

# No severe ozone depletion in the tropical stratosphere in recent decades

Jayanarayanan Kuttippurath<sup>1,\*</sup>, Gopalakrishna Pillai Gopikrishnan<sup>1</sup>, Rolf Müller<sup>2</sup>, Sophie Godin-Beekmann<sup>3</sup>, Jerome Brioude<sup>4</sup>

<sup>1</sup> CORAL, Indian Institute of Technology Kharagpur, Kharagpur–721302, India

5 <sup>2</sup> Institute of Energy and Climate Research (IEK-7), Forschungszentrum Jülich, Jülich, Germany

<sup>3</sup> LATMOS, Sorbonne Université, UVSQ, CNRS, Paris, France

<sup>4</sup> Laboratoire de l'Atmosphère et des Cyclones (LACy), UMR 8105, Météo France/CNRS/Université de La Réunion, St Denis de La Réunion, France

10 *Correspondence to:* J. Kuttippurath ([jayan@coral.iitkgp.ac.in](mailto:jayan@coral.iitkgp.ac.in))

## Abstract.

Stratospheric ozone is an important constituent of the atmosphere. Significant changes in its concentrations have great consequences for the environment in general and for ecosystems, in particular. Here, we analyse ground-based, ozonesonde and satellite ozone measurements to examine the ozone depletion, and the spatiotemporal trends in ozone [in the tropics](#) during the past five decades (1980–2020). The amount of column ozone in the tropics is relatively small (250–270 DU) compared to high and mid-latitudes (Northern Hemisphere 275–425 DU; Southern Hemisphere 275–350 DU). In addition, the tropical total ozone trend is very small ( $\pm 0-0.2$  DU yr<sup>-1</sup>) as estimated for the period 1998–2022. No observational evidence is found regarding the indications or signatures of severe stratospheric ozone depletion in the tropics in contrast to a recent claim. Finally, current understanding and observational evidence do not provide any support for the possibility of an ozone hole occurring outside Antarctica today with respect to the present-day stratospheric halogen levels.

## 1 Introduction

Ozone is a triatomic molecule, and 90% of its atmospheric abundance is located in the stratosphere, roughly from 10 to 50 km above the ground (*e.g.* Cicerone, 1987). Stratospheric ozone is chemically produced in the tropical stratosphere around 25–35 km and transported to the middle and high latitudes. Therefore, stratospheric ozone mixing ratios are highest in the tropics and decrease towards the polar regions (London, 1992; Coldewey-Egbers et al., 2020). In general, the production of ozone is effective at low latitudes, and thus ozone mixing ratios at middle and high latitudes are smaller than those in the tropics. However, the ozone column, which is the integrated concentration of ozone from the surface to the top of the atmosphere (about 100 km), increases with latitude towards the poles, as its column amount is determined by atmospheric transport, which vertically [propagate](#) downwards at middle and high latitudes (*e.g.* Staehelin et al., 2001). As ozone absorbs ultraviolet radiation (UV-B radiation, 280–320 nm), a decrease in its atmospheric concentration would facilitate more UV

incidence on the Earth's surface. This is a great concern as UV-B radiation is harmful for life on the Earth (*e.g.* Bernhard et al., 2020).

35 Since the late 1970s, ozone in the Antarctic lower stratosphere has shown a dramatic seasonal decrease, which is driven by anthropogenic halogens (Farman et al., 1985). Understanding of stratospheric ozone chemistry, model simulations and measurements (*e.g.* Tuck et al., 1989; Pyle et al., 1994) showed that the decline in ozone was due to the occurrence of polar stratospheric clouds (PSCs) in winter on which the inactive halogens are converted into active forms, that catalytically destroy ozone in the presence of sunlight during spring (*e.g.* Solomon, 1986; Crutzen and Arnold, 1986; Poole and McCromick, 1988). The depletion of ozone deepened in the 1980's and peaked in the 1990s. The ozone loss in the Antarctic lower stratosphere is severe because of the unusual meteorology there, in particular winter/spring periods with very low 40 temperatures and the formation of a polar vortex that effectively isolates the mid-latitude air from polar air. For strong polar ozone loss to occur, it is essential that high levels of active chlorine are maintained up to spring (August, September and October in the Antarctic) (Müller et al., 2018). However, ozone loss in other regions, including the Arctic, never reach similar and widespread low levels as that during Antarctic spring. Note that occasionally localised atmospheric dynamics can result in short lived small areas with low column ozone or mini ozone holes (McCormack and Hood, 1997; Millán and 45 Manney, 2017).

The change in globally averaged annual total column ozone (TCO) in the mid-1990s with respect to pre-ozone hole (pre-1980) levels is about 5%, but about 17% in Antarctica, and the global TCO remains stable since the 2000s (Ball et al., 2019; Weber et al., 2018, 2022). The upper stratospheric decrease in ozone (4–8%) was induced by the increase in chlorine loading from 1980 to the late 1990s (*e.g.* Steinbrecht et al., 2017), but ozone has been steadily increasing thereafter due to the 50 reduction in stratospheric halogens (WMO, 2018; Steinbrecht et al., 2017). The decrease in upper stratospheric temperature caused by the increase in atmospheric CO<sub>2</sub> slows down the ozone loss catalytic reactions, which has also helped to increase ozone there. On the other hand, Godin-Beekmann et al. (2022) shows a 1–3%/dec reduction in the lower stratospheric ozone of both mid-latitudes and tropics since 2000. There are also studies indicating a significant reduction in ozone loss rates in Antarctica (Solomon et al., 2016; Kuttippurath and Nair, 2017; Pazmino et al., 2018), but statistically significant positive 55 trends are not detected in other regions (WMO, 2018).

In contrast to the mid-latitudes, ozone loss in the tropics is very small, and available analyses also show very small or nonsignificant trends (Randel et al., 2011; Heue et al., 2016; Lelieveld and Dentener, 2000; Staehelin and Poberaj, 2008; Thompson et al., 2021; Bogner et al., 2022). However, recently Lu (2022) claimed severe ozone depletion in the tropical stratosphere by using TOST (Trajectory mapped Ozonesonde dataset for the Stratosphere and Troposphere) data for the 60 period 1960–2010. The study claimed that there is even an ozone hole, which is seven times larger than the Antarctic ozone hole. Furthermore, the ozone hole in the tropics according to that study would be currently increasing and would be a great threat to life in the region. Chipperfield et al. (2022) in response showed that there is no robust, credible observational

evidence for tropical ozone depletion. Also, the satellite and ground-based observations show that there is only 3–5% decrease in the tropical lower stratospheric ozone, which is far lower than that reported by Lu (2022). Chipperfield et al. (2022) further observe that the number of ozonesonde profiles used by Lu (2022) is very few, which has an impact on the smoothing method used for generating the TOST data. Since the SHADOZ ozonesonde network was established in the 1990s, there have only been continuous ozone measurements since this period in the Southern Hemisphere (SH), which are inadequate to claim a year-around large ozone hole in the tropics prior to 1990. Although, the reprocessing (*i.e.* ensure high quality in the ozonesonde measurement system by following the consensus-based operating procedures and reprocessing guidelines established by ozonesonde experts around the world) has greatly enhanced the ozone data, these profiles were not considered in TOST. Furthermore, the cosmic ray driven electron-induced ozone loss in the tropics are ill-constructed, as it requires clouds like polar stratospheric clouds (PSCs), which are not present in the tropical lower stratosphere (Lu, 2010). The satellite and modelled CFC-12 data also do not support the lower stratospheric ozone depletion in the tropics (Hoffmann et al., 2014), suggesting that the results of Lu (2022) are flawed. Therefore, we present an in-depth investigation of tropical stratospheric ozone and its trend based on various ground-based, satellite and reanalysis data for the past five decades.

## 2 Data and Methods

### 2.1 GOZCARDS and SWOOSH ozone profile data

Global OZone Chemistry and Related trace gas Data records for the Stratosphere (GOZCARDS v2.2) is a bias-corrected merged satellite-based stratospheric ozone dataset for the period 1979–2018. These data are produced by combining measurements from different satellites, such as Atmospheric Chemistry Experiment Fourier Transform Spectrometer (ACE-FTS) on SCISAT, Stratospheric Aerosol and Gas Experiment (SAGE) I, its successor SAGE II, Halogen Occultation Experiment (HALOE) and Microwave Limb Sounder (MLS) on Upper Atmosphere Research Satellite (UARS) and Aura, by using SAGE II data as the primary reference. These data contain ozone mixing ratios and standard error for the altitude range of 215–0.21 hPa in 10° latitude bins. The GOZCARDS data are in good agreement with other satellite and ground-based ozone measurements. The GOZCARDS data do not show any upturn of more than 0.5–1%, which makes them suitable for global ozone trend analysis. More details can be found in Froidevaux et al. (2015).

Stratospheric Water and OzOne Satellite Homogenised (SWOOSH) version 2 is a merged data from different limb-sounding satellite instruments; SAGE II, SAGE III, HALOE, UARS MLS and Aura MLS. The primary SWOOSH data are the zonally averaged monthly-mean time series of ozone mixing ratios at pressure levels between 316 and 1 hPa. These data are available from 1980 to date on 2.5°, 5°, and 10° zonal mean grids. The measurements are homogenised by applying corrections calculated from the measurements taken during the overlap period of those instruments. The bias in different satellite data used for SWOOSH is mostly within 0.2 ppmv with respect to ozonesondes (Davis et al., 2016).

## 2.2 SBUV and GSG merged TCO data

95 Solar Backscatter Ultraviolet (SBUV) Merged Ozone Data Set (MOD v8.6) provides the longest available satellite-based  
time series of profile and TCO from a single instrument type for the period 1970–2013 (except a 5-year gap in the 1970s).  
Data from nine independent SBUV-type instruments are included in the record. Although modifications in instrument design  
were made in the evolution from the Nimbus-4 Backscattered Ultraviolet instrument to the modern SBUV ([SBUV 2](#)) type,  
the basic principle of measurement and retrieval algorithm remain the same; lending consistency to this data record  
100 compared to those based on measurements using different instrument types (Frith et al., 2018). The SBUV zonal mean ozone  
profiles agree within 10%, mostly within 5%, when compared to ground-based and other satellite measurements (Kramarova  
et al., 2013; DeLand et al., 2012).

The [merged](#) GOME/SCIAMACHY/GOME2 (GSG) TCO dataset is a comprehensive compilation of measurements from  
three satellite instruments: Global Ozone Monitoring Experiment (GOME), Scanning Imaging Absorption Spectrometer for  
105 Atmospheric Chartography (SCIAMACHY) and GOME-2 (Lerot et al., 2014). By combining data from multiple  
instruments, GSG offers improved coverage and temporal continuity from 1995 to date (Weber et al., 2018). The ozone  
retrievals are based on the University of Bremen weighing function DOAS (WFDOAS) v4 algorithm (Coldewey-Egbers et  
al., 2005). These data are in good agreement with the World Ozone and UV Data Centre (WOUDC) ozonesonde  
measurements, with an average bias of 2–3% for the zonal and global averaged values (Fioletov et al., 2002).

