- 1 Investigation of spatial and temporal variability in lower tropospheric ozone 2 from RAL Space UV-Vis satellite products Richard J. Pope^{1,2}, Brian J. Kerridge^{3,4}, Richard Siddans^{3,4}, Barry G. Latter^{3,4}, Martyn P. Chipperfield^{1,2}, Wuhu 3 Feng^{1,5}, Matilda A. Pimlott¹, Sandip S. Dhomse^{1,2}, Christian Retscher⁶ and Richard Rigby^{1,7} 4 5 1: School of Earth and Environment, University of Leeds, Leeds, United Kingdom 6 2: National Centre for Earth Observation, University of Leeds, Leeds, United Kingdom 7 3: Remote Sensing Group, STFC Rutherford Appleton Laboratory, Chilton, United Kingdom 8 4: National Centre for Earth Observation, STFC Rutherford Appleton Laboratory, Chilton, United Kingdom 9 5: National Centre for Atmospheric Science, University of Leeds, Leeds, United Kingdom 10 6: European Space Agency, ESRIN, Frascati, Italy 11 7: Centre for Environmental Modelling and Computation, University of Leeds, Leeds, United Kingdom 12 Revised version for Atmospheric Chemistry and Physics 13 Correspondence to: Richard J. Pope (r.j.pope@leeds.ac.uk) **Key Points** 14 15 The RAL Space profile retrieval algorithm for ultraviolet-visible nadir sounders has good vertical 16 sensitivity to retrieve lower tropospheric column ozone (LTCO₃). 17 OMI, SCIAMACHY and GOME-1 have suitably stable LTCO₃ records in comparison to ozonesondes 18 and are merged to form the first long-term satellite LTCO₃ record (1996-2017). 19 Comparison of 5-year averages for 1996-2000 and 2013-2017 suggests a significant LTCO₃ increase 20 21 (3.0 to 5.0 DU) in the tropics/sub-tropics over the satellite-era. 22 Abstract: 23 Ozone is a potent air pollutant in the lower troposphere and an important short-lived climate forcer (SLCF) in 24 the upper troposphere. Studies using satellite data to investigate spatiotemporal variability of troposphere 25 ozone (TO_3) have predominantly focussed on the tropospheric column metric. This is the first study to 26 investigate long-term spatiotemporal variability in lower tropospheric column ozone (LTCO₃, surface-450 hPa 27 sub-column) by merging multiple European Space Agency - Climate Change Initiative (ESA-CCI) products 28 produced by the Rutherford Appleton Laboratory (RAL) Space. We find that in the LTCO₃, the degrees of 29 freedom of signal (DOFS) from these products varies with latitude range and season and is up to 0.8,
- 30 indicating that the retrievals contain useful information on lower TO₃. The spatial and seasonal variation of
- 31 the RAL Space products are in good agreement with each other but there are systematic offsets of up to 3.0-
- 32 5.0 DU between them. Comparison with ozonesondes shows that the Global Ozone Monitoring Experiment
- 33 (GOME-1, 1996-2003), the SCanning Imaging Absorption spectroMeter for Atmospheric
- 34 CartograpHY (SCIAMACHY, 2003-2010) and the Ozone Monitoring Instrument (OMI, 2005-2017) have stable
- 35 LTCO₃ records over their respective periods, which can be merged together. However, GOME-2 (2008-2018)
- 36 shows substantial drift in its bias with respect to ozonesondes. We have therefore constructed a robust
- 37 merged dataset of LTCO₃ from GOME-1, SCIAMACHY and OMI between 1996 and 2017. Comparing the
- 38 LTCO₃ differences between the 1996-2000 and 2013-2017 5-year averages, we find sizeable positive
- increases (3.0-5.0 DU) in the tropics/sub-tropics, while in the northern mid-latitudes, we find small scale
- 40 differences in LTCO₃. Therefore, we conclude that there has been a substantial increase in tropical/sub-

41 tropical LTCO₃ during the satellite-era, which is consistent with tropospheric column ozone (TCO₃) records
42 from overlapping time-periods (e.g. 2005-2016).

43 **1. Introduction**

44 Tropospheric ozone (TO_3) is a short-lived climate forcer (SLCF) and, is the third most important greenhouse 45 gas (GHG; e. g. Myhre et al., 2013). TO₃ is also a hazardous air pollutant with adverse impacts on human 46 health (WHO, 2018) and the biosphere (e.g. agricultural and natural vegetation; Sitch et al., 2007). Since the 47 pre-industrial (PI) period, anthropogenic activities have increased the atmospheric loading of ozone (O_3) 48 precursor gases, most notably nitrogen oxides (NO_x) and methane (CH_4), resulting in a substantial increase in 49 TO₃ of 25-50% since 1900 (Gauss et al., 2006; Lamarque et al., 2010; Young et al., 2013). The PI to present day (PD) radiative forcing (RF) from TO₃ is estimated by the Intergovernmental Panel on Climate Change 50 51 (IPCC) to be 0.47 Wm⁻² (Forster et al., 2021) with an uncertainty range of 0.24-0.70 Wm⁻².

- 52 During the satellite-era, with a number of missions since 2000, extensive records of TO₃ have been
- produced, e.g. by the European Space Agency Climate Change Initiative (ESA-CCI; ESA, 2019). However, the large overburden of stratospheric O₃, coupled with the different vertical sensitivities and sources of error
- 55 associated with observations in different wavelength regions (e.g. Eskes and Boersma 2003; Ziemke et al.,
- 56 2011; Miles et al., 2015) contributes to large-scale spatiotemporal inconsistencies between the records
- 57 (Gaudel et al., 2018). Various studies (e.g. Heue et al. 2016; Pope et al., 2018; Ziemke et al. 2019) analysing
- TO₃ trends usually focussed on one or two instruments. The work by Gaudel et al. (2018) was part of the
- 59 Tropospheric Ozone Assessment Report (TOAR), which represented a large global effort to understand
- spatiotemporal patterns and variability in TO₃. Gaudel et al., (2018) analysed ozonesondes and multiple
- polar orbiting-nadir viewing satellite products and reported large-scale discrepancies in the spatial
 distribution, magnitude, direction and significance of the TCO₃ trends. While the satellite records did cover
- 63 slightly different time periods, they were unable to provide any definitive reasons for these discrepancies
- 64 beyond briefly suggesting that differences in measurement techniques and retrieval methods were likely to
- 65 be causing the observed spatial inconsistencies. Another factor introducing inconsistencies is the assumed
- tropopause height for the different products. Some products used the World Meteorological Organisation
- 67 (WMO) definition of "the first occurrence of the 2 K/km lapse-rate" while some others e.g. integrated the 0-
- 68 6 km and 6-12 km sub-columns to derive the tropospheric column. The use of different a priori products
- 69 within the retrieval scheme will have also provided inconsistencies.
- 70 The vertical sensitivity of each product (function of measurement technique and retrieval methodology)
- used by Gaudel et al. (2018) has a substantial impact on which part of the troposphere (and stratosphere)
- 72 the O₃ signal is weighted towards. The vertical sensitivity can be referred to as the "averaging kernel" (AK),
- 73 which provides the relationship between perturbations at different levels in the retrieved and true profiles
- 74 (Rodgers, 2000; Eskes and Boersma, 2003). As the instruments' vertical sensitivities differ, they are likely to
- 75 be influenced differently by processes controlling TO₃ temporal variability in different layers of the
- troposphere (e.g. lower troposphere influenced more by precursor emissions vs. the upper troposphere
- subject more to the influence from stratospheric-tropospheric exchange). Therefore, the differing vertical
- 78 sensitivities, and thus the TO₃ they are retrieving, could be driving the inconsistencies in reported TCO₃
- 79 trends between products.
- 80 While many studies have previously focussed on TCO₃ (e.g. Gaudel et al. (2018); Ziemke et al. (2019)),
- 81 several nadir-viewing ultraviolet-visible (UV-Vis) sounders can retrieve TO₃ between the surface to 450 hPa
- 82 (i.e. lower tropospheric column O₃, LTCO₃). The retrieval scheme from the Rutherford Appleton Laboratory
- 83 (RAL) Space exploits information from the O₃ Huggins bands (325-335 nm), as well as the Hartley band (270-

- 307nm), to retrieve high quality LTCO₃ and was selected for the ESA-CCI and EU Copernicus Climate Change 84
- 85 Service. As a result, the RAL Space LTCO₃ products (and equivalent from other providers) are valuable
- 86 resources to investigate global and regional O₃-related air quality (e.g. Richards et al., 2013; Pope et al.,
- 87 2018; Russo et al., 2023).

88 In this study, we explore the spatiotemporal variability of LTCO₃ from several UV-Vis sounders produced by 89 RAL Space. While Gaudel et al., (2018) used a range of UV-Vis and infrared (IR) TCO₃ products, including the 90 RAL Space Ozone Monitoring Instrument (OMI) product, we focus here on several RAL Space UV-Vis 91 products. Here, we aim to explore the consistencies between them, their vertical sensitivities, $LTCO_3$ stability 92 against ozonesonde records and suitability for long-term trend analysis. In our manuscript, Section 2 93 discusses the satellite/ozonesonde datasets used, Section 3 presents are results, while Section 4 summarises 94 our conclusions and discussion points.

95 2. Methodology and Datasets

96 2.1. Datasets

97 The four RAL Space UV-Vis satellite products investigated here are from OMI, the Global Ozone Monitoring 98 Experiment – 1 (GOME-1), GOME-2 and the SCanning Imaging Absorption spectroMeter for Atmospheric 99 CartograpHY (SCIAMACHY), all of which were developed as part of the ESA-CCI project (Table 1). GOME-1, 100 GOME-2, SCIAMACHY and OMI flew on ESA's ERS-2, MetOp-A, ENVISAT and NASA's Aura satellites in sun-101 synchronous low Earth polar orbits with local overpass times of 10.30, 9.30, 10.00 and 13.30, respectively. 102 They are all nadir viewing with spectral ranges which include the 270-350 nm range used for ozone profile 103 retrieval. The spatial footprints of the respective instruments at nadir are 320 km × 40 km, 80 km × 40 km, 104 240 km × 30 km and 24 km × 13 km (Boersma et al., 2011; Miles et al., 2015; Shah et al., 2018). The scheme 105 established by RAL Space to retrieve height-resolved O_3 profiles with tropospheric sensitivity (Miles et al., 106 2015) was applied to all of these satellite instruments. The scheme is based on the optimal estimation (OE) 107 approach of Rogers et al., (2000) and provides state-of-the-art retrieval sensitivity to lower TO₃, which is 108 described in detail by Miles et al., (2015) and by Keppens et al., (2018). The differences between the retrieval 109 versions (i.e. fv214 and fv300) in Table 1 are primarily linked to the instrument types where GOME-1, GOME-110 2 and SCIAMACHY are across-track scanning instruments while OMI uses a 2-D array detector. For this work, 111 the data were filtered for good quality retrievals whereby the geometric cloud fraction was <0.2, the lowest 112 sub-column O_3 value was > 0.0, the solar zenith angle < 80.0°, the convergence flag = 1.0 and the normalised 113 cost function was < 2.0. These filters also remove OMI pixels influenced by the OMI row anomaly (Torres et 114 al., 2018), so there is reduced OMI data coverage over the record. However, we find this has minimal impact 115 on our results with substantial proportions of data (e.g. millions of retrievals per year at the start and end of 116 the OMI record) available for analysis in our study.

117

2.2. Ozonesondes and Application of Satellite Averaging Kernels

118 To help understand the impact of the satellite AKs on retrieved LTCO₃ and stability of the satellite 119 instruments listed in Table 1 over time, we use ozonesonde data between 1995 and 2019 from the World 120 Ozone and Ultraviolet Radiation Data Centre (WOUDC), the Southern Hemisphere ADditional Ozonesondes 121 (SHADOZ) project and from the National Oceanic and Atmospheric Administration (NOAA). Keppens et al., 122 (2018) undertook a detailed assessment of the ESA-CCI TO₃ data sets, including the RAL UV-Vis profile data 123 sets used in this study (mostly older versions though) using ozonesondes. They found that the RAL LTCO₃ 124 products typically had a positive bias of about 40%, apart from OMI which was closer to 10%. On the global 125 scale, tropospheric drift in GOME-1 and OMI over time was approximately -5% and 10% per decade, 126 respectively. However, GOME-2 and SCIAMACHY had significant tropospheric drift trends of approximately 127 40% per decade. The recent Copernicus Product Quality Assessment Report (PQAR) Ozone Products Version

- 128 2.0b (Copernicus, 2021) undertook a more recent assessment of nadir ozone profiles using the level 3
- products from RAL listed in **Table 1.** They found that in the troposphere, OMI/GOME-1 and
- 130 SCIMACHY/GOME-2 had biases of -20% and 10%. GOME-1 tropospheric drift was deemed to be insignificant
- 131 (-10% to 5% per decade), while GOME-2 and SCIAMACHY had a significant drift of 30% and 20% per decade,
- respectively. OMI also had an insignificant tropospheric drift of 10% per decade.

133 In this study, for comparisons between ozonesonde profiles and satellite retrievals, each ozonesonde profile 134 was spatiotemporally co-located to the closest satellite retrieval. Here, all the retrievals within 6 hours of the 135 ozonesonde launch were subsampled and then the closest retrieval in space (i.e. within 500 km) was taken 136 for the final co-located one. Therefore, there was one satellite retrieval for every ozonesonde profile to help reduce the spatiotemporal sampling difference errors. Here, ozonesonde O₃ measurements were rejected if 137 138 the O₃ or pressure values were unphysical (i.e. < 0.0), if the O₃ partial pressure > 2000.0 mPa or the O₃ value 139 was set to 99.9, and whole ozonesonde profiles were rejected if at least 50% of the measurements did not 140 meet these criteria. These criteria are similar to those applied by Keppens et al., (2018) and Hubert et al., 141 (2016). To allow for direct like-for-like comparisons between the two quantities, accounting for the vertical 142 sensitivity of the satellite, the instrument AKs were applied the ozonesonde profiles. Here, each co-located 143 ozonesonde profile (in volume mixing ratio) was used to derive ozone sub-columns (in number density) on

144 the satellite pressure grid. The application of the AKs for the UV-Vis instruments was done using **Equation 1**:

153

 $sonde_{AK} = AK. (sonde_{int} - apr) + apr$ (1)

where *sonde_{AK}* is the modified ozonesonde sub-column profile (Dobson units, DU), *AK* is the averaging kernel matrix, *sonde_{int}* is the sonde sub-column profile (DU) on the satellite pressure grid and *apr* is the apriori subcolumn amount (DU). Here, the ozonesonde profile, on its original pressure grid (typically in units of ppbv or mPa) are converted into ozone sub-columns between each pair of measurement levels. These sub-columns are then aggregated up to the larger sub-columns (e.g. the LTCO₃ range is between the surface and 450 hPa) on the coarser satellite pressure grid.

152 **3. Results**

3.1. Satellite Vertical Sensitivity

154 Figure 1 represents average AKs for all the instruments listed in Table 1 for 2008 (1998 for GOME-1) in the 155 northern (NH) and southern (SH) hemispheres between the equator and 60°S & N. Of the four RAL Space 156 products, OMI O₃ profiles appear to contain the most information with degrees of freedom of signal (DOFS) 157 of 5.0 or above for the full atmosphere. Here, the DOFS represents the number of independent pieces of 158 information on the vertical profile in the retrieval (i.e. the sum of the AK diagonal). SCIAMACHY has the 159 lowest sensitivity with average DOFS ranging between 4.12 and 4.64. The DOFS tends to be larger in NH for 160 all the products, though there is no clear pattern in the seasonality (i.e. January vs. July). In terms of $LTCO_3$, 161 OMI again has greater sensitivity than the others with average hemispheric and seasonal DOFS ranging 162 between 0.63 and 0.68. For GOME-1 (GOME-2), the LTCO₃ DOFS range between 0.37 and 0.50 (0.39 and 163 0.46). SCIAMACHY LTCO₃ DOFS range between 0.44 and 0.52. Therefore, while SCIAMACHY has the lowest 164 overall information on the full atmospheric ozone, it has reasonably good information in the LTCO₃, as do 165 the other instruments. These results are robust given the large number of retrievals (N) that have been used to derive the average AKs (i.e. N > 65,000 in all cases). 166

167 While Figure 1 provides spatial average information on LTCO₃ DOFS, Figure 2 shows spatial maps for

- 168 December-January-February (DJF) and June-July-August (JJA) over the respective instrument records. The
- 169 largest LTCO₃ DOFs occur over the ocean ranging between approximately 0.4 and 0.6 for GOME-1, GOME-2
- and SCIAMACHY, while OMI has larger ocean values between 0.7 and 0.8. Over land, the LTCO₃ DOFS tend to

be lower and between 0.3 and 0.5 for GOME-1, GOME-2 and SCIAMACHY. Again, OMI has larger values on
land of between 0.4 and 0.7. Depending on the hemispheric season, the summer-time (JJA in NH and DJF in
SH) LTCO₃ DOFS are larger for each instrument. Overall, OMI (GOME-2) retrievals contain the largest (lowest)
amount of information on LTCO₃.

175 The impact of the satellite vertical sensitivity is further investigated by co-locating the products with the 176 merged ozonesonde data set, over their respective mission periods (globally and in the NH and SH) and the 177 AKs applied to assess the impact on the ozonesondes (Figure 3). For all the instruments, there are suitable 178 samples sizes (N > 1000 in all cases) of co-located retrievals and derived ozonesonde LTCO₃. In the case of GOME-1, the global distribution has a 25th-75th percentile (25 75%) range of approximately 8.0 to 20.0 DU 179 180 and a median of 14.0 DU. The apriori 25_75% range and median values are 16.0 to 22.0 and 19.0 DU. These 181 substantial differences between retrieved and apriori values confirm there is sensitivity in the GOME-1 182 retrieval to lower tropospheric ozone. It can be seen from Equation 1 that if a satellite instrument had 183 perfect sensitivity at all levels (i.e. AK=1), there would be no change in co-located ozonesonde LTCO₃ 184 distribution when the AKs are applied. However, given AK values are less than 1.0 in Figure 1, leading to the 185 DOFS of approximately 0.5, there is a shift in the median value towards the apriori from approximately 21.0 to 19.0 DU. The corresponding ozonesonde 10th-90th percentile (10 90%) range of 13.0 to 26.0 DU expanded 186 187 to 12.0 to 27.0 DU. Therefore, the application of the AKs to the ozonesondes actually increases the range of 188 observed values. In the NH, the GOME-1 median (25 75% range) is 14.0 (4.0-24) DU while the apriori median 189 (25_75% range) is 21.0 (18.0-23.0) DU. The ozonesonde median (25_75% range) is 22.0 (19.0-25.0) DU while 190 application of the AKs yields values of 19.0 (16.0-24.0) DU. In the SH, the GOME-1 median (25 75% range) is 191 12.0 (8.0-17.0) DU while the apriori median (25 75% range) is 14.0 (12.0-16.0) DU. The ozonesonde median 192 (25_75% range) is 12.0 (11.0-17.0) DU while application of the AKs yields values of 12.0 (6.0->40.0) DU. In 193 comparison, GOME-2 shows a similar response though the shift in LTCO₃ value between the apriori and 194 satellite is smaller. This makes sense given the lower vertical sensitivity of GOME-2. In the SH, the application 195 of the AKs to the ozonesondes yields a very large range in the percentiles. It is likely that the South Atlantic 196 Anomaly (SAA – i.e. where charged particles directly impact UV detectors increasing dark-current noise, 197 which in turn reduces the number of retrievals from all UV sensors, notably both GOME-1 and GOME-2; 198 Keppens et al., 2018), given the typically larger values and signal corruption, is driving the large response in 199 the ozonesonde+AKs range.

200 For OMI, the global distribution has a median (25_75% range) of 17.0 (13.0-25.0) DU yielding a substantial 201 shift from the apriori median (25_75% range) of 18.0 (16.0-22.0) DU. In the NH, the satellite median (25_75% 202 range) is 18.0 (13.0-25.0) DU and the apriori median (25_75% range) value is 20.0 (17.0-23.0) DU. In the SH, 203 the satellite median (25 75% range) is 14.0 (10.0-22.0) DU and the apriori median (25 75% range) value of 204 15.0 (13.0-19.0) DU. When the AKs are applied to the ozonesondes there is typically an increase in the 205 median LTCO₃ and range by approximately 3.0-4.0 DU. This increase in LTCO₃ when the OMI AKs are applied 206 to the ozonesondes contrasts with the other satellite instruments. While the vertical smearing from the 207 stratosphere would intuitively be expected to increase the tropospheric layer retrieval, and thus the AK 208 adjustment to decrease the ozonesonde value, in the case of OMI there is a negative excursion in the AKs 209 into the lowermost stratosphere (see Figure 1), so the opposite occurs. For SCIAMACHY, a similar 210 relationship occurs to that of GOME-1 and GOME-2 with a shift of the satellite LTCO₃ median away from the 211 apriori by 1.0-3.0 DU and an increase the in 25_75% range by 10.0-15.0 DU. Apart from the SH, the 212 application of the AKs to the ozonesondes shifts the LTCO₃ median by 2.0-3.0 DU but the 25_75% range 213 remains similar. Overall, there is shift in the satellite LTCO₃ median value away from the apriori with an 214 increase in the 25_75% and 10_90% ranges. A similar pattern occurs in multiple cases between the

ozonesondes and the ozonesondes+AKs. Therefore, all the instruments have reasonable vertical sensitivity in
 LTCO₃ with substantial perturbations from the apriori and to the satellite LTCO₃ distribution.

217 **3.2.** Lower Tropospheric Column Ozone Seasonality

Multiple studies have investigated the seasonality of TO₃ from space observing large biomass burning and lightning induced O₃ in the South Atlantic (Ziemke et al., 2006; Ziemke et al., 2011; Pope et al., 2020), enhanced summertime TO₃ over the Mediterranean (Richards et al., 2013), TO₃ over large precursor regions such as China and India (Verstraeten et al., 2015) and the enriched northern hemispheric background O₃ during springtime (Ziemke et al., 2006). Here, we compare the long-term seasonal (DJF and JJA) spatial distributions of RAL Space LTCO₃ products (**Figure 4**).

- OMI and GOME-2 LTCO₃ have regions of consistency (e.g. JJA NH enhanced background TO₃, between 20.0
- DU and 30.0 DU, and the Mediterranean TO₃ peak, >25.0 DU), but the SAA interferes with the signal of the
- 226 biomass burning induced secondary O₃ formation from Africa and South America. However, for OMI, this
- ozone plume ranges between 23.0 and 27.0 DU (18.0 and 20.0 DU) in DJF (JJA). There are also clear LTCO $_3$
- hotspots over anthropogenic regions (e.g. eastern China and northern India) peaking at over 25.0 DU in JJA.
- The GOME-1 LTCO₃ spatial patterns are consistent with that of OMI and GOME-2, but there is a systematic low bias relative to OMI and GOME-2 in the absolute LTCO₃ of 3.0 DU to 7.0 DU, depending on geographical
- low bias relative to OMI and GOME-2 in the absolute LTCO₃ of 3.0 DU to 7.0 DU, depending on geographica
 location (e.g. 20.0-22.0 DU over northern India for GOME-2 and OMI, while 16-18 DU for GOME-1). These
- differences in the GOME-1 and GOME-2/OMI LTCO₃ seasonal averages are likely to be at least partly due to
- underlying LTCO₃ tendencies between the respective instrument time periods. This is investigated further in
- 234 Section 3.4. The SCIAMACHY spatial pattern and absolute LTCO₃ values are more consistent with OMI and
- 235 GOME-2. Moreover, SCIAMACHY shows limited sensitivity to the SAA and resolves the biomass burning /
- lightning O₃ sources detected by OMI over South America, South Atlantic and Africa (18.0-20.0 DU in JJA).
- However, especially in the NH in DJF, there appears to be regions of latitudinal banding in the LTCO₃ spatial
- patterns (e.g. 0°-30°N), which are not observed (or to the same extent) as the other UV-Vis sounders.
- Overall, GOME-2 and OMI are in good agreement spatially and seasonally with similar absolute LTCO₃ values.
 In DJF and JJA, OMI appears to be 2.0-3.0 DU lower and larger than GOME-2, respectively. This is reasonable
- 241 given the similar temporal records they cover (2005-2017 vs. 2007-2018). SCIAMACHY has similar spatial-
- seasonal patterns but has systematically larger (3.0-5.0 DU) DJF values in comparisons to OMI and GOME-2.

The satellite LTCO₃ seasonality is consistent with that of the ozonesondes. Here, the median (25th percentile, 75th percentile) ozonesonde LTCO₃ values for the NH in DJF, NH in JJA, SH in DJF and SH in JJA are 18.0 (15.7, 20.0) DU, 20.8 (16.7, 24.6) DU, 10.8 (8.2, 14.8) DU and 14.4 (12.1, 16.3) DU, respectively. Therefore, the NH LTCO₃ values are larger than those in the SH and the JJA LTCO₃ values are larger than the DJF equivalent. All of which are consistent with the four instrument LTCO₃ seasonal distributions.

248

3.3. Satellite Instrument Temporal Stability

249 For accurate assessment of satellite LCTO₃ temporal variability, there needs to be insignificant drift over 250 time, whereas bias which is constant over time can be tolerated. The most appropriate data set with which 251 to assess satellite long-term drifts is that of the ozonesonde record, albeit that it has certain limitations 252 potentially including temporal changes in accuracy (Stauffer et al., 2020) as well as geographical coverage. 253 Figure 5 shows annual time series of the satellite-ozonesonde (with AKs applied) median biases for three 254 latitude bands: 90°-30°S, 30°S-30°N and 30-90°N. The hatched pixels show where the biases are non-255 substantial, defined as the 25_75% difference range intersecting with zero. For GOME-1, the mean bias (MB) 256 is -5.34, -3.21 and -0.90 DU for the three regions, respectively. For the 30-90°N region, several years show 257 substantial biases of -6.0 to -3.0 DU. The two other latitude bands have few substantial years but in the

- tropical band, both 2002 and 2003 show substantial biases of approximately -5.0 DU. To assess the stability
- of the instruments with time, a simple linear least-squares fit was performed with regional trends of -0.32, -
- 260 0.98* and -0.03 DU/yr. A substantial trend (shown by an asterisk) has a p-value < 0.05 as defined as
- 261 $|M/\sigma_M| > 2.0$ (e.g. Pope et al., 2018), where M and σ_M are the linear trend and trend uncertainty,
- respectively. While, the 30-90°N region had a sizable systematic bias, it was stable with time, as was the bias
- for the 90°-30°S region. However, the 2002 and 2003 biases in the 30S°-30°N region gave rise to a substantial
 drift in the GOME-1 record.
- 265 For GOME-2, the record MB is 1.91, -5.05 and 1.64 DU for the respective latitude bands, all of which have 266 substantial bias trends at 0.62*, -0.70* and 0.22* DU/yr. Therefore, the GOME-2 LTCO₃ records from this processing run are not stable and cannot be used further in the study. SCIAMACHY has regional mean biases 267 268 of 1.33, 4.47 and 2.81 DU. In the 30-90°N region, the bias is non-substantial. While there are substantial 269 biases peaking at 3.0-5.0 DU in the 90°-30°S region, neither region has a critical drift trend. The largest 270 substantial biases are in the 30S°-30°N region (>5.0 DU) for 2006 to 2008. While the positive trend of 0.21 271 DU/yr is insignificant, we do not use the SCIAMACHY data in later years when harmonising the LTCO₃ records 272 (section 3.4). OMI has MBs of -5.16, -2.91 and -0.41 DU with only a few of the year-latitude pixels having 273 substantial biases peaking at -6.0 to -3.0 DU in the 30-90°N region. The resulting bias trends are -0.12, 0.22 274 and -0.10 DU/yr, which all have p-values > 0.05. Therefore, GOME-1, OMI and SCIAMACHY were deemed 275 suitable LTCO₃ records for use in this study.

3.4. Lower Tropospheric Column Ozone Merged Record

- 277 The RAL Space products cover the full period between 1996 and 2017. Therefore, there is the opportunity to 278 merge and harmonise these records to produce a long-term record to look at the spatiotemporal variability 279 of LTCO₃. From Figure 5, the OMI record appears to be stable with time globally, providing a suitable data set 280 between 2005 and 2017. The GOME-2 record appears not to be sufficiently stable across its record (2008-281 2018), so is not included in subsequent analysis. The GOME-1 record covers 1996 to 2010, but given the loss 282 of geographical coverage due to the onboard tape recorder failing in June 2003 (van Roozendael, 2012), a 283 true global average is only available between 1996 and 2003. Figure 5 shows that GOME-1 bias with respect 284 to the ozonesonde record is not stable in the tropics but this is predominantly driven by instrument-285 ozonesonde differences in 2003. Therefore, 2003 is also dropped leaving the GOME-1 global record between 286 1996 and 2002. The GOME-1 tropical bias for 2002 is similar to that of 2003 (-5.0 DU) but the biases for the 287 other latitude bands are less distinct. The regional average LTCO₃ values for 2002 in Figure 5 are also 288 comparable to neighbouring years (e.g. 2000 and 2001). SCIAMACHY also does not have a full year of data 289 for 2002, so we have included the GOME-1 2002 data in our analysis.
- 290 While OMI (2005-2017) and GOME-1 (1996-2002) now cover a large proportion of the global record, there is 291 still a systematic difference between them. Different UV-Vis instruments can have inconsistencies in their 292 retrieved products (e.g. van der A et al., (2006), Heue et al., (2016)) and often require a systematic 293 adjustment to create a harmonised record. Here, there is overlap in the raw records between 2005 and 2010 294 for GOME-1 and OMI. The GOME-1 record does have large missing data gaps globally, but for the mid-295 latitude and tropical latitude bands, there is sufficient sampling to inter-compare the two records. Therefore, 296 for each swath, the nearest OMI retrieval is co-located to that of GOME-1, but has to be within 250 km. The 297 local overpass times are different (i.e. GOME-1 10.30 and OMI 13.30) but within approximately 3-hours, so 298 the diurnal cycle impacts are likely to be of a secondary order and we are confident in merging the records. 299 Based on the co-located OMI and GOME-1 data, we derived long-term latitude-month offset which are 300 added to GOME-1 (1996-2002) to harmonise the records. The was done using latitudinal bins of 60°S-30°S, 301 30°S-30°N and 30°N-60°N. Given the lack of GOME-1 data outside of 60°S-60°N due to the failure of the

302 GOME-1 tape recorder in June 2003, there was insufficient data to derive offsets, the high-latitudes data is 303 excluded in the following sections. Where there was good spatial coverage from GOME-1 between 2005 and 304 2010, once the offset had been applied, gridded OMI and GOME-1 where data existed for both, on a pixel by 305 pixel basis, were averaged together.

306 For 2003 and 2004, we use the SCIAMACHY spatial fields to gap fill the record. Figure 5 shows that 307 SCIAMACHY had some substantially large biases compared to the ozonesondes in 2006, 2007 and 2008 but 308 was reasonable for other years. Therefore, we use the global distributions from SCIAMCHY for both years 309 but scale them to expected values between 2002 and 2005. This is achieved by getting the globally weighted 310 (based on surface area) LTCO₃ average for GOME-1 (2002 with GOME-1 with the OMI offset applied) and 311 OMI (2005) and the SCIAMACHY (2003-2004). Based on the difference between 2002 and 2005, an annual 312 linear global scaling is applied in 2003 and 2004 for the SCIAMACHY spatial fields. Thus, we have developed a 313 harmonised LCTO₃ record between 1996 and 2017. Examples of the harmonised data for Europe and East 314 Asia are shown in Figure 6. Overall, there is non-linear variability in the two regional time-series where red 315 and blue show the GOME-1 and OMI LTCO₃ time series and then black shows where they have been merged 316 (including SCIAMACHY for 2003 and 2004). For Europe (East Asia), the seasonal cycle ranges between 10.0 317 (13.0) and 30.0 (27.0) DU, respectively, with annual average values between 18.0 (18.0) and 22.0 (21.0) DU.

318 **3.5. Lower Tropospheric Column Ozone Temporal Variability**

319 The harmonised RAL Space data set can now be used to investigate decadal scale spatiotemporal variability 320 in LTCO₃. Figure 7 shows the global long-term (1996-2017) average in LTCO₃ and the 5-year average 321 anomalies for 1996-2000, 2005-2009 and 2013-2017. In the long-term average (Figure 7a), there is clear SH 322 to NH LTCO₃ gradient with background values of 13.0-17.0 DU and 20-23.0 DU, respectively. There are 323 hotspots over East Asia, the Middle East/Mediterranean and northern India of 24.0-25.0 DU. The largest SH 324 LTCO₃ values (20.0-22.0 DU) are between 30-15°S spanning southern Africa, the Indian Ocean and Australia. 325 Minimum LTCO₃ values (<12.0 DU) are over the Himalayas (due to topography) and the tropical oceans. As 326 shown in Figure 2, there is sufficient information (e.g. LTCO₃ DOFS mostly > 0.5) in the tropics and mid-327 latitudes for the instruments used to form the merged LTCO₃ data. This provides confidence in this merged 328 LTCO₃ record for long-term temporal analysis. Note, the SAA has been masked out in all the panels. The 329 1996-2000 anomaly map (Figure 7b) shows values to be similar (i.e. -1.0 to 1.0 DU) with respect to the 1996-330 2017 mean between 30°N and 60°N. A similar relationship occurs at approximately 30°S. However, in tropics 331 and NH sub-tropics (15°S to 30°N), the anomalies are more negative, ranging between approximately -3.0 332 and -1.0 DU. The green polygon-outlined regions show where the 1996-2000 LTCO₃ average represents a 333 substantial difference (p-value < 0.05) from the long-term average. This is based on the Wilcoxon rank test 334 (WRT), which is the nonparametric counterpart of the Student t-test that relaxes the constraint on normality 335 of the underlying distributions (Pirovanoet al., 2012). As well as this tropical band, the 60-45°S band shows 336 anomalies of a similar magnitude. In the 2005-2009 anomaly map (Figure 7c), there are widespread, though 337 non-substantial, anomalies of -1.5.0 to 0.0 DU. There are small clusters of substantial anomalies (e.g. 338 southern Africa at -2.0 to -1.0 DU and over the Bering Sea between 1.0 and 2.0 DU) but with limited spatial 339 coherence. In the 2013-2017 anomaly map (Figure 7d), they remain small LTCO₃ anomalies in the northern 340 mid-latitudes (-1.0 to 1.0 DU). A similar pattern occurs in the southern sub-tropics and mid-latitudes, though 341 the anomalies are larger peaking at 1.5 DU around 60-45°S (some have p-values < 0.05). However, in the 342 tropics and sub-tropics (15°S-30°N), there are positive anomalies of 1.0 to 2.0 DU throughout the region, 343 peaking at 2.0-2.5 DU over Africa.

Overall, these anomalies suggest there has been limited change in LTCO₃, between 1996 and 2017, in the NH
 mid-latitudes (e.g. as can be seen for Europe and East Asia in Figure 6). Unfortunately, the SAA masks any

346 useful information on LTCO₃ over South America, but generally there has been a moderate LTCO₃ increase in 347 the SH mid-latitudes. The largest and most substantial changes have been in the tropics and sub-tropics (i.e. 348 15°S to 30°N) switching from sizeable negative anomalies (-2.0 to -1.0 DU) in the 1996-2000 LTCO₃ average 349 to positive anomalies (1.0-2.0 DU) in the 2013-2017 LTCO₃. Figure 8 shows the difference between the 2013-350 2017 and 1996-2000 averages. Over the tropics/sub-tropics (15°S-30°N), the largest increases (p-values < 351 0.05) of 3.0 to 5.0 DU occur peaking Africa, India and South-East Asian (>5.0 DU). Thus, showing a large-scale increase in tropical LTCO₃ between 1996 and 2017. In the NH mid-latitudes, the absolute LTCO₃ differences 352 353 are relatively small (-1.0 to -1.5 DU) but there are consistent, though some negative differences (generally -354 2.0 and -1.0 DU) are over North America and Russia. In the SH mid-latitudes, there has been moderate 355 increase in LTCO₃ of 2.0-3.5 DU. However, southern Africa shows more localised decreases of up to 3.0 DU 356 and non-substantial differences at 30°S across the Indian Ocean. The ozonesondes are consistent with 357 satellite 1996-2000 and 2013-2017 average LTCO₃ differences. In the tropics, the majority of ozonesonde 358 sites show increases between these two periods ranging between 0.5 and 5.0 DU. Over Europe (i.e. northern 359 mid-latitudes), the ozonesonde LTCO₃ differences range between -0.5 and 0.5 DU suggesting limited LTCO₃ 360 change over time.

361 3.6. Long-term LTCO₃ Trends

362 In line with TOAR-II, we have added additional metrics on the temporal change in LTCO₃ over the merged 363 instrument record. Here, we have calculated the linear trends in LTCO₃ in 15° latitude bins between 60°S-364 60°N along with the 95% confident range and associated p-values (see Table 2). In the tropical latitudes 365 (15°S-30°N), all the linear trends show substantial increasing trends (2.89-4.12 DU/decade) between 1996 366 and 2017; all with p-values tending to 0.0. This is consistent with the LTCO₃ positive differences (3.0-5.0 DU) 367 between the 1996-2000 and 2013-2017 averages (Figure 8). In the northern mid-latitudes (30-60°N), there 368 are smaller positive trends (1.33 and 0.49 DU/decade) but the 95% confidence values intersect with 0.0 and 369 have larger p-values. Again, this is consistent with the near-zero differences between the 1996-2000 and 370 2013-2017 averages (Figure 8). In the southern mid-latitudes (30-60°S), the trends are substantially positive 371 (1.85 and 4.49 DU/decade) with near-zero p-values. Again, this is consistent with the substantial differences 372 (2.0-4.0 DU) between the 1996-2000 and 2013-2017 averages. The 15-30°S trend is small at 0.94 DU/decade 373 with a moderate p-value of 0.35, indicating this not to be a substantial trend.

4. Discussion and Conclusions

374

375 Multiple studies have used satellite records to investigate change in TCO₃ in recent decades. Gaudel et al., 376 (2018) used a range of UV-Vis and IR TCO₃ products between 2005 and 2016. The UV-Vis sounders generally 377 show substantial positive trends (0.1-0.8 DU/yr) in the tropics/sub-tropics and a mixed response in the mid-378 latitudes. The IR instruments typically showed significant decreasing trends (-0.5 to -0.2 DU/yr) in 379 background regions and isolated regions of substantial TCO₃ enhancements. Ziemke et al., (2019) used a 380 long-term merged record of TCO₃ from the Total Ozone Mapping Spectrometer (TOMS) and Ozone 381 Monitoring Instrument/Microwave Limb Sounder (OMI-MLS) between 1979 and 2016. Over this period, they 382 found significant increases of TCO_3 of 1.5 to 6.5 DU, especially over India and East Asia. Heue et al., (2016) 383 used a long-term tropical TCO₃ record (GOME, SCIAMACHY, OMI, GOME-2A and GOME-2B) finding 384 significant increases (0.5-2.0 DU/decade) over central Africa and the South Atlantic. However, the study by 385 Wespes et al., (2018) from IASI (an IR sounder) indicated that TCO₃ decreased between 2008 and 2017 by -386 0.5 to -0.1 DU/yr. Gaudel et al., (2018) reported similar TCO₃ tendencies using two IASI products (IASI-FORLI 387 and IASI-SOFRID). However, Boynard et al., (2018) and Wespes et al., (2018) report a step-change in 2010 in 388 the IASI-FORLI O₃ data which could influence observed long-term trends. Therefore, studies using IR 389 products available to TOAR-I and Wespes (2018) are no longer considered reliable.

- In this study, for the first time we analysed long-term changes in LTCO₃ using a merged satellite UV-Vis
 sounder record. Overall, we found that LTCO₃ was lower (by 1.0-3.0) in the tropics between 1996 and 2000
 in comparison to the long term average (i.e. 1006, 2017). Similar LTCO, values exist between the 2005, 2000.
- in comparison to the long-term average (i.e. 1996-2017). Similar LTCO₃ values exist between the 2005-2009
- and long-term averages, while the 2013-2017 average shows substantially larger tropical values (1.0-2.5 DU)
 than the long-term average. Therefore, this tropical increase (3.0-5.0 DU) in LTCO₃ between 1996 and 2017 is
- consistent with other reported increases in TCO₃. A similar consistency is found in the NH mid-latitudes, with
- minimal changes in LTCO₃ observed here and in trends in TCO₃ reported in Gaudel et al., (2018) and Ziemke
- et al., (2019). Sizable LTCO₃ increases in the SH mid-latitudes are also consistent with Gaudel et al., (2018)
 and Ziemke et al., (2019), though they differ from IASI retrieved TCO₃ trends as reported by Wespes et al
 (2018). Overall, the long-term changes in LTCO₃ reported here and the literature TCO₃ trends from satellite
- 400 UV products are comparable in regard to latitude dependence and direction. It therefore seems that the
 401 positive tendencies in TCO₃ reported in the literature from UV soundings over the satellite-era are associated
 402 with, and could be driven by, changes occurring in LTCO₃.
- 403 For future work, a detailed study is required to disentangle the reported TCO₃ and LTCO₃ trends reported by
- 404 UV-Vis and IR sounders, which would benefit from satellite level-2 data produced from level-1 data sets
 405 which are more uniform over time along with other improvements. This can potentially be done also by
- which are more uniform over time along with other improvements. This can potentially be done also by
 using a 3D atmospheric chemistry model (ACM) to investigate the changes in lower and upper tropospheric
- 407 ozone, and application of the satellite AKs (i.e. the vertical sensitivity of the different satellite products) to
- the model from the different sounders to establish how satellite vertical sensitivity potentially changes the
- simulated TO₃ tendency of the model. An ACM would also be a useful tool to help diagnose the importance
- 410 of LTCO₃ contributions to the TCO₃ tendencies, and which processes might be driving any spatiotemporal
- 411 changes (e.g. surface emissions, atmospheric chemistry/surface deposition, stratospheric-tropospheric O₃
- 412 exchanges etc.). Finally, together with improved, extended reprocessed versions of the data sets used in this
- 413 study, the launch of the Sentinel 5 Precursor (S5P) satellite (in October 2017) can be used to extend the
- 414 merged data record of LTCO₃, along with new polar orbiting platforms such as Sentinel-5 and IASI-NG
 415 instruments on future EUMETSAT MetOp-Second Generation satellites.

416 Acknowledgements:

- 417 This work was funded by the UK Natural Environment Research Council (NERC) by providing funding for the
- 418 National Centre for Earth Observation (NCEO, award reference NE/R016518/1) and funding from the
- 419 European Space Agency (ESA) Climate Change Initiative (CCI) post-doctoral fellowship scheme (contract
- 420 number 4000137140). Anna Maria Trofaier (ESA Climate Office) provided support and advice throughout the
- 421 fellowship.

422 Conflicting Interests

423 The authors declare that they have no conflicts of interest.

424 Data Availability

- 425 The RAL Space satellite data is available via the NERC Centre for Environmental Data Analysis (CEDA) Jasmin
- 426 platform subject to data requests. The RAL Space satellite data will be uploaded to the Zenodo open access
- 427 portal (https://zenodo.org/) if this manuscript is accepted for publication in ACP after the peer-review
- 428 process. The ozonesonde data for WOUDC, SHADOZ and NOAA is available from <u>https://woudc.org/</u>,
- 429 <u>https://tropo.gsfc.nasa.gov/shadoz/</u>and <u>https://gml.noaa.gov/ozwv/ozsondes/</u>.

430 Author Contributions

- 431 RJP, MPC and BJK conceptualised and planned the research study. RJP and MAP analysed the satellite data
- 432 provided by RAL Space (BJK, RS, BGL) with support from BJK, RS and BGL. MPC, SD and CR provided scientific

advice, while WF and RR provided technical support. RJP prepared the manuscript with contributions fromall co-authors.

435 **References:**

- Boersma KF, et al. 2011. An improved tropospheric NO₂ column retrieval algorithm for the Ozone Monitoring
 Instrument. Atmospheric Measurement Techniques, 4, 1905–1928, doi: 10.5194/amt-4-1905-2011.
- Copernicus. 2021. Product Quality Assessment Report (PQAR) Ozone products Version 2.0b, Issued by
 BIRA-IASB/Jean-Christopher Lambert, Ref: C3S_D312b_Lot2.2.1.2_202105_PQAR_O3_v2.0b.
- 440 ESA. 2019. Climate Change Initiative. http://cci.esa.int/ozone (last accessed 02/05/2023).
- Eskes HJ and Boersma KF. 2003. Averaging kernels for DOAS total column satellite retrievals. *Atmospheric Chemistry and Physics*, **3**, 1285–1291, doi: 10.5194/acp-3-1285-2003.
- Forster P, et al. 2021. Climate Change 2021: The Physical Science Basis. Contribution of Working Group I to
 the Sixth Assessment Report of the Intergovernmental Panel on Climate
 Change, doi:10.1017/9781009157896.009.
- Gaudel A, et al. 2018. Tropospheric Ozone Assessment Report: Present day distribution and trends of
 tropospheric ozone relevant to climate and global atmospheric chemistry model evaluation. *Elementa*, 6
 (39), 1-58, doi: 10.1525/elementa.291.
- 449 Gauss M, et al. 2006. Radiative forcing since preindustrial times due to ozone change in the troposphere and
- 450 the lower stratosphere. *Atmospheric Chemistry and Physics*, **6**, 575-599, doi: 10.5194/acp-6-575-2006.
- Heue KP, et al. 2016. Trends of tropical tropospheric ozone from 20 years of European satellite
 measurements and perspectives for the Sentinel-5 Precursor. *Atmospheric Measurement Techniques*, 9,
 5037-5051, doi: 10.5194/amt-9-5037-2016.
- Hubert D, et al. 2016. Ground-based assessment of the bias and long-term stability of 14 limb and
 occultation ozone profile data records. *Atmospheric Measurement Techniques*, 9, 2497-2534, doi:
 10.5194/amt-9-2497-2016.
- Keppens A, et al. 2018. Quality assessment of the Ozone_cci Climate Research Data Package (release 2017) –
 Part 2: Ground-based validation of nadir ozone profile data products. *Atmospheric Measurement Techniques*, 11, 3769-3800, doi: 10.5194/amt-11-3769-2018.
- Lamarque JF, et al. 2010. Historical (1850-2000) gridded anthropogenic and biomass burning emissions of
 reactive gases and aerosols: methodology and application. *Atmospheric Chemistry and Physics*, 10, 70177039, doi: 10.5194/acp-10-7017-2010.
- 463 Miles GM, et al. 2015. Tropospheric ozone and ozone profile retrieved from GOME-2 and their validation.
 464 *Atmospheric Measurement Techniques*, **8**, 385-398, doi: 10.5194/amt-8-385-2015.
- Myhre G, et al. 2013. Anthropogenic and Natural Radiative Forcing, in: Climate Change 2013: The Physical
 Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental
 Panel on Climate Change Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA,
- 468 659–740.
 469 Pope RJ, et al. 2018. Widespread changes in UK air quality observed from space. *Atmospheric Science Letters*,
- 470 **19:e817**, doi: 10.1002/asl.817.
- 471 Pope RJ, et al. 2020. Substantial Increases in Eastern Amazon and Cerrado Biomass Burning-Sourced
- Tropospheric Ozone. *Geophysical Research Letters*, **47 (3)**, e2019GL084143, doi: 10.1029/2019GL084143.
- 473 Pirovano G, et al. 2012. Investigating impacts of chemistry and transport model formulation on model
- 474 performance at European scale. *Atmospheric Environment*, **59**, 93-109, doi:
- 475 10.1016/j.atmosenv.2011.12.052.

- 476 Richards NAD, et al. 2013. The Mediterranean summertime ozone maximum: global emission sensitivities
 477 and radiative impacts. *Atmospheric Chemistry and Physics*, **13**, 2231-2345, doi: 10.5194/acp-13-2331-2013.
- 478 Rodgers, C.D. 2000. Inverse methods for atmospheric sounding: Theory and practice. New Jersey, USA:479 World Science.
- 480 Russo, M.R., Kerridge, B.J., Abraham, N.L., et al. 2003. Seasonal, interannual and decadal variability of
- tropospheric ozone in the North Atlantic: comparison of UM-UKCA and remote sensing observations for
 2005-2018. *Atmospheric Chemistry and Physics*, 23 (11), 6169-6196, doi: 10.5194/acp-23-6169-2023.
- 483 Shah S, et al. 2018. Evaluation of SCIAMACHY Level-1 data versions using nadir ozone profile retrievals in the 484 period 2003-2011, *Atmospheric Measurement Techniques*, **11**, 2345-2360, doi: 10.5194/amt-11-2345-2018.
- 485 Sitch S, et al. 2007. Indirect radiative forcing of climate change through ozone effects on the land-carbon 486 sink. *Nature*, **448**, 791-794, doi: 10.1038/nature06059.
- Stauffer RM, et al. 2020. A post-2013 Dropoff in Total Ozone at a Third of Global Ozonesonde Stations:
 Electrochemical Concentration Cell Instrument Artefacts?, *Geophysical Research Letters*, 47 (11),
 e2019GL086791, doi: 10.1029/2019GL086791.
- Torres, O., Bhartia, P.K., Jethva, H. and Ahn, C. 2018. Impact of the ozone monitoring instrument row
 anomaly on the long-term record of aerosol products. *Atmospheric Measurement Techniques*, **11**, 27012715, doi: 10.5194/amt-11-2701-2018.
- van der A, et al. 2006. Detection of the trend and seasonal variation in tropospheric NO₂ over China. *Journal*of *Geophysical Research*, **11**, D12317, doi: 10.1029/2005JD006594.
- Van Roozendael M, et al. 2012. Sixteen years of GOME/ERS-2 total ozone data: The new direct-fitting GOME
 Data Processor (GDP) version 5—Algorithm description. *Journal of Geophysical Research: Atmospheres,* 117,
 D03305, doi: 10.1029/2011JD016471.
- Verstraeten WW, et al. 2015. Rapid increase in tropospheric ozone production and export from China. *Nature Geoscience*, **8**, 690-695, doi: 10.1038/NGEO2493.
- Wespes C, et al. 2018. Decrease in tropospheric levels in the Northern Hemisphere observed by IASI. *Atmospheric Chemistry and Physics*, **18**, 6867-6885, doi: 10.5194/acp-18-6867-2018.
- Young PJ, et al. 2013. Pre-industrial to end 21st century projections of tropospheric ozone from the
 Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP). *Atmospheric Chemistry and Physics*, 13, 2063-2090, doi: 10.5194/acp-13-2063-2013.
- Ziemke JR, et al. 2006. Tropospheric ozone determined from Aura OMI and MLS: Evaluation of
 measurements and comparison with the Global Modelling Initiative's Chemical Transport Model, *Journal of*
- 507 *Geophysical Research*, **111** (D19303), doi: 10.1029/2006JD007089.
- Ziemke JR, et al. 2011. A global climatology of tropospheric and stratospheric ozone derived from Aura
 OMI/MLS measurements, *Atmospheric Chemistry and Physics*, **11**, 9237-9251, doi: /10.5194/acp-11-92372011.
- Ziemke JR, et al. 2019. Trends in global tropospheric ozone inferred from a composite record of
 TOMS/OMI/MLS/OMPS satellite measurements and the MERRA-2 GMI simulation. *Atmospheric Chemistry and Physics*, **19**, 3257-3269, doi: 10.5194/acp-19-3257-2019.
- 514
- 515
- 516
- 210
- 517
- 518

519 Figures & Tables:

-				
Data Provider	Satellite Profile	Product Link	Data	Data Size
	Products & Version		Range	
RAL Space	OMI–fv214	http://www.ceda.ac.uk/	2004-2018	1442 GB
RAL Space	GOME-2A-fv300	http://www.ceda.ac.uk/	2007-2019	1007 GB
RAL Space	GOME-1-fv301	http://www.ceda.ac.uk/	1995-2011	703 GB
RAL Space	SCIAMACHY-fv300	http://www.ceda.ac.uk/	2002-2012	718 GB

Table 1: List of RAL Space level-2 satellite ozone profile data sets.

Latitude Band	LTCO₃ Trend (DU/decade) (95% Confidence Interval)	LTCO₃ Trend (ppbv/decade) (95% Confidence Interval)	p-values
60°S ≤ Latitude < 45°S	4.49 (2.51, 6.48)	10.37 (5.79, 14.95)	0.00
45°S ≤ Latitude < 30°S	1.85 (0.11, 3.59)	4.27 (0.26, 8.28)	0.03
30°S ≤ Latitude < 15°S	0.94 (-1.05, 2.93)	2.17 (-2.42, 6.76)	0.35
15°S ≤ Latitude < 0°	2.89 (1.27, 4.52)	6.68 (2.94, 10.43)	0.00
0° ≤ Latitude < 15°N	3.93 (3.13, 4.72)	9.06 (7.23, 10.89)	0.00
15°N ≤ Latitude < 30°N	4.12 (3.25, 4.97)	9.50 (7.51, 11.48)	0.00
30°N ≤ Latitude < 45°N	1.33 (-0.34, 3.01)	3.08 (-0.78, 6.95)	0.11
45°N ≤ Latitude < 60°N	0.49 (-1.14, 2.13)	1.14 (-2.64, 4.91)	0.55

Table 2: $LTCO_3$ trends (DU/decade and ppbv/decade) for latitude bands (15° bins) between 60°S and 60°N.

523 The 95% confidence intervals of the trends are shown in brackets. The trend p-values are also shown.



Figure 1: Average averaging kernels (AKs) for the instruments listed in **Table 1** for the northern and southern hemispheres ($60^{\circ}S-60^{\circ}N$) in January and July of 2008 (1998 for GOME-1). The average degrees of freedom of signal (DOFS) is shown as is DOFS LTCO₃ which represents the DOFS in the lower tropospheric column ozone

 $(LTCO_3)$. N represents the number of retrievals in each average AK average.



5360.00.20.30.50.70.81.0537Figure 2: Seasonal distributions of LTCO3 degrees of freedom of signal (DOFS) in DJF and JJA for GOME-1,538GOME-2, OMI and SCIAMACHY averaged over the full record for each instrument.





547 size.



Figure 4: Seasonal distributions of LTCO₃ in December-January-February (DJF) and June-July-August (JJA) for
 OMI, GOME-1, GOME-2 and SCIAMACHY averaged over the full record for each instrument.



-15.0-10.0-5.00.05.010.015.0555Figure 5: Latitudinal-annually varying satellite-sonde, with AKs applied, LTCO3 (DU) median (50^{th} percentile)556biases. Hatched regions show where the spread in the 25^{th} and 75^{th} percentiles intersect with 0.0. The mean557bias (MB) and trend are for the full time series of each hemisphere. The * for the trend term indicates that it558has a p-value < 0.05. The latitude bands are $90-30^\circ$ S, 30° S- 30° N and $30-90^\circ$ N.



Figure 6: Examples of the merged LTCO₃ (DU) data set for Europe and East Asia. The GOME-1, OMI and
 merged time series are shown in red, blue and black, respectively. The merged record also includes globally
 scaled LTCO₃ data from SCIAMACHY for 2003 and 2004. Dashed lines represent the annual averages and the

576 monthly mean time-series are solid lines.



578 **Figure 7**: LTCO₃ (DU) merged data set from GOME-1 (1996-2002), SCIAMACHY (2003-2004) and OMI (2005-

579 2017). a) 1996-2017 long-term average, b) 1996-2000 average anomaly, c) 2005-2009 average anomaly and 580 d) 2013-2017 average anomaly. Anomalies are relative to the long-term average (panel a). Green polygon-

d) 2013-2017 average anomaly. Anomalies are relative to the long-term average (panel a). Green polygon outlined regions show significant anomalies (95% confidence level and where the absolute anomaly > 1.0 DU)

582 from the long-term average using the Wilcoxon Rank Test. White/grey pixels are where the South Atlantic

583 Anomaly influence on retrieved LTCO₃ has been masked out.



Figure 8: LTCO3 (DU) merged data set from GOME-1 (1996-2002), SCIAMACHY (2003-2004) and OMI (2005-2017) where the difference between the 2013-2017 average and 1996-2000 average is shown. Green polygon-outlined regions show substantial differences (95% confidence level and where the absolute difference > 1.0 DU) using the Wilcoxon Rank Test. Grey pixels are where the South Atlantic Anomaly influence on retrieved LTCO₃ has been masked out. Circles show differences in ozonesonde LTCO₃ (DU) over the same time periods as the merged satellite record.