fig. 7, but for CCM data from EMAC-FUB model.











Fig. 10. Same as Fig. 9, but for CCM data from EMAC-FUB model.







