Response to Referee #2: 1

3 Responses to individual comments are given here below:

5 [Overall the authors have done a goodly amount of change to the paper as the marked revision shows. The paper is well written and with current figures that are both legible and concise for the paper. The revision 6 7 now includes discussion in various places of the jump/drift with IASI that was not in the original draft. 8 They have also re-done their trend analyses in accordance to this jump/drift in IASI.]

- 9
- We thank the referee for her/his appreciation for the changes made in the manuscript in order to address all the two referees' comments. 10
- 11 12

13

14 15

16

17

18

2

4

[In the revision the authors have added discussion of a downward "jump" in tropospheric ozone measurements around September 2010 for IASI which they mention causes an artificial overall downward trend or drift of about -2.8 DU/decade in the NH. This is new to the revision which mentions this jump several times including the Conclusions section as possibly affecting the calculated trends. In an effort to account for this jump the authors have modified their regression trend model by including two different constants in the regression for the two separate time periods, before and after September 2010. An argument is made that the jump-related drift of -2.8 DU does not explain the larger negative trends in summer of ~-5

- 19 DU/decade measured by IASI.
- 20 The authors state that the reason for the downward jump in tropospheric ozone from IASI is not clear. The reference list includes Keppens et al. and Boynard et al. papers that discuss a detected negative drift in IASI 21
- 22 tropospheric ozone. Both are related papers using IASI and are in preparation/under review for this same
- issue; the two papers are mentioned specifically in the revision in regards to the drift/jump in IASI ozone. 23
- 24 The revision states that Boynard et al. (this issue) describes the IASI drift as being caused largely by a
- 25 downward discontinuity "jump" in the IASI data around September 2010. An earlier paper published by
- Boynard et al. (2016) shows in their Figure 15 evidence of this jump and a persistent downward drift/trend 26
- 27 in IASI tropospheric ozone relative to ozonesondes in both the NH and SH extra-tropics. The downward
- 28 drift (including jump) for IASI tropospheric ozone relative to the ozonesondes indicated by Boynard et al.

29 (2016) was never discussed in their 2016 paper. The current revision references the Boynard and Keppens

- 30 papers that are under review for this same special issue regarding the IASI jump/drift.]
- 31 A discontinuity was not clearly enough demonstrated in the paper of Boynard et al. (2016). Furthermore,
- that paper reports the validation of the previous FORLI-O₃ product (FORLI-v20140922), while Boynard et 32 al. (under review, this issue) validates the last FORLI-O₃ product (FORLI-v20151009) which is the one 33
- 34 used in this manuscript and his companion paper (Wespes et al., 2017). As a consequence, only the results
- 35 reported in Boynard et al (under review, this issue) should be considered for the present O₃ trend analysis.
- 36
- 37 [The authors state that tropospheric ozone for IASI has one piece of information that corresponds to ground-
- 38 to-300 hPa. The authors define this as middle-low troposphere (MLT) ozone. They mention that the upper
- 39 level 300 hPa tends to minimize influence from stratospheric ozone in the retrievals. The revision still states
- 40 that the significant negative trends in the SH are hard to explain, and mention that stratospheric ozone
- influence may be a large reason for this band structure of negative MLT trends throughout the SH year-41
- round. There will be questions from readers regarding the very nature of IASI nadir retrievals in resolving 42

tropospheric ozone, especially how much tailing influence from ozone above 300 hPa (includingstratosphere) there is in the MLT measurements, especially in the extra-tropics.]

45 In a previous paper (Wespes et al., 2016), we have specifically quantified the stratospheric contributions

46 into the ground-300 hPa O₃ columns as function of the latitudes and the periods. Values from that study

47 have already been referenced here precisely to discuss critically the possible impact of the stratosphere in

48 the tropospheric O_3 trends. We don't see how to improve on this without duplicating this earlier paper.

49

50 [Papers listed in my first review describing zero or positive trends measured in extra-tropical tropospheric

- 51 ozone are not included in the revision. There are several reasons stated for not referencing them in the 52 revision such as issues of MLT versus UT or differences in the vertical resolution of the measurements.
- 53 The author's response is that including reference to these is beyond the scope of the present paper. The

54 Petetin et al. (2016) paper (the diurnal cycle paper) that I mentioned in my first review used

55 MOZAIC+IAGOS aircraft measurements over Frankfurt and showed statistically significant increases in

56 ozone throughout the troposphere from ground to 300 hPa (i.e., MLT). Regarding the TOAR, another basic

57 issue for the satellite measurements including IASI is their short records for doing trend analyses and that

58 their time periods are generally quite different.]

59 We have indeed carefully looked at the paper of Petetin et al. (2016) which is of high relevance. However,

60 considering the recent decline observed in the anthropogenic O₃ precursors in the N.H. (since 2010-2011),

61 the results presented in their Figures 5 and S6-9 are not comparable with ours. They show "relative

63 hence, they are not representative of the period analyzed here on global scale. For this reason, we have

65 have been reported from independent existing datasets including MOZAIC and IAGOS.

66

67 Moreover, we would like to stress that significant efforts have been devoted in order to disentangle trend from other dynamical effects by applying a dedicated MLR model to the IASI data and that it has been 68 69 shown achievable even from a relatively short period of measurements (Wespes et al., 2016). While other methods are also valuable, we feel that a proper comparison can only be made against studies that have 70 71 used similar MLR. In that respect, we now provide in the revised manuscript additional discussion based 72 on the results of Leventidou et al., (under review), Heue et al. (2016) and Ebojie et al. (2016) which have 73 applied MLR to the data from UV sounders. From these papers, one can easily see the challenge in 74 homogenizing independent datasets and reaching consensus in determining tropospheric O₃ trends. 75 Nevertheless, significant negative trends are interestingly found at the global scale (~-4 to -8 DU/dec; Ebojie et al., 2016) and in the tropics over the Oceanic regions (~ -1 to -2.5 DU/dec; Leventidou et al., 76 under review) while significant positive trends are observed over Africa (Leventidou et al., under review). 77 78 These results are comparable with those reported in our manuscript, despite the different studied periods

79 (1995-2015 in Leventidou et al. and 2003-2011 in Ebojie et al., 2016). These papers have been added in the

80 revised manuscript.

81

82 In order not to oversell the results, we have also made clearer in the revised manuscript (abstract, section

- **83** 4.3 and conclusion) that, at this stage, no consensus in terms of O_3 trends can be easily reached in the
- troposphere from the available measurements (UV or IR satellites, O_3 sondes, aircrafts, ground-based
- 85 measurements,...) for the reasons already mentioned in the manuscript (time-varying instrumental biases,

89

90 [There appears to be some questions regarding the IASI MLT ozone measurements themselves for 91 evaluating trends. The drift for IASI tropospheric ozone is a bit disturbing as it is rather large and not 92 explainable from either the current study or those of Keppens et al. or Boynard et al. that are related IASI 93 papers also in review in this same issue. The negative trends throughout much of the NH and SH for IASI 94 MLT ozone appear to be in contradiction to zero or positive trends measured from other independent data 95 sources (aircraft, ozonesonde, satellite), albeit of differing (usually longer) time records and not specifically 96 calculated for ground-to-300hPa as IASI. The authors attribute negative trends in the NH as possibly due 97 to reductions in emissions in recent years, particularly over N. America and Europe. The authors state that 98 the negative trends in the SH are hard to explain, but possibly of stratospheric origin.]

99 See our responses to the two previous comments about reconciling trend with previous studies and the100 stratospheric contribution into the S.H. trends.

101

102 We acknowledge that the origin of the artificial drift is still unclear and under investigation at present. 103 However, we feel that the inclusion of a second constant in the MLR model is a pretty robust way to deal 104 with this issue and, hence, that the jump should not interfere with the trends. The fact that this correction 105 did not change the broad results gives somewhat more confidence in the retrieved O_3 trend.

106

107 [Given over 9 years of measurements from IASI for detecting decadal changes in global tropospheric ozone 108 (main theme of the paper), it would seem important to compare decadal changes in IASI MLT ozone 109 directly with decadal changes in other independent data products in the paper such as station ozonesondes 110 or IAGOS aircraft ozone. This paper is going to raise some doubts with readers as to the IASI trend results 111 given the current unknowns with the data. There is really not enough 1-1 comparison evidence presented 112 from other independent measurements to test validity of the IASI trend results.]

We certainly agree that such a 1-1 comparison would be of great interest. However it is definitively not straightforward as it would deserve setting careful criteria on co-locations and involve a detailed investigation of the respective vertical sensitivities. Moreover, differences in spatio-temporal samplings

 $\label{eq:should be also investigated.} \ Attempts have been made to apply a MLR model on the same O_3 sondes dataset$

117 (smoothed by the IASI averaging kernels) as the one used for the validation in Boynard et al. (this issue).

118 Unfortunately, the regression residual errors were found too large to retrieve significant trend or dynamical

119 covariates because of the weak temporal sampling of the dataset. We feel and we would like the reviewer

and the editor to agree that such a 1-1 comparison is outside the scope of this paper, which again, exploitsthe global coverage of IASI to extract relevant trends.

122

We also would like to stress that the validation of FORLI- O_3 in Boynard et al. (this issue) with O_3 sondes highlights a jump but not a persistent downward drift, as it has been confirmed by calculating the drift

separately over the periods before or after the jump (cfr our responses to previous comments to referees #1

126

and #2).

128	Finally, we would like to notify the reviewer on the fact that such an evaluation of the different IASI-O ₃
129	products against O ₃ sondes and daily IAGOS commercial aircraft profiles is being initiated by the IASI data
130	providers (ULB/LATMOS, LA and LISA) in collaboration with the IAGOS data providers and the TOAR-
131	climate leaders (A. Gaudel and O. Cooper).
132	
133	
134	
135	
136	
137	
138	
139	
140	
141	
142	
143	
144	
145	
146	
147 148	
148 149	
149	
151	
152	
153	
154	
155	
156	
157	
158	
159	
160	
161	
162	
163	
164	
165	
166	
167	

- 168 List of relevant changes made in the manuscript:
- 169

170 Abstract:

- L. 23-26: "Despite that no consensus in terms of tropospheric O₃ trends is currently reached from the available independent datasets (UV or IR satellites, O₃ sondes, aircrafts, ground-based measurements...) for the reasons that are discussed in the text, this finding is consistent with..."
- 175

176 Section 4.1:

- L. 236-272: "On the contrary, the tropical Pacific region exhibits significant negative trends that are similar to those reported from UV sounders in Ebojie et al. (2016) and in Leventidou et al. (in review) over previous periods, while Heue et al. (2016) mainly reports significant positive trend over that region."
- L. 285-286: "Significant positive trends over South-East Asia have also been reported from UV sounders over previous periods (e.g. Ebojie et al., 2016)."
- L. 303-304: "Significant negative change in tropospheric O₃ over these regions were also
 reported in Ebojie et al. (2016)."
- 185

186 Section 4.3:

- L. 425 428: "If reconciling the trend biases between the datasets by applying the vertical sensitivity of each measurement type to a common platform, as proposed in the TOAR-climate assessment report is beyond the scope of this study and if, at this stage, there is no consensus in determining tropospheric O₃ trends, the improvement in using a MLR instead of a SLR model ..."
- 192

193 Conclusions:

- L. 589-594 : "Currently, no consensus in terms of O₃ trends in the troposphere is reached from the available measurements (UV or IR satellites, O₃ sondes, aircrafts, ground-based measurements,...) for several reasons (time-varying instrumental biases, differences in the methodology used for calculating trends, in the measurement period, in the upper boundary of the O₃ columns, in the retrieval algorithm, in the spatio-temporal sampling, in the vertical sensitivity of the instrument,...) (Section 4.3; the TOAR-climate report – Gaudel et al., in review). However, determination, with IASI ..."
- 201
- 202 **References:**
- Ebojie, F., Burrows, J. P., Gebhardt, C., Ladstätter-Weißenmayer, A., von Savigny, C.,
 Rozanov, A., Weber, M., and Bovensmann, H.: Global tropospheric ozone variations from
 205 2003 to 2011 as seen by SCIAMACHY, Atmos. Chem. Phys., 16, 417-436,
 doi:10.5194/acp-16-417-2016, 2016.