## 110 2.3 SHADOZ, WOUDC and TOST ozonesonde data

Southern Hemisphere ADditional OZonesondes (SHADOZ) is a project designed to measure the vertical profiles of ozone  
from a number of tropical stations using ozonesondes, which started in 1998. These measurements make use of  
Electrochemical Concentration Cell (ECC) sondes. The ECC instrument has a gas-sampling pump connected to the ozone  
sensor to a radiosonde for data telemetry (Komhyr, 1995). The accuracy of ozonesonde measurements are better than 5%. A  
115 detailed description of these data is given in Thompson et al. (2017). Table S1 lists location of [the SHADOZ stations](#).

We also use the WOUDC ECC ozonesonde data for the period 1980–2022. The ECC ozonesonde is interfaced with a  
radiosonde, which transmits the data, including ozone, atmospheric pressure, temperature and relative humidity. The  
measurements in WOUDC were performed mainly with VIZ radiosondes during the period 1980–1991, followed by RS-80  
[radiosondes](#) until 2009 and the iMet radiosondes thereafter. The VIZ radiosondes use a hypsometer for pressure  
120 measurements, and they have an accuracy of  $\pm 0.2$  hPa at altitudes above 20 hPa (Conover and Stroud, 1958). The RS-80  
radiosondes are paired with electronic boards, which are capable of transmitting data every 7 seconds. The ECC  
ozonesondes have a precision of about 3–5% and an absolute accuracy of about 10% (Tarasick et al., 2019; Smit et al.,  
2007). However, the advanced versions (v2) have improved electronic components that transmit data every second. The i-

Met radiosondes are equipped with a GPS receiver that measures the geometric altitude, in addition to atmospheric pressure  
125 (Johnson et al., 2018).

TOST is a global 3-dimensional height-resolved ozone dataset, derived from WOUDC ozone sounding records across the globe using trajectory mapping. These data are spatially interpolated using 96-hour forward and backward trajectories calculated using the HYSPLIT v 4.8 model at each 1 km altitude from the surface for a number of locations. The National Centres for Environmental Prediction/ (NCEP) meteorological data are used to drive the trajectory model. The bias of TOST  
130 data is about 10% or less, but there are larger biases in the upper troposphere lower stratosphere (UTLS) region and in areas with sparse measurements. Furthermore, the precision and accuracy of TOST data further depend on the HYSPLIT model and meteorology used for its simulations. A detailed description of the TOST data (variable used: trop\_strat\_zbith\_mean) is given in Tarasick et al. (2019) and Chipperfield et al. (2022).

#### **2.4 TROPOMI, OMI, OMPS and TOMS ozone column data**

135 Tropospheric Ozone Monitoring Instrument (TROPOMI) utilises a combination of spectral bands in the UV and visible wavelength ranges (270–850 nm), specifically designed to capture the absorption features of ozone in the Earth's atmosphere. By measuring the intensity of sunlight reflected or scattered by the atmosphere, TROPOMI can retrieve precise information on the TCO amount. With its high spatial resolution (7x5 km), TROPOMI provides global measurements and detects the ozone depletion events (Inness et al., 2019). In general, the retrieval of TCO from TROPOMI employing the  
140 GODFIT algorithm has an accuracy of about 1% (Spurr et al., 2021).

Ozone Mapping and Profile Suite (OMPS) is one among the five instruments on-board the Suomi National Polar-orbiting Partnership (Suomi NPP) satellite, which is designed to measure TCO. The spectrometer uses the backscattered solar radiances each 0.42 nm between 300 to 380 nm, with 1 nm spectral resolution. The swath of OMPS is approximately 50x2800 km<sup>2</sup>, with a field of view (FoV) of 0.27° along track and 110° across track. These measurements have a negative  
145 bias of 2–4% compared to reference products (Flynn et al., 2014).

Ozone Monitoring Instrument (OMI) has been key in providing accurate measurements of TCO from 2004 onwards. By applying the advanced Ultraviolet (UV) and Visible (VIS) spectrometry techniques, OMI captures sunlight scattered by the Earth's atmosphere to determine ozone concentrations (Levelt et al., 2006). It operates in two wavelength ranges: 270–370 nm in UV and 350–500 nm in VIS. The spectral resolution is 0.45 nm for UV and 0.63 nm for VIS. Its retrieval algorithm  
150 (DOAS) processes the spectral information to derive TCO values. Its high spatial resolution (25x25 km) enables detailed mapping of the global distribution of TCO with a bias less than 6% in the tropics and mid-latitudes (Huang et al., 2018).

The Total Ozone Mapping Spectrometers (TOMS) are a series of instruments designed to measure TCO. Here, we use TCO measurements from TOMS aboard Nimbus-7 (N7) and Earth Probe (EP) covering the period from 1979 to 2004 (McPeters et

al., 1998). TOMS employs a single monochromator and a scanning mirror to sample the backscattered solar ultraviolet radiation at 3° intervals along a line perpendicular to the orbital plane. EP-TOMS employs six discrete wavelengths ranging from 309 to 360 nm, using triangular slit functions with a nominal 1 nm bandwidth. The estimated uncertainty of TOMS data is about 3.3%, and there is a bias of 1–2% among the ozone data from different TOMS platforms (Kroon et al., 2008).

## 2.6 Methods

We have estimated the long-term trends in ozone by applying the linear method using two sets of measurements. It defines a two-sided alternative hypothesis, for which the slope of regressed line is non-zero. The standard error of the slope is estimated using the assumption of residual normality, and statistical significance of the trend is estimated by finding the p-value derived from the Wald-Test with t-distribution. We have considered the slope to be statistically significant if its p-value is < 0.05 (95% CI).

We use a multiple linear regression (MLR) model for computing the trends, which estimates the long-term change in ozone, which is driven by different processes and are represented here as the explanatory variables. The proxy variables include El Nino Southern Oscillation (ENSO), quasi-biennial oscillation at 10 hPa (QBO<sub>1</sub>), quasi-biennial oscillation at 30 hPa (QBO<sub>2</sub>), 11-year solar cycle (SF), stratospheric Aerosol Optical Depth (sAOD) and the independent linear trend terms (L<sub>pre</sub> and L<sub>post</sub>) to evaluate the change before and after the peak in ODSs in the stratosphere (Godin-Beekmann et al., 2022). The standardised and normalised (standard deviation = 1 and mean = 0) time series of ozone is regressed using the following equation:

$$y(z, t) = C_1(z, t) \cdot QBO_1(t) + C_2(z, t) \cdot QBO_2(t) + C_3(z, t) \cdot ENSO(t) + C_4(z, t) \cdot SF(t) + C_5(z, t) \cdot sAOD(t) + (C_6(z, t) + C_7(z, t)(t - t_1)) \cdot L_{pre}(t) + (C_8(z, t) + C_9(z, t)(t - t_1)) \cdot L_{post}(t) + C_{10}(z, t) \cdot Gap(t) + \varepsilon(z, t)$$

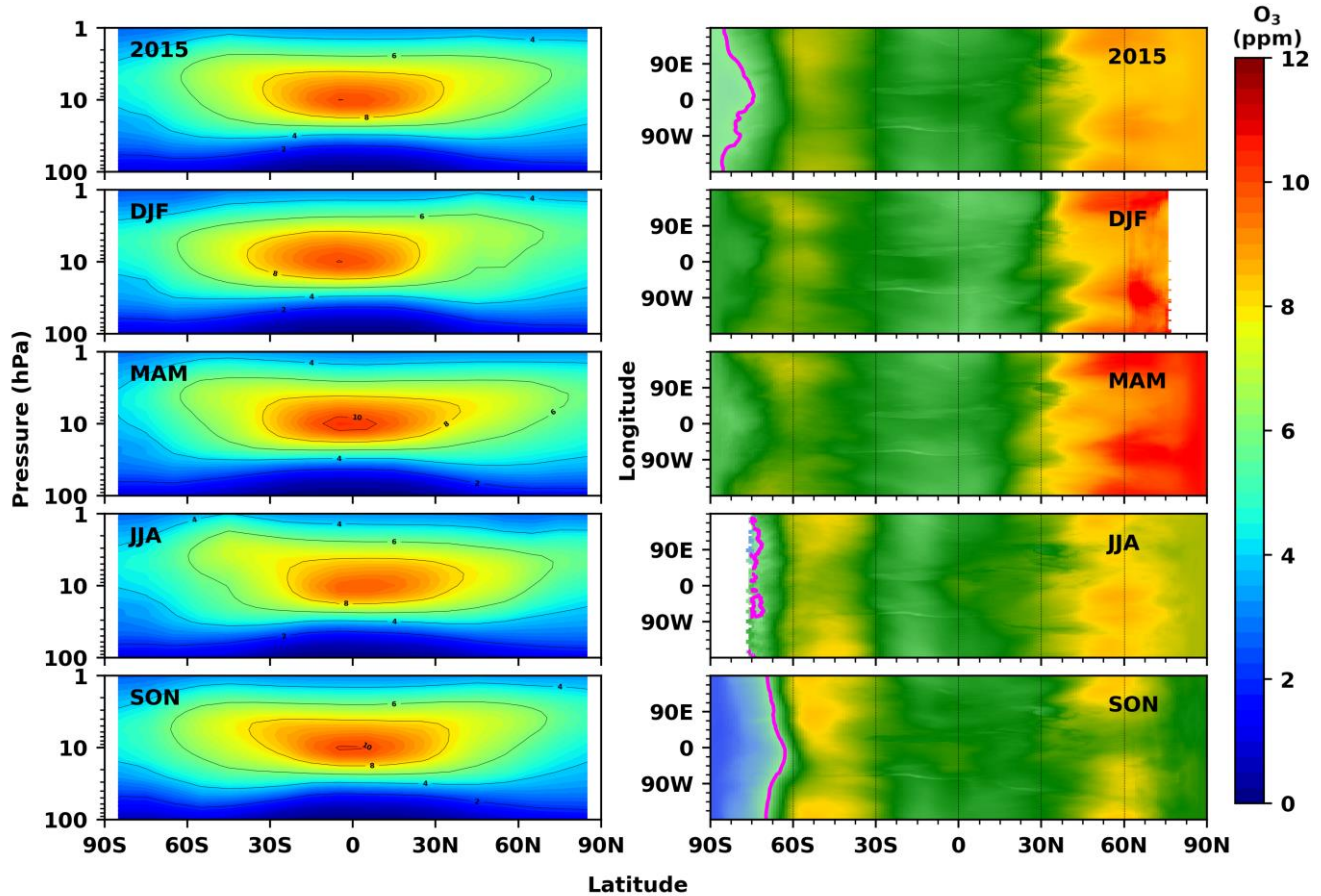
Where *Gap* is the value representing the turnaround period,  $y(z, t)$  is the ozone time series at different  $z$  altitude levels,  $C_1$  to  $C_{10}$  are the fitted coefficients,  $t_1$  is January 1997,  $t_2$  is January 2000 and  $\varepsilon$  is the residual term.

