1 A Global Ozone Climatology from Ozone Soundings via Trajectory Mapping:

2 A Stratospheric Perspective

- J. J. Liu^{1,2,3}, D. W. Tarasick¹, V. E. Fioletov¹, C. McLinden¹, T. Zhao⁴, S. Gong¹, C. Sioris¹, J. J.
 Jin⁵, G. Liu⁶, and O. Moeini⁷
- ⁵ ¹Science and Technology Branch, Environment Canada, 4905 Dufferin Street, Downsview,
- 6 Ontario, Canada M3H 5T3
- ⁷ ²Department of Geography and Planning, University of Toronto, 100 St. George Street, Toronto,
- 8 Ontario, Canada M5S 3G3
- ⁹ ³Nanjing University, Nanjing, Jiangsu, China, 210093
- ⁴Nanjing University of Information Science & Technology, Nanjing, Jiangsu, China, 210044
- ⁵Universities Space Research Association, and Global Modeling and Assimilation Office,
- 12 Goddard Space Flight Center, NASA, Greenbelt, MD, USA
- ⁶Space Sciences Laboratory, University of California, Berkeley, California, USA, 94720
- ⁷Department of Earth and Space Science & Engineering, York University,
- 15 4700 Keele Street, Toronto, Ontario, M3T 1P3
- 16 Submitted to Atmospheric Chemistry and Physics
- 17 12 June 2013
- 18 ¹Corresponding author:
- 19 Department of Geography and Planning, University of Toronto,
- 20 100 St. George Street, Room 5047, Toronto, Ontario, Canada M5S 3G3
- 21 E-mail:janejj.liu@utoronto.ca, Tel: (416) 946-8257, FAX: (416) 946-3886

22 Abstract

23 This study explores a domain-filling trajectory approach to generate a global ozone climatology 24 from relatively sparse ozonesonde data. Global ozone soundings comprising 51,898 profiles at 25 116 stations over 44 years (1965-2008) are used, from which forward and backward trajectories 26 are calculated from meteorological reanalysis data, to map ozone measurements to other 27 locations and so fill in the spatial domain. The resulting global ozone climatology is archived monthly for five decades from the 1960s to the 2000s on a grid of $5^{\circ} \times 5^{\circ} \times 1$ km (latitude, 28 29 longitude, and altitude), from the surface to 26 km altitude. It is also archived yearly from 1965 30 to 2008. The climatology is validated at 20 selected ozonesonde stations by comparing the actual 31 ozone sounding profile with that derived through trajectory mapping of ozone sounding data 32 from all stations except the one being compared. The two sets of profiles are in good agreement, 33 both individually with correlation coefficients (r) between 0.975 and 0.998 and root mean square 34 (RMS) differences of 87 to 482 ppbv, and overall with r = 0.991 and an RMS of 224 ppbv. The 35 ozone climatology is also compared with two sets of satellite data, from the Satellite Aerosol and 36 Gas Experiment (SAGE) and the Optical Spectrography and InfraRed Imager System (OSIRIS). 37 The ozone climatology compares well with SAGE and OSIRIS data in both seasonal and zonal 38 means. The mean differences are generally quite small, with maximum differences of 20% above 39 15 km. The agreement is better in the northern hemisphere, where there are more ozonesonde 40 stations, than in the southern hemisphere; it is also better in the middle and high latitudes than in 41 the tropics where reanalysis winds are less accurate. This ozone climatology captures known 42 features in the stratosphere, as well as seasonal and decadal variations of these features. 43 Compared to current satellite data, it offers more complete high latitude coverage as well as a 44 much longer record. The climatology shows clearly the depletion of ozone from the 1970s to the

45 mid 1990s and ozone recovery in the lower stratosphere in the 2000s. When this climatology is

46 used as the upper boundary condition in an Environment Canada operational chemical forecast

- 47 model, the forecast is improved in the vicinity of the upper troposphere–lower stratosphere
- 48 (UTLS) region. As this ozone climatology is neither dependent on *a priori data* nor
- 49 photochemical modeling, it provides independent information and insight that can supplement
- 50 satellite data and model simulations of stratospheric ozone.

51 1 Introduction

52 Ozone is an important trace gas in the atmosphere, playing a significant role in atmospheric 53 chemical, dynamical, and radiative processes. In the stratosphere, emissions of human-produced 54 ozone depleting substances led to a substantial ozone decline that has been reversed by the 55 implementation of the Montreal Protocol in 1987 and its subsequent amendments (WMO, 2003, 56 2007). Identifying stratospheric ozone recovery and assessing the impact of ozone changes on 57 climate are important issues that have attracted considerable attention (e.g., Randel and Wu, 58 1999; Weatherhead and Andersen, 2006; Waugh et al., 2009; Eyring et al., 2010; Ziemke and 59 Chandra, 2012). Stratospheric ozone can also impact chemistry and dynamics in the troposphere, 60 both via direct exchange and the Brewer-Dobson circulation (e.g., Holton et al., 1995; Stohl et 61 al., 2003) and via effects on planetary wave propagation and global climate (Baldwin and 62 Dunkerton, 2001; Gillett and Thompson, 2003).

63 An understanding of the distribution of stratospheric ozone and its long-term changes is a 64 critical step to assessing the interactions between ozone variability and climate change. Satellite 65 observations have advantages of consistent quality and global coverage, but suffer from biases 66 between different, or successive instruments and limited vertical or horizontal resolution 67 depending on the measurement approach. Considerable effort has been made previously to 68 develop ozone climatologies, typically combining satellite measurements from one or more 69 instruments with other data sources, often ozonesondes. These datasets are either meridionally 70 and vertically resolved (two dimensions in latitude and altitude, e. g., Fortuin and Kelder, 1998; 71 Lamsal et al., 2004; McPeters et al., 2007; Randel and Wu, 2007; Hassler et al., 2008; Jones et 72 al., 2009; McLinden et al., 2009; McPeters and Labow, 2012) or of ozone column that is 73 horizontally resolved (two dimensions in latitude and longitude, e. g., Ziemke et al., 2005, 2011). Stratospheric ozone climatologies in two or three dimensions have also been developed from
chemistry-climate model simulations (e.g., Eyring et al., 2010; SPARC CCMVal, 2010).
Although considerable advances have been made in the past decade to improve model
representations of ozone transport and chemistry, uncertainties remain with regard to
parameterizations, radiation transport schemes, and simulation of the Brewer-Dobson circulation,
as well as prediction of the late-spring breakup of the Antarctic vortex and of the Antarctic ozone
hole (e. g., SPARC CCMVal, 2010; Forster et al. 2011; Wang and Waugh, 2012).

81 In this study, we explore a different approach to develop a long-term ozone climatology 82 from global ozonesonde data for satellite and model *a priori*, satellite and climate model data 83 validation, troposphere-stratosphere exchange, and atmospheric climate-chemistry interaction 84 research. Ozone soundings have the advantages of long-term and quasi-global coverage, high accuracy, and high vertical resolution but are sparse in space and time. Given ozone's long 85 lifetime of weeks to months in the lower stratosphere (e.g., Jacob, 1999; Liu and Ridley, 1999), 86 87 a measurement of ozone mixing ratio at one place and time generally provides a good estimate of 88 ozone mixing ratio in that same air parcel several days before or after. It is therefore possible to 89 extend sparse ozonesonde measurements and to fill the gaps in the spatial domain by trajectory 90 calculations, assuming the ozone mixing ratio along each trajectory path is constant. This is a 91 technique that has been used successfully with other stratospheric data (Sutton et al., 1994; 92 Newman and Schoeberl, 1995; Morris et al., 2000), and with ozone data in both the stratosphere 93 and troposphere. Stohl et al. (2001) used trajectory statistics to extend one year of MOZAIC 94 (Measurement of Ozone and Water Vapour by Airbus In-service Aircraft) ozone measurements 95 into a 4-season ozone climatology at 10° longitude by 6° latitude and three vertical heights. 96 Trajectory mapping has also been used to fill gaps in satellite MLS (Microwave Limb Sounder)

97 ozone data in the stratosphere in order to calculate the tropospheric residual ozone column
98 (Schoeberl et al., 2007). Tarasick et al. (2010) developed high resolution (1°×1°×1 km in
99 latitude, longitude, and altitude) tropospheric ozone fields for North America from ozonesonde
100 data from the INTEX (Intercontinental Transport Experiment) and ARCTAS (Arctic Research of
101 the Composition of the Troposphere from Aircraft and Satellites) campaigns. This has been
102 recently extended to global tropospheric ozonesonde data by Liu et al. (2013).

Our study is aimed at (1) developing a global long-term ozone climatology from ozonesondes in 3 dimensions (latitude, longitude, and altitude), (2) validating this dataset against independent ozonesondes and satellite observations, (3) characterizing the stratospheric ozone distribution and its decadal and seasonal variabilities, and (4) illustrating applications of this dataset in providing evidence of ozone recovery and in improving simulations of tropospheric ozone.

109 In the following, Section 2 describes the methodology of trajectory mapping and the 110 input ozonesonde data. Section 3 describes the validation of the trajectory-mapped dataset with 111 ozonesonde data and the comparison between this dataset and satellite datasets from SAGE (the 112 Satellite Aerosol and Gas Experiment) and OSIRIS (the Optical Spectrography and InfraRed 113 Imager System). In Section 4, the variation of stratospheric ozone at different time scales is 114 characterized. Section 5 discusses the timing of stratospheric ozone recovery, based on this 115 ozone climatology, and shows the improvement in the simulation of tropospheric ozone in an air 116 quality forecast model when this climatology is used as the upper boundary condition. 117 Conclusions are provided in Section 6.

118

119 **2 Methods**

120 **2.1 Global Ozonesonde Data**

121 The global ozonesonde data used herein were acquired from the World Ozone and Ultraviolet 122 Radiation Data Centre (WOUDC), which is one of five World Data Centres of the Global 123 Atmosphere Watch programme of the World Meteorological Organization. The WOUDC 124 operates a scientific archive, providing ozone data to the international scientific community 125 (Wardle et al., 1998; Hare et al. 2007). Table 1 describes these stations, including their 126 geolocations (latitude, longitude, and elevation), operation periods, and the total number of 127 profiles used in this study. The stations' geolocations are illustrated in Figure 1a, showing their 128 non-uniform distribution around the globe. There are more stations in the northern hemisphere 129 (NH) than in the southern hemisphere (SH), and more stations over Europe and North America 130 than over Asia. Large gaps appear over China and eastern Russia. The total number of available 131 stations is 47 in the 1970s (Figure 1b). This number gradually increases with time. In the 2000s, 132 there are 57 stations (Figure 1c) with more coverage in the tropics, South Asia, East Asia, the 133 Arctic, and the Antarctic. Overall, we employ 51,898 ozone soundings globally at 111 stations, 134 in different periods (Table 1) over 44 years from 1965 to 2008.