207	-	Heue, KP., Coldewey-Egbers, M., Delcloo, A., Lerot, C., Loyola, D., Valks, P., and van
208		Roozendael, M.: Trends of tropical tropospheric ozone from 20 years of European satellite
209		measurements and perspectives for the Sentinel-5 Precursor, Atmos. Meas. Tech., 9, 5037-
210		5051, doi:10.5194/amt-9-5037-2016, 2016. Leventidou, E., Weber, M., Eichmann, KU., and Burrows, J. P.: Harmonisation and trends
211 212	-	of 20-years tropical tropospheric ozone data, Atmos. Chem. Phys. Discuss.,
212		https://doi.org/10.5194/acp-2017-815, in review, 2017.
214		
215		
216		
217		
218		
219		
220		
221		
222		
223		
224		
225		
226		
227		
228		
229		
230		
231		
232 233		
233		
234		
235		
237		
238		
239		
240		
241		
242		
243		
244		
245		

Decrease in tropospheric O₃ levels of the Northern Hemisphere observed by IASI

Catherine Wespes¹, Daniel Hurtmans¹, Cathy Clerbaux^{1,2}, Anne Boynard² and Pierre-François
 Coheur¹

- ¹Spectroscopie de l'Atmosphère, Service de Chimie Quantique et Photophysique, Faculté des
- 252 Sciences, Université Libre de Bruxelles (ULB), Bruxelles, Belgique
- ²LATMOS/IPSL, UPMC Univ. Paris 06 Sorbonne Universités, UVSQ, CNRS, Paris, France
- 254 255

256 Abstract

In this study, we describe the recent changes in the tropospheric ozone (O_3) columns measured by 257 the Infrared Atmospheric Sounding Interferometer (IASI) onboard the Metop satellite during the 258 first 9 years of the IASI operation (January 2008 to May 2017). Using appropriate multivariate 259 regression methods, we discriminate significant linear trends from other sources of O₃ variations 260 captured by IASI. The geographical patterns of the adjusted O₃ trends are provided and discussed 261 on the global scale. Given the large contribution of the natural variability in comparison with that 262 of the trend (25-85% vs 15- 50%, respectively) to the total O_3 variations, we estimate that 263 additional years of IASI measurements are generally required to detect the estimated O₃ trends 264 with a high precision. Globally, additional 6 months to 6 years of measurements, depending on the 265 regions and the seasons, are needed to detect a trend of [5] DU/decade. An exception is interestingly 266 found during summer at mid-high latitudes of the North Hemisphere (N.H.; ~ 40°N-75°N) where 267 268 the large absolute fitted trend values (~[0.5] DU/yr on average) combined with the small model residuals (~10%) allow the detection of a band-like pattern of significant negative trends. Despite 269 270 that no consensus in terms of tropospheric O₃ trends is currently reached from the available independent datasets (UV or IR satellites, O3 sondes, aircrafts, ground-based measurements,...) 271 272 for the reasons that are discussed in the text, tThis finding is consistent with the reported decrease 273 in O_3 precursor emissions in recent years, especially in Europe and US. The influence of continental pollution on that latitudinal band is further investigated and supported by the analysis 274 of the O₃-CO relationship (in terms of correlation coefficient, regression slope and covariance) 275 that we found to be the strongest at northern mid-latitudes in summer. 276

277

278 **1 Introduction**

279

 O_3 plays a key role throughout the whole troposphere where it is produced by the photochemical oxidation of carbon monoxide (CO), non-methane volatile organic compounds (NMVOCs) and methane (CH₄) in the presence of nitrogen oxides (NO_x) (e.g. Logan et al., 1981). O₃ sources in the troposphere are the in situ photochemical production from anthropogenic and natural precursors, and the downwards transport of stratospheric O₃. Being a strong pollutant, a major reactive species and an important greenhouse gas in the upper troposphere, O₃ is of highest interest for air quality, atmospheric chemistry and radiative forcing studies. Thanks to its long lifetime (several weeks) relatively to transport timescales in the free troposphere (Fusco and Logan, 2003), O₃ also contributes to large-scale transport of pollution far from source regions with further impacts on global air quality (e.g. Stohl et al., 2002; Parrish et al., 2012) and climate. Monitoring and understanding the time evolution of tropospheric O₃ at a global scale is, therefore, crucial to apprehend future climate changes. Nevertheless, a series of limitations make O₃ trends particularly challenging to retrieve and to interpret.

293

294 Since the 1980s, while the O_3 precursors anthropogenic emissions have increased and shifted 295 equatorward in the developing countries (Zhang et al., 2016), extensive campaigns and routine in situ and remote measurements at specific urban and rural sites have provided long-term but sparse 296 datasets of tropospheric O₃ (e.g. Cooper et al., 2014 and references therein). Ultraviolet and Visible 297 298 (UV/VIS) atmospheric sounders onboard satellites provide tropospheric O₃ measurements with a much wider coverage, but they result either from indirect methods (e.g. Fishman et al., 2005) or 299 300 from direct retrievals which are limited by coarse vertical resolution (Liu et al., 2010). All these 301 datasets also suffer from a lack of homogeneity in terms of measurement methods (instrument and 302 algorithm) and spatio-temporal samplings (e.g. Doughty et al., 2011; Heue et al., 2016; Leventidou 303 et al., in review). Those limitations, in addition to the large natural inter-annual variability (IAV) 304 and decadal variations in tropospheric O₃ levels (due to large-scale dynamical modes of O₃ variations and to changes in stratospheric O_3 , in stratosphere-troposphere exchanges, in precursor 305 emissions and in their geographical patterns), introduce strong biases in trends determined from 306 independent studies and datasets (e.g. Zbinden et al., 2006; Thouret et al., 2006; Logan et al., 307 308 2012; Parrish et al., 2012 and references therein). As a consequence, determining accurate trends requires a long period of high density and homogeneous measurements (e.g. Payne et al., 2017). 309

310

Such long-term datasets are now becoming obtainable with the new generation of nadir-looking and polar-orbiting instruments measuring in the thermal infrared region. In particular, about one decade of O₃ profile measurements, with a good sensitivity in the troposphere independently from the layers above, is now available from the IASI (Infrared Atmospheric Sounding Interferometer) sounder aboard the European Metop platforms, allowing to monitor regional and global variations in tropospheric O₃ levels (e.g. Dufour et al., 2012; Safieddine et al., 2013; Wespes et al., 2016).

317

In this study, we examine the tropospheric O_3 changes behind the natural IAV as measured by 318 IASI over January 2008-May 2017. To that end, we use the approach described in Wespes et al. 319 (2017), which relies on a multi-linear regression (MLR) procedure, for accurately differentiating 320 trends from other sources of O_3 variations; the latter being robustly identified and quantified in 321 that companion study. In Section 2, we briefly review the IASI mission and the tropospheric O₃ 322 product, and we shortly describe the multivariate models (annual or seasonal) that we use for fitting 323 the daily O₃ time series. In Section 3, after verifying the performance of the MLR models over the 324 325 available IASI dataset, we evaluate the feasibility to capture and retrieve significant trend parameters, apart from natural O_3 dependencies, by performing trend sensitivity studies. In Section 4, we present and discuss the global distributions of the O_3 trends estimated from IASI in the troposphere. The focus is given in summer over and downwind anthropogenic polluted areas of the N.H. where the possibility to infer significant trends from the first ~9 years of available IASI measurements is demonstrated. Finally, the O_3 -CO correlations, enhancement ratios and covariance are examined for characterizing the origin of the air masses in regions of positive and negative trends.

- 333
- 334 335

2 IASI O₃ measurements and multivariate regression

336 The IASI instrument is a nadir-viewing Fourier transform spectrometer that records the thermal infrared emission of the Earth-atmosphere system between 645 and 2760 cm⁻¹ from the polar Sun-337 synchronous orbiting meteorological Metop series of satellites. Metop-A and -B have been 338 successively launched in October 2006 and September 2012. The third and last launch is planned 339 340 in 2018 with Metop-C to ensure homogeneous long-term IASI measurements. The measurements are taken every 50 km along the track of the satellite at nadir and over a swath of 2200 km across 341 342 track, with a field of view of four simultaneous footprints of 12 km at nadir, which provides global coverage of the Earth twice a day (at 9:30 AM and PM mean local solar time). The instrument 343 presents a good spectral resolution and a low radiometric noise, which allows the retrieval of 344 numerous gas-phase species in the troposphere (e.g. Clerbaux et al., 2009, and references therein; 345 Hilton et al., 2012; Clarisse et al., 2011). 346

347

348 In this paper, we use the FORLI-O₃ profiles (Fast Optimal Retrievals on Layers for IASI processing chain set up at ULB; v20151001) retrieved from the IASI-A (aboard Metop-A) daytime 349 350 measurements (defined with a solar zenith angle to the sun $< 80^{\circ}$) which are characterized by a 351 good spectral fit (determined here by quality flags on biased or sloped residuals, suspect averaging kernels, maximum number of iteration exceeded,...) and which correspond to clear or almost-clear 352 353 scenes (a filter based on a fractional cloud cover below 13% has been applied; cfr Clerbaux et al., 2009; Hurtmans et al., 2012). These profiles are characterized by a good vertical sensitivity in the 354 troposphere and the stratosphere (e.g. Wespes et al., 2017). The FORLI algorithm relies on a fast 355 radiative transfer and retrieval methodology based on the Optimal Estimation Method (Rodgers, 356 2000) and is fully described in Hurtmans et al. (2012). The FORLI-O₃ profiles, which are retrieved 357 at 40 constant vertical layers from surface up to 40 km and an additional 40-60 km one, have 358 359 already undergone thorough characterization and validation exercises (e.g. Anton et al., 2011; Dufour et al., 2012; Gazeaux et al., 2012; Hurtmans et al., 2012; Parrington et al., 2012; Pommier 360 et al., 2012; Scannell et al., 2012; Oetjen et al., 2014; Boynard et al., 2016; Wespes et al., 2016; 361 Keppens et al. 2017; Boynard et al., 2017). They demonstrated a good degree of accuracy, of 362 precision and of vertical sensitivity with no instrumental drift, to capture the large-scale dynamical 363 modes of O₃ variability in the troposphere independently from the layers above (Wespes et al., 364 365 2017), with the possibility to further differentiate long-term O_3 changes in the troposphere (Wespes et al., 2016). Note, however, that a drift in the N.H. MLT O₃ over the whole IASI dataset is reported
in Keppens et al. (this issue) and Boynard et al. (this issue) from comparison with O₃ sondes. This
drift (~2.8DU/dec in the N.H.) is shown in Boynard et al. (this issue) to result from a discontinuity
("jump" as called in Boynard et al., this issue) in September 2010 in the IASI O₃ time series, for
reasons that are unclear at present. Furthermore, the drift strongly decreases (<|1| DU/dec on
average) after the "jump" and it becomes even non-significant for most of the stations (significant
positive drift is also found for some stations) over the periods before or after the jump, separately.