## 3. Results and Discussion

### 3.1 Ozone variability and trends across the latitudes

Figure 1 (left) shows the latitudinal distribution of zonal mean stratospheric ozone from the merged satellite data (GOZCARDS) averaged for the period 1984–2021. The data show high ozone mixing ratios (10–11 ppm at 25–35 km) in the tropics (30° N–30° S), which decrease toward the high latitudes (2–5 ppm at 25–35 km). Since the production of ozone is higher in the tropics, ozone mixing ratios are highest in the tropical middle stratosphere. As the intensity of atmospheric

180 transport is different with seasons, there are also analogous changes in ozone distribution across the latitudes and altitudes. The seasonal variability of ozone is minimal in the tropics and very high in the polar regions with respect to the latitudinal distribution of sunlight and variability of the dynamical processes. Therefore, the seasonal averages show comparatively high ozone in summer and spring, and relatively lower ozone in autumn and winter in the tropics. Since the winter transport is stronger, the ozone values in the northern hemispheric middle and high latitudes are comparatively higher during this  
 185 period (*e.g.* see the 4 ppm contour). Relatively lower ozone values are found in the mid-latitudes (*e.g.* 6–7 ppm at 10 hPa), but the lowest in the polar regions (3–4 ppm at 100 hPa). The smaller wintertime ozone values in the polar lower stratosphere (1–3 ppm) indicate the seasonal ozone loss there (*e.g.* Randel and Cobb, 1994; Chipperfield et al., 2015).



190 Figure 1: Left: Latitudinal distribution of ozone mixing ratios in ppm averaged for the period 1984–2021 and throughout the seasons, as derived from the GOZCARDS data. Right: The global seasonal and annual distribution of Total Column Ozone (TCO in DU) for 2015 as measured by Ozone Mapping and Profiler Suite (OMPS) shows the region of ozone hole (Magenta line), *i.e.* TCO less than 220 DU. The white spaces are data gaps. Here, DJF is December, January and February; MAM is March, April and May; JJA is June, July and August; SON is September, October and November.

Figure 1 (right) shows the seasonal distribution of TCO across the latitudes for 2015 measured by OMPS. In contrast to mixing ratios, the TCO distribution shows high values in the northern high latitudes in winter and spring, and very low values in the SH spring. The Antarctic ozone hole is clearly visible in austral spring, but the analysis for the boreal spring is masked by the data gaps. However, a reduction of 50–60 DU, which is the average TCO loss expected in a normal cold Arctic winter, in 0°–50° E and 100°–130° E around 70° N, is clearly captured (Goutail et al., 2005). The seasonal variation of ozone in the tropical latitudes is very small, but the SH mid-latitudes show high values in winter and NH in spring, as the Brewer Dobson Circulation (BDC) is stronger in winter and spring (Lin and Fu, 2013). Here, we have used the data from OMPS for the year 2015 to show the changes in TCO, since there was a pronounced Antarctic ozone hole in that year.

Figure 2 shows the TCO averaged over the tropics for the period 1978–2022, which is within 250–280 DU in this time period from all available measurements and reanalysis datasets. We also observe a decrease in peak TCO in the tropics during the period 1995–1999 (around 255 DU) when compared to the previous and following years (> 255 DU). However, there is an increase in TCO post-1997 and there is no significant difference (10–15 DU) in TCO among different datasets during the entire period. Furthermore, the bias in measurements from different instruments is within 5–10 DU, which shows that the data are robust and there is no substantial loss of ozone in the tropics during the period 1979–2022. The tropical column ozone is never below 220 DU. We have also computed the trends in TCO using MERRA-2, ERA-5 and satellite data (combined SBUV and OMPS measurements). The satellite-based estimates show significant negative trends ( $-0.076 \pm 0.028$  DU yr<sup>-1</sup> and  $-0.093 \pm 0.059$  DU yr<sup>-1</sup>) in the pre- and post-1997 periods, whereas the reanalysis data show nonsignificant trends in both periods. Conversely, the GSG (GOME–SCIAMACHY–GOME 2) data yield nonsignificant positive trends in the post-2000 period.

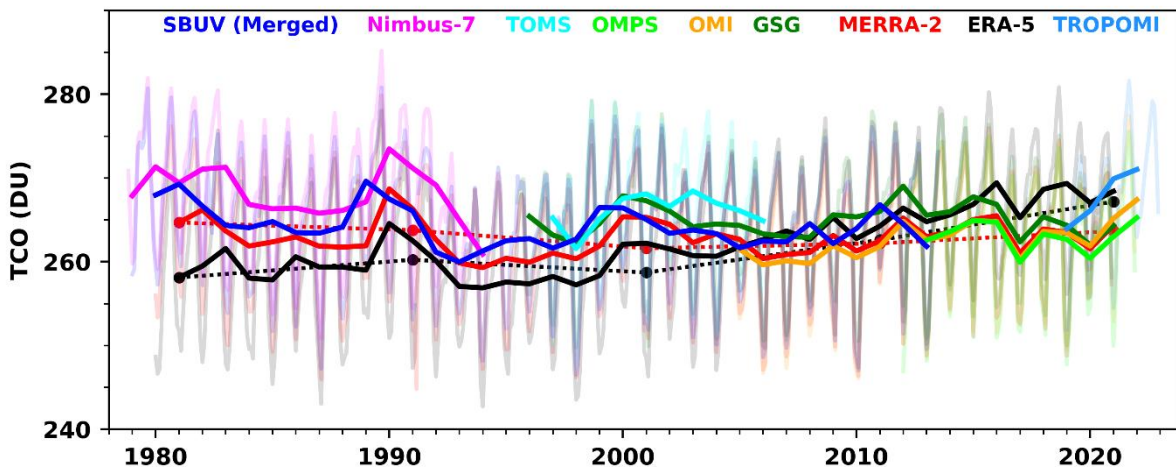


Figure 2: The distribution of Total Column Ozone (TCO in DU) averaged over the tropics (30° S–30° N) from different satellites for the period 1978–2022. The light lines show the monthly distribution, whereas dark lines show the annually averaged value of TCO. The dotted



line shows the decadal distribution of TCO from MERRA-2 and ERA-5. The peak in 1991 may be driven by the Mount Pinatubo volcanic eruption.

We have also estimated the trends in ozone in the stratosphere using the SWOOSH (Fig. 3) and GOZCARDS (Fig. S1) data for the period 1984–2022, and the trends are statistically nonsignificant (at the 95 CI) at most altitudes in both datasets. The SWOOSH estimates for the period 1984–1997 show nonsignificant, but high negative trends of about  $-0.035$  ppm  $\text{yr}^{-1}$  in the upper stratosphere and  $-0.015$  ppm  $\text{yr}^{-1}$  in the middle and lower stratosphere. Some regions also show nonsignificant positive trends ( $0.03$ – $0.04$  ppm  $\text{yr}^{-1}$ ) such as the lower stratosphere in all seasons (but DJF and JJA in GOZCARDS and these are significant). The negative trends indicate the impact of high amounts of stratospheric halogens during the period 1984–1997. In contrast, the estimates for the period 1998–2022 show nonsignificant positive trends ( $0.01$ – $0.025$  ppm  $\text{yr}^{-1}$ ) throughout the stratosphere across the seasons. The positive trends in other latitudes and altitudes are mostly within  $0.01$ – $0.02$  ppm  $\text{yr}^{-1}$ , and are significant. The highest among these trends ( $0.025 \pm 0.01$  ppm  $\text{yr}^{-1}$ ) are found in NH and SH low-latitude mid-stratosphere (above 10 hPa) in March, April, May (MAM).

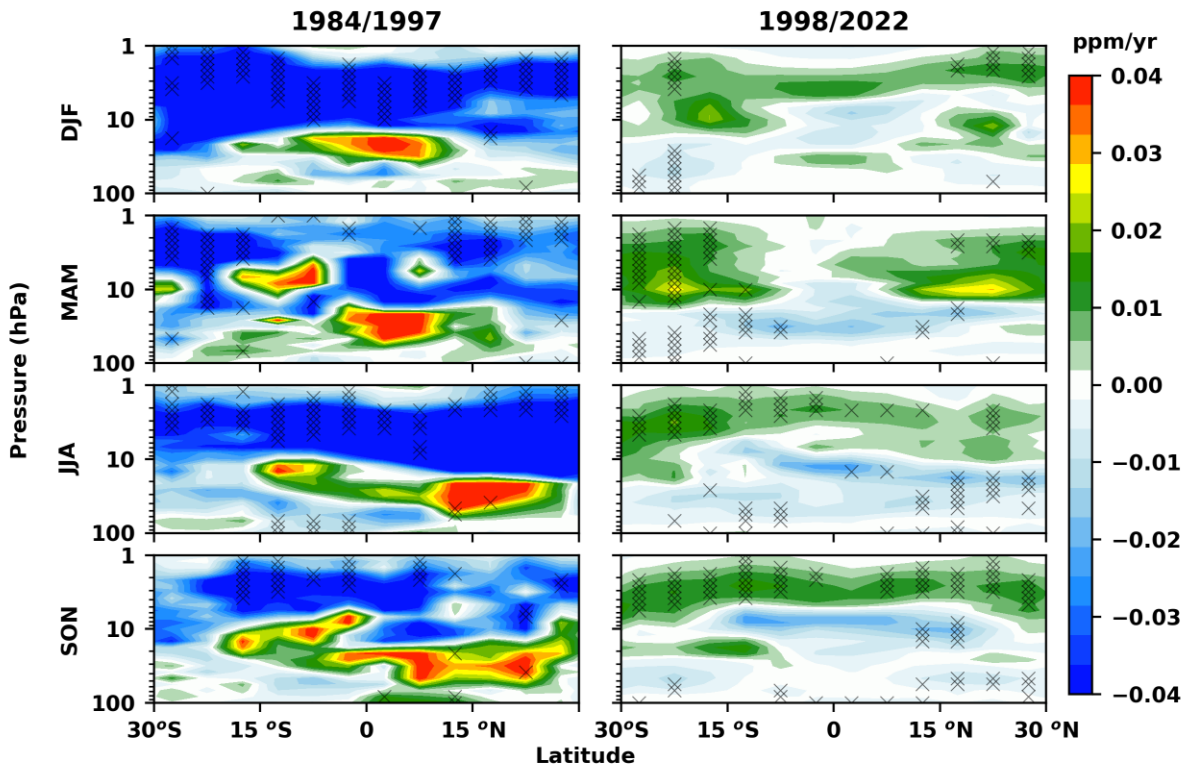


Figure 3: Trends in mixing ratio of ozone estimated for each season using the SWOOSH data for the periods of 1984–1997 and 1998–2022. The stippled regions are statistically significant at the 95% CI. Here, DJF is December, January and February; MAM is March, April and May; JJA is June, July and August; SON is September, October and November.