135 Most of the profiles are from the electrochemical concentration cell (ECC)-type 136 ozonesonde, which was introduced in the early 1970s and adopted by a majority of stations in the 137 global network by the early 1980s. Virtually all the data in the most recent decade are from ECC 138 sondes. The remainder are from Brewer-Mast (BM) sondes (currently still in use at one site), the 139 Japanese KC96 sonde, and the Indian sonde. Prior to the early 1990s, three stations in Europe 140 (Praha, Lindenberg and Legionowo) flew the GDR sonde. A majority of the data before 1980 is 141 from BM sondes or similar (both the GDR and Indian sondes are similar in design to the BM 142 sonde). A small amount of data is available in the early 1960s from carbon-iodine sondes

(similar to the KC sondes) and from Regener sondes (which operated by the chemiluminescentreaction of ozone with luminol).

145 When properly prepared and handled, ECC ozonesondes have a precision of 3-5% (1- σ) 146 and an absolute accuracy of about 5% in the stratosphere (Kerr et al., 1994; Smit et al., 2007; Deshler et al., 2008; Liu et al., 2009). The ozone sensor response time (e^{-1}) of about 25 seconds 147 148 gives the sonde a vertical resolution of about 100 metres for a typical balloon ascent rate of 4 149 m/s. Two types of ECC ozonesondes are in current use, the 2Z model manufactured by EnSci 150 Corp. and the 6A model manufactured by Science Pump, with minor differences in construction 151 and some variation in recommended concentrations of the potassium iodide sensing solution and 152 of its phosphate buffer. The maximum variation in stratospheric response resulting from these 153 differences is likely of the order of 2-3% (Smit et al., 2007).

The precision of other sonde types is somewhat poorer, at about 5-10% (Kerr et al., 1994; Smit et al., 1996). In early intercomparisons (Attmannspacher and Dütsch, 1970, 1981) the Indian and the GDR sonde showed significantly poorer precision than other sonde types.

157 The largest systematic differences between sondes are in the lower stratosphere, where 158 the BM and GDR sondes gave readings about 5-10% lower than the ECC and KC sondes in early 159 intercomparisons (Attmannspacher and Dütsch, 1970; 1981). In later intercomparisons (Kerr et 160 al., 1994; Smit et al., 1996; Deshler et al., 2008) the KC sondes have shown a low bias of about 161 5% in the lower stratosphere, and the BM has generally shown a low bias as well. The Indian 162 sonde has generally shown little bias in the lower stratosphere, but somewhat lower precision 163 than the other non-ECC sondes. Regener sondes were used regularly for only a brief period in 164 the 1960s, as they showed somewhat erratic response (Hering and Dütsch, 1965).

In the middle and upper stratosphere below 26 km differences in sonde response are small. Overall, between the tropopause and 26 km the precision of the various sonde types in recent decades is generally within $\pm 5\%$, and any systematic biases between them or compared to other ozone sensing techniques are smaller than $\pm 5\%$ (World Climate Research Programme, 169 1998).

170

171 **2.2 Trajectory Calculation and Global Ozone Mapping**

172 To prepare for trajectory modelling, the ozone mixing ratio profile for each sounding was

173 integrated into 26 vertical levels at 1-kilometre resolution from sea level. Ozone measurements

above 26 km were excluded because of their higher uncertainties (e.g., Fioletov et al., 2006).

175 Trajectories for all 26 levels were calculated using the Hybrid Single-Particle Lagrangian

176 Integrated Trajectory (HYSPLIT) model (version 4.9) (Draxler and Hess, 1998, Draxler et al.,

177 2012), available from the Air Resources Laboratory, National Oceanic and Atmospheric

178 Administration (NOAA) (http://www.arl.noaa.gov /HYSPLIT info.php). HYSPLIT is driven by

the reanalysis meteorological data from the National Centers for Environmental

180 Prediction/National Center for Atmospheric Research (NCEP/NCAR) (Kalnay et al., 1996).

181 These data are available from 1965 to the present. Both forward and backward trajectories for 4

182 days (96 hours) were calculated for 51,898 ozone soundings and the trajectory positions were

183 stored.

Ozone mixing ratios in the 26 levels from each sounding were assigned to the corresponding trajectory positions along the forward and backward paths every 6 hours for 4 days (32 positions for each level). Then, the trajectory-mapped data were binned at intervals of 5° latitude and 5° longitude, at each 1-km altitude, and averaged. This bin size is less than the typical ozone correlation length in the stratosphere of about 1000-1500 km (Liu et al., 2009).

Two different altitude coordinates were employed for this binning, and so two sets of maps were produced, one whose vertical coordinate is altitude above sea level, and the other altitude above ground level. A global ozone climatology of monthly means for each altitude was generated for each decade from the 1960s to the 2000s. The climatology is also archived yearly from 1965 to 2008 at the same horizontal and vertical resolutions. Figure 1 illustrates the improved spatial coverage if trajectory mapping is used. The trajectory mapping greatly spreads out the ozone information along the trajectory paths, increasing the spatial domain to include most of the globe.

196 Most atmospheric models using assimilated meteorological data produce tropopause 197 heights that are more accurate than climatology. Using an ozone climatology that includes a 198 climatological tropopause with such models can produce, in effect, spurious stratospheric 199 intrusions as air in the upper troposphere (when the actual tropopause is high) is assigned 200 stratospheric values of ozone (e.g., Makar et al., 2010). The opposite effect can occur when the 201 actual tropopause is low. For this reason, we generated three ozone climatology datasets. The 202 first dataset is a conventional climatology that uses ozonesonde data in both troposphere and 203 stratosphere, while in the second and third dataset, the stratospheric ozone climatology and 204 tropospheric ozone climatology are generated separately with trajectories originating exclusively 205 from the stratosphere and the troposphere, respectively. The explicitly separated troposphere and 206 stratosphere, which overlap in the upper troposphere-lower stratosphere (UTLS) region, are 207 intended to be used for model *a priori* where an atmospheric model determines the tropopause 208 location itself. This paper focuses on characterization and validation of the first and second 209 datasets, while the tropospheric climatology is discussed in a separate paper (Liu et al., 2013). 210

211 **3. Validation and Comparison with Independent Ozone Observations**

212 **3.1 Comparison with Ozonesonde Profiles**

213 To test the trajectory methodology, the actual ozone profile at selected stations was compared 214 with that produced by the trajectory mapping using the ozone soundings at all stations except the 215 one being compared. Figure 2 shows examples of such comparison in the NH winter and summer months (January and July) in 1990s at four stations: Boulder (40.1° N, 105.3° W, 1689 216 m), Hohenpeissenberg (47.8° N, 11.0° E, 975 m), Uccle (50.8° N, 4.0° E, 100 m), and Eureka 217 218 (80.0° N, 86.2° W, 310 m). The paired profiles shown are in reasonable agreement. Seasonal 219 variation is well captured in the trajectory-derived ozone profiles. The correlation coefficient (r)220 between the two datasets is between 0.950 and 0.999. Biases are between -170 and +80 ppbv, 221 while RMS differences range from 60 to 340 ppbv. 222 Globally, twenty stations were selected (Table 2, Figure 1a), with a range of latitude from 223 71° S (Neumayer) to 80° N (Eureka), and longitude from 171° W (Samoa) to 178° E (Fuji), and 224 in altitude from near sea level (Tahiti) to 1745 m (Nairobi). There are 10 stations each in the 225 northern and southern hemispheres, while the number of stations is 9 and 11, respectively, in the 226 eastern and western hemisphere.

In Figure 3, the decadal monthly mean ozone abundance from ozonesonde data at selected stations is compared with that from trajectory mapping without input from the station being tested, for all the 20 stations in the 1990s and 2000s. Overall, the correlation coefficient between the two datasets is 0.99. The overall bias is negligible, at 1.2 ppbv, while the RMS difference is 224 ppbv. For individual stations, the correlation coefficient ranges between 0.975 and 0.999 and the RMS differences between 87 and 482 ppbv.

233

3.2 Comparison with SAGE Ozone Data

235 The satellite instrument SAGE (McCormick et al., 1989) was designed to measure ozone in the 236 stratosphere where most ozone resides. The SAGE mission expanded over 20 years starting from 237 SAGE I onboard AEM2 (Application Explorer Mission) in 1979. SAGE I was in operation until 238 1981. SAGE II, on board ERBS (Earth Radiation Budget Satellite), was in operation from 1984-239 2005. SAGE uses the highly accurate solar occultation technique and has a vertical resolution of 240 less than 1 km. Ozone is derived from measurements in the spectral range of 0.385-1.020 µm 241 (Cunnold et al., 1989). In the absence of clouds, SAGE can detect some layers in the middle and 242 upper troposphere after a significant improvement in the algorithms (Kent et al., 1993). Wang et 243 al. (2002) compared SAGE II with ozonesonde data and found that SAGE tends to slightly 244 overestimate ozone (less than 5%) between 15 and 20 km altitude and to systematically 245 underestimate ozone in the troposphere by approximately 30% between 2 and 8 km altitude. 246 This is consistent with the comparisons we make with the trajectory-mapped data in Figure 4 247 below.

This study uses SAGE I version 7.0 and SAGE II version 6.2 (Wang et al., 2006). Vertical profiles are reported in number density at 1-km resolution from 0.5 km to 60.5 km. Each profile is matched with temperature and pressure from NMC/NCEP (National Meteorological Centre/National Centre for Environment Protection) to derive ozone mixing ratio from the surface to 26 km (McLinden et al., 2009). The comparison with SAGE data is conducted here in two ways.

Firstly, profiles from the ozone climatology were compared with SAGE and ozonesonde data vertically. About 1400 SAGE profiles that coincide with ozonesonde profiles, from 1979 to 2005, were extracted. Each pair of SAGE and ozonesonde profiles was matched with the profile from the ozone climatology in the corresponding month, decade, and $5^{\circ} \times 5^{\circ}$ latitude-longitude 258 bin. That is, the comparisons between ozonesonde and SAGE profiles were made at the same 259 location and time, while ozonesonde profiles were compared with those from the ozone 260 climatology at the same location, month, and decade. Figure 4 shows the vertical distribution of 261 the mean relative differences in the 1990s in NH summer (JJA) and winter (DJF) by four 262 latitudinal zones. Above 15 km, mean differences from the ozone climatology are mostly <10%, 263 as are SAGE-sonde differences. Below 15 km, the ozone climatology is closer to ozonesondes 264 than SAGE. The distribution of the bias for vertical profiles of SAGE is consistent with previous 265 studies (e.g., Wang et al., 2002; Fioletov et al., 2006). The comparison between the climatology 266 and ozonesondes is better in the summer than in the winter, and better at higher latitudes $(45-90^{\circ})$ N, 45-90° S) than at lower latitudes (0-45° N, 0-45° S). This appears to be largely the case for 267 268 SAGE as well. To some extent, the difference profiles of the climatology from sondes in Figure 269 4 are similar to the corresponding SAGE-sonde difference profiles, which may indicate that they 270 are largely due to sampling issues with the sondes.