373

For the purpose of this work, we focus on a tropospheric column ranging from ground to 300 hPa 374 375 (called MLT – Middle-Low Troposphere – in this study) that includes the altitude of maximum sensitivity of IASI in the troposphere (usually between 4 and 8 km altitude), which limits as much 376 as possible the influences of the stratospheric O₃ and that was shown in Wespes et al. (2017) to 377 exhibit independent deseasonalized anomalies/dynamical processes from those in the stratospheric 378 layers. The stratospheric contribution into the tropospheric O₃ columns have been previously 379 estimated in Wespes et al. (2016) as ranging between 30% and 65% depending on the region and 380 the season with the smallest contribution as well as the largest sensitivity in the northern mid-381 latitudes in spring-summer where the O₃ variations, hence, mainly originate from the troposphere. 382 383 We use almost the same MLR model (in its annual or its seasonal formulation) as the one developed in the companion paper (see Eq.1 and 2; Section 2.2 in Wespes et al., 2017), which 384 includes a series of geophysical variables in addition to a linear trend (LT) term. In order to take 385 account of the observed "jump" properly, we modified the previously used MLR model so that the 386 constant term is split into two components covering the periods before and after the September 387 2010 "jump", separately. The MLR which is performed using an iterative stepwise backward 388 389 elimination approach to retain the most relevant explanatory variables (called "proxies") at the end of the iterations (e.g. Mäder et al., 2007) is applied on the daily IASI O₃ time series. The main 390 391 selected proxies used to account for the natural variations in O₃ are namely the QBO (Quasi-Biennial Oscillation), the NAO (North Atlantic Oscillation) and the ENSO (El Niño-Southern 392 Oscillation) (cfr Table 1 in Wespes et al. (2017) for the exhaustive list of the used proxies). Their 393 associated standard errors are estimated from the covariance matrix of the regression coefficients 394 and are corrected to take into account the uncertainty due to the autocorrelation of the noise 395 residual (see Eq. 3 in Wespes et al. (2016)). The common rule that the regression coefficients are 396 significant if they are greater in magnitude than 2 times their standard errors is applied (95% 397 confidence limits defined by 2σ level). The MLR model was found to give a good representation 398 of the IASI O₃ records in the troposphere over 2008-2016, allowing us to identify/quantify the 399 400 main O₃ drivers with marked regional differences in the regression coefficients. Time-lags of 2 and 4 months for ENSO are also included hereafter in the MLR model to account for a large but 401 delayed impact of ENSO on mid- and high latitudes O₃ variations far from the Equatorial Pacific 402 where the ENSO signal originates (Wespes et al., 2017). 403 404

405 **3 Regression performance and sensitivity to trend**

In this section, we first verify the performance of the MLR models (annual and seasonal; in terms of residual errors and variation explained by the model) to globally reproduce the time evolution of O_3 records over the entire studied period (January 2008 – May 2017). Based on this, we then

- 410 investigate the statistical relevance for a trend study from IASI in the troposphere by examining
- 411 the sensitivity of the pair IASI-MLR to the retrieved LT term.
- 412

Figure 1 presents the seasonal distributions of tropospheric O_3 measured by IASI averaged over January 2008 – May 2017 (left panels), along with the root-mean-squared error of the seasonal

regression fit (*RMSE*, in DU; middle panels) and the contribution of the fitted seasonal model into

416 the IASI O₃ time series (in %; right panels), calculated as $\frac{\sigma(O_3^{\text{Fitted}_model}(t))}{\sigma(O_3(t))}$ where σ is the standard

deviation relative to the regression models and to the IASI O_3 time series. These two statistical 417 parameters help to evaluate how well the fitted model explains the variability in the IASI O_3 418 419 observations. The seasonal patterns of O_3 measurements are close to those reported in Wespes et al. (2017) for a shorter period (see Section 2.1 and 3.1 in Wespes et al. (2017) for a detailed 420 description of the distributions) and they clearly show, for instance, high O₃ values over the highly 421 populated areas of Asia in summer. The distributions from Fig.1 show that the model reproduces 422 between 35% and 90% of the daily O₃ variation captured by IASI and that the residual errors varies 423 424 between 0.01 DU and 5 DU (i.e. the *RMSE* relative to the IASI O_3 time series are of ~15% on 425 global average and vary between 10% in the N.H. in summer and 30% in specific tropical regions). On an annual basis (data not shown), the model explains a large fraction of the variation in the 426 IASI O₃ dataset (from ~45% to ~85%) and the *RMSE* are lower than 4.5 DU everywhere (~3 DU 427 on the global average). The relative *RMSE* is less than 1% in almost all situations indicating the 428 429 absence of bias.

430

431 The seasonal distributions of the contribution to the total variation in the MLT from the adjusted 432 harmonic terms and explanatory variables, which account for the "natural" variability, and from 433 the LT term are shown in Fig. 2 (left and right panels, respectively). The grey areas in the LT panels refer to the LT terms rejected by the stepwise backward elimination process. The crosses 434 indicate that the trend estimate in the grid cell is non-significant in the 95% confidence limits (2σ 435 level) when accounting for the autocorrelation in the noise residual at the end of the elimination 436 procedure. While the large influence of the seasonal variations and of the main drivers - namely 437 ENSO, NAO and QBO - on the IASI O₃ records has been clearly attested in Wespes et al. (2017), 438 we demonstrate in Fig.2 that the LT also contributes considerably to the O₃ variations detected by 439 440 IASI in the troposphere. The LT contribution generally ranges from 15% to 50%, with the largest values (~30-50%) being observed at mid-high latitudes in the S.H. (30°S-70°S) and in the N.H. 441 (~45°N-70°N) in summer. In the S.H., they are associated with the smallest tropospheric O₃ 442 columns (Fig.1; left panels) and the smallest natural contributions (<25%; left panels), while in the 443

444 N.H. summer, they interestingly correspond to large MLT O₃ columns, large natural contributions 445 (\sim 50-60%) and the smallest *RMSE* (<12 % or <3 DU). From the annual regression model, the 446 natural variation and the trend contribute respectively for 30-85% and up to 40% to the total 447 variation in the MLT.

448 In Fig.3, we further investigate the robustness of the estimated trends by performing sensitivity tests in regions of significant trend contributions (e.g. in the N.H. mid-latitudes in summer; cfr 449 Fig.2). The ~9-year time series of IASI O₃ daily averages (dark blue) along with the results from 450 the seasonal regression model with and without the LT term included in the model (light blue and 451 orange lines, respectively) are represented in the top row panel for one specific location (Fig.3a 452 and b; highlighted by a blue circle in the JJA panel in Fig.4). The second row panel provides the 453 deseasonalised IASI (dark blue line) and fitted time series (calculated by subtracting the adjusted 454 seasonal cycle from the time series) resulting from the adjustment with and without the LT term 455 included in the MLR model (light blue and orange lines, respectively). The differences between 456 the fitted models with and without LT are shown in the third rows (pink lines). They match fairly 457 well the adjusted trend over the IASI period (3^d row panel, grey lines; the trend and the *RMSE* 458 values are also indicated) and the adjustment without LT leads to larger residuals (e.g. 459 RMSE JJA W/0 LT=3.37 DU vs. RMSE JJA with LT=3.21 DU in summer). This result demonstrates the 460 possibility to capture trend information from ~9 years of IASI-MLR with only some compensation 461 effects by the other explanatory variables, contrary to what was observed when considering a 462 shorter period of measurements or a lesser temporal sampling (i.e. monthly dataset; e.g. Wespes 463 464 et al., 2016). It is also worth to mention that the O₃ changes calculated over the whole IASI dataset in summer are larger than the RMSE of the model residuals (increase of 5.39 ± 1.86 DU vs RMSE 465 of 3.21 DU), underlying the statistical relevance of trend estimates. 466

467 The robustness of the adjusted trend is verified at the global scale in Fig.4 which represents the seasonal distributions of the relative differences in the RMSE with and without the LT included in 468 the MLR model, calculated as $[(RMSE_{w/o} LT - RMSE_{with} LT)/RMSE_{with} LT \times 100]$ (in %). An 469 increase in the *RMSE* when excluding LT from the MLR is observed almost everywhere in regions 470 of significant trend contributions (Fig.2), especially in mid-high latitudes of the S.H. and of the 471 N.H. in summer where it reaches 10%. This result indicates that adjusting LT improves the 472 performance of the model and, hence, that a trend signal is well captured by IASI at a regional 473 scale in the troposphere. From the annual model, the increase in the RMSE only reaches 5% at 474 mid-high latitudes of the S.H. (data not shown). In regions of weak or non-significant trend 475 contribution (see crosses in Fig.2), no improvement is logically found. 476

- 477
- 478 **4 O₃ trend over 2008-2017**
- 479 **4.1 Annual and seasonal trends**
- 480

The annual and the seasonal distributions of the fitted LT terms which are retained in the annual 481 482 and the seasonal MLR models by the stepwise elimination procedure are respectively represented in Fig. 5 (a) and (b) (in DU/yr). Generally, the mid-high latitudes of both hemispheres and, more 483 particularly, the N.H. mid-latitudes in summer reveal significant negative trends, while the tropics 484 are mainly characterized by non-significant or weak significant trends. Even if trends in emissions 485 have already been able to qualitatively explain measured tropospheric O₃ trends over specific 486 regions, the magnitude and the trend estimates considerably vary between independent 487 measurement datasets (e.g. Cooper et al., 2014; the TOAR report-Tropospheric Ozone 488 Assessment Report: Present day distribution and trends of tropospheric ozone relevant to climate 489 490 and global atmospheric chemistry model evaluation, Lead Authors: A. Gaudel and O.R. Coopercoordinated by the International Global Atmospheric Chemistry Project and available on 491 http://www.igacproject.org/activities/TOAR - Gaudel et al., in review and submitted to Elementa; 492 and references therein) for the reasons discussed in Section 1 and they are not 493 reproduced/explained by model simulations (e.g. Jonson et al., 2006; Cooper et al., 2010; Logan 494 495 et al., 2012; Wilson et al., 2012; Hess et al., 2013; and references therein). As a result, comparing/reconciling the adjusted trends with independent measurements, even on a qualitative 496 497 basis, remains difficult. Nevertheless, several of the statistically significant features observed in Fig.5 show, interestingly, qualitative consistency with respect to recent published findings: 498

499

- The S.H. tropical region extending from the Amazon to tropical eastern Indian Ocean 500 seems to indicate a general increase with, for example, a DJF trend of $\sim 0.23\pm0.18$ DU/yr 501 (i.e. 2.09±1.70 DU over the IASI measurement period), despite the large IAV in the MLT 502 503 which characterizes the tropics and which likely explains the high frequency of nonsignificant trends. Enhanced O_3 levels over that region have already been analysed for 504 505 previous periods (e.g. Logan et al., 1985, 1986; Fishman et al., 1991; Moxim et al., 2000; Thompson et al., 2000, 2007; Sauvage et al., 2006, 2007; Heue et al., 2016; Ebojie et al., 506 507 2016; Archibald et al., 2017; Leventidou et al., in review). For instance, the large O₃ enhancement of ~0.36±0.25 DU/yr (i.e. 3.3±2.3 DU over the whole IASI period) stretching 508 from southern Africa to Australia over the north-east of Madagascar during the austral 509 winter-spring likely originates from large IAV in the subtropical jet-related stratosphere-510 troposphere exchanges which have been found to primarily contribute to the tropospheric 511 O₃ trends over that region (Liu et al., 2016; 2017). Nevertheless, this finding should be 512 mitigated by the fact that the trend value in the S.H. tropics is of the same magnitude as the 513 514 *RMSE* of the regression residuals (~2-4.5 DU; see Fig.1).

515On the contrary, the tropical Pacific region exhibits significant negative trends that are516similar to those reported from UV sounders in Ebojie et al. (2016) and in Leventidou et al.517(in review) over previous periods, while Heue et al. (2016) mainly reports significant518positive trend over that region.