The GOZCARDS data also show similar trends, but those in the upper and middle stratosphere are slightly lower than that in SWOOSH in all seasons, about 0.1-0.2 ppmv yr<sup>-1</sup> in 1984–1997. However, the trend in the middle stratosphere in DJF is slightly higher at 15°–30° S in GOZACARDS during the pre-1997 period. The trend computed for the post-1997 period is very similar and those in the lower and middle stratosphere are nonsignificant in both datasets. Therefore, we have examined the difference between GOZCARDS and SWOOSH ozone, which is shown in Fig. S2. In general, GOZCARDS shows relatively higher values in the middle stratosphere (25–35 km) until 2004, which is the UARS MLS period. However, GOZCARDS shows slightly lower values with SWOOSH during the HALOE period, from 1991 to 2004, within 0.5 ppmv. The agreement between both datasets is excellent in the lower and upper stratosphere, and throughout the stratosphere during 2004–2020, within 0.1 ppmv. Our results are consistent with those of Szelag et al. (2020), as they also find significant negative trends in the lower stratosphere (up to -3% yr<sup>-1</sup>), but positive trends in the middle and upper stratosphere in spring and summer in the tropics.

### 3.2 Tropical ozone variability and trends

Our analyses (Figs. 3, S1, S3 and S5) show that there was substantial ozone loss in the 1984–1997 period at all latitudes and seasons, which is consistent in all the satellite based (GOZCARDS and SWOOSH) and reanalysis (ERA-5 and MERRA-2; see Supplementary File, Table S2) data used in this study. Neutral O<sub>3</sub> trends are also found in the tropics, and is consistent in both ozone profile and TCO measurements. Recently Lu (2022) claimed that there is strong ozone loss that he refers to as an “ozone hole” in the tropics in the past decades (1990–2020), which is reported to be present in all seasons and increasing in size day by day. The author further argues that this “ozone hole” is similar to that in Antarctica and even the chemical mechanisms causing it were the same. However, there are serious concerns about that particular study and the so-called tropical “ozone hole”. First, the data Lu (2022) used are mainly from the pre-satellite era and these data have plenty of gaps in the tropical region (Chipperfield et al., 2022). For instance, Fig. 4 shows the data used by Lu (2022), in which there are large data gaps in the tropical latitudes in all three decades (1960s, 1970s and 1980s). These data gaps are in the middle stratosphere for 1960 and 1980, but in the entire lower and middle stratosphere for 1970. The ozone values in the tropics are about 20–40 ppbv and there is hardly any significant change in tropical ozone from 1960 to 2010. Note that there is no signature of an ozone hole in Antarctica in this data (not shown), which also illustrates the problem of TOST data in accurately representing stratospheric ozone. In brief, very small values are observed in TOST in the tropics and the data gaps make it not suitable for statistical analysis. Second, the low ozone value region in the tropics is known to the scientific community for long (London, 1992) and the reason for this is the tropical upwelling branch of BDC that carries air with low ozone to the lower stratosphere (10–20 km).

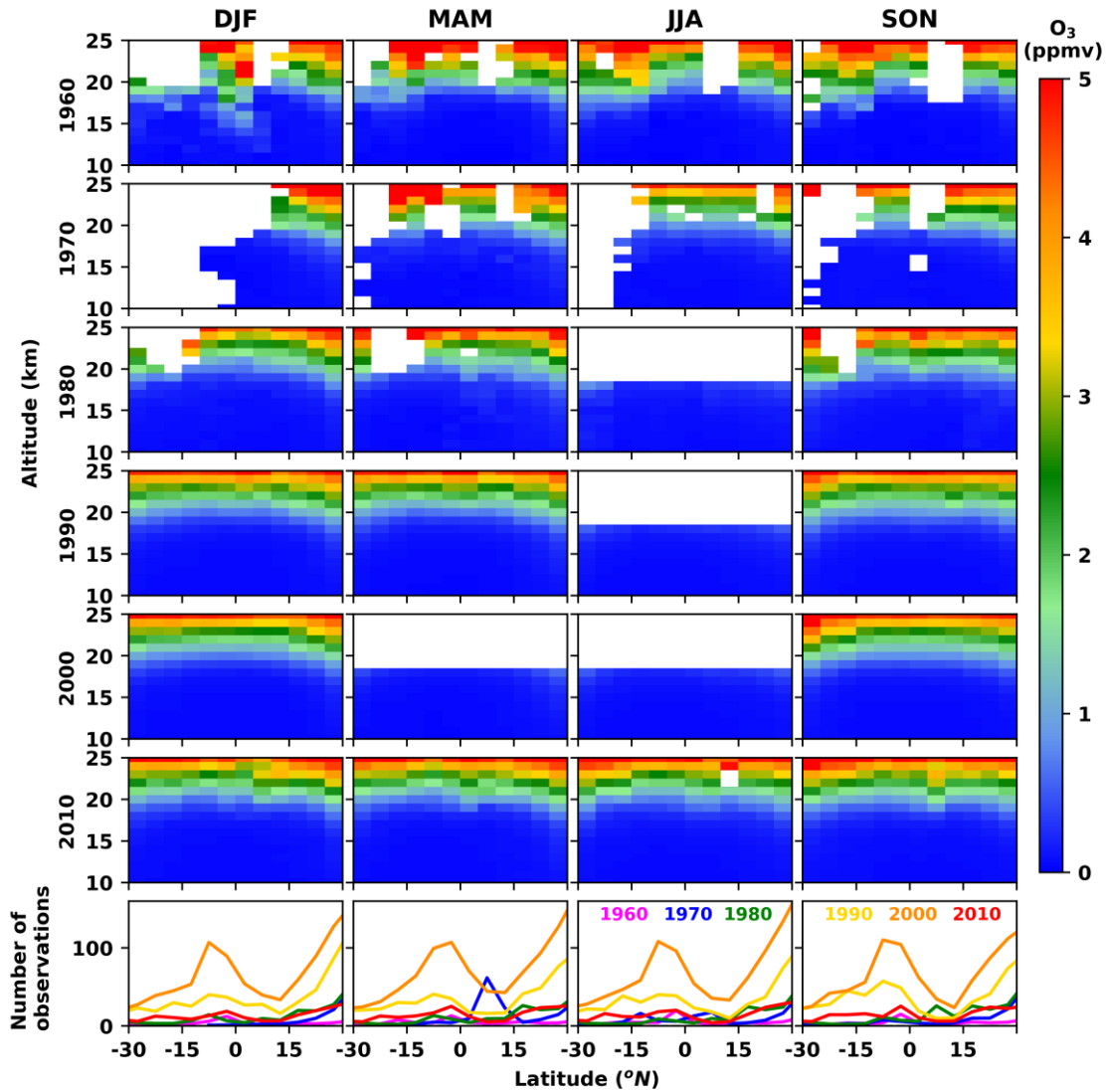
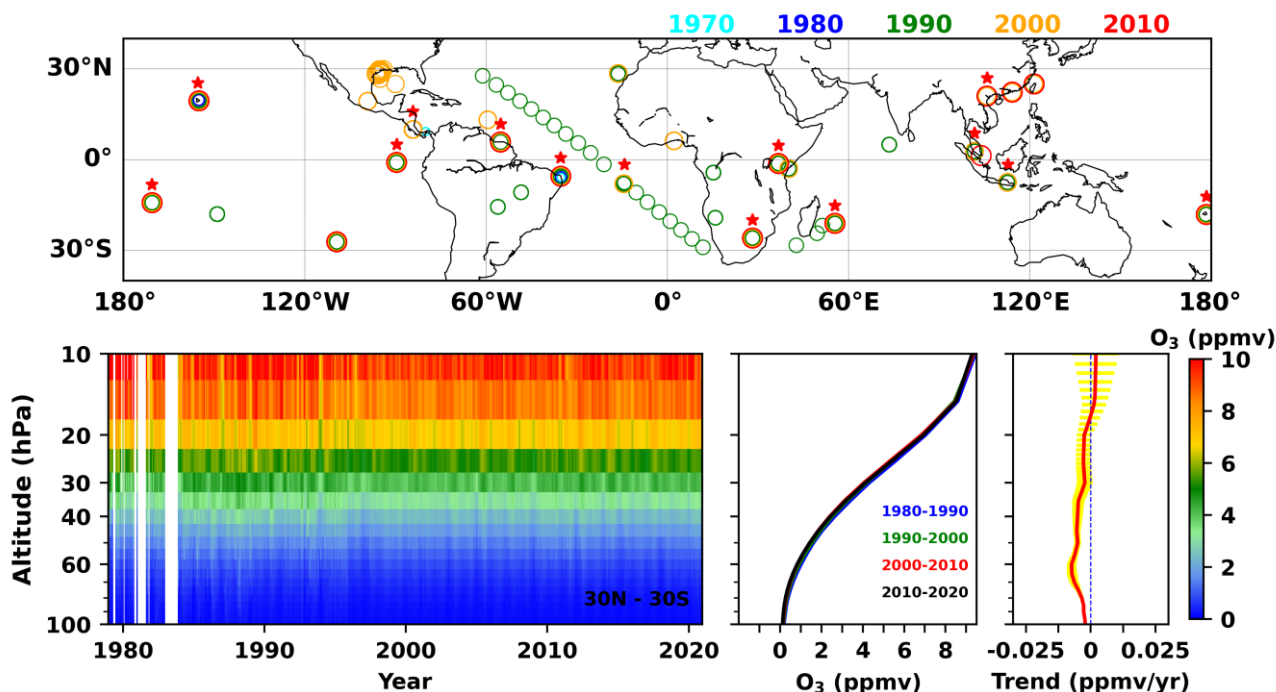


Figure 4: Average of vertical distribution of ozone from the Trajectory-mapped Ozonesonde dataset for Troposphere and Stratosphere (TOST) in each decade from 1960 to 2010. White areas indicate data gaps. Here, DJF is December, January and February; MAM is March, April and May; JJA is June, July and August; SON is September, October and November. The bottom panel shows the number of ozonesonde observations at 19 km for each decade.

We have used all ozonesonde measurements available in the tropics from WOUDC to further examine the ozone values (Fig. 5). As expected very small values are observed in the tropical lower stratosphere, approximately 2 ppm. The decadal change of ozone is also very small (middle panel) in the past four decades, and the long-term analysis shows nonsignificant trends, at about  $0.01 \pm 0.008$  ppm yr<sup>-1</sup> for all three latitude bands (0–30° N, 0–30° S and 30°–30° N/S).

270 We have also applied the MLR method to find the trend in ozone by using the SWOOSH and GOZCARDS data. The estimated trends are nonsignificant at most altitudes during the period 1984–1997 (Fig. S7). Both data show a statistically significant decline of ozone in the upper stratosphere (5–1 hPa) during the period 1984–1997. The upper stratosphere shows a negative trend of around  $-0.035 \text{ ppm yr}^{-1}$ , and the middle and lower stratosphere show a negative trend of around  $-0.015 \text{ ppm yr}^{-1}$ . Although much of these tropical regions have noticeable positive trends ( $0.03\text{--}0.04 \text{ ppm yr}^{-1}$ ), they are  
 275 nonsignificant. However, the trend estimated for the period 1998–2022 suggests that ozone is increasing ( $0.025\text{--}0.05 \text{ ppm yr}^{-1}$ ) in the stratosphere across the seasons (Figs. S8 and S9), except in the lower-latitude lower stratosphere where the values are slightly negative ( $-0.01 \pm 0.002 \text{ ppm yr}^{-1}$ ) and are statistically significant.



280 Figure 5. Top panel: Locations of the ozonesonde stations in the tropics. The Southern Hemisphere Additional OZonesondes (SHADOZ) stations are marked with a red star. Bottom left panel: Ozone profiles from WOUDC ozonesonde averaged for the tropics. Bottom Middle Panel. Mean ozone distribution over each decade in the tropics. Bottom right panel: The yearly averaged ozone trends for the period 1980–2020.

Furthermore, we have collocated the SHADOZ measurements to the nearest grids of TOST data and estimated the linear trends and bias of TOST at 15–35 km. The decadal mean of SHADOZ data does not show any significant change in ozone concentrations, except above 30–32 km, which can also be due to balloon measurement errors at these altitudes (Fig. S10).  
 285 The trend estimated for the individual SHADOZ stations exhibit either significant positive trends of about  $0.01 \pm 0.005$

ppmv yr<sup>-1</sup> or significant negative trends of about 0.01–0.035 ppm yr<sup>-1</sup> in the lower stratosphere (below 25 km). The middle stratospheric trends are neutral or positive at the SH stations.

290 The bias in TOST data, that were used by Lu (2022), estimated using the collocated SHADOZ measurements in the tropics are shown in Fig. 6. The TOST data show hardly any bias below 20 km at most stations, but a low bias (1–1.5 ppmv) above that at all stations, except Nairobi, Hilo and Irena, where the TOST data show higher bias of about 1–1.5 ppmv. This is one of the reasons for the low ozone found in the study by Lu (2022). In addition, the comparison between TOST and satellite data (GOZCARDS and SWOOSH) shows that TOST is biased low by 0.1–0.45 ppmv in the lower stratosphere, which increases with altitude (Fig. S11). Also, the ozone transported vertically from the tropical tropopause to the stratosphere  
295 usually is characterised by very low ozone values. Therefore, the low tropical ozone values are driven by dynamics (Telford et al., 2009; Chipperfield et al., 2018).

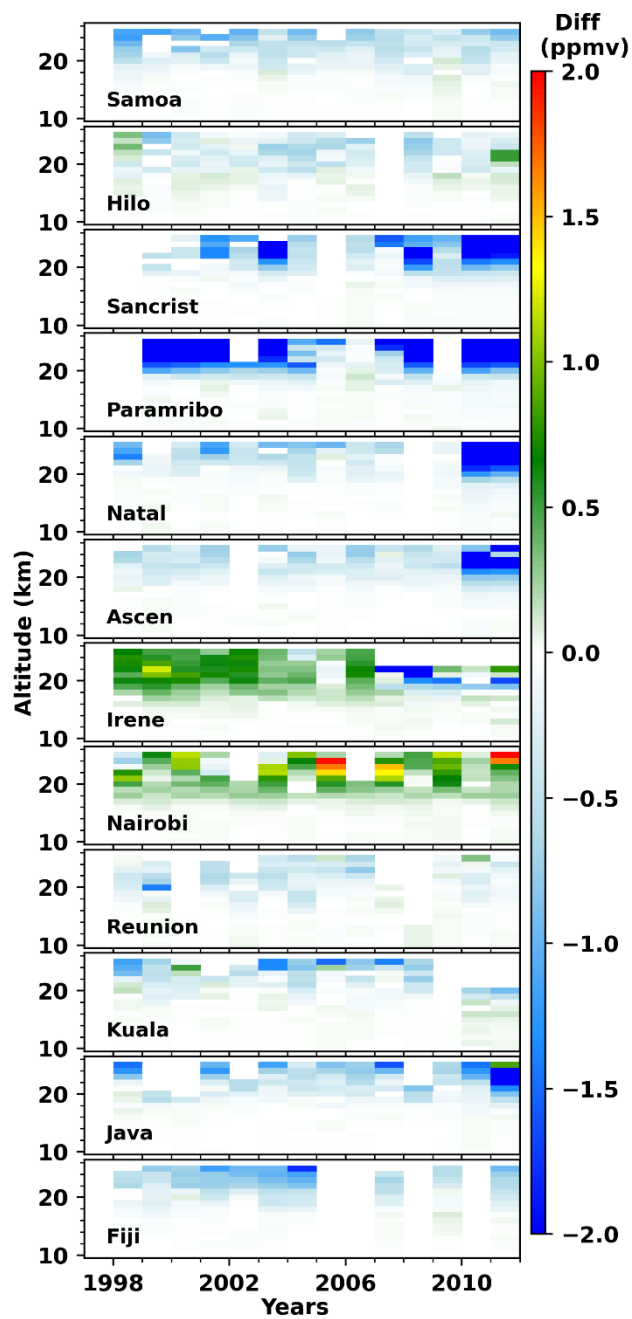


Figure 6: The bias in TOST data (TOST - SHADOZ in ppmv) calculated using collocated measurements from SHADOZ measurements for the period 1998–2012.

300 Third, Lu (2022) used the percentage change in ozone to define the “ozone hole”, which is not a good metric to show how  
much ozone is present in a region. Rather, an ozone hole definition (*i.e.* ozone values below 220 DU) should be based on the  
amount of ozone present in a region, not relative to some other decade or a period. Apart from that, ozone loss is a seasonal  
process in the polar regions and therefore the comparison must be made with respect to the period of ozone loss with respect  
305 relative to previous decades in that region. Fourth, the amount of TCO in the tropical region was never below 220 DU and  
there is a slight increase in ozone in the stratosphere and troposphere after the year 2005 (see Fig. 2). Additionally, Lu (2022)  
incorrectly assigns tropical altitudes above 10 km to the stratosphere, but troposphere extends up to 16–18 km there (Seidel  
et al., 2001), in which very low ozone can be found over the tropical Pacific due to vertical transport of clean boundary layer  
air by convection (Kley, 1997). Lu (2022) therefore incorrectly claims that Polvani et al. (2017) and Newton et al. (2018)  
310 had reported very low ozone values in the tropical lower stratosphere. Polvani et al. (2017) only discusses ozone at 70 hPa  
(18 km) and higher, whereas Newton et al. (2018) attribute the low ozone to “uplift of almost-unmixed boundary-layer air”  
to altitudes of 100–150 hPa (14–17 km). Therefore, no TCO measurements show values below 220 DU, but all depict a  
small increase in ozone after 2005, in contrast to the claim made by Lu (2022). Five, the formation of polar vortex and PSCs  
are key to ozone loss in the polar winter and spring. Formation of PSC particles is also required for the cosmic-ray-driven  
315 electron-induced reaction (CRE) mechanism put forward by Lu (2022). However, no such phenomena are reported for the  
tropical stratosphere; indeed, there is no evidence for ice particles in the tropical stratosphere in measurements (Zou et al.,  
2022; Chipperfield et al., 2022). Therefore, no such heterogeneous ozone loss is observed in the low latitudes and there is no  
basis for the CRE theory (Groß and Müller, 2011). Finally, it is already well established a couple of decades ago based on  
all then available measurements that the trends in tropical stratospheric ozone are largely absent or minimal at best for the  
320 period 1979–1997 (Staehelin et al., 2001), which is neither acknowledged nor discussed in Lu (2022).

### 3.3 Reasons for the lower values of ozone in the tropics

We also replicated the analysis made by Lu (2022), in addition to a detailed analysis by Chipperfield et al. (2022) with the  
same TOST data, and find the following issues with Lu's claim on tropical ozone loss. (i) The TOST data Lu (2022) used are  
sparse in the tropical latitudes in the troposphere and stratosphere in all three previous decades of 1960s, 1970s and 1980s  
325 (see Fig. 4, top three panels and Fig. S12). Although the values are very small (20–30 ppb), which is expected there, the data  
cannot be subtracted from another dataset with gaps in them. One cannot claim any scientific process with interpolated data  
with huge gaps in them, as shown here. (ii) As opposed to Lu's statement of continuous decline, we find a slight increase or  
no significant change in ozone from 1980 to the next decades in various independent datasets.

The tropical stratospheric ozone has increased at least by 10–20 ppb in the past decades according to our analysis of a wide  
330 range of available data, in contrast to Lu's claim that, the so-called tropical “ozone hole” were expanding. The recent  
strengthening of BDC has reduced the ozone values in the tropical stratosphere, which is reflected in the analysis of ozone

for recent decades (Butchart et al., 2006). Due to the accelerated motion of air in the tropics, the time for photochemical production of ozone is reduced, which is another reason for the declining trend in ozone there (Avalone and Prather, 1996). The enhanced ozone transport to the middle latitudes further reduces ozone in the lower stratosphere (Wargan et al., 2018).  
335 In addition to the changes in relative strength of upper and lower branches of the BDC (Butchart et al., 2006; Keeble et al., 2018; Abalos et al., 2019), the increase in halogen containing short-lived species as there are no regulations or policies to curb them (Hossaini et al., 2015; Villamayor et al., 2023), widening of extratropical troposphere (Zubov et al., 2013; Bognar et al., 2022), increased aerosol loading (Andersson et al., 2015), and unexpected emissions of CFC-11 (Fleming et al., 2020) and inorganic iodine (Cuevas et al., 2018; Karagodin-Doyennel et al., 2021) could also decrease tropical lower stratospheric  
340 ozone. There is also a study suggesting that the reduction in solar activity might reduce ozone in the tropical regions (Arsenovic et al., 2018). However, trend detection in the tropical latitudes is difficult due to the large dynamical variability there, as also found by Stone et al. (2018). Note that the warming of tropical upper troposphere causes a sharp temperature gradient between tropics and mid-latitudes, which would push the jet, and thus lift the tropopause. This, in turn, produces enhanced meridional transport between the regions (tropics to mid-latitudes) through the lower branch of BDC, and is  
345 projected to continue through the turn of the century. Henceforth, tropical lower stratospheric ozone is also expected to decline further in the coming decades (Zubov et al., 2013). In brief, the change in tropical ozone presented in Lu (2022) is mostly due to the issues in the data used in his study, and the lower values of ozone in the troposphere are driven by dynamics. In the tropics, there are no new ozone loss processes and certainly there is no “ozone hole” formed, as claimed.

Apart from these arguments, the claim by Lu (2022) regarding the lower ozone values and its impact is based on the volume  
350 (molar) mixing ratios in the tropical lower stratosphere. However, the ozone peak is around 30–35 km at these latitudes when we consider volume mixing ratios (molar mixing ratios), and hence, the analyses of Lu (2022) miss the major part of tropical ozone. When we examine the column values, they are never below 220 DU and there is no big threat from UV radiation.

#### 4. Conclusions

355 The analyses of stratospheric ozone in the tropics presented here show a consistent picture of ozone evolution in the past four decades. There is no significant loss or increase of tropical stratospheric ozone although slightly negative trends are found during the period of 2000–2020. Recent studies have suggested that the negative trends in the tropical upwelling region are caused by dynamical processes; including the increase in the speed of BDC. This is clearly pictured in the time series of tropical ozone in recent years. The long-term trend in tropical TCO for the period (1998–2022) also shows no notable  
360 difference from the past decades. Lu (2022)’s claim is solely based on one decadal dataset, which has only few profiles (see Fig. S12) and the dataset is available only for the lower stratosphere. On the other hand, here we have analysed a set of satellite, balloon-borne ground-based and reanalyses data to examine tropical ozone, and find that the claims are not properly based on measurements or model simulations, and the data Lu (2022) used are inadequate to analyse tropical stratospheric



ozone. In addition, there is no such threat as Lu (2022) claimed due to the slight negative trends in ozone in the past two  
365 decades (1998–2022) as these changes are driven by stratospheric dynamics. In summary, there is no tropical “ozone hole”  
and the evidence provided by Lu (2022) for such a phenomenon is seriously flawed.

*Data availability.* TOST data is available via <https://woudc.org/archive/products/ozone/vertical-ozone-profile/ozonesonde/1.0/tost/>, GOZCARDS and MERRA–2 data are available on <https://disc.gsfc.nasa.gov/>, ERA-5 data are available on <https://cds.climate.copernicus.eu/cdsapp#!/dataset/reanalysis-era5-single-levels?tab=overview>, SHADOZ is  
370 available via <https://tropo.gsfc.nasa.gov/shadoz/>, OMPS TCO is available at <https://ozonewatch.gsfc.nasa.gov/>, WOUDC data are available: <https://woudc.org/home.php>, SBUV MOD is available at [https://acd-ext.gsfc.nasa.gov/Data\\_services/merged/](https://acd-ext.gsfc.nasa.gov/Data_services/merged/), SWOOSH data is available at <https://csl.noaa.gov/groups/csl8/swoosh/>, TROPOMI data is available at <https://sentinel.esa.int/web/sentinel/user-guides/sentinel-5p-tropomi>, GSG data is available at <http://www.iup.uni-bremen.de/UVSAT/datasets/merged-wfdoas-total-ozone>

375 *Authorship contributions.* JK: Conceptualization, Methodology, Validation, Formal analysis, Investigation, Resources, Data Curation, Visualization, Writing - Original Draft, Writing - Review & Editing. GSG: Software, Validation, Formal analysis, Data Curation, Visualization, Writing - Review & Editing. RM: Methodology, Formal analysis, Data Curation, Writing - Review & Editing. SGB: Formal analysis, Investigation, Data Curation, Writing - Review & Editing. JB: Formal analysis, Investigation, Data Curation, Writing - Review & Editing.

380 *Competing interests.* JK, RM, and SGB are editors of ACP, otherwise, there is no competing interest.

*Acknowledgements.* We thank the Chairman, CORAL and the Director, Indian Institute of Technology Kharagpur, for providing the facility for this study. The authors thank M. P. Chipperfield, Anne Thompson and L. Froidevaux for their comments and suggestions on the draft. GSG acknowledges the Prime Ministers Research Fellowship from the Ministry of Education, GoI for his Ph.D. at IIT KGP.

385 *Financial Support.* This study did not receive any project funding.

## References

- Abalos, M., Polvani, L., Calvo, N., Kinnison, D., Ploeger, F., Randel, W., and Solomon, S.: New Insights on the Impact of Ozone-Depleting Substances on the Brewer-Dobson Circulation, *J. Geophys. Res. Atmos.*, 124, 2435–2451, <https://doi.org/10.1029/2018jd029301>, 2019.
- 390 Andersson, S. M., Martinsson, B. G., Vernier, J.-P., Friberg, J., Brenninkmeijer, C. A. M., Hermann, M., van Velthoven, P. F. J., and Zahn, A.: Significant radiative impact of volcanic aerosol in the lowermost stratosphere, *Nat. Commun.*, 6, <https://doi.org/10.1038/ncomms8692>, 2015.

- 395 Arsenovic, P., Rozanov, E., Anet, J., Stenke, A., Schmutz, W. and Peter, T. Implications of potential future grand solar minimum for ozone layer and climate. *Atmos. Chem. Phys.*, 18(5), pp.3469-3483. <https://doi.org/10.5194/acp-18-3469-2018>, 2018
- Avallone, L. M. and Prather, M. J.: Photochemical evolution of ozone in the lower tropical stratosphere, *J. Geophys. Res. Atmos.*, 101, 1457–1461, <https://doi.org/10.1029/95jd03010>, 1996.
- Ball, W. T., Alsing, J., Staehelin, J., Davis, S. M., Froidevaux, L., and Peter, T.: Stratospheric ozone trends for 1985–2018: sensitivity to recent large variability, *Atmos. Chem. Phys.*, 19, 12731–12748, <https://doi.org/10.5194/acp-19-12731-2019>, 400 2019.
- Bernhard, G. H., Fioletov, V. E., Groß, J.-U., Ialongo, I., Johnsen, B., Lakkala, K., G. L. Manney, R. Mueller, and T. Svendby: Record-breaking increases in Arctic solar ultraviolet radiation caused by exceptionally large ozone depletion in 2020. *Geophysical Research Letters*, 47, e2020GL090844. <https://doi.org/10.1029/2020GL090844>, 2020.
- 405 Bognar, K., Tegtmeier, S., Bourassa, A., Roth, C., Warnock, T., Zawada, D., and Degenstein, D.: Stratospheric ozone trends for 1984–2021 in the SAGE II–OSIRIS–SAGE III/ISS composite dataset, *Atmos. Chem. Phys.*, 22, 9553–9569, <https://doi.org/10.5194/acp-22-9553-2022>, 2022.
- Butchart, N., Scaife, A. A., Bourqui, M., de Grandpré, J., Hare, S. H. E., Kettleborough, J., Langematz, U., Manzini, E., Sassi, F., Shibata, K., Shindell, D., and Sigmond, M.: Simulations of anthropogenic change in the strength of the Brewer–Dobson circulation, *Clim. Dyn.*, 27, 727–741, <https://doi.org/10.1007/s00382-006-0162-4>, 2006.
- 410 Chipperfield, M. P., Chrysanthou, A., Damadeo, R., Dameris, M., Dhomse, S. S., Fioletov, V., Frith, S. M., Godin-Beekmann, S., Hassler, B., Liu, J., Müller, R., Petropavlovskikh, I., Santee, M. L., Stauffer, R. M., Tarasick, D., Thompson, A. M., Weber, M., and Young, P. J.: Comment on “Observation of large and all-season ozone losses over the tropics” [*AIP Adv.* 12, 075006 (2022)], *AIP Adv.*, 12, <https://doi.org/10.1063/5.0121723>, 2022.
- 415 Chipperfield, M. P., Dhomse, S. S., Feng, W., McKenzie, R. L., Velders, G. J. M., and Pyle, J. A.: Quantifying the ozone and ultraviolet benefits already achieved by the Montreal Protocol, *Nat. Commun.*, 6, <https://doi.org/10.1038/ncomms8233>, 2015.
- Cicerone, R. J.: Changes in Stratospheric Ozone, *Science*, 237, 35–42, <https://doi.org/10.1126/science.237.4810.35>, 1987.
- 420 Coldewey-Egbers, M., Weber, M., Lamsal, L. N., de Beek, R., Buchwitz, M., and Burrows, J. P.: Total ozone retrieval from GOME UV spectral data using the weighting function DOAS approach, *Atmos. Chem. Phys.*, 5, 1015–1025, <https://doi.org/10.5194/acp-5-1015-2005>, 2005.
- Conover, W.C. and Stroud, W.G. A high-altitude radiosonde hypsometer. *J. Atmos. Sci.*, 15(1), pp.63-68, [https://doi.org/10.1175/1520-0469\(1958\)015%3C0063:AHARH%3E2.0.CO;2](https://doi.org/10.1175/1520-0469(1958)015%3C0063:AHARH%3E2.0.CO;2). 1958
- Crutzen, P. J. and Arnold, F.: Nitric acid cloud formation in the cold Antarctic stratosphere: a major cause for the springtime ‘ozone hole,’ *Nature*, 324, 651–655, <https://doi.org/10.1038/324651a0>, 1986.

- 425 Cuevas, C. A., Maffezzoli, N., Corella, J. P., Spolaor, A., Vallelonga, P., Kjær, H. A., Simonsen, M., Winstrup, M., Vinther, B., Horvat, C., Fernandez, R. P., Kinnison, D., Lamarque, J.-F., Barbante, C., and Saiz-Lopez, A.: Rapid increase in atmospheric iodine levels in the North Atlantic since the mid-20th century, *Nat. Commun.*, 9, <https://doi.org/10.1038/s41467-018-03756-1>, 2018.
- Davis, S. M., Rosenlof, K. H., Hassler, B., Hurst, D. F., Read, W. G., Vömel, H., Selkirk, H., Fujiwara, M., and Damadeo, R.: The Stratospheric Water and Ozone Satellite Homogenized (SWOOSH) database: a long-term database for climate studies, *Earth Syst. Sci. Data*, 8, 461–490, <https://doi.org/10.5194/essd-8-461-2016>, 2016.
- 430 DeLand, M. T., Taylor, S. L., Huang, L. K., and Fisher, B. L.: Calibration of the SBUV version 8.6 ozone data product, *Atmos. Meas. Tech.*, 5, 2951–2967, <https://doi.org/10.5194/amt-5-2951-2012>, 2012.
- Farman, J. C., Gardiner, B. G., and Shanklin, J. D.: Large losses of total ozone in Antarctica reveal seasonal ClO<sub>x</sub>/NO<sub>x</sub> interaction, *Nature*, 315, 207–210, <https://doi.org/10.1038/315207a0>, 1985.
- 435 Fioletov, V. E.: Global and zonal total ozone variations estimated from ground-based and satellite measurements: 1964–2000, *J. Geophys. Res. Atmos.*, 107, <https://doi.org/10.1029/2001jd001350>, 2002.
- Fleming, E. L., Newman, P. A., Liang, Q., and Daniel, J. S.: The Impact of Continuing CFC-11 Emissions on Stratospheric Ozone, *J. Geophys. Res. Atmos.*, 125, <https://doi.org/10.1029/2019jd031849>, 2020.
- 440 Flury, T., Wu, D. L., and Read, W. G.: Variability in the speed of the Brewer–Dobson circulation as observed by Aura/MLS, *Atmos. Chem. Phys.*, 13, 4563–4575, <https://doi.org/10.5194/acp-13-4563-2013>, 2013.
- Flynn, L., Long, C., Wu, X., Evans, R., Beck, C. T., Petropavlovskikh, I., McConville, G., Yu, W., Zhang, Z., Niu, J., Beach, E., Hao, Y., Pan, C., Sen, B., Novicki, M., Zhou, S., and Seftor, C.: Performance of the Ozone Mapping and Profiler Suite (OMPS) products, *J. Geophys. Res. Atmos.*, 119, 6181–6195, <https://doi.org/10.1002/2013jd020467>, 2014.
- 445 Frith, S. M., Kramarova, N. A., Stolarski, R. S., McPeters, R. D., Bhartia, P. K., and Labow, G. J.: Recent changes in total column ozone based on the SBUV Version 8.6 Merged Ozone Data Set, *J. Geophys. Res. Atmos.*, 119, 9735–9751, <https://doi.org/10.1002/2014jd021889>, 2014.
- Froidevaux, L., Anderson, J., Wang, H.-J., Fuller, R. A., Schwartz, M. J., Santee, M. L., Livesey, N. J., Pumphrey, H. C., Bernath, P. F., Russell, J. M., III, and McCormick, M. P.: Global Ozone Chemistry And Related trace gas Data records for the Stratosphere (GOZCARDS): methodology and sample results with a focus on HCl, H<sub>2</sub>O, and O<sub>3</sub>, *Atmos. Chem. Phys.*, 15, 10471–10507, <https://doi.org/10.5194/acp-15-10471-2015>, 2015.
- 450 Godin-Beekmann, S., Azouz, N., Sofieva, V. F., Hubert, D., Petropavlovskikh, I., Effertz, P., Ancellet, G., Degenstein, D. A., Zawada, D., Froidevaux, L., Frith, S., Wild, J., Davis, S., Steinbrecht, W., Leblanc, T., Querel, R., Tourpali, K., Damadeo, R., Maillard Barras, E., Stübi, R., Vigouroux, C., Arosio, C., Nedoluha, G., Boyd, I., Van Malderen, R., Mahieu, E., Smale, D., and Sussmann, R.: Updated trends of the stratospheric ozone vertical distribution in the 60° S–60° N latitude range based on the LOTUS regression model, *Atmos. Chem. Phys.*, 22, 11657–11673, <https://doi.org/10.5194/acp-22-11657-2022>, 2022.

- 460 Goutail, F., Pommereau, J.-P., Lefèvre, F., van Roozendael, M., Andersen, S. B., Kåstad Høiskar, B.-A., Dorokhov, V., Kyrö, E., Chipperfield, M. P., and Feng, W.: Early unusual ozone loss during the Arctic winter 2002/2003 compared to other winters, *Atmos. Chem. Phys.*, 5, 665–677, <https://doi.org/10.5194/acp-5-665-2005>, 2005.
- Groß, J.-U. and Müller, R.: Do cosmic-ray-driven electron-induced reactions impact stratospheric ozone depletion and global climate change? *Atmos. Environ.*, 45, 3508–3514, <https://doi.org/10.1016/j.atmosenv.2011.03.059>, 2011.
- 465 Heue, K.-P., Coldewey-Egbers, M., Delcloo, A., Lerot, C., Loyola, D., Valks, P., and van Roozendael, M.: Trends of tropical tropospheric ozone from 20 years of European satellite measurements and perspectives for the Sentinel-5 Precursor, *Atmos. Meas. Tech.*, 9, 5037–5051, <https://doi.org/10.5194/amt-9-5037-2016>, 2016.
- Hoffmann, L., Hoppe, C.M., Müller, R., Dutton, G.S., Gille, J.C., Griessbach, S., Jones, A., Meyer, C.I., Spang, R., Volk, C.M. and Walker, K.A. Stratospheric lifetime ratio of CFC-11 and CFC-12 from satellite and model climatologies. *Atmos. Chem. Phys.*, 14(22), pp.12479-12497. <https://doi.org/10.5194/acp-14-12479-2014>, 2014
- 470 Hossaini, R., Chipperfield, M. P., Montzka, S. A., Rap, A., Dhomse, S., and Feng, W.: Efficiency of short-lived halogens at influencing climate through depletion of stratospheric ozone, *Nat. Geosci.*, 8, 186–190, <https://doi.org/10.1038/ngeo2363>, 2015.
- Huang, G., Liu, X., Chance, K., Yang, K., and Cai, Z.: Validation of 10-year SAO OMI ozone profile (PROFOZ) product using Aura MLS measurements, *Atmos. Meas. Tech.*, 11, 17–32, <https://doi.org/10.5194/amt-11-17-2018>, 2018.
- 475 Inness, A., Flemming, J., Heue, K.-P., Lerot, C., Loyola, D., Ribas, R., Valks, P., van Roozendael, M., Xu, J., and Zimmer, W.: Monitoring and assimilation tests with TROPOMI data in the CAMS system: near-real-time total column ozone, *Atmos. Chem. Phys.*, 19, 3939–3962, <https://doi.org/10.5194/acp-19-3939-2019>, 2019.
- Johnson, B., Schnell, R.C., Cullis, P., Sterling, C.W., Jordan, A.F. and Petropavlovskikh, I.V. December. Data Homogenization Results from Three NOAA Long-term ECC Ozone Sonde Records. In AGU Fall Meeting Abstracts (Vol. 480 2018, pp. A31I-2959), <https://ui.adsabs.harvard.edu/abs/2018AGUFM.A31I2959J/abstract>, 2018
- Karagodin-Doyennel, A., Rozanov, E., Sukhodolov, T., Egorova, T., Saiz-Lopez, A., Cuevas, C. A., Fernandez, R. P., Sherwen, T., Volkamer, R., Koenig, T. K., Giroud, T., and Peter, T.: Iodine chemistry in the chemistry–climate model SOCOL-AERv2-I, *Geosci. Model Dev.*, 14, 6623–6645, <https://doi.org/10.5194/gmd-14-6623-2021>, 2021.
- 485 Keeble, J., Brown, H., Abraham, N. L., Harris, N. R. P., and Pyle, J. A.: On ozone trend detection: using coupled chemistry–climate simulations to investigate early signs of total column ozone recovery, *Atmos. Chem. Phys.*, 18, 7625–7637, <https://doi.org/10.5194/acp-18-7625-2018>, 2018.
- Kley, D.: Tropospheric chemistry and transport. *Science*, 276(5315), pp.1043-1044, 1997.
- Komhyr, W. D., Barnes, R. A., Brothers, G. B., Lathrop, J. A., and Opperman, D. P.: Electrochemical concentration cell ozonesonde performance evaluation during STOIC 1989, *J. Geophys. Res. Atmos.*, 100, 9231–9244, 490 <https://doi.org/10.1029/94jd02175>, 1995.

- Kramarova, N. A., Frith, S. M., Bhartia, P. K., McPeters, R. D., Taylor, S. L., Fisher, B. L., Labow, G. J., and DeLand, M. T.: Validation of ozone monthly zonal mean profiles obtained from the version 8.6 Solar Backscatter Ultraviolet algorithm, *Atmos. Chem. Phys.*, 13, 6887–6905, <https://doi.org/10.5194/acp-13-6887-2013>, 2013.
- Kroon, M., Veeffkind, J. P., Sneep, M., McPeters, R. D., Bhartia, P. K., and Levelt, P. F.: Comparing OMI-TOMS and OMI-DOAS total ozone column data, *J. Geophys. Res. Atmos.*, 113, <https://doi.org/10.1029/2007jd008798>, 2008.
- 495
- Kuttippurath, J. and Nair, P. J.: The signs of Antarctic ozone hole recovery, *Sci. Rep.*, 7, <https://doi.org/10.1038/s41598-017-00722-7>, 2017.
- Lelieveld, J. and Dentener, F. J.: What controls tropospheric ozone? *J. Geophys. Res. Atmos.*, 105, 3531–3551, <https://doi.org/10.1029/1999jd901011>, 2000.
- 500
- Lerot, C., Van Roozendaal, M., Spurr, R., Loyola, D., Coldewey-Egbers, M., Kochenova, S., van Gent, J., Koukouli, M., Balis, D., Lambert, J. -C., Granville, J., and Zehner, C.: Homogenized total ozone data records from the European sensors GOME/ERS-2, SCIAMACHY/Envisat, and GOME-2/MetOp-A, *J. Geophys. Res. Atmos.*, 119, 1639–1662, <https://doi.org/10.1002/2013jd020831>, 2014.
- Levelt, P. F., van den Oord, G. H. J., Dobber, M. R., Malkki, A., Huib Visser, Johan de Vries, Stammes, P., Lundell, J. O. V., and Saari, H.: The ozone monitoring instrument, *IEEE Trans. Geosci. Remote Sensing*, 44, 1093–1101, <https://doi.org/10.1109/tgrs.2006.872333>, 2006.
- 505
- Lin, P. and Fu, Q.: Changes in various branches of the Brewer-Dobson circulation from an ensemble of chemistry climate models, *J. Geophys. Res. Atmos.*, 118, 73–84, <https://doi.org/10.1029/2012jd018813>, 2013.
- Liu, G., Liu, J., Tarasick, D. W., Fioletov, V. E., Jin, J. J., Moeini, O., Liu, X., Sioris, C. E., and Osman, M.: A global tropospheric ozone climatology from trajectory-mapped ozone soundings, *Atmos. Chem. Phys.*, 13, 10659–10675, <https://doi.org/10.5194/acp-13-10659-2013>, 2013.
- 510
- London, J. and Liu, S. C.: Long-term tropospheric and lower stratospheric ozone variations from ozonesonde observations, *J. Atmos. Terr. Phys.*, 54, 599–625, [https://doi.org/10.1016/0021-9169\(92\)90100-y](https://doi.org/10.1016/0021-9169(92)90100-y), 1992.
- Lu, Q.-B.: Cosmic-ray-driven electron-induced reactions of halogenated molecules adsorbed on ice surfaces: Implications for atmospheric ozone depletion and global climate change, *Phys. Rep.*, 487, 141–167, <https://doi.org/10.1016/j.physrep.2009.12.002>, 2010.
- 515
- Lu, Q.-B.: Observation of large and all-season ozone losses over the tropics, *AIP Advan.*, 12, <https://doi.org/10.1063/5.0094629>, 2022.
- Millan, L. and Manney, G.: An assessment of Ozone Mini-holes Representation in Reanalyses Over the Northern Hemisphere. *Atmospheric Chemistry and Physics Discussions*, pp.1-22, 2017
- 520
- McCormack, J.P. and Hood, L.L.: The frequency and size of ozone “mini-hole” events at northern midlatitudes in February. *Geophysical research letters*, 24(21), pp.2647-2650, 1997

- McPeters, R.: Earth probe total ozone mapping spectrometer (TOMS) data product user's guide (Vol. 206895). National Aeronautics and Space Administration, Goddard Space Flight Center, 525 <https://ntrs.nasa.gov/api/citations/19990019486/downloads/19990019486.pdf>, 1998.
- Müller, R., Grooß, J.-U., Zafar, A. M., Robrecht, S., and Lehmann, R.: The maintenance of elevated active chlorine levels in the Antarctic lower stratosphere through HCl null cycles, *Atmos. Chem. Phys.*, 18, 2985–2997, <https://doi.org/10.5194/acp-18-2985-2018>, 2018.
- Newton, R., Vaughan, G., Hints, E., Filus, M.T., Pan, L.L., Honomichl, S., Atlas, E., Andrews, S.J. and Carpenter, L.J.: 530 Observations of ozone-poor air in the tropical tropopause layer. *Atmospheric chemistry and physics*, 18(7), pp.5157-5171, 2018
- Pazmiño, A., Godin-Beekmann, S., Hauchecorne, A., Claud, C., Khaykin, S., Goutail, F., Wolfram, E., Salvador, J., and Quel, E.: Multiple symptoms of total ozone recovery inside the Antarctic vortex during austral spring, *Atmos. Chem. Phys.*, 18, 7557–7572, <https://doi.org/10.5194/acp-18-7557-2018>, 2018.
- 535 Polvani, L.M., Wang, L., Aquila, V. and Waugh, D.W.: The impact of ozone-depleting substances on tropical upwelling, as revealed by the absence of lower-stratospheric cooling since the late 1990s. *Journal of Climate*, 30(7), pp.2523-2534, 2017.
- Poole, L. R. and McCormick, M. P.: Polar stratospheric clouds and the Antarctic ozone hole, *J. Geophys. Res. Atmos.*, 93, 8423–8430, <https://doi.org/10.1029/jd093id07p08423>, 1988.
- Pyle, J. A., Harris, N. R. P., Farman, J. C., Arnold, F., Braathen, G., Cox, R. A., Faucon, P., Jones, R. L., Megie, G., O'Neill, 540 A., Platt, U., Pommereau, J. -P., Schmidt, U., and Stordal, F.: An overview of the EASOE Campaign, *Geophys. Res. Lett.*, 21, 1191–1194, <https://doi.org/10.1029/94gl00004>, 1994.
- Randel, W. J. and Cobb, J. B.: Coherent variations of monthly mean total ozone and lower stratospheric temperature, *J. Geophys. Res. Atmos.*, 99, 5433–5447, <https://doi.org/10.1029/93jd03454>, 1994.
- Randel, W. J. and Thompson, A. M.: Interannual variability and trends in tropical ozone derived from SAGE II satellite data 545 and SHADOZ ozonesondes, *J. Geophys. Res. Atmos.*, 116, <https://doi.org/10.1029/2010jd015195>, 2011.
- Seidel, D.J. and Randel, W.J. Recent widening of the tropical belt: Evidence from tropopause observations. *Journal of Geophysical Research: Atmospheres*, 112(D20), 2007
- Smit, H.G., Straeter, W., Johnson, B.J., Oltmans, S.J., Davies, J., Tarasick, D.W., Hoegger, B., Stubi, R., Schmidlin, F.J., Northam, T. and Thompson, A.M.: Assessment of the performance of ECC-ozonesondes under quasi-flight conditions in the 550 environmental simulation chamber: Insights from the Juelich Ozone Sonde Intercomparison Experiment (JOSIE). *Journal of Geophysical Research: Atmospheres*, 112(D19), 2007
- Solomon, S.: Progress towards a quantitative understanding of Antarctic ozone depletion, *Nature*, 347, 347–354, <https://doi.org/10.1038/347347a0>, 1990.
- Solomon, S., Ivy, D. J., Kinnison, D., Mills, M. J., Neely, R. R., III, and Schmidt, A.: Emergence of healing in the Antarctic 555 ozone layer, *Science*, 353, 269–274, <https://doi.org/10.1126/science.aae0061>, 2016.

Solomon, S., Garcia, R. R., Rowland, F. S., and Wuebbles, D. J.: On the depletion of Antarctic ozone, *Nature*, 321, 755–758, <https://doi.org/10.1038/321755a0>, 1986.

SPARC: Chapter 4: Overview of Ozone and Water vapour [https://www.sparc-climate.org/wp-content/uploads/sites/5/2022/01/04\\_S-RIP\\_Report\\_Ch04.pdf](https://www.sparc-climate.org/wp-content/uploads/sites/5/2022/01/04_S-RIP_Report_Ch04.pdf), 2017

560 Spurr, R., Loyola, D., Heue, K.P., Van Roozendaal, M. and Lerot, C.: S5P/TROPOMI Total Ozone ATBD. Deutsches Zentrum für Luftund Raumfahrt (German Aerospace Center), Weßling, Germany, Tech. Rep. S5P-L2-DLR-ATBD-400A, 2021

Stahelin, J., and Poberaj, C. S.: Long-term Tropospheric Ozone Trends: A Critical Review, *Climate Variability and Extremes during the Past 100 Years*, 271–282, [https://doi.org/10.1007/978-1-4020-6766-2\\_18](https://doi.org/10.1007/978-1-4020-6766-2_18), 2008.

565 Stahelin, J., Harris, N. R. P., Appenzeller, C., and Eberhard, J.: Ozone trends: A review, *Rev. Geophys.*, 39, 231–290, <https://doi.org/10.1029/1999rg000059>, 2001.

Steinbrecht, W., Froidevaux, L., Fuller, R., Wang, R., Anderson, J., Roth, C., Bourassa, A., Degenstein, D., Damadeo, R., Zawodny, J., Frith, S., McPeters, R., Bhartia, P., Wild, J., Long, C., Davis, S., Rosenlof, K., Sofieva, V., Walker, K., Rahpoe, N., Rozanov, A., Weber, M., Laeng, A., von Clarmann, T., Stiller, G., Kramarova, N., Godin-Beekmann, S.,  
570 Leblanc, T., Querel, R., Swart, D., Boyd, I., Hocke, K., Kämpfer, N., Maillard Barras, E., Moreira, L., Nedoluha, G., Vigouroux, C., Blumenstock, T., Schneider, M., García, O., Jones, N., Mahieu, E., Smale, D., Kotkamp, M., Robinson, J., Petropavlovskikh, I., Harris, N., Hassler, B., Hubert, D., and Tummon, F.: An update on ozone profile trends for the period 2000 to 2016, *Atmos. Chem. Phys.*, 17, 10675–10690, <https://doi.org/10.5194/acp-17-10675-2017>, 2017.

Stone, K. A., Solomon, S., and Kinnison, D. E.: On the Identification of Ozone Recovery, *Geophys. Res. Lett.*, 45, 5158–  
575 5165, <https://doi.org/10.1029/2018gl077955>, 2018.

Szelağ, M. E., Sofieva, V. F., Degenstein, D., Roth, C., Davis, S., and Froidevaux, L.: Seasonal stratospheric ozone trends over 2000–2018 derived from several merged data sets, *Atmos. Chem. Phys.*, 20, 7035–7047, <https://doi.org/10.5194/acp-20-7035-2020>, 2020.

Tarasick, D. W., Carey-Smith, T. K., Hocking, W. K., Moeini, O., He, H., Liu, J., Osman, M. K., Thompson, A. M.,  
580 Johnson, B. J., Oltmans, S. J., and Merrill, J. T.: Quantifying stratosphere-troposphere transport of ozone using balloon-borne ozonesondes, radar windprofilers and trajectory models, *Atmos. Environ.*, 198, 496–509, <https://doi.org/10.1016/j.atmosenv.2018.10.040>, 2019.

Telford, P., Braesicke, P., Morgenstern, O., and Pyle, J.: Reassessment of causes of ozone column variability following the eruption of Mount Pinatubo using a nudged CCM, *Atmos. Chem. Phys.*, 9, 4251–4260, <https://doi.org/10.5194/acp-9-4251-2009>,  
585 2009, 2009.

Thompson, A. M., Stauffer, R. M., Wargan, K., Witte, J. C., Kollonige, D. E., and Ziemke, J. R.: Regional and Seasonal Trends in Tropical Ozone from SHADOZ Profiles: Reference for Models and Satellite Products, *J. Geophys. Res. Atmos.*, 126, <https://doi.org/10.1029/2021jd034691>, 2021.

590 Thompson, A. M., Witte, J. C., Sterling, C., Jordan, A., Johnson, B. J., Oltmans, S. J., Fujiwara, M., Vömel, H., Allaart, M.,  
Piters, A., Coetzee, G. J. R., Posny, F., Corrales, E., Diaz, J. A., Félix, C., Komala, N., Lai, N., Ahn Nguyen, H. T., Maata,  
M., Mani, F., Zainal, Z., Ogino, S., Paredes, F., Penha, T. L. B., da Silva, F. R., Sallons-Mitro, S., Selkirk, H. B., Schmidlin,  
F. J., Stübi, R., and Thiongo, K.: First Reprocessing of Southern Hemisphere Additional Ozonesondes (SHADOZ) Ozone  
Profiles (1998–2016): 2. Comparisons With Satellites and Ground-Based Instruments, *J. Geophys. Res. Atmos.*, 122,  
<https://doi.org/10.1002/2017jd027406>, 2017.

595 Tuck, A. F., Watson, R. T., Condon, E. P., Margitan, J. J., and Toon, O. B.: The planning and execution of ER-2 and DC-8  
aircraft flights over Antarctica, August and September 1987, *J. Geophys. Res. Atmos.*, 94(D9), 11181– 11222,  
[doi:10.1029/JD094iD09p11181](https://doi.org/10.1029/JD094iD09p11181), 1989.

Villamayor, J., Iglesias-Suarez, F., Cuevas, C.A. et al.: Very short-lived halogens amplify ozone depletion trends in the  
tropical lower stratosphere, *Nat. Clim. Chang.*, 13, 554–560, <https://doi.org/10.1038/s41558-023-01671-y>, 2023

600 Wargan, K., Orbe, C., Pawson, S., Ziemke, J. R., Oman, L. D., Olsen, M. A., Coy, L., and Emma Knowland, K.: Recent  
Decline in Extratropical Lower Stratospheric Ozone Attributed to Circulation Changes, *Geophys. Res. Lett.*, 45, 5166–5176,  
<https://doi.org/10.1029/2018gl077406>, 2018.

Weber, M., Coldewey-Egbers, M., Fioletov, V. E., Frith, S. M., Wild, J. D., Burrows, J. P., Long, C. S., and Loyola, D.:  
605 Total ozone trends from 1979 to 2016 derived from five merged observational datasets – the emergence into ozone recovery,  
*Atmos. Chem. Phys.*, 18, 2097–2117, <https://doi.org/10.5194/acp-18-2097-2018>, 2018.

Weber, M., Arosio, C., Coldewey-Egbers, M., Fioletov, V.E., Frith, S.M., Wild, J.D., Tourpali, K., Burrows, J.P. and  
Loyola, D.: Global total ozone recovery trends attributed to ozone-depleting substance (ODS) changes derived from five  
merged ozone datasets. *Atmospheric Chemistry and Physics*, 22(10), pp.6843-6859, 2022.

610 WMO (World Meteorological Organization). Scientific Assessment of Ozone Depletion: 2022, GAW Report No. 278, 509  
pp.; WMO: Geneva, 2022.

WMO (2018): Braesicke, P., Neu, J.L., Fioletov, V.E., Godin-Beekmann, S., Hubert, D., Petropavlovskikh, I., Shiotani, M.,  
Sinnhuber, B.M., Ball, W., Chang, K.L. and Damadeo, R., 2018. Update on Global Ozone: Past, Present, and Future,  
Chapter 3 in WMO Scientific Assessment of Ozone Depletion (2018).

615 Zou, L., Griessbach, S., Hoffmann, L., and Spang, R.: A global view on stratospheric ice clouds: assessment of processes  
related to their occurrence based on satellite observations, *Atmos. Chem. Phys.*, 22, 6677–6702, <https://doi.org/10.5194/acp-22-6677-2022>, 2022.

Zubov, V., Rozanov, E., Egorova, T., Karol, I., and Schmutz, W.: Role of external factors in the evolution of the ozone layer  
and stratospheric circulation in 21st century, *Atmos. Chem. Phys.*, 13, 4697–4706, <https://doi.org/10.5194/acp-13-4697-2013>, 2013.

620

---