Secondly, decadal monthly mean from SAGE were gridded at the same 3-dimensional resolution as the ozone climatology ($5^{\circ} \times 5^{\circ} \times 1$ km in latitude, longitude, and altitude). The two datasets, for the same time period, are compared by grid cell, along with ozonesonde data. Figure 5 compares the three datasets by season, taking data at grid cells where all the data are available in the 1990s and the 2000s.

In general, the trajectory-derived ozone climatology agrees well with ozonesondes (left panels) and the degree of scatter shows little variation with season. The scatter is due to imperfections in the wind fields, as well as measurement error in the original sonde data. The SAGE-ozonesonde correlation appears reasonably good for the four seasons (middle panels), although the degree of scatter is larger than that for the ozone climatology. The ozone climatology correlates well with SAGE data in all seasons (right panels) with RMS errors that are about the same as the SAGE-sonde comparison (middle panels). Most notable is that the leftmost panels show much less scatter (smaller RMS errors and higher r) in all seasons than the two comparisons involving SAGE data.

285 The same data points are divided into six 30° latitude zones (Figure 6). The climatology 286 is closely correlated with independent ozonesondes, with r > 0.98 in all cases. The SAGE-287 ozonesonde correlations (Figure 6, right panels) show more scatter than the ozone climatology-288 ozonesonde correlations in all six latitude zones. As in Figure 5, the ozone climatology-SAGE 289 correlations (not shown) are similar to the SAGE-sonde correlations. Table 3 lists statistics for 290 the ozone climatology and SAGE data in NH winter and summer seasons and 3 altitudes in the 291 1990s and the 2000s. Both datasets capture the altitude-dependent seasonal variations. The ozone 292 climatology and SAGE data are well-correlated at 14.5 km with $r \sim 0.9$. The correlation becomes smaller at higher altitudes. At 19.5 km, the correlation coefficient is ~0.8 for both winter and 293 294 summer in the 1990s and the 2000s. At 24.5 km the correlation is poorer, reflecting the fact that 295 the RMS/Mean is lower. The mean ozone mixing ratio from the climatology dataset is higher 296 than SAGE at 14.5 km (similar to the SAGE-sonde bias in Figure 4) but shows little bias at 297 higher altitudes. The RMS difference between the two datasets decreases with altitude from 298 about 30% at 14.5 km to about 15-20% at 24.5 km.

Figure 7 shows the relative difference (= $[O_{3(Clim)} - O_{3(SAGE)}]$ /mean $[O_{3(Clim)}, O_{3(SAGE)}]$, in %) in NH winter and summer in the 1990s at 19.5 km and 24.5 km. The comparison covers parts of the globe where both datasets are available. At 19.5 km, the ozone climatology shows significant positive bias relative to SAGE data in several regions, including the region downwind of the east coast of the United States and over the Atlantic ocean in the southern hemisphere and the 304 Antarctic. There are also evident negative biases over mountains in North and South America, 305 Greenland and the Tibetan region. The magnitude of differences is large at 19.5 km (ranging 306 from -100% to 100%), in part perhaps because the ozone gradient with altitude is largest here 307 (Figure 2). At 24.5 km, differences are smaller, ranging from -40% to 40%. Differences are 308 smaller in summer than in winter (Table 3). While the negative biases over mountains may be 309 due to error in the vertical motion fields of the trajectory model, the positive biases noted do not 310 appear to be related to topography. A comparison of individual sonde stations to SAGE (not 311 shown) suggests that they may be due to individual station biases, promulgated by the trajectory 312 mapping.

313

314 **3.3 Comparison with OSIRIS Ozone Data**

315 OSIRIS is onboard the Swedish/Canadian/French/Finnish Odin satellite, which was launched 316 into a polar, sun-synchronous orbit at 600 km in February 2001 (Murtagh et al., 2002; Llewellyn 317 et al., 2004). OSIRIS can scan the Earth's limb at tangent heights from 10 to 70 km, covering 318 82.2° S to 82.2°N. The spectrograph measures scattered sunlight from 280-800 nm, with ~1 nm 319 spectral resolution. The field of view is about 1.288 arc min in the vertical direction, equivalent 320 to \sim 1 km height at the tangent point. The overall accuracy is within 10% between 15 to 35 km 321 altitudes (von Savigny et al., 2003). The mean bias of OSIRIS from coincident SAGE 322 measurements is less than 2% between 18 km to 53 km (Degenstein et al., 2009). 323 From 2002 to 2010, 5,900,889 data points from OSIRIS were matched with ozone values 324 from this ozone climatology, each pair being in the same month in the 2000s at the same location within the range of a grid ($5^{\circ} \times 5^{\circ} \times 1$ km in latitude, longitude, and altitude). The 5,900,889 pairs 325 326 of data were compared by season (Figure 8) and by latitudinal zone (Figure 9). Overall, the two

327 datasets agree reasonably well (Figures 8a, 9a and 9b). By season, the two datasets agree better 328 above 20 km (relative difference < 10%) than lower altitudes (Figure 8a). The agreement is best 329 in NH summer and fall (maximum relative difference < 5%) and poorest in winter (maximum 330 relative difference < 20%) (Figure 8a). Both datasets can generally capture profile variation with 331 season in both magnitude and profile shape (Figures 8b and 8c). The largest relative difference 332 ($\approx 20\%$) appears in winter over the UTLS region where ozone values from OSIRIS are smaller than those from this ozone climatology (Figures 8a-8c). However, it should be noted that the 333 334 very strong gradients of ozone concentration with height in this region imply that an altitude bias 335 of as little as 100 m (which could be introduced by the response time of the ascending 336 ozonesonde) could account for this difference. Examined by latitude zone (Figures 9a and 9b), 337 the largest relative differences (again, $\sim 20\%$) are found below 20 km, in the 0-45° latitude range 338 in both hemispheres. The climatology is closer to OSIRIS from 45° to 90°N, where there is 339 greater density of sonde data, with a relative difference less than 7%. Global mean profiles for 340 different latitudinal zones are very similar to each other in the two datasets, with higher ozone in 341 higher northern and southern latitudes at the same altitude (Figures 9c and 9d).

342

343 4 Global Ozone Distribution

344 **4.1 Long-term Climatology**

Figure 10 shows the latitude-altitude distribution of the ozone climatology averaged from the 1970s to the 2000s. The trajectory mapped ozone climatology can generally capture the steep ozone changes (<100 to >500 ppbv) in the vicinity of the tropopause over all latitudes. Near 25 km, the ozone climatology is lower at the poles than at lower latitudes. This distribution is

349 similar to that from SAGE data (e.g., Fioletov, 2008).

350 The mean ozone distribution from the 1970s to the 2000s at 19.5 km is shown in Figure 351 11a. In order to fill the remaining gaps ($\sim 1\%$) and to reduce small-scale "noise", the 352 climatological average values obtained from the mapping are fitted to a linear combination of 353 spherical functions (Figure 11b). Ozone concentration increases with latitude from ~500 ppbv in 354 the tropics to ~4000 ppbv at the poles, reflecting the influence of the Brewer-Dobson circulation. 355 Moreover, the ozone climatology shows significant longitudinal variation. In the northern 356 hemisphere, Figure 11 shows a wave-like pattern with high ozone over the northernmost parts of 357 North America and eastern Asia, as also found by Stohl et al. (2001), who attributed this feature 358 to the quasi-permanent troughs over the continents (e.g., Fleagel and Businger, 1980). In the 359 southern hemisphere, ozone appears higher over the Antarctic continent.

360

361 **4.2 Decadal Variability**

362 The distribution of ozone at 19.5 km is shown by decade from the 1970s to the 2000s in Figure 363 12. The large gaps in the tropics and in the middle latitudes of the southern hemisphere in the 364 1970s are largely filled in by the SHADOZ program (Southern Hemisphere ADditional 365 OZonesondes, Thompson et al., 2003a, 2003b, 2007) starting from the 1990s (Figures 1b and 366 1c). The global ozone distribution shows some consistent features in the four decades, as 367 described in section 4.1 for the long-term mean. The decadal variability is striking in both 368 hemispheres. Table 4 shows the decadal ozone variation in January and July for the globe, the 369 Arctic (60-90° N), and the Antarctic (60-90° S), averaged by only taking values from the grid 370 cells where ozone data are available in all the four decades. All regions show ozone depletion 371 from the 1970s to the 1990s and ozone recovery in the 2000s. The decadal variation is larger in 372 the winter season than in the summer season in both hemispheres, presumably because chlorine 373 is activated in polar winter. At the 25 km level, the variation is even larger (not shown).

375 4.3 Seasonal Variability

Figure 13 shows the seasonal variability of ozone in the 1990s at 19.5 km, demonstrating that 376 377 there are sufficient data to separate the seasonal variability. The seasonal variation at this level 378 resembles the variation in total ozone (e.g., Fioletov, 2008). The latitudinal gradient (lower 379 ozone in the equator and higher ozone in the middle and high latitudes) is well captured with a 380 maximum in spring and a minimum in fall. In the northern hemisphere, the spring ozone 381 maximum over the Canadian Arctic is evident. Another high ozone region in the winter and 382 spring is located over northernmost eastern Asia. This is more obvious in the 2000s (not shown), 383 because of ozone recovery in the stratosphere. Both maxima can be also partly seen in the SAGE 384 data up to 60° N (not shown). In the southern hemisphere, ozone starts to accumulate over the 385 South Pole in the SH summer (DJF), gradually increases in abundance and area in SH fall 386 (MAM) and reaches a maximum in the SH winter (JJA). In the SH spring (SON), with 387 heterogeneous destruction of ozone, ozone over the South Pole reaches a minimum, leaving a 388 band of high ozone between 30° and 60° S. This feature is reflected in the dip in the ozone 389 contours at 30°-55° S in Figure 10 above 20 km. It should be regarded with caution, however, as 390 there are relatively few ozone sounding stations in this latitude range (Figure 1), and so most of 391 the information comes via relatively long trajectories, with correspondingly larger uncertainties 392 (see also Figure 14). In addition, the lack of radiosounding stations in this region must also 393 increase the relative uncertainty in the winds in the NCEP database that are used to calculate 394 trajectories. Nevertheless, we note that a similar dip in the ozone contours is seen at 40°-60° N, 395 where ozonesonde data are most abundant. The smoothed ozone fields shown in the right panels 396 of Figure 13 are gap-filled and have reduced small-scale variation and reduced extrema. Seasonal 397 variability in the 2000s (not shown) is similar to that in the 1990s.

400

399 4.4 Data Coverage and Standard Errors

401 coverage at 19.5 km averaged over 12 months is 34% in the 1970s, increasing to 52% in the 402 1980s and to \sim 70% in the 1990s and the 2000s (Table 5). The coverage at higher or lower 403 altitudes is about the same as at this altitude. The average number of samples is around 20-25 per 404 grid cell and there are no large differences in this number between decades or among layers. The 405 averaged standard error is about 6-7% of the mean at 19.5 km and about 4-5% of the mean at 406 24.5 km (Note that standard error is calculated simply from all data points in a grid cell. 407 Depending on the wind speed, some cells may contain more than one value from an individual 408 trajectory, and so the standard error calculation will be biased low). Seasonally, most missing 409 data are in low latitudes (Figure 14). In the NH summer, data are also missing over northeast 410 Asia. In the SH summer (DJF), there are large gaps over South America and the Atlantic Ocean 411 and some regions in the eastern Pacific Ocean. As there are more stations in North America and 412 Europe (Figure 1), the number of samples per cell is the largest over these regions. In the NH 413 winter, there is a band at mid-latitudes (40-70° N) with a large number of samples. 414 Consequently, the standard error of the mean is low in the region. In the southern hemisphere 415 between 40-70° S, with fewer stations, the standard errors are higher, due to the smaller number 416 of samples. This ozone climatology was generated with 4-day forward and backward trajectories. 417 We also explored 6-day trajectories and found a general increase in data coverage and a decrease 418 in the standard error with the 6-day runs. For example, at 19.5 km in January, the coverage 419 increases from 68% to 80% and the mean standard error decreases from 4.8% to 4.7%. Future

As shown in Figure 12, the trajectory-mapped coverage is improved after the 1980s. The

420 studies may use additional data sources (e.g., SAGE and OSIRIS data) and improved trajectory

421 methods (e.g., using a variable time step in the stratosphere and troposphere) to eliminate data422 gaps and enhance accuracy.

423

424 **5** Applications

425 In the following we present two examples of applications of the ozone climatology.

426 **5.1 Stratospheric Ozone Recovery**

427 Following the implementation of the Montreal Protocol and its amendments, there has been

428 considerable interest in tracking the expected recovery of stratospheric ozone, including the dates

429 when ozone may return to its 1960 and 1980 levels and to full ozone recovery, i.e., no longer

430 influenced by human-produced ozone depleting substances (ODSs) (e.g., Weatherhead and

431 Andersen, 2006; Shepherd and Jonsson, 2008; Waugh et al., 2009; Eyring et al., 2010; Ziemke

432 and Chandra, 2012). Ozone recovery remains a challenging issue because of the complexity of

433 the natural variability of ozone, as well as the complex impact of greenhouse-gas induced

434 climate change and the removal of atmospheric ODSs owing to the Montreal Protocol. Eyring et

435 al. (2010) analyzed results from 17 chemistry-climate models and predicted that global

436 stratospheric ozone will return to its 1980 and 1960 levels by ~2025 and ~2040 respectively.

437 Ziemke and Chandra (2012) suggested an earlier recovery of stratospheric ozone on the basis of

438 their analysis of OMI (Ozone Monitoring Instrument) satellite data. Some studies have suggested

that ozone recovery would be larger at higher latitudes than lower latitudes (Weatherhead and

440 Andersen, 2006; Ziemke and Chandra, 2012).

Here we analyze ozone time series in the stratosphere using the yearly averaged ozone
climatology. Only data from grid points where ozone values are available in all years from 1970
to 2007 are employed to avoid biases due to data sampling. Using this criterion, the northern
hemisphere from 30° N to 90° N is a region with data coverage exceeding 70%. Data coverage in

other regions is somewhat lower. Figure 15 shows the annual area-weighted mean ozone 445 between 30° N and 90° N at 21.5 km (~50 hPa) and 24.5 km (~40 hPa), along with 446 447 corresponding 3-year running means. Ozone depletion before the mid-1990s is evident at both 448 levels (as well as in between, not shown). Turnaround at both levels is in 1996. By 2005, ozone 449 was about half-way to returning to its pre-1980 value at both altitudes. This figure is quite 450 similar to Figure 16a of Ziemke and Chandra (2012) which also shows ozone recovery starting 451 from the mid-1990s. The shape of the ozone variation in Figure 15a is also similar to that in 452 Waugh et al. (2009, in their Figure 2c) in a simulation. Weatherhead and Andersen (2006) 453 merged total column ozone data from TOMS (the Total Ozone Mapping Spectrometer) and 454 SBUV/2 (the Solar Backscattered Ultraviolet) satellite instruments and found the lowest ozone 455 values during 1993-1997 over 30° N to 80° N, which also is similar to our results. Their results 456 imply faster recovery in the northern part of the northern hemisphere than is indicated by some 457 atmospheric models (Weatherhead and Andersen, 2006). Figure 15 supports this conclusion, from an independent ozone data source. 458

459

460 5.2 Upper Boundary Conditions for Tropospheric Ozone Simulations in Chemistry Models 461 Previous studies have demonstrated the importance of upper boundary conditions to accurate 462 modeling of tropospheric ozone in regional chemical forecast models (Tarasick et al., 2007; 463 Makar et al., 2010, and references therein). The Canadian Global Environmental Multiscale 464 meteorology model (GEM) has been coupled with the Modelling Air quality and CHemistry 465 (MACH) system (Talbot et al., 2009) to operationally predict tropospheric ozone in a global uniform 1°×1° horizontal resolution. The GEM-MACH global model was used for the CalNex 466 467 (California Research at the Nexus of Air Quality and Climate Change) field campaign in 2010. 468 The forecasted ozone fields during the campaign from May 10 to June 20, 2010 were evaluated

469 by applying upper boundary conditions with three different ozone climatologies, including 470 monthly averaged data from HALOE (HALogen Occultation Experiment), SAGE-II, and this 471 ozone climatology for the 2000s (note that this climatology does not include ozonesonde data 472 from 2010). For this initial experiment, rather than use the stratospheric and tropospheric 473 climatologies separately, as suggested in Section 2.2, the complete stratosphere-troposphere 474 climatology was used, with a dynamic adjustment of the ozone climatology to the model-475 predicted tropopause height, as employed by Makar et al. (2010). The ozone profiles from the 476 three GEM-MACH global simulations were averaged from May 10 to June 20 and compared 477 with independent ozonesondes for the same period at six CalNex campaign stations (Figure 16). 478 Although modeled ozone for heights above 15 km are generally in good agreement, large 479 discrepancies can be found in the UTLS region (8-15 km) between the ozonesondes and the 480 simulations. Using the ozonesondes as reference, the simulated ozone profiles in the upper 481 troposphere and over the UTLS region are generally improved in shape and magnitude when this 482 climatology is used, compared to those using the satellite climatologies. The improvement 483 appears best at Joshua Tree, Point Reves, and Point Sur over 5-15 km, where HALOE and SAGE 484 tends to overestimate and underestimate the ozone value, respectively. At Shasta, Trinidad Head, 485 and Kelowna (all further north), the profiles are better in the troposphere with the new 486 ozonesonde climatology, but in the UTLS all the model profiles are too smooth compared to the 487 ozonesondes (black line), suggesting that there is dynamical activity in the UTLS that is not 488 well-represented in the model, or that the use of single stratosphere-troposphere climatologies 489 tends unduly to smooth the sharp change in mixing ratio at the tropopause. 490

491 **6** Conclusions

492 A new ozone climatology (1960s-2000s) in 3 dimensions (latitude, longitude, and altitude) has 493 been developed based on the global ozonesonde record and trajectory statistics. The input data 494 comprise 51898 ozone soundings at 111 stations over 44 years (1965-2008), obtained from the 495 WOUDC. Forward and backward trajectories are performed for 4 days each from each sounding, 496 driven by NCEP reanalysis data. The resulting global ozone climatology is archived monthly for 497 five decades from the 1960s to the 2000s with a grid size of $5^{\circ} \times 5^{\circ} \times 1$ km (latitude, longitude, and 498 altitude). The ozone climatology is also archived yearly from 1965 to 2008 at the same 499 horizontal and vertical resolutions.

500 This climatology dataset is tested at 20 selected stations by comparing the actual ozone 501 sounding profile with that derived through the trajectory technique, using the ozone soundings at 502 all stations except the one being tested. The two sets of profiles are in good agreement with an 503 overall correlation coefficient of 0.99 and an RMS error of 224 ppbv (Figure 3). By individual 504 station, the correlation coefficient ranges between 0.975 and 0.998 and RMS error between 87 505 and 482 ppbv. The ozone climatology is also compared with two sets of satellite data, from 506 SAGE and OSIRIS, in the stratosphere. Although agreement is generally quite good (within 507 10%), there are larger biases in the UTLS, especially over mountains and in areas where 508 ozonesonde measurements are particularly sparse.

The comparison with ozonesondes and satellite measurements provides us some confidence that the trajectory-mapping approach is an effective tool for interpolating sparse ozonesonde measurements. The comparison also assesses where uncertainties and limitations lie. Overall, we have more confidence in this climatology over the northern hemisphere than over the southern hemisphere, and in the middle and high latitudes than in the tropics. At the defined grid system of $5^{\circ} \times 5^{\circ} \times 1$ km (latitude, longitude, altitude), there are enough 516 1990s when SHADOZ data became available. However, there are still gaps, about 25% after the
517 1990s and mainly over the tropical oceans, which are interpolated with a spherical function
518 smoothing algorithm (Figure 13).

The ozone climatology in the long-term mean (1970s-2000s) can capture the general features in the ozone distribution, such as the sharp gradient of ozone in the vicinity of the tropopause, the latitudinal variation of tropopause height, and the ozone gradient between low and high latitudes (Figures 10 and 11). Seasonal variability of ozone in both hemispheres, as well as in the Arctic and the Antarctic, is captured (Figure 13).

524 The ozone climatology shows clearly the depletion of ozone in the northern middle and high latitudes from the 1970s to the 1990s and ozone recovery in the 2000s (Figure 15), implying 525 526 possible faster recovery of ozone in the region than predicted by most atmospheric models. This 527 is consistent with analyses of satellite data from TOMS and SBUV/2 by Weatherhead and 528 Andersen (2006) and from OMI by Ziemke and Chandra (2012). Our ozone data, however, are 529 from an independent source and a record that is twice as long as the satellite record. 530 This ozone climatology has been used as the initial and upper boundary conditions for a 531 simulation of tropospheric ozone using the operational chemical forecast model at Environment 532 Canada. The model performance is improved notably in the vicinity of the tropopause (Figure

533 16).

Since the 1990s, trajectory mapping of atmospheric species has been explored with some success, suggesting that the assumptions and principles for this approach are valid. The degree of success depends on the credibility of the driving winds and available measurements of the species of interest. The results of this study can provide insight for trajectory mapping of other species having similar or longer lifetime. In the future, this work can be improved by including 539 more ozone measurement from different platforms, such as MOZAIC and SAGE. Longer

- 540 trajectories could be employed in the stratosphere, and possibly better-resolved wind fields.
- This ozone climatology is latitudinally, longitudinally, and vertically resolved and it can reveal longitudinal variation in the stratosphere that two-dimensional ozone climatologies cannot show. It covers higher latitudes than current satellite data, and also a rather longer time period. As it is neither dependent on *a priori data* nor photochemical modeling, it provides independent information and insights that can supplement satellite data and model simulations and enhance our understanding of stratospheric ozone.
- 547

548 Acknowledgements

549 The global ozone sounding data were obtained from the World Ozone and Ultraviolet Radiation

550 Data Center (<u>http://www.woudc.org</u>) operated by Environment Canada, Toronto, Ontario,

551 Canada, under the auspices of the World Meteorological Organization. We thank many whose

dedication makes such a dataset possible. The SAGE and OSIRIS teams are appreciated for their

datasets for validation. We acknowledge the trajectory model HYSPLIT (Hybrid Single Particle

554 Lagrangian Integrated Trajectory Model) from the NOAA Air Resources Laboratory

555 (<u>http://www.arl.noaa.gov/ready.html</u>), driven by the NCEP/NCAR reanalysis data from the

556 NOAA Physical Sciences Division (<u>http://www.esrl.noaa.gov/psd/data/reanalysis</u>

557 <u>/reanalysis.shtml</u>). The first author is grateful to the Natural Sciences and Engineering Research

558 Council of Canada (NSERC) and Environment Canada for a research fellowship.

560 **References**

- 561 Attmannspacher, A., and Dütsch, H. U.: International ozone sonde intercomparison at the
- 562 Observatory Hohenpeissenberg, Ber. Dtsch. Wetterdienstes, 120, 1-85, 1970.
- 563 Attmannspacher, A., and Dütsch, H. U.: Second international ozone sonde intercomparison at
- the Observatory Hohenpeissenberg, Ber. Dtsch. Wetterdienstes, 157, 1-64, 1980.
- Baldwin, M. P., and Dunkerton, T. J.: Stratospheric harbingers of anomalous weather regimes,
 Science, 294, 581-584, 2001.
- 567 Côté, J., Gravel, S., Méthot, A., Patoine, A., and Roch, M.: The operational CMC-MRB Global
- 568 Environmental Multiscale (GEM) model, part I: Design considerations and formulation, Mon.
- 569 Wea. Rev., 126(6), 1373-1395, 1998.
- 570 Cunnold D. M., et al.: Validation SAGEII ozone measurements, J. Geophy. Res. 94, 8447-8460,
 571 1989.
- 572 Degenstein, D. A., Bourassa, A. E., Roth, C. Z., and Llewellyn, E. J.: Limb scatter ozone
- 573 retrieval from 10 to 60 km using a multiplicative algebraic reconstruction technique, Atmos.
- 574 Chem. Phys., 9, 6521–6529, doi:10.5194/acp-9-6521-2009, 2009.
- 575 Deshler, T., Mercer, J., Smit, H. G. J., Stuebi, R., Levrat, G., Johnson, B. J., Oltmans, S. J.,
- 576 Kivi, R., Thompson, A. M., Witte, J., Davies, J., Schmidlin, F. J., Brothers, G., and Sasaki T.:
- 577 Atmospheric comparison of electrochemical cell ozoneso ndes from different manufacturers,
- and with different cathode solution strengths: The Balloon Experiment on Standards for
- 579 Ozonesondes, J. Geophys. Res., 113, D04307, doi:10.1029/2007JD008975, 2008.
- 580 Draxler, R. R., and Hess, G. D.: An overview of the HYSPLIT_4 modeling system for
- trajectories, dispersion and deposition, Aust. Met. Mag., 47, 295–308, 1998.
- 582 Draxler, R., Stunder, B., Rolph, G., Stein, A., Taylor, A.: HYSPLIT_4 User's Guide, 2012.

- 583 Eyring, V., et al.: Multimodel projections of stratospheric ozone in the 21st century, J. Geophys.
- 584 Res., 112, D16303, doi:10.1029/2006JD008332, 2007.
- 585 Fioletov, V. E.: D. Ozone Climatology, Trends, and Substances that Control Ozone,
- 586 ATMOSPHERE-OCEAN 46, 39–67 doi:10.3137/ao.460103, 2008.
- 587 Fioletov, V. E., Tarasick, D. W., and Petropavlovskikh, I.: Estimating ozone variability and
- 588 instrument uncertainties from SBUV(/2), ozonesonde, Umkehr, and SAGE II measurements:
- 589 Short-term variations, J. Geophys. Res., 111, D02305, doi:10.1029/2005JD006340, 2006.
- 590 Fleagle, R. G. and Businger, J. A.: An Introduction to Atmospheric Physics, 2nd ed., Academic,
- 591 San Diego, Calif., 1980.
- 592 Forster, P. M, et al.: Evaluation of radiation scheme performance within chemistry climate
- 593 models, J. Geophys. Res., 116, D10302, doi:10.1029/2010JD015361, 2011.
- 594 Fortuin, J., and Kelder, H.: An ozone climatology based on ozonesonde and satellite
- 595 measurements, J. Geophys. Res., 103, 31,709-31,733, 1988.
- 596 Gillett, N.P., and Thompson, D. W. J.: Simulation of recent Southern Hemisphere climate
- 597 change, Science, 302, 273-275, 2003.
- 598 Hare, E. W., Carty, E. J., Fioletov, V., and Wardle, D. I.: User Guide to the WMO/GAW World
- 599 Ozone Data Centre, Version 3.0, Environment Canada, 2007.
- 600 Hassler, B., Bodeker, G. E., and Dameris, M.: Technical Note: A new global database of trace
- 601 gases and aerosols from multiple sources of high vertical resolution measurements, Atmos.
- 602 Chem. Phys., 8, 5403–5421, doi:10.5194/acp-8-5403-2008, 2008.
- 603 Hering, W. S. and Dütsch, H. U.: Comparison of chemiluminescent and electrochemical
- ozonesonde observations, J. Geophys. Res., 70(22), 5483–5490,
- 605 doi:10.1029/JZ070i022p05483, 1985.

- 606 Holton, J. R., Haynes, P. H., McIntyre, M. E., et al.: Stratosphere-troposphere exchange. Rev.
- 607 Geophys, 33(4): 403-439, 1995.
- Jacob, D. J.: Introduction to Atmospheric Chemistry, Princeton University Press: Princeton, NJ,1999.
- 510 Jones, A., Urban, J., Murtagh, D. P., Eriksson, P. S., Brohede, Haley, C., Degenstein,
- D., Bourassa, A., von Savigny, C., Sonkaew, T., Rozanov, A., Bovensmann, H., and
- Burrows, J.: Evolution of stratospheric ozone and water vapour time series studied with
- 613 satellite measurements, Atmos. Chem. Phys., 9, 6055-6075, 2009.
- Kalnay, E., Kanamitsu, M., Kistler, R., Collins, W., Deaven, D., Gandin, L., et al.: The
- 615 NCEP/NCAR 40-year reanalysis project. Bulletin of the American Meteorological Society, 77,
- 616 437–471, 1996.
- 617 Kent et al.: A model for the separation of cloud and aerosols in SAGEII occultation data, J.
- 618 *Geophys. Res.* 98, 20,725-20735,1993.
- 619 Kerr, J. B., Fast, H., McElroy, C. T., Oltmans, S. J., Lathrop, J. A., Kyro, E., Paukkunen, A.,
- 620 Claude, H., Köhler, U., Sreedharan, C. R., Takao, T., and Tsukagoshi, Y.: The 1991 WMO
- 621 international ozonesonde intercomparison at Vanscoy, Canada, Atmosphere-Ocean, 32, 685-622 716, 1994.
- 623 Lamsal, L., Weber, M., Tellmann, S., and Burrows, J.: Ozone column classified climatology of
- ozone and temperature profiles based on ozonesonde and satellite data, J. Geophys. Res., 109,
- 625 D20304, doi:10.1029/2004JD004680, 2004.
- 626 Liu, S., and Ridley, B.: Tropospheric Ozone, In Atmospheric Chemistry and Global Change,
- 627 edited by Brasseur, G. P., J. J. Orlando, G. S. Tyndall, pp. 645-686, Oxford University Press,
- 628 1999.

- 629 Liu, G, Tarasick, D. W., Fioletov, V. E., Sioris, C. E., and Rochon, Y. J.: Ozone correlation
- 630 lengths and measurement uncertainties from analysis of historical ozonesonde data in North
- 631 America and Europe, J. Geophys. Res., 114, D04112, doi:10.1029/2008JD010576, 2009.
- 632 Liu, G, Liu, J. J., Tarasick, D. W., Fioletov, V. E., Jin, J. J., Najafabadi, O., Liu, Xiong, and
- 633 Sioris, C.E.: A global tropospheric ozone climatology from trajectory-mapped ozone
- soundings, submitted to Atmospheric Chemistry and Physics Discussion, 13, 11473–11507,
- 635 2013.
- Llewellyn, E. et al.: The OSIRIS instrument on the Odin spacecraft, Can. J. Phys., 82, 411–422,
 2004.
- Makar, P. A., Gong, W., Mooney, C., Zhang, J., Davignon, D., Samaali, M., Moran, M. D., He,
- H., Tarasick, D. W., Sills, D., and Chen, J.: Dynamic adjustment of climatological ozone
- boundary conditions for air-quality forecasts, Atmos. Chem. Phys., 10, 8997-9015,
- 641 doi:10.5194/acp-10-8997-2010, 2010.
- 642 McCormick, M. P.; Zawodny, J. M., Veiga, R. E., Larsen, J. C., Wang, P. H.: An overview of
- sage I and II ozone measurements, Planet. Space Sci. 37, 1567-1586, 1989.
- 644 McLinden, C. A., Tegtmeier, S., and Fioletov, V.: Technical Note: A SAGE-corrected SBUV
- zonal-mean ozone data set, Atmos. Chem. Phys. 9, 7963-7972, 2009.
- 646 Morris, G. A., Gleason, J. F., Ziemke, J., and Schoebert, M. R.: Trajectory mapping: A tool for
- 647 validation of trace gas observations, J. Geophy. Res., 105(D14), 17,875–17,894, 2000.
- 648 Murtagh, D., Frisk, U., Merino, F., Ridal, M., Jonsson, A., Stegman, J., Witt, G., Eriksson, P.,
- Jimenez, C., Megie, G., de la Noe, J., Ricaud, P., Baron, P., Pardo, J. R., Hauchcorne, A.,
- 650 Llewellyn, E. J., Degenstein, D. A., Gattinger, R. L., Lloyd, N. D., Evans, W. F. J., McDade,
- I. C., Haley, C. S., Sioris, C., von Savigny, C., Solheim, B. H., McConnell, J. C., Strong, K.,

- 652 Richardson, E. H., Leppelmeier, G. W., Kyrola, E., Auvinen, H., and Oikarinen, L.: An
- overview of the Odin atmospheric mission, Can. J. Phys., 80, 309–319, 2002.
- Newman, P. A. and Schoeberl, M. R.: A reinterpretation of the data from the NASA
- 655 Stratosphere-Troposphere Exchange Project, Geophys. Res. Lett., 22(18), 2501–2504, 1995.
- Randel, W. and Wu, F.: Cooling of the Arctic and Antarctic Polar stratospheres due to ozone
- 657 depletion, J. Climate, 12, 1467-1479, 1999.
- Randel, W. and Wu, F.: A stratospheric ozone profile data set for 1979-2005: variability, trends,
- and comparisons with column ozone data, J. Geophys. Res., 111, D06313,
- dio:10.1029/2006JD007339, 2007.
- 661 Schoeberl, M. R., Ziemke, J. R., Bojkov, B., Livesey, N., Duncan, B., Strahan, S., and others: A
- trajectory-based estimate of the tropospheric ozone column using the residual method, J.
- 663 Geophys. Res., 112, D24S49, doi:10.1029/2007JD008773, 2007.
- 664 Shepherd, T. G. and Jonsson, A. I.: On the attribution of stratospheric ozone and temperature
- changes to changes in ozone-depleting substances and well-mixed greenhouse gases, Atmos.
- 666 Chem. Phys., 8,1435–1444, 2008.
- 667 Smit, H. G. J., Sträter, W., Helten, M., Kley, D., Ciupa, D., Claude, H. J., Köhler, U., Hoegger,
- B., Levrat, G., Johnson, B., Oltmans, S. J., Kerr, J. B., Tarasick, D. W., Davies, J., Shitamichi,
- 669 M., Srivastav, S. K., and Vialle, C.: JOSIE: The 1996 WMO international intercomparison of
- ozonesondes under quasi-flight conditions in the environmental chamber at Jülich, in
- Atmospheric Ozone: Proceedings of the Quadrennial O₃ Symposium, l'Aquila, Italy, edited by
- R. D. Bojkov and G. Visconti, pp. 971-974, Parco Sci. e Tecnol. d'Abruzzo, Italy, 1996.
- 673 Smit, H. G. J., Straeter, W., Johnson, B., Oltmans, S., Davies, J., Tarasick, D. W., Hoegger, B.,
- 674 Stubi, R., Schmidlin, F., Northam, T., Thompson, A., Witte, J., Boyd I., and Posny, F.:
- Assessment of the performance of ECC-ozonesondes under quasi-flight conditions in the

- 676 environmental simulation chamber: Insights from the Juelich Ozone Sonde Intercomparison
- 677 Experiment (JOSIE), J. Geophys Res., 112, D19306, doi:10.1029/2006JD007308, 2007.

678 SPARC CCMVal, SPARC CCMVal Report on the Evaluation of Chemistry-Climate Models, V.

- Eyring, T. G. Shepherd, D. W. Waugh (Eds.), SPARC Report No. 5, WCRP-X, WMO/TD-No.
- 680 X, 2010.
- 681 Stohl, A., James, P., Forster, C., and Spichtinger, N.: An extension of Measurement of Ozone
- and Water Vapour by Airbus-In-service aircraft (MOZAIC) ozone climatologies using
- trajectory statistics, J. Geophys. Res., 106, D21, 27,757-27,768, doi:10.1029/2001JD000749,
- 684 2001.
- 685 Stohl, A., Bonasoni, P., Cristofanelli, P., Collins, W., Feichter, J., Frank, A., Forster, C.,
- 686 Gerasopoulos, E., Gäggeler, H., James, P., Kentarchos, T., Kromp-Kolb, H., Krüger, B.,
- Land, C., Meloen, J., Papayannis, A., Priller, A., Seibert, P., Sprenger, M., Roelofs, G. J.,
- 688 Scheel, H. E., Schnabel, C., Siegmund, P., Tobler, L., Trickl, T., Wernli, H., Wirth, V., Zanis,
- 689 P., and Zerefos, C.: Stratosphere-troposphere exchange: a review, and what we have learned
- 690 from STACCATO, J. Geophys. Res., 108(D12), 8516, doi:10.1029/2002JD002490, 2003.
- 691 Sutton, R. T., Maclean, H., Swinbank, R., O'Neill, A., and Taylor, F. W.: High-resolution
- 692 stratospheric tracer fields estimated from satellite observations using Lagrangian trajectory
- 693 calculations, J. Atmos. Sci., 51, 2995-3005, 2004.
- Talbot, D.; Moran, M. D., Bouchet, V., Crevier, L.-P., Menard, S., Kallaur, A.: Development of
- a New Canadian Operational Air Quality Forecast Model. In Air Pollution Modeling and Its
- Application XIX; Borrego, C., Miranda, A.I., Eds.; Springer: Berlin, Germany, pp. 470–478,
- *697* 2009.
- 698 Tarasick, D. W., Moran, M. D., Thompson, A. M., Carey-Smith, T., Rochon, Y., Bouchet, V. S.,
- 699 Gong, W., Makar, P. A., Stroud, C., Ménard, S., Crevier, L.-P., Cousineau, S., Pudykiewicz,

- J. A., Kallaur, A., Moffet, R., Ménard, R., Robichaud, A., Cooper, O. R., Oltmans, S. J.,
- 701 Witte, J. C., Forbes, G., Johnson, B. J., Merrill, J., Moody, J. L., Morris, G., Newchurch, M.
- J., Schmidlin, F. J., and Joseph, E.: Comparison of Canadian Air Quality Forecast Models
- with tropospheric ozone profile measurements above mid-latitude north america during the
- 704 IONS/ICARTT campaign: evidence for stratospheric input, J. Geophys. Res.,
- 705 112, No. D12, D12S22, doi:10.1029/2006JD007782, 2007.
- 706 Tarasick, D. W., Jin, J. J., Fioletov, V. E., Liu, G., Tompson, A. M., Oltmans, S. J., Liu, J.,
- 707 Sioris, C. E., Liu, X., Cooper, O. R., Dann, T., and Thouret, V.: High-resolution tropospheric
- ozone fields for INTEX and ARCTAS from IONS ozonesondes, J. Geophys. Res., 115,
- 709 D20301, doi:10.1029/2009JD012918, 2010.
- 710 Thompson, A. M., Witte, J. C., McPeters, R. D., Oltmans, S. J., Schmidlin, F. J., Logan, J. A.,
- Fujiwara, M., Kirchhoff, V. W. J. H., Posny, F., Coetzee, G. J. R., Hoegger, B., Kawakami,
- S., Ogawa, T., Johnson, B. J., Vömel, H., and Labow, G.: Southern Hemisphere Additional
- 713 Ozonesondes (SHADOZ) 1998-2000 tropical ozone climatology 1. Comparison with Total
- 714 Ozone Mapping Spectrometer (TOMS) and ground-based measurements, J. Geophys. Res.,
- 715 108, D2, 8238, doi: 10.1029/2001JD000967, 2003a.
- 716 Thompson, A. M., Witte, J. C., Oltmans, S. J., Schmidlin, F. J., Logan, J. A., Fujiwara, M.,
- 717 Kirchhoff, V. W. J. H., Posny, F., Coetzee, G. J. R., Hoegger, B., Kawakami, S., Ogawa, T.,
- 718 Fortuin, J. P. F., and Kelder, H. M.: Southern Hemisphere Additional Ozonesondes
- 719 (SHADOZ) 1998-2000 tropical ozone climatology 2. Tropospheric variability and the zonal
- 720 wave-one, J. Geophys. Res., 108, D2,8241, doi: 10.1029/2002JD002241, 2003b.
- 721 Thompson, A. M., Witte, J. C., Smit, H. G. J., Oltmans, S. J., Johnson, B. J., Kirchhoff, V. W. J.
- H., and Schmidlin, F. J.: Southern Hemisphere Additional Ozonesondes (SHADOZ) 1998-
- 723 2004 tropical ozone climatology: 3. Instrumentation, station-to-station variability, and

- evaluation with simulated flight profiles, J. Geophys. Res., 112, D03304,
- 725 doi:10.1029/2005JD007042, 2007.
- von Savigny, C., Haley, C. S., Sioris, C. E., McDade, I. C., et al.: Stratospheric ozone profiles
- retrieved from Limb scattered sunlight radiance spectra measured by the OSIRIS instrument
- 728 on the Odin satellite, Geophys. Res. Lett., 30, 1755, doi:10.1029/2002GL016401,2003.
- 729 Wang, H. J., Cunnold, D. M., Thomason, L. W., Zawodny, J. M., and Bodeker, G. E.:
- Assessment of SAGE version 6.1 ozone data quality, J. Geophys. Res., 107, 4691, doi:
- 731 10.1029 / 2002JD002418, 2002.
- 732 Wang. L. and Waugh, D. W.: Chemistry-climate model simulations of recent trends in lower
- stratospheric temperatures and stratospheric residual circulation. J. Geophys. Res., 117,
- 734 D09109, doi:10.1029/2011JD017130, 2012.
- 735 Wardle, D.I., Hare, E. W., Carty, E. J., and Fioletov, V. E.: The Guide to the WMO/GAW
- 736 World Ultraviolet Radiation Data Centre (WUDC) Version 5.1, Atmospheric Environment
- 737 Service, Environment Canada, 1998.
- 738 Waugh, D. W., Oman, L., Kawa, S. R., Stolarski, R. S., Pawson, S., Douglass, A. R., Newman,
- P. A., and Nielsen J. E.: Impacts of climate change on stratospheric ozone recovery, Geophys.
- 740 Res. Lett., 36, L03805, doi:10.1029/2008GL036223, 2009.
- 741 Weatherhead, E. C. and Andersen, S. B.: The search for signs of recovery of the ozone layer,
- 742 Nature, 441, 39–45, doi:10.1038/nature04746, 2006.
- 743 WMO(World Meteorological Organization): Scientific Assessment of Ozone Depletion: 2002,
- Global Ozone Research and Monitoring Project--Report No. 47, Geneva, Switzerland, 498 pp.,
- 745 2003.

- 746 WMO (World Meteorological Organization): Scientific Assessment of Ozone Depletion: 2006,
- Global Ozone Research and Monitoring Project --Report No. 50, Geneva, Switzerland, 572
 pp., 2007.
- 749 World Climate Research Programme: SPARC/IOC/GAW Assessment of Trends in the Vertical
- 750 Distribution of Ozone, Stratospheric Processes and Their Role in Climate, World Meteorol.
- 751 Organ. Global Ozone Res. Monit. Proj. Rep. 43, Geneva, Switzerland, 1998.
- 752 Ziemke, J. R., Chandra, S., and Bhartia, P. K.: A 25-year data record of atmospheric ozone from
- 753 TOMS Cloud Slicing: Implications for trends in stratospheric and tropospheric ozone, J.
- 754 Geophys. Res., 110, D15105, doi:10.1029/2004JD005687, 2005.
- 755 Ziemke, J. R., Chandra, S., Labow, G. J., Bhartia, P. K., Froidevaux, L., and Witte, J. C.: A
- global climatology of tropospheric and stratospheric ozone derived from Aura OMI and MLS
- 757 measurements, Atmos. Chem. Phys., 11, 9237–9251, 2011.
- 758 Ziemke, J. R. and Chandra, S.: Development of a climate record of tropospheric and
- stratospheric column ozone from satellite remote sensing: evidence of an early recovery of
- 760 global stratospheric ozone, Atmos. Chem. Phys., 12, 5737-5753, 2012.

Station	Station Name	Latitude	Longitude	Altitude	Start	End	No.
ID		(°)	(°)	(m)	Year	Year	Profiles
255	Ainsworth	42.6	-100	789	1986	1986	7
229	Albrook	9	-79.6	66	1980	1980	20
18	Alert	82.5	-62.4	127	1987	2008	1068
111	Amundsen-Scott	-90	0	2820	1965	1987	272
348	Ankara	40	32.9	896	1994	2001	167
328	Ascension Island	-8	-14.4	91	1990	2005	415
199	Barrow	71.3	-156.6	11	1974	1974	3
104	Bedford	42.5	-71.3	80	1969	1971	77
181	Berlin	52.5	13.4	50	1966	1973	350
197	Biscarrosse/Sms	44.4	-1.2	18	1976	1983	359
67	Boulder	40.1	-105.3	1689	1979	1996	556
338	Bratts Lake	50.2	-104.7	592	2003	2008	229
329	Brazzaville	-4.3	15.3	314	1990	1992	82
394	Broadmeadows	-37.7	144.9	108	1999	2003	219
72	Byrd	-80	-119.5	1528	1965	1966	59
38	Cagliari	39.3	9.1	4	1968	1980	419
108	Canton Island	-2.8	-171.7	3	1965	1965	31
20	Caribou	46.9	-68	192	1981	1981	1
444	Cheju	33.5	126.5	300	2001	2001	13
224	Chilca	-12.5	-76.8	-1	1975	1975	3
138	Christchurch	-43.5	172.6	34	1965	1965	25
77	Churchill	58.8	-94.1	35	1973	2008	1431
198	Cold Lake	54.8	-110.1	702	1977	1981	66
236	Coolidge Field	17.3	-61.8	10	1976	1976	7
334	Cuiaba	-15.6	-56.1	990	1992	1992	21
450	Davis	-68.6	78	16	2003	2005	61
316	De Bilt	52.1	5.2	9.5	1994	2006	650

Table 1. Ozonesonde stations, including each station's station ID number, geolocation, the

764	number of	profiles us	sed in this	study, as	well as	measurement	period.

238	Denver	39.8	-104.9	1611	1977	1977	1
441	Easter Island	-27.2	-109.4	62	1995	1997	75
21	Edmonton	53.6	-114.1	766	1970	2008	1562
456	Egbert	44.2	-79.8	253	2003	2008	185
213	El Arenosillo	37.1	-6.7	41	1977	1983	20
335	Etosha Pan	-19.2	15.9	1100	1992	1992	16
315	Eureka	80	-86.2	310	1992	2008	1077
105	Fairbanks	64.8	-147.9	138	1965	1965	47
438	Fiji	-18.1	178.3	6	1997	2005	247
203	Ft. Sherman	9.3	-80	57	1977	1977	16
228	Gimli	50.6	-97.1	228	1980	1985	31
76	Goose Bay	53.3	-60.4	40	1965	2008	1774
237	Great Falls	47.5	-111.4	1118	1977	1977	4
40	Haute Provence	43.9	5.7	674	1981	1997	61
477	Heredia	10	-84.1	1176	2006	2006	69
109	Hilo	19.6	-155.1	11	1965	2001	679
99	Hohenpeissenberg	47.8	11	975	1966	2007	4147
344	Hong Kong	22.3	114.2	66	2000	2007	310
418	Huntsville	34.7	-86.6	196	1999	2003	160
303	Iqaluit	63.8	-68.6	20	1991	1992	30
437	Java	-7.6	112.7	50	1998	2006	264
265	Irene	-25.9	28.2	1524	1990	2006	346
336	Isfahan	32.5	51.4	1550	1999	2008	101
404	Jokioinen	60.8	23.5	103	1995	1998	99
439	Kaashidhoo	5	73.5	1	1999	1999	54
7	Kagoshima	31.6	130.6	157.5	1969	2005	841
457	Kelowna	49.9	-119.4	456	2003	2008	234
225	Kourou	5.3	-52.7	4	1974	1974	3
149	La Paz	-16.5	-68	3420	1965	1965	10
436	La Reunion Island	-21.1	55.5	24	1998	2006	232
256	Lauder	-45	169.7	370	1986	2006	1270

254	Laverton	-37.9	144.8	21	1984	1999	384
221	Legionowo	52.4	21	96	1979	2008	1404
43	Lerwick	60.1	-1.2	80	1992	2001	551
174	Lindenberg	52.2	14.1	112	1975	2008	2074
235	Long View	32.5	-94.8	103	1976	1976	2
29	Macquarie Island	-54.5	159	6	1994	2003	343
308	Madrid / Barajas	40.5	-3.7	650	1994	2008	489
400	Maitri	-70.5	11.5	223.5	1994	1998	94
448	Malindi	-3	40.2	-6	1999	2006	87
233	Marambio	-64.2	-56.6	196	1988	2006	348
466	Natal	-5.4	-35.3	32	2002	2007	177
227	Mcdonald Observatory	30.7	-90.9	2081	1969	1969	6
88	Mirny	-66.6	93	30	1989	1991	114
190	Naha	26.2	127.7	27	1989	2008	706
175	Nairobi	-1.3	36.8	1745	1996	2006	398
219	Natal	-5.9	-35.2	32	1979	2000	215
323	Neumayer	-70.7	-8.3	42	1992	2008	1198
10	New Delhi	28.5	77.2	247.5	1984	2006	146
280	Novolasarevskaya	-70.8	11.9	110	1985	1991	393
89	Ny Alesund	78.9	11.9	242.5	1990	2006	1541
210	Palestine	31.8	-95.7	121	1975	1985	163
156	Payerne	46.5	6.6	491	1968	2007	4776
217	Poker Flat	65.1	-147.5	357.5	1979	1982	40
187	Poona	18.6	73.9	559	1984	2003	112
333	Porto Nacional	-10.8	-48.4	240	1992	1992	15
242	Praha	50	14.5	304	1979	2008	1242
131	Puerto Montt	-41.5	-72.8	5	1965	1965	5
24	Resolute	74.7	-95	40	1966	2008	1649
297	S. Pietro Capofiume	44.7	11.6	11	1984	1993	98
191	Samoa	-14.3	-170.6	82	1995	2006	435
434	San Cristobal	-0.9	-89.6	8	1998	2006	271

239	San Diego	32.8	-117.2	72.5	1977	1977	2
234	San Juan	18.5	-66.1	17	1976	1976	6
401	Santa Cruz	28.4	-16.3	36	1996	2003	320
12	Sapporo	43.1	141.3	19	1969	2008	1004
443	Sepang Airport	2.7	101.7	17	1998	2007	235
132	Sofia	42.8	23.4	588	1982	1991	239
231	Spokane	47.7	-117.4	576	1976	1976	7
101	Syowa	-69	39.6	22	1966	2008	1284
432	Tahiti	-18	-149	2	1995	1999	168
95	Taipei	25	121.5	25	2000	2001	64
14	Tateno / Tsukuba	36.1	140.1	31	1968	2008	1299
157	Thalwil	46.8	8.5	515	1966	1968	183
205	Thiruvananthapuram	8.5	77	60	1984	2006	165
460	Thule	76.5	-68.7	57	1991	2003	248
65	Toronto	43.8	-79.5	198	1976	1994	16
445	Trinidad Head	40.8	-124.2	55	1999	2001	109
53	Uccle	50.8	4.4	100	1965	2007	4648
318	Valentia	51.9	-10.3	14	1994	2008	352
257	Vanscoy	52.1	-107.2	510	1990	2004	57
107	Wallops Island	37.9	-75.5	13	1970	2008	1280
64	Washington	39	-77.5	84	1965	1966	89
194	Yorkton	51.3	-102.5	504	1975	1978	72

Station ID	Station Name	Latitude (°)	Longitude (°)	Altitude (m)
328	Ascension Island	-8	-14.4	91
67	Boulder	40.1	-105.3	1689
315	Eureka	80	-86.2	310
438	Fiji	-18.1	178.3	6
76	Goose Bay	53.3	-60.4	40
109	Hilo	19.6	-155.1	11
99	Hohenpeissenberg	47.8	11	975
344	Hong Kong	22.3	114.2	66
265	Irene	-25.9	28.2	1524
336	Isfahan	32.5	51.4	1550
256	Lauder	-45	169.7	370
190	Naha	26.2	127.7	27
175	Nairobi	-1.3	36.8	1745
219	Natal	-5.9	-35.2	32
323	Neumayer	-70.7	-8.3	42
191	Samoa	-14.3	-170.6	82
434	San Cristobal	-0.9	-89.6	8
401	Santa Cruz	28.4	-16.3	36
432	Tahiti	-18	-149	2
53	Uccle	50.8	4.4	100

Table 2. Geolocation of the 20 stations for validation.

Table 3. Comparison of the ozone climatology and SAGE in the 1990s and the 2000s at three772altitudes in NH winter (DJF) and summer (JJA), respectively. All the correlation coefficients (r)773are significant at a significance level <1% (p<0.01). O_{3(Clim)} and O_{3(SAGE)} are the ozone mixing774ratios for the climatology and SAGE data, respectively, averaged over grid cells where both775data are available (the number of grid cells is indicated as N). RMS is the root mean square776difference between the climatology and SAGE data; and RMS/Mean is the ratio of the RMS to

14.5 km		19.5	19.5 km		24.5 km	
DJF	JJA	DJF	JJA	DJF	JJA	
1990s						
328	313	1736	1584	4763	4677	
312	286	1747	1640	4632	4711	
1.05	1.09	0.99	0.97	1.03	0.99	
104	87	413	337	714	508	
33	29	24	21	15	11	
0.94	0.95	0.85	0.86	0.31	0.62	
1967	1908	1633	1544	1636	1579	
2000s						
323	310	1698	1580	4950	4817	
306	286	1702	1624	4651	4737	
1.06	1.08	1.00	0.97	1.06	1.02	
116	105	405	337	819	679	
37	35	24	21	17	14	
0.93	0.92	0.87	0.87	0.32	0.46	
1930	1904	1630	1496	1584	1618	
	14.5 DJF 1990s 328 312 1.05 104 33 0.94 1967 2000s 323 306 1.06 116 37 0.93 1930	14.5 km DJF JJA 1990s 328 328 313 312 286 1.05 1.09 104 87 33 29 0.94 0.95 1967 1908 2000s 323 306 286 1.06 1.08 116 105 37 35 0.93 0.92 1930 1904	14.5 km 19.5 DJF JJA DJF 1990s 328 313 1736 312 286 1747 1.05 1.09 0.99 104 87 413 33 29 24 0.94 0.95 0.85 1967 1908 1633 2000s 2000s 2000s 310 1698 306 286 1702 1.06 1.08 1.00 116 105 405 37 35 24 0.93 0.92 0.87 1930 1904 1630	14.5 km $19.5 km$ DJFJJADJFJJA1990s32831317361584312286174716401.051.090.990.9710487413337332924210.940.950.850.861967190816331544 $2000s$ 32331016981580306286170216241.061.081.000.97116105405337373524210.930.920.870.871930190416301496	14.5 km $19.5 km$ 24.5 DJFJJADJFJJADJF $1990s$ 328 313 1736 1584 4763 312 286 1747 1640 4632 1.05 1.09 0.99 0.97 1.03 104 87 413 337 714 33 29 24 21 15 0.94 0.95 0.85 0.86 0.31 1967 1908 1633 1544 1636 $2000s$ 323 310 1698 1580 4950 306 286 1702 1624 4651 1.06 1.08 1.00 0.97 1.06 116 105 405 337 819 37 35 24 21 17 0.93 0.92 0.87 0.87 0.32 1930 1904 1630 1496 1584	

777 the mean of $O_{3(Clim)}$ and $O_{3(SAGE)}$.

781 **Table 4**. Decadal variability of ozone mixing ratio at 19.5 km averaged over the globe, the

Arctic (60-90° N), and the Antarctic (60-90° S) from the 1970s to the 2000s in January and July.

783 The mean is area-weighted and is computed taking only values from grid cells where ozone data

are available in all four decades. The change is relative to the mean in %.

The G		be	The Arctic		The An	tarctic
Decade	Ozone	Change	Ozone	Change	Ozone	Change
	(ppbv)	(%)	(ppbv)	(%)	(ppbv)	(%)
	January					
1970s	2665	8.0	3578	9.8	2772	2.9
1980s	2534	2.7	3308	1.5	2753	2.2
1990s	2297	-6.9	3044	-6.6	2592	-3.8
2000s	2375	-3.8	3108	-4.6	2656	-1.4
Mean	2468		3260		2693	
	July					
1970s	2066	8.8	2364	7.6	3519	7.6
1980s	1905	0.3	2179	-0.9	3323	1.7
1990s	1820	-4.2	2085	-5.2	3096	-5.3
2000s	1805	-4.9	2164	-1.5	3137	-4.0
Mean	1899		2198		3269	

785

Table 5. Standard error of the mean (SE/mean, in %), coverage (in % of the total grid cells), and
the number of samples per grid (in N/grid) by decade at 19.5 and 24.5 km, respectively.

		19.5 km			24.5 km	
	SE/Mean	Coverage		SE/Mean	Coverage	
Decade	(%)	(%)	N/grid	(%)	(%)	N/grid
1970s	6.3	33.5	20	3.9	30	22
1980s	7.1	51.7	18	5.0	46.1	19
1990s	6.1	74.3	26	3.9	72.8	26
2000s	6.4	72.4	23	4.0	72.8	22



Fig. 1. Global distribution of ozonesonde stations (green and red dots) used in this study: (a) all stations from the 1960s to the 2000s, (b) in the 1970s, and (c) in the 2000s. Green dots in (a) indicate stations used for validation (see Table 2). In (b) and (c), the station locations are overlaid with the coverage of ozone climatology (in green) generated in this study at 19.5 km in the 1970s and the 2000s, respectively (see Figure 12 for the ozone values).



Fig. 2. Ozone profiles at Boulder, Hohenpeissenberg, Uccle, and Eureka: decadal monthly means in January (left panels) and July (right panels) in the 1990s. The actual sounding profiles are labeled as Sonde, while the profiles from trajectories without input from the station being tested are labeled as Traj. The error bar is 4 times the standard error of the mean (equivalent to 95% confidence limits on the averages).



Fig. 3. Correlation (*r*) between ozone mixing ratios from ozonesondes and from trajectories, without input of the stations being tested, for 20 stations in the 1990s and the 2000s. The dashed line shows the 1:1 axis. The correlation is significant at the 1% level (p < 0.01). N denotes the number of data points and RMS the root mean square error. Y represents trajectory-derived ozone mixing ratio and X ozone mixing ratio from ozonesondes.



Fig. 4. Mean relative difference between the trajectory-derived ozone climatology and ozonesonde data (in red) for latitude zones $45-90^{\circ}$ N, $0^{\circ}-45^{\circ}$ N, 45° S -0° , and $90-45^{\circ}$ S in NH winter (DJF, left panels) and summer (JJA, right panels) in the 1990s. Coincident SAGE and ozonesonde profiles are compared in green. See text for details.



Fig. 5. Correlation (r) of ozone mixing ratio between trajectory-derived, SAGE, and ozonesonde data for the 1990s and 2000s for NH winter (1st row), spring (2nd row), summer (3rd row), and winter (4th row). The dashed line shows the 1:1 axis. N denotes the number of data points and RMS the root mean square error. All correlations are significant at a level better than 1%.



Fig. 6. Correlation (r) of ozone mixing ratio between trajectory-derived and ozonesonde data (left panels) and between SAGE and ozonesonde data (right panels) for the 1990s and 2000s for different latitude zones. The dashed line shows the 1:1 axis. N denotes the number of data points and RMS the root mean square error. All correlations are significant at a level better than 1%.



Fig. 7. Relative difference in ozone mixing ratio between the ozone climatology and SAGE data in NH winter (DJF, left panels) and summer (JJA, right panels) in the 1990s at 19.5 km (upper panels) and 24.5 km (lower panels). The comparison is made when both datasets are available at a cell and is expressed as the percentage difference of climatology minus SAGE divided by their mean.



Fig. 8. (a) Altitude variation in relative difference between the trajectory-derived ozone climatology and OSIRIS data in four seasons, (b) Global mean ozone profile from the ozone climatology by season. (c) Global mean ozone profile from OSIRIS by season (NH spring: MAM, summer: JJA, fall: SON, winter: DJF).



Fig. 9. Relative difference between the trajectory-derived ozone climatology and OSIRIS data in four latitude zones (a) in the northern hemisphere and (b) in the southern hemisphere, (c) global mean ozone profile from the ozone climatology in four latitude zones: $90-45^{\circ}$ S, 45° S- 0° , $0^{\circ}-45^{\circ}$ N, and $45-90^{\circ}$ N, and (d) global mean ozone profile from OSIRIS in the four zones.



Fig. 10. Long-term latitude-altitude distribution of the ozone climatology (1970s-2000s).



(b) Gap-filled and Smoothed Ozone (ppbv), 1970s-2000s, 19.5 km



Fig. 11. (a) Long-term ozone climatology (1970s-2000s) at 19.5 km with gaps (white areas). (b) The gaps are filled and the ozone field is smoothed by a spherical function-based interpolation.



Fig. 12. Decadal variation in ozone at 19.5 km for (a) the1970s, (b) the 1980s, (c) the 1990s and (d) the 2000s. White areas indicate missing data.



Fig. 13. Ozone fields with gaps (white areas) in the NH winter (DJF), spring (MAM), summer (JJA), and fall (SON) at 19.5 km in the 1990s (left panels) and the corresponding smoothed ozone fields (right panels).

(a) O₃ SE/Mean (%), 1990s, DJF, 19.5 km



(c) O₃ SE/Mean (%), 1990s, MAM, 19.5 km



(e) O₃ SE/Mean (%), 1990s, JJA, 19.5 km





80

60

40

20

0

-20

-40

-60

-80

-150

0

-100

5

-50

Latitude (Deg.)

(b) Number of Samples, 1990s, DJF, 19.5 km



(d) Number of Samples, 1990s, MAM, 19.5 km



(f) Number of Samples, 1990s, JJA, 19.5 km





Fig. 14. The standard error of the mean (left panels) and number of samples (right panels) in the NH winter (DJF), spring (MAM), summer (JJA), and fall (SON) in the 1990s at 19.5 km.



Fig. 15. Annual mean time series at (a) 21.5 km (~50 hPa) and (b) 24.5 km (~40 hPa) averaged over 30-60°N. The solid line is a 3-year running mean and the dashed line indicates the average ozone level before 1980. The area-weighted average is taken using data only from the grid cells where ozone data are available in all years.



Fig. 16. Ozone model performance compared with measurements at six ozonesonde sites during the 2010 CalNex field study, using data from HALOE, SAGE II and this new ozone climatology, respectively, as upper boundary conditions.