The trends over the South-East Asia are mostly non-significant and vary by season. In 520 spring-summer, some grid cells in India, in mainland China and eastwards downwind 521 China exhibit significant positive trends reaching ~0.45 DU/yr (i.e. ~4.2 DU over the IASI 522 measurement period). This tends to indicate that the tropospheric O_3 increases which have 523 been shown to mainly result from a strong positive trend in the Asian emissions over the 524 past decades (e.g. Zhao et al., 2013; Cooper et al., 2014; Zhang et al., 2016; Cohen et al., 525 2017; Tarasick et al., 2017; and references therein) but also from a substantial change in 526 the stratospheric contribution (Verstraeten et al., 2015) persists through 2008-2017 despite 527 the recent decrease in O₃ precursor emissions recorded in China after 2011 (e.g. Duncan et 528 529 al., 2016; Krotkov et al., 2016; Miyazaki et al., 2017; Van der A et al. 2017). This would 530 indicate that this decrease is probably too recent/weak to recover the 2008 O_3 levels over 531 the entire region. Significant positive trends over South-East Asia have also been reported from UV sounders over previous periods (e.g. Ebojie et al., 2016). Note, however, that this 532 finding has to be taken carefully given the large model residuals (*RMSE* of \sim 2-4 DU; cfr 533 Section 3, Fig.1) over that region. Finally, the large uncertainty in trend estimates over the 534 South-East Asia might reflect the large IAV in the biomass-burning emissions and 535 lightning NO_x sources, in addition to the recent changes in emissions. 536

The mid- and high latitudes of the S.H. show clear patterns of negative trends, all over the 538 year and in a more pronounced manner during winter-spring, with larger amplitudes than 539 those of the *RMSE* values ($\sim 0.33 \pm 0.14$ DU/vr on average in the 35°S-65°S band; i.e. a 540 trend amplitude of $\sim |3.1| \pm 1.3$ DU over the studied period vs a RMSE value of ~ 2.5 DU). 541 542 These significant negative trends in the S.H. are hard to explain but, considering the stratospheric contribution into the tropospheric columns (natural and artificial due to the 543 544 limited IASI vertical sensitivity) in the mid-high latitudes of the S.H. (~40-60%; see supplementary materials in Wespes et al., 2016) and the negative significant trends 545 previously reported from IASI in the UTLS/low stratosphere in the 30°S-50°S band, they 546 547 could be in line with those derived by Zeng et al. (2017) in the UTLS for a clean rural site of the S.H. (Lauder, New Zealand), which mainly originate from increasing troppause 548 549 height and O_3 depleting substances. Significant negative change in tropospheric O_3 over 550 these regions were also reported in Ebojie et al. (2016).

537

551

In the N.H., a band-like pattern of negative trends is observed in the 40°N-75°N latitudes 552 covering Europe and North America, especially during summer. Averaged annual trend of 553 -0.31±0.17 DU/yr and summer trend of -0.47±0.22 DU/yr (i.e. -2.87±1.57 DU and -554 4.36±2.02 DU, respectively, from January 2008 to May 2017) are estimated in that 555 latitudinal band. These trend values are significantly larger than the RMSE of the MLR 556 557 model (<3.5 DU in JJA; cfr Section 3, Fig.1). Interestingly, both the annual and summer 558 trends are amplified relative to the ones calculated in the mid-latitudes of the N.H. over the 2008-2013 period of IASI measurements (-0.19±0.05 DU/yr and -0.30±0.10 DU/yr for the 559 14

annual and the summer trends, respectively, calculated in the 30°N-50°N band; see Wespes 560 et al. (2016)). This finding is line with previous studies which point out a possible leveling 561 off of tropospheric O₃ in summer due to the decline of anthropogenic O₃ precursor 562 emissions observed since 2010-2011 in North America, in Western Europe and also in 563 some regions of China (e.g. Cooper et al., 2010; 2012; Logan et al., 2012; Parrish et al., 564 2012; Oltmans et al., 2013; Simon et al., 2015; Ebojie et al., 2016; Archibald et al., 2017; 565 Miyazaki et al., 2017). It even goes a step further by suggesting a possible decrease in the 566 tropospheric O₃ levels. Archibald et al. (2017) recently reported a net decrease of ~5% in 567 the global anthropogenic NO_x emissions in the 30°N-90°N latitude band, which is 568 consistent with the annual significant negative trend of -0.27±0.15 DU/yr for O₃ estimated 569 from IASI in that band. We should also note that, even if these latitudes are characterized 570 by the lowest stratospheric contribution (~30-45%; see supplementary materials in Wespes 571 et al., 2016), it might partly mask/attenuate the variability in the tropospheric O_3 levels. 572

574 575

4.2 Expected year for trend detection

573

576 In this section, we further verify that it is indeed possible to infer, from the studied IASI period, 577 the significant negative trend derived in the 40° N-75°N band in summer (~|0.5| DU/yr on average, 578 see Section 4.1) by determining the expected year from which such a trend amplitude would be 579 detectable at a global scale. This is achieved by estimating the minimum duration (with probability 580 0.90) of the IASI O₃ measurements that would be required to detect a trend of a specified 581 magnitude, and its 95% confidence level, following the formalism developed in Tiao et al. (1990) 582 and in Weatherhead et al. (1998):

$$N^* \approx \left[\frac{3.3 \cdot \sigma_{\varepsilon}}{\left|\tau_{yr}\right|} \cdot \sqrt{\frac{1+\Phi}{1-\Phi}}\right]^{2/3} \qquad \text{Eq (1)}$$

584

$$CL_{N^*} = [N^* \cdot e^{-B}; N^* \cdot e^{+B}]$$
 Eq (2)

Where N^* is the number of the required years, σ_{ϵ} is the standard deviation of the autoregressive 585 noise residual \mathcal{E}_t , τ_{vr} is the magnitude of the trend per year, Φ is the lag-1 autocorrelation of the 586 noise. The magnitude of the variation and of the autocorrelation in the noise residuals are taken 587 into account for a better precision on the trend estimate. Given that large variance (σ_{e}^{2}) and large 588 positive autocorrelation Φ of the noise induce small signal-to-noise ratio and long trend-like 589 segments in the dataset, respectively, these two parameters increase the number of years that would 590 be required for detecting a specified trend. CL_{M^*} is the 95% confidence limits which is not 591 symmetric around N^* and depends on B, an estimated uncertainty factor calculated as 592

593 $\frac{4}{3\sqrt{D}}\sqrt{\frac{1+\Phi}{1-\Phi}}$, with *D* the number of days in the IASI datasets, which accounts for the uncertainty

in Φ (the uncertainty in σ_{ε} being negligible given that only a few years of data are needed to 594 estimate it; cfr Weatherhead et al. (1998)). As a result, based on the available IASI-A and proxies 595 datasets and assuming that the MLR model used in this study is accurate, we estimate, in Fig. 6 (a) 596 and (b), the expected year when an O₃ trend amplitude of |5| DU per decade (i.e. $\tau_{yr} = |0.5|$ DU/yr 597 which corresponds to the averaged absolute value of the fitted negative trends in the N.H. summer; 598 599 see Fig.5b) is detectable, and its associated maximal confidence limit, respectively. The results in Fig. 6 clearly demonstrates the possibility to infer, from the available IASI dataset, such significant 600 trends in the mid- and high latitudes of the N.H. in summer and fall (trend detectable from ~2016-601 2017 with an uncertainty of ~6-9 months; cfr Fig.6b). On the contrary, the tropical regions and the 602 N.H. in winter-spring would require additional ~ 6 months to 6 years of measurements to detect 603 an amplitude of |0.5| DU/yr (trend significant only from ~ 2017 – 2023 or after depending on the 604 location and the season). Note also that the strongest negative trends (up to -0.85 DU/yr, i.e. τ_{yr} = 605 [0.85] DU/yr, see Fig.5b) observed in specific regions of the N.H. mid-latitudes would only require 606 ~6 years of IASI measurements for being detected. The mid- and high latitudes of the S.H. would 607 require the shortest period of IASI measurement for detecting a specified trend, with only ~7 years 608 \pm 6 months of IASI measurements to detect a |0.5| DU/yr trend (trend detectable from ~2015). That 609 band-like pattern in the S.H. corresponds to the region with the weakest IAV and contribution from 610 large-scale dynamical modes of variability in the IASI MLT columns (see Section 3, Fig.1 and 2), 611 which translates into small σ_{ϵ}^2 and Φ . Note however that an additional few months of IASI data 612 are required to confirm the smaller negative trend of ~-0.35 DU/yr measured on average in the 613 S.H. (see Fig.5; a period ~9 years ± 6 months being necessary to detect a trend amplitude of |3.5|614 DU/dec). Given that large σ_{ε} means large noise residual in the IASI data, the regions of short or 615 long required measurement period coincide, as expected, well with the small or high RMSE values 616 of the regression residuals (see Section 3, Fig. 1). 617

618

619 The regions of the longest measurement periods required in the tropics for a trend detection (up to 620 ~16 years of IASI data) correspond to known patterns of widespread high O₃: (a) above intense biomass burnings in Amazonia and eastwards across tropical Atlantic (Logan et al., 1986; Fishman 621 et al., 1991; Moxim et al., 2000; Thompson et al., 2000, 2007; Sauvage et al., 2007), (b) eastwards 622 Africa across the South Indian Ocean which is subject to large variations in the stratospheric 623 influences during the winter-spring austral period (JJA-SON) (Liu et al., 2016; 2017), (c) 624 Eastwards Africa across the North Indian Ocean to India likely due to large lightning NO_x 625 emissions above central Africa during the wet season associated with the northeastward jet 626 627 conducting a so-called "O₃ river" (Tocquer et al., 2015) and (d) above regions of positive ENSO 628 "chemical" effect in Equatorial Asia/Australia and eastwards above northern and southern tropical regions (Wespes et al., 2016) explained by reduced rainfalls and biomass fires during El Niño 629

conditions (e.g. Worden et al., 2013). In fact, most of these patterns (a, b and d) are closely 630 connected with strong El- Niño events which extend the duration of the dry season and cause 631 severe droughts, producing intense biomass burning emissions, for instance, over South America 632 (e.g. Chen et al., 2011; Lewis et al., 2011) and South Asia/Australia (e.g. Oman et al., 2013; Valks 633 et al., 2014; Ziemke et al., 2015), and which alter the tropospheric circulation by increasing the 634 transport of stratospheric O₃ into the troposphere (e.g. Voulgarakis et al., 2011; Neu et al., 2014) 635 and the transport of biomass burning air masses to the Indian Ocean (Zhang et al., 2012). In 636 637 summary, these large-scale indirect ENSO-related variations in tropospheric O_3 and the lightning NO_x impact on O₃, which are not accounted for in the MLR by specific representative proxies, are 638 misrepresented in the regression models. They induce large noise residuals, i.e. large σ_{ε} , and, 639 hence, extends the time period needed to detect a trend of any given magnitude. 640

641

Figure 6, finally, suggests that a long duration is also required, especially in summer, above and east of China to quantify the anthropogenic impact on the local changes in the MLT: additional 3 ± 1.5 years or 5 ± 1.5 years for a given |5| or |3.5| DU/dec trend are respectively calculated. This result could be explained by large perturbations in the MLT columns induced by recent decreases after decades of almost constant increases in surface emissions in China (e.g. Cohen et al., 2017; Miyazaki et al., 2017).

- 648
- 649 4.3 Multi-linear vs single linear model
- 650

651 Even if MLR have already been used for extracting trends in stratospheric and total O₃ columns 652 (e.g. Mäder et al., 2007; Frossard et al., 2013; Rieder et al., 2013; Knibbe et al., 2014), single linear 653 regressions (SLR) without discriminating the natural (chemical and dynamical) factors describing 654 the O_3 variability are still commonly used (e.g. Cooper et al., 2014; the TOAR-climate report – 655 Gaudel et al., in review Tropospheric Ozone Assessment Report: Present day distribution and trends of tropospheric ozone relevant to climate and global atmospheric chemistry model 656 evaluation, Lead Authors: A. Gaudel and O.R. Cooper submitted to Elementa.; and references 657 therein). They are, however, suspected to contribute to trend biases retrieved between independent 658 measurements. In addition to the time-varying instrumental biases, trend biases can also be related 659 660 to differences in the spatial and the temporal samplings (e.g. Doughty et al., 2011; Saunois et al., 661 2012; Lin et al., 2015), in the measurement period, in the upper boundary of the O₃ columns, in the algorithm and in the vertical sensitivity of the measurements. The latter artificially alters the 662 characteristics of the sounded layer by contaminations from the above and the below layers leading 663 to a mixing of the trend but also of the natural characteristics originating from these different layers 664 (e.g. troposphere and stratosphere). The differences in the studied period, in the tropopause 665 definition and in the spatio-temporal sampling might also imply differences in the natural influence 666 on the measured O₃ variations. While the impact of the natural contribution is taken into account 667 668 in the MLR model, it might introduce an additional bias in the trend determined from SLR, making

further challenging to compare trends estimated from a series of inhomogeneous independent 669 670 measurements. Substantial effort in homogenizing independent tropospheric O₃ column (TOCs) 671 datasets have been performed in the TOAR-climate assessment report (Gaudel et al., in reviewsubmitted to Elementa), but large SLR trend biases remain between the TOAR datasets, in 672 673 particular, between the satellite datasets where the lack of homogeneity in terms of troppause 674 calculation (same tropopause definition but different temperature profiles are used), of instrument 675 vertical sensitivities and of spatial sampling has been specifically pointed as possible causes for 676 the trend divergence.

677

678 If **R**reconciling the trend biases between the datasets by applying the vertical sensitivity of each measurement type to a common platform, as proposed in the TOAR-climate assessment report is 679 beyond the scope of this study, and if, at this stage, there is no consensus in determining 680 tropospheric O₃ trends, but the improvement in using a MLR instead of a SLR model for 681 determining more accurate/realistic trends is explored here by comparing the seasonal distributions 682 of the trends estimated from MLR (see Fig. 5 (b) in Section 4.1.) and from SLR (presented in 683 Fig.7). Note that the constant term in the SLR is split into two components (covering the periods 684 before and after the September 2010 "jump") to take account of the observed "jump" (see Section 685 2). The highest differences in the fitted trends derived from the two methods are found in the 686 tropics and in some regions of the mid-latitudes of the N.H. They likely result from overlaps 687 between the LT term and other covariates. For instance, the regions with high significant SLR 688 trends (~0.3-0.6 DU/yr over the tropical western and middle Pacific) during the period extending 689 from September to May match the regions with strong El Niño/Southern Oscillation influence (cfr 690 691 Fig. 8 and 12 in Wespes et al., 2016). On the contrary, the MLR model lends generally weak significant negative or non-significant trends in the Pacific Niño region during that period and it 692 693 would even need additional ~3 to 4 years of IASI measurements for detecting the fitted SLR trends 694 (see Section above). The effect of ENSO in overestimating the fitted SLR trend is further illustrated on Fig. 8 which represents the time series of O₃ observed by IASI and adjusted by the annual MLR 695 696 model (top row) along with the deseasonalized times series (middle row) and the fitted SLR and MLR trends (bottom row). The fitted signal of the ENSO proxy from the MLR regression 697 (calculated as $x_i X_{norm,i}$ following Wespes et al. (2017)) is also represented (bottom row). That 698 699 example clearly shows that the ENSO influence is considerably compensated by the adjustment of 700 the linear trend in the SLR regression (annual trend of -0.48±0.06 DU/yr from SLR vs -0.23±0.16 DU/yr from MLR for that example). Finally, differences between the SLR and the MLR models 701 are also observed in the region with strong positive NAO influence over the Icelandic/Arctic region 702 during MAM (see Wespes et al. (2016) for a description of the NAO-related O₃ changes). On the 703 contrary, the sub-tropical S.H. exhibit similar seasonal patterns of negative trends from both the 704 SLR and the MLR. It results from the weak natural IAV and contributions in tropospheric O₃ above 705 706 that region (see Section 3, Fig. 1 and 2), which, hence, limits the compensation effects. 707

708 In summary, despite the fact that considering a long period of measurements is usually 709 recommended in SLR study to overcome the dynamical cycles and, hence, to help in 710 discriminating their influences from trends, we show that, considering that some dynamics have 711 irregular or no particular periodicity (e.g. NAO, ENSO), it is not accurate enough. Furthermore, 712 accurate satellite measurements of tropospheric O_3 at a global scale are quite recent, limiting the 713 period of available and comparable datasets (e.g. Payne et al., 2017). As a consequence, we support 714 here that using a reliable multivariate regression model based on geophysical parameters and adapted for each specific sounded layer is a robust method for differentiating a "true" trend from 715 any other sources of variability and, hence, that it should help in resolving trend differences 716 717 between independent datasets.

719 **4.4 Continental influence**

720

718

721 In this section, we use the capabilities of IASI to simultaneously measure O_3 and CO in order (1) 722 to differentiate tropospheric and stratospheric air masses, (2) to identify the regions influenced by the continental export/intercontinental transport of O₃ pollution and (3) to evaluate that continental 723 influence on tropospheric O₃ trends as observed by IASI. Similar tracer correlations between CO 724 and O₃ have already been used to give insight into the photochemical O₃ enhancement in air 725 pollution transport (e.g. Parrish et al., 1993; Bertschi et al., 2005). However, there are only a few 726 studies using global satellite data for this purpose (Zhang et al., 2006; Voulgarakis et al., 2010; 727 728 Kim et al., 2013) and the analysis of the O₃-CO relationship for better understanding the origin of 729 O₃ trends in the troposphere has, to the best of our knowledge, never been explored.

730

Fig.9a and 9b show the seasonal patterns of the O_3 -CO correlations (referred as R_{O3-CO}) and of the 731 dO_3/dCO regression slopes calculated in the troposphere (from the surface to 300 hPa) over the 732 733 studied IASI period (January 2008 – May 2017). The dO₃/dCO regression slopes, which represent 734 the so-called O_3 -CO enhancement ratio, is used to evaluate the photochemical O_3 production in continental outflow regions. The R_{O3-CO} and the dO_3/dCO distributions are similar and clearly 735 show regional and seasonal differences in the strength of the O₃-CO relationships. The patterns of 736 737 positive and negative correlations allows to discriminate the outflow regions characterized by 738 photochemical O₃ production from precursors (including CO) or CO destruction (both identified by positive R_{03-C0}) from the regions characterized by O₃ loss (chemical destruction or surface 739 deposition) or by strong stratospheric contaminations (both identified by negative R_{03-CO}). 740 Negative R_{O3-CO} and dO_3/dCO are measured at high latitudes of both hemispheres all over the year, 741 but more specifically at high latitudes of the S.H. in summer-fall (with R_{O3-CO} <-0.25 on averages 742 in DJF and MMA). Given that high latitudes experience more O₃ destruction than the low latitudes 743 due to a lack of sunlight, the negative correlations for the high latitude regions might also reflect 744 745 air masses originating from/characterizing the stratosphere due to natural intrusion or to artificial 746 mixing with the troposphere introduced by the limited vertical sensitivity of IASI in the highest latitudes (stratospheric contribution varying between ~40% and 65%; see supplementary materials 747 19 in Wespes et al., 2016). These processes are likely at the origin of the band-like pattern of negative trends in the S.H. discussed in Sections 3 and 4.1. Negative R_{O3-CO} and dO_3/dCO are also found above the Caribbean, the Arabic Peninsula and the North Indian Ocean in JJA/SON and the South Atlantic in DJF. They are in line with Kim et al. (2013) and they likely reflect the influence of lightning NO_x which produce O₃ but also OH oxidizing CO (e.g. Sauvage et al., 2007; Labrador et al., 2004).

754

755 Strong positive correlations are identified in both R_{O3-CO} and dO_3/dCO patterns over the tropical regions and for mid-latitudes of both Hemispheres during the peak of photochemistry in summer. 756 757 Maxima ($R_{O3-CO} > 0.8$ and $dO_3/dCO > 0.5$) are detected in continental pollution outflow regions in the N.H., especially downwind South-East Asia and over the South Africa/Amazonia/South 758 Atlantic region. These O₃-CO correlation patterns from IASI are fully consistent with those 759 measured by TES (Zhang et al., 2006; 2008; Voulgarakis et al., 2010) and by OMI/AIRS (Kim et 760 al., 2013), which have been interpreted with global CTM's as originating from Asian pollution 761 influence and from combustion sources including biomass burning, respectively. The high positive 762 R_{03-CO} found in JJA at mid-latitudes of the N.H. are detected in a lesser extent in DJF reflecting 763 the decreasing photochemistry. It is also worth pointing out in Fig. 9 the clear hemispheric 764 differences in the R_{O3-CO} and dO_3/dCO values at mid-high latitudes and, more particularly, the shift 765 of positive R_{O3-CO} and dO_3/dCO towards higher latitudes of the N.H. during summer (e.g. R_{O3-CO} = 766 0.24 in summer vs 0.038 in spring in the 35°N-55°N band). As a consequence, these results suggest 767 that the band-like pattern of negative trends measured by IASI in summer might substantially 768 reflect the continental pollution influence and, hence, that it could result from the decline of 769 770 anthropogenic O₃ precursor emissions. Nevertheless, interpreting O₃-CO correlations in the free 771 troposphere, especially in photochemically aged pollution plumes far from the emission sources 772 towards the highest latitudes, remains complicated by the mixing of the continental combustion 773 outflow with stratospheric air masses, in addition to the background dynamic and photochemistry (e.g. Liang et al., 2007). 774

775

776 Finally, we also provide in Fig.9c the seasonal patterns of O₃-CO covariances (COV_{O3-CO}). They confirm the band-like pattern of the weak natural variation captured in the S.H. mid-latitudes (see 777 778 Sections 3 and 4.1) and help identifying the region downwind East China, the northern midlatitudes outflow region and the South tropical region as the ones with the highest pollution 779 780 variability, in addition to the strongest O₃-CO correlations. To conclude, the particularly strong positive O₃-CO relationship in terms of R_{O3-CO}, dO₃/dCO and COV_{O3-CO} measured over and 781 downwind North-East India/East China in summer in comparison with the ones measured 782 downwind East US and over Europe indicate that South-East Asia experiences the most of the 783 intense pollution episodes of the N.H. with the largest O₃-CO variability (COV_{03-CO} > 40×10^{33} 784 mol².cm⁻⁴) and the largest O₃ enhancement ($dO_3/dCO > 0.5$) over the last decade. The strong O₃-785 CO relationship in that region is associated with the significant increase that is detected in the IASI 786

 O_3 levels downwind East of Asia (see Section 4.1) despite the net decrease in O_3 precursor emissions recorded in China after 2011 (e.g. Cohen et al., 2017; Miyazaki et al., 2017).

789

790 **5** Conclusions

791

792 In this study, we have explored, for the first time, the possibility to infer significant trends in tropospheric O₃ from the first ~10 years (January 2008 – May 2017) of IASI daily measurements 793 at a global scale. To this end, MLR analyses have been performed by applying a multivariate 794 regression model (annual and seasonal formulations), including a linear trend term in addition to 795 796 chemical and dynamical proxies, on gridded mean tropospheric ozone time series. This work follows on the analysis of the main dynamical drivers of O₃ variations measured by IASI, which 797 was recently published in Wespes et al. (2017). We have first verified the performance of the MLR 798 models in explaining the variations in daily time series over the whole studied period. In particular, 799 we have shown that the model reproduces a large part of the O_3 variations (>70%) with small 800 residuals errors (*RMSE* of $\sim 10\%$) at northern latitudes in summer. We have then performed O₃ 801 trend sensitivity tests to verify the possibility to capture trend characteristics independently from 802 natural variations. Despite the weak contribution of trends to the total variation in the MLT O₃ 803 columns at a global scale, the results demonstrate the possibility to discriminate significant trends 804 from the explanatory variables, especially in summer at mid-high latitudes of the N.H. (~45°N-805 70°N) where the contribution and the sensitivity of trends are the largest (contribution of ~30-50% 806 and a ~10% increase in the RMSE excluding the LT in the model). We then focused on the 807 interpretation of the global trend estimates. We have found an interesting significant positive 808 809 trends in the S.H. tropical region extending from the Amazon to the tropical eastern Indian Ocean and over South-East Asia in spring-summer which should however be carefully considered given 810 811 the high RMSE of the regression residuals in these regions. The MLR analysis reveals a band-like 812 pattern of high significant negative trends in the N.H. mid-high latitudes in summer (-0.47±0.16 DU/yr on average in the 45°N-70°N band). The statistical significance of such trend estimates is 813 further verified by estimating, based on the autocorrelation and on the variance of the noise 814 residuals, the minimum number of years of IASI measurements that are required to detect a trend 815 of a [5] DU/dec magnitude. The results clearly demonstrate the possibility to determine such a trend 816 amplitude from the available IASI dataset and the used MLR model at northern mid-high latitudes 817 in summer, while much larger measurement periods are necessary elsewhere. In particular, the 818 regions with the longest required period, in the tropics, highlight a series of known processes that 819 are closely related to the El-Niño dynamic, which underlies the lack of associated 820 parameterizations in the MLR model. The importance of using reliable MLR models in 821 understanding large-scale O₃ variations and in determining trends is further explored by comparing 822 the trends inferred from MLR and from SLR, the latter being still commonly used by the 823 international community. The comparison has clearly highlighted the gain of MLR in attributing 824 the trend-like segments in natural variations, such as ENSO, to the right processes and, hence, in 825 826 avoiding misinterpretation of "apparent" trends in the measurement datasets. Nevertheless, it is 21

worth noting that there could be a possible impact of the sampling (because of the cloud and quality 827 828 filters applied) and of the jump in September 2010 that has been identified in the IASI dataset (see Section 2), in both MLR and SLR trends. Finally, by exploiting the simultaneous and vertically-829 resolved O₃ and CO measurements from IASI, we have provided and used the O₃-CO correlations 830 in the troposphere to help determining the origins/characteristics of patterns of negative or positive 831 trends. The distributions have allowed us to identify, in particular, strong positive O₃-CO 832 correlations, regression slopes and covariance in the N.H. mid-latitudes and northward during 833 summer, which suggests a continental pollution influence in the N.H. band-like pattern of high 834 significant negative trends recorded by IASI and, hence, a direct effect of the policy measures 835 taken to reduce emissions of O₃ precursor species. 836

837

This study supports overall the importance of using (1) high density and long term homogenized 838 satellite records, such as those provided by IASI, and (2) complex models with predictor functions 839 that describe the O₃-regressors dependencies for a more accurate determination of trends in 840 tropospheric O₃ - as required by the scientific community, e.g. in the Intergovernmental Panel on 841 Climate Change (IPCC, 2013) - and for further resolving trend biases between independent 842 843 datasets (Payne et al., 2017; the TOAR-climate assessment report – Gaudel et al., in review). 844 Currently, no consensus in terms of O₃ trends in the troposphere is reached from the available 845 measurements (UV or IR satellites, O3 sondes, aircrafts, ground-based measurements,...) for several reasons (time-varying instrumental biases, differences in the methodology used for 846 847 calculating trends, in the measurement period, in the upper boundary of the O₃ columns, in the 848 retrieval algorithm, in the spatio-temporal sampling, in the vertical sensitivity of the instrument,...) 849 (Section 4.3; the TOAR-climate report – Gaudel et al., in review). However, Ddetermination, with IASI, of robust trends in tropospheric O_3 at the global scale will be achievable in the near future 850 851 by merging the homogeneous O₃ profiles from the three successive instruments onboard Metop-A 852 (2006); -B (2012) and -C (2018) platforms and from the IASI-Next Generation instrument onboard the Metop Second Generation series of satellites (Clerbaux and Crevoisier, 2013; Crevoisier et al., 853 854 2014). A long record of tropospheric O_3 measurements will be also assured by the Cross-track Infrared Sounder (CrIS) onboard the Joint Polar Satellite System series of satellites. 855

857 Acknowledgments

IASI has been developed and built under the responsibility of the Centre National d'Etudes
Spatiales (CNES, France). It is flown onboard the Metop satellites as part of the EUMETSAT
Polar System. The IASI L1 data are received through the EUMETCast near real time data
distribution service. We acknowledge the financial support from the ESA O₃-CCI, Copernicus O₃C3S and Eumetsat AC SAF projects. The research in Belgium is also funded by the Belgian State
Federal Office for Scientific, Technical and Cultural Affairs and the European Space Agency (ESA
Prodex IASI Flow and AC SAF).

865

856

867 Figure captions

868

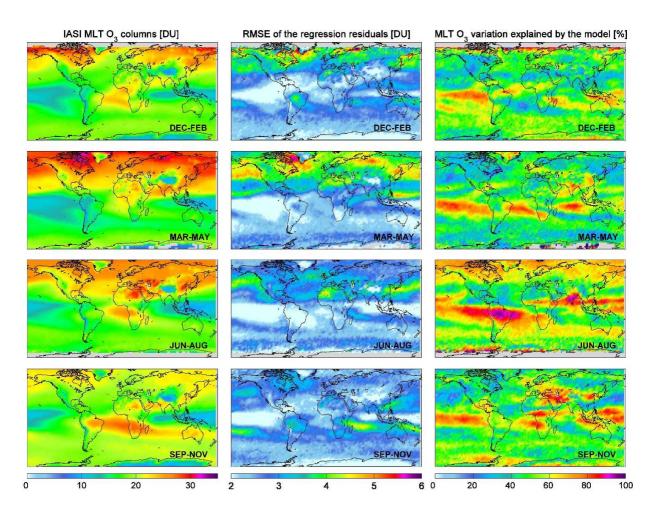


Fig.1. Seasonal distribution of O₃ tropospheric columns (in DU, integrated from ground to 300hPa) measured by IASI and averaged over January 2008 – May 2017 (left panel), of the *RMSE* of the regression fits (in DU, middle panel) and of the fraction of the variation in IASI data explained by the regression model, calculated as $[100 \times (\sigma(O_3^{Fitted_mod\,el}(t))/\sigma(O_3(t)))]$ (in %, right panel). Data are averaged over a 2.5°x2.5° grid box.

875

869

876

877

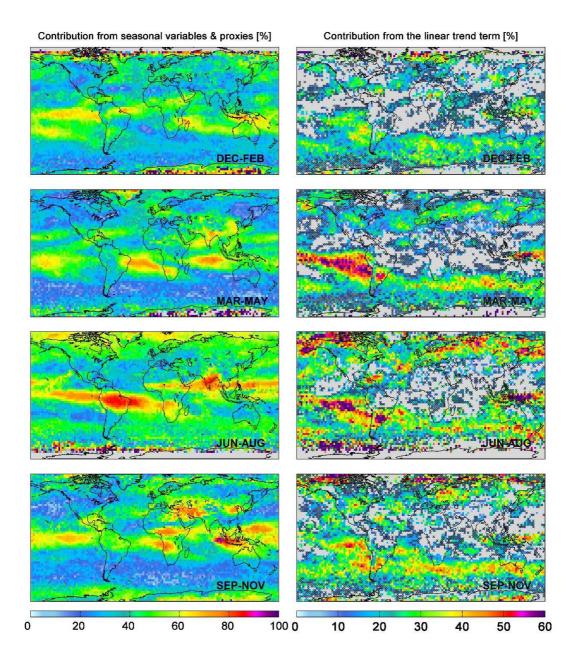
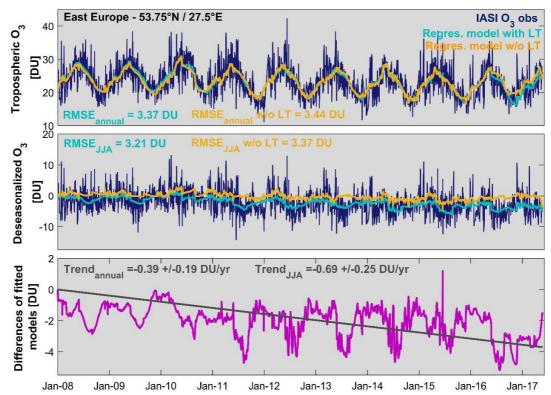
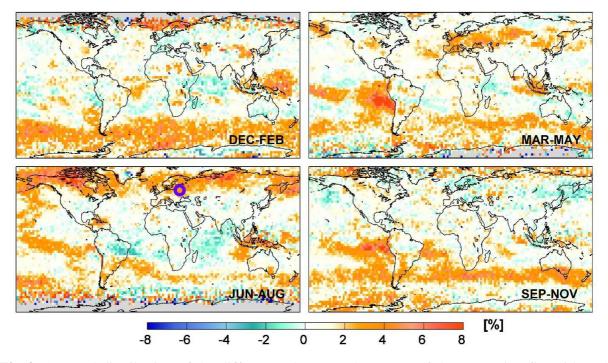


Fig.2. Seasonal distributions of the contribution from the seasonal and explanatory variables into 881 IASI **O**₃ variations estimated the 882 as $\left|100\times\sigma\left(\sum_{n=1;j=2}^{4;m} [a_n; b_n; x_j] \left[\cos(n\omega t); \sin(n\omega t); X_{norm,j}\right]\right) / \sigma(O_3(t))\right] \text{ (in \%, left panels) and of the}\right|$ 883 contribution from the linear trend calculated as $\left[100 \times \sigma(x_{j=1} \cdot trend) / \sigma(O_3(t))\right]$ (in %, right panels). 884 The grey areas and crosses refer to the non-significant grid cells in the 95% confidence limits (2σ 885 886 level). Note that the scales are different. 887



888

Fig.3. Examples of daily time series of IASI O₃ measurements (dark blue) and of the fitted seasonal regression models with (light blue) and without (orange) the linear term in the troposphere (1st row). Daily time series of the deseasonalised O₃ (observations and regression models; 2^d row) and of the difference of the fitted models with and without the linear trend term as well as the adjusted annual trend (pink and grey lines, respectively; 3^d row) (given in DU). The *RMSE* (annual and for the JJA period in DU) and the trend values (annual and for the JJA period in DU/yr) are also indicated.



898

Fig.4. Seasonal distribution of the differences between the *RMSE* of the regression fits with and without linear trend term $[(RMSE_w/o_LT - RMSE_with_LT)/RMSE_with_LT \times 100]$ (in %). The blue circles in the JJA panel refer to the case presented in Fig.3.

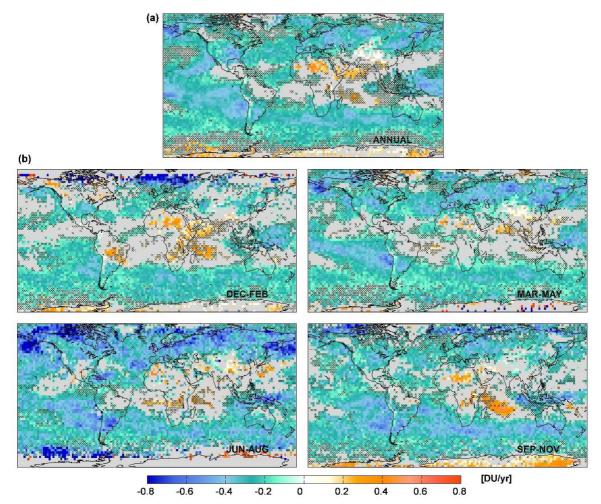
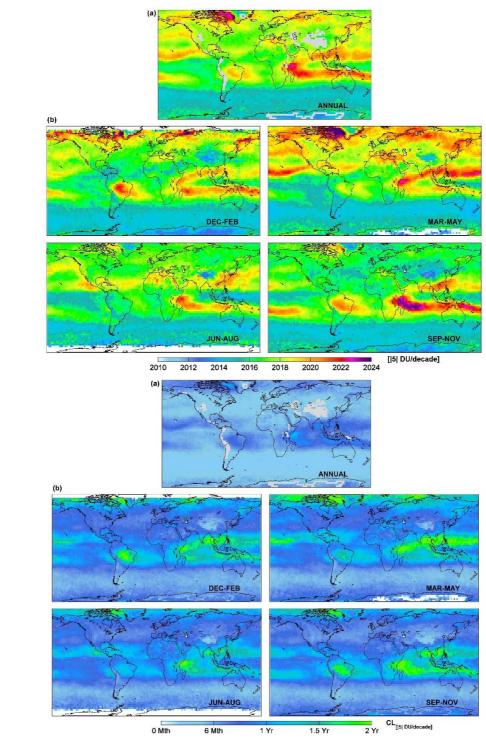
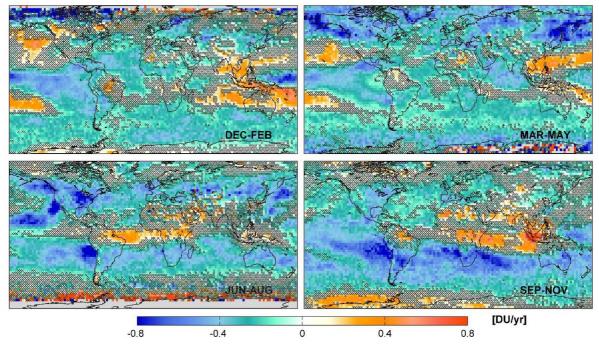


Fig.5. (a) Annual and (b) seasonal distributions of the adjusted trends in DU/yr from the multi-linear regression models. The grey areas and crosses refer to the non-significant grid cells in the 95% confidence limits (2 σ level).





- given trend of |5| DU per decade starting at the beginning of the studied period (20080101) and (b)
- 915 associated maximal confidence limits from the annual (top panel) and the seasonal (bottom panels)
- 916 regression models.



917

Fig.7. Seasonal distributions of the fitted linear term trends (given in DU/yr) derived from a single

919 linear regression model. The crosses refer to the non-significant grid cells in the 95% confidence

920 limits (2σ level).

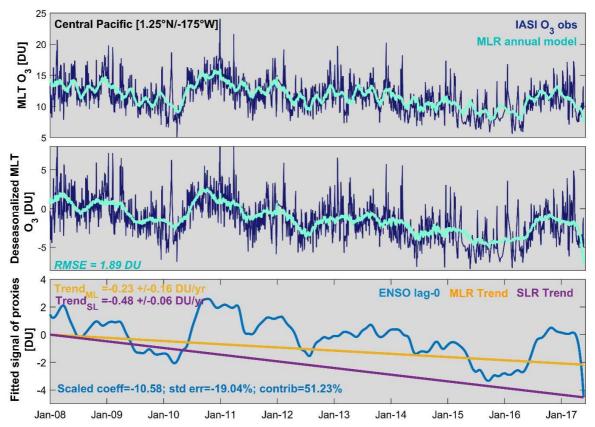


Fig.8. Daily time series of O₃ measured by IASI and adjusted by the multivariate annual regression model (top row and middle row for the deseasonalized O₃), along with the adjusted trends derived from the single and the multivariate linear regressions (SLR and MLR) and of the fitted signal of ENSO proxy (one of the main retained proxies in the multivariate regression model) calculated as $[x_j X_{norm,j}]$ (bottom row) over the equatorial central Pacific (negative ENSO "dynamical" effect) (given in DU). The *RMSE* of the multivariate regression fit and the fitted SLR and MLR trend values are also indicated.

- 930
- 931
- 932
- 933
- 934
- 935

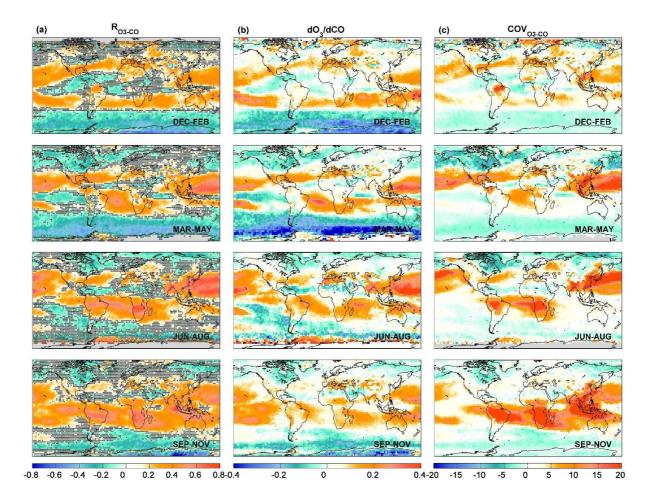


Fig.9. Global distributions of (a) the correlation coefficients (R_{O3-CO}), (b) the regression slope (dO_3/dCO in mol.cm⁻²/mol.cm⁻²) and (c) the covariances (COV_{O3-CO} in 10^{33} mol.cm⁻²×mol.cm⁻²) of daily median IASI tropospheric O_3 and CO over January 2008 – May 2017. Data are averaged over a 2.5°x2.5° grid box. Crosses in R_{O3-CO} panels (a) refer to the non-significant grid cells in the 95% confidence intervals (2σ level).

954 **References**

Anton, M., D. Loyola, C. Clerbaux, M. Lopez, J. Vilaplana, M. Banon, J. Hadji-Lazaro, P. Valks,
N. Hao, W. Zimmer, P. Coheur, D. Hurtmans, and L. Alados-Arboledas: Validation of the MetopA total ozone data from GOME-2 and IASI using reference ground-based measurements at the
Iberian peninsula, Remote Sensing of Environment, 115, 1380-1386, 2011.

960

Archibald, A., Y. Elshorbany et al.: Tropospheric Ozone Assessment Report: Critical review of
the present-day and near future tropospheric ozone budget, Elem. Sci. Anth., in-review, 2017.

Bertschi, I. T. and Jaffe, D. A.: Long-range transport of ozone, carbon monoxide, and aerosols to
the NE Pacific troposphere during the summer of 2003: Observations of smoke plumes from Asian
boreal fires, J. Geophys. Res., 110(D5), D05303, doi:10.1029/2004JD005135, 2005.

- 967
 968 Boynard, A., D. Hurtmans, M. Koukouli, et al.: Seven years of IASI ozone retrievals from FORLI:
 969 validation with independent total column and vertical profile measurements, Atmos. Meas. Tech.,
 970 9, 4327-4353, 2016.
- 971

Boynard, A., D. Hurtmans, K. Garane, F. Goutail, J. Hadji-Lazaro, M. E. Koukouli, C. Wespes,
A. Keppens, J.-P. Pommereau, A. Pazmino, D. Balis, D. Loyola, P. Valks, S. Hassinen, P.-F.
Coheur and C. Clerbaux, Validation of the IASI FORLI/Eumetsat O₃ products using satellite
(GOME-2), ground-based (Brewer-Dobson, SAOZ) and ozonesonde measurements, in
preparation for this QOS special issue.

- Chen, Y., Randerson, J. T., Morton, D. C., DeFries, R. S., Collatz, G. J., Kasibhatla, P. S., iglio,
 L., Jin, Y., and Marlier, M. E.: Forecasting Fire Season Severity in South America Using Sea
 Surface emperature Anomalies, Science, 334, 787–791, doi:10.1126/science.1209472, 2011.
- 981

Clarisse, L., Y. R'Honi ; P.-F. Coheur ; D. Hurtmans, and C. Clerbaux : Thermal infrared nadir
observations of 24 atmospheric gases. Geophysical Research Letters, 38, L10802, 2011.

- 984
 985 Clerbaux, C., A. Boynard, L. Clarisse, M. George, J. Hadji-Lazaro, H. Herbin, D. Hurtmans, M.
 986 Pommier, A. Razavi, S. Turquety, C. Wespes, and P.-F. Coheur: Monitoring of atmospheric
 987 composition using the thermal infrared IASI/MetOp sounder, Atmos. Chem. Phys., 9, 6041-6054,
 988 2009.
- 988 989

990 Clerbaux C. and C. Crevoisier: New Directions: Infrared remote sensing of the troposphere from991 satellite: Less, but better, Atmospheric Environment, 72, 24-26, 2013.

- 992
- Crevoisier, C., Clerbaux, C., Guidard, V., Phulpin, T., Armante, R., Barret, B., Camy-Peyret, C.,
 Chaboureau, J.-P., Coheur, P.-F., Crépeau, L., Dufour, G., Labonnote, L., Lavanant, L., HadjiLazaro, J., Herbin, H., Jacquinet-Husson, N., Payan, S., Péquignot, E., Pierangelo, C., Sellitto, P.,
- and Stubenrauch, C.: Towards IASI-New Generation (IASI-NG): impact of improved spectral
- 997 resolution and radiometric noise on the retrieval of thermodynamic, chemistry and climate
- 998 variables, Atmos. Meas. Tech., 7, 4367-4385, 2014.

- 999
 1000 Cohen, Y., et al.: Climatology and long-term evolution of ozone and carbon monoxide in the UTLS
 1001 at northern mid-latitudes, as seen by IAGOS from 1995-2013, ACPD, in-review, 2017.
 1002
- 1003 Cooper, O., D. Parrish, A. Stohl, M. Trainer, P. Nédélec, V. Thouret, J.-P. Cammas, S. Oltmans,
 1004 B. Johnson and D. Tarasick: Increasing springtime ozone mixing ratios in the free troposphere
 1005 over western North America, Nature, 463, 344–348, doi:10.1038/nature08708, 2010.
- 1006
- 1007 Cooper, O. R., R.-S. Gao, D. Tarasick, T. Leblanc, and C. Sweeney: Long-term ozone trends at
 1008 rural ozone monitoring sites across the United States, 1990–2010, *J. Geophys. Res.*, *117*, D22307,
 1009 doi:10.1029/2012JD018261, 2012.
 1010
- 1011 Cooper, O. R., D. D. Parrish, J. Ziemke, N. V. Balashov, M. Cupeiro, I. E. Galbally, S. Gilge, L.
 1012 Horowitz, N. R. Jensen, J.-F. Lamarque, V. Naik, S. J. Oltmans, J. Schwab, D. T. Shindell, A. M.
 1013 Thompson, V. Thouret, Y. Wang, R. M. Zbinden: Global distribution and trends of tropospheric
 1014 ozone: An observation-based review, Elementa: Science of the Anthropocene, 2, 000029,
 1015 doi:10.12952/journal.elementa.000029, 2014.
- 1017 Doughty, D. C., Thompson, A. M., Schoeberl, M. R., Stajner, I., Wargan, K., and Hui, W. C. J.: 1018 An intercomparison of tropospheric ozone retrievals derived from two Aura instruments and North America in 2006, 1019 measurements in western J. Geophys. Res., 116. 1020 D06303,doi:10.1029/2010JD014703, 2011.
- 1021

- Dufour, G., M. Eremenko, A. Griesfeller, B. Barret, E. LeFlochmoen, C. Clerbaux, J. HadjiLazaro, P.-F. Coheur, and D. Hurtmans: Validation of three different scientific ozone products
 retrieved from IASI spectra using ozonesondes, Atmos. Meas. Tech., 5, 611-630, 2012.
- Duncan, B. N., Lamsal, L. N., Thompson, A. M., Yoshida, Y., Lu, Z., Streets, D. G., Hurwitz, M.
 M., and Pickering, K. E.: A space-based, high-resolution view of notable changes in urban
 NOx pollution around the world (2004-2005), J. Geophys. Res. 121, 976–96, 2016.
- Ebojie, F., Burrows, J. P., Gebhardt, C., Ladstätter-Weißenmayer, A., von Savigny, C., Rozanov,
 A., Weber, M., and Bovensmann, H.: Global tropospheric ozone variations from 2003 to 2011 as
 seen by SCIAMACHY, Atmos. Chem. Phys., 16, 417-436, doi:10.5194/acp-16-417-2016, 2016.
- 1033
- Fishman, J., K. Fakhruzzaman, B. Cros and D. Nganda: Identification of widespread pollution in
 the southern-hemisphere deduced from satellite analyses, Science, 252, 1693-1696, 1991.
- Fishman, J., Creilson, J. K., Wozniak, A. E., and Crutzen, P. J.: Interannual variability of
 stratospheric and tropospheric ozone determined from satellite measurements, J. Geophys. Res.,
 1039 110, D20306, doi:10.1029/2005JD005868, 2005.
- 1040
- Frossard, L., H.E. Rieder, M. Ribatet, J. Staehelin, J. A, Maeder, S. Di Rocco, A. C. Davison, T.
 Pete.: On the relationship between total ozone and atmospheric dynamics and chemistry at mid-

latitudes – Part 1: Statistical models and spatial fingerprints of atmospheric dynamics and
chemistry, Atmos. Chem. Phys., 13, 147–164, doi:10.5194/acp-13-147-2013, 2013.

Fusco, A. C., and J. A. Logan: Analysis of 1970–1995 trends in tropospheric ozone at northern
hemisphere midlatitudes with the GEOSCHEM model, J. Geophys. Res., 108(D15), 4449,
doi:10.1029/2002JD002742, 2003.

1049

1060

1065

1068

1073

1045

1050 Gaudel, A., O. R. Cooper, G. Ancellet, B. Barret, A. Boynard, J. P. Burrows, C. Clerbaux, P.-F. Coheur, J. Cuesta, E. Cuevas, S. Doniki, G. Dufour, F. Ebojie, G. Foret, O. Garcia, M. J. Granados-1051 1052 Muñoz, J. Hannigan, F. Hase, B. Hassler, G. Huang, D. Hurtmans, D. Jaffe, N. Jones, P. Kalabokas, B. Kerridge, S. Kulawik, B. Latter, T. Leblanc, E. Le Flochmoën, W. Lin, J. Liu, X. 1053 Liu, E. Mahieu, A. McClure-Begley, J. Neu, M. Osman, M. Palm, H. Petetin, I. Petropavlovskikh, 1054 1055 R. Querel, N. Rahpoe, A. Rozanov, M. G. Schultz, J. Schwab, R. Siddans, D. Smale, M. Steinbacher, H. Tanimoto, D. Tarasick, V. Thouret, A. M. Thompson, T. Trickl, E. Weatherhead, 1056 1057 C. Wespes, H. Worden, C. Vigouroux, X. Xu, G. Zeng, J. Ziemke: Tropospheric Ozone Assessment Report: Present-day distribution and trends of tropospheric ozone relevant to climate 1058 and global atmospheric chemistry model evaluation, submitted to Elementa. 1059

- Gazeaux, J., C. Clerbaux, M. George, J. Hadji-Lazaro, J. Kuttippurath, P.-F. Coheur, D. Hurtmans,
 T. Deshler, M. Kovilakam, P. Campbell, V. Guidard, F. Rabier, and J.-N. Thepaut:
 Intercomparison of polar ozone profiles by IASI/Metop sounder with 2010 concordiasi
 ozonesonde observations, Atmos. Meas. Tech., 5, 7923-7944, 2012.
- Hess, P.G. and R. Zbinden: Stratospheric impact on tropospheric ozone variability and trends:
 1990–200, Atmos. Chem. Phys., 13, 649–674, 2013.
- 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,
 doi:10.5194/amt-9-5037-2016, 2016.
- Hilton, F., R. Armante, T. August, et al. : Hyperspectral Earth Observation from IASI: Five Years
 of Accomplishments, Bulletin of the American Meteorological Society, vol. 93, issue 3, pp. 347370, 2012.
- Hurtmans, D., P. Coheur, C.Wespes, L. Clarisse, O. Scharf, C. Clerbaux, J. Hadji-Lazaro, M.
 George, and S. Turquety: FORLI radiative transfer and retrieval code for IASI, Journal of
 Quantitative Spectroscopy and Radiative Transfer, 113, 1391-1408, 2012.
- 1081

- Intergovernmental Panel on Climate Change, Climate Change 2013: The Physical Science Basis.
 Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel
 on Climate Change, edited by T. F. Stocker et al., pp. 164–270, Cambridge Univ. Press,
 Cambridge, U. K., and New York, 2013.
- 1086

Jonson, J. E., Simpson, D., Fagerli, H., and Solberg, S.: Can we explain the trends in European
ozone levels?, Atmos. Chem. phys., 6, 51–66, doi:10.5194/acp-6-51-2006, 2006.

1089

Keppens, A., J.-C. Lambert, J. Granville, D. Hubert, T. Verhoelst, S. Compernolle, B. Latter, B.
Kerridge, R. Siddans, A. Boynard, J. Hadji-Lazaro, C. Clerbaux, C. Wespes, D. R. Hurtmans, P.F. Coheur, J. van Peet, R. van der A, K. Garane, M. E. Koukouli, D. S. Balis, A. Delcloo, R. Kivi,
R. Stübi, S. Godin-Beekmann, M. Van Roozendael, C. Zehner: Quality assessment of the
Ozone_cci Climate Research Data Package (release 2017): 2. Ground-based validation of nadir
ozone profile data products, in preparation for this QOS special issue.

1096

Kim, P. S., D. J. Jacob, X. Liu, J. X.Warner, K. Yang, K. Chance, V. Thouret, and P. Nedelec:
Global ozone–CO correlations from OMI and AIRS: constraints on tropospheric ozone sources,
Atmos. Chem. Phys., 13, 9321–9335, 2013.

- Knibbe J. S., R. J. van der A, and A. T. J. de Laat: Spatial regression analysis on 32 years of total
 column ozone data, Atmos. Chem. Phys., 14, 8461–8482, 2014.
- 1103

1100

Krotkov, N. A., McLinden, C. A., Li, C., Lamsal, L. N., Celarier, E. A., Marchenko, S. V., Swartz,
W. H., Bucsela, E. J., Joiner, J., Duncan, B. N., Boersma, K. F., Veefkind, J. P., Levelt, P. F.,
Fioletov, V. E., Dickerson, R. R., He, H., Lu, Z., and Streets, D. G.: Aura OMI observations of
regional SO₂ and NO₂ pollution changes from 2005 to 2015, Atmos. Chem. Phys., 16, 4605-4629,
doi:10.5194/acp-16-4605-2016, 2016.

1109

1113

1117

Labrador, L. J., von Kuhlmann, R., and Lawrence, M. G.: Strong sensitivity of the global mean
OH concentration and the tropospheric oxidizing efficiency to the source of NOx from lightning,
Geophys. Res. Lett., 31, L06102, doi:10.1029/2003GL019229, 2004.

- Leventidou, E., Weber, M., Eichmann, K.-U., and Burrows, J. P.: Harmonisation and trends of 20years tropical tropospheric ozone data, Atmos. Chem. Phys. Discuss., https://doi.org/10.5194/acp2017-815, in review, 2017.
- Lewis, S. L., Brando, P. M., Phillips, O. L., van der Heijden, G. M. F., and Nepstad, D.: The 2010
 Amazon Drought, Science, 331, 554–554, doi:10.1126/science.1200807, 2011.
- Liang, Q., Jaegle, L., Hudman, R. C., Turquety, S., Jacob, D. J., Avery, M. A., Browell, E. V.,
 Sachse, G. W., Blake, D. R., Brune, W., Ren, X., Cohen, R. C., Dibb, J. E., Fried, A., Fuelberg,
 H., Porter, M., Heikes, B. G., Huey, G., Singh, H. B., andWennberg, P. O.: Summertime influence
 of Asian pollution in the free troposphere over North America, J. Geophys. Res., 112, D12S11,
 doi:10.1029/2006JD007919, 2007.
- 1126
- Lin, M., L. W. Horowitz, O. R. Cooper, D. Tarasick, S. Conley, L. T. Iraci, B. Johnson, T. Leblanc,
 I. Petropavlovskikh, and E. L. Yates: Revisiting the evidence of increasing springtime ozone
 mixing ratios in the free troposphere over western North America, Geophys. Res. Lett., 42, 8719–
 8728, doi:10.1002/2015GL0653112015.
- 1131

- Liu, X., P. K. Bhartia, K. Chance, R. J. D. Spurr, and T. P. Kurosu: Ozone profile retrievals from
 the Ozone Monitoring Instrument, Atmos. Chem. Phys., 10, 2521–2537, 2010.
- 1134
- Liu, J., Rodriguez, J. M., Thompson, A. M., Logan, J. A., Douglass, A. R., Olsen, M. A., Steenrod,
- 1136 S. D., and Posny, F.: Origins of tropospheric ozone interannual variation over Reunion: A model
- 1137 investigation, J. Geophys. Res.-Atmos., 121, 521–537, doi:10.1002/2015jd023981, 2016.
- 1138
- Liu, J., et al.: Causes of interannual variability over the southern hemispheric tropospheric ozone
 maximum, Atmos. Chem. Phys., 17, 3279–3299, 2017.
- 1141
- Logan, J. A., M. J. Prather, S. C. Wofsy, and M. B. McElroy: Tropospheric chemistry: A global
 perspective, J. Geophys. Res., 86 (NC8), 7210–7254, doi:10.1029/JC086iC08p07210, 1981.
- Logan, J. A.: Tropospheric Ozone: Seasonal behaviour, Trends, and Anthropogenic Influence, J.
 Geophys. Res., 90(D6), 10 463–10 482, 1985.
- 1147

- Logan, J. A., and V. W. J. H. Kirchhoff: Seasonal variations of tropospheric ozone at Natal, Brazil,
 J. Geophys. Res., 91(D7), 7875–7881, doi:10.1029/JD091iD07p07875, 1986.
- Logan, J. A., Staehelin, J., Megretskaia, I. A., Cammas, J.-P., Thouret, V., Claude, H., De Backer,
 H., Steinbacher, M., Scheel, H.-E., Stübi, R., Fröhlich, M., and Derwent, R.: Changes in ozone
 over Europe: Analysis of ozone measurements from sondes, regular aircraft (MOZAIC) and alpine
 surface sites, J. Geophys. Res., doi:10.1029/2011JD016952, 2012.
- Mäder, J. A., J. Staehelin, D. Brunner, W.A. Stahel, I. Wohltmann, and T. Peter: Statistical
 modelling of total ozone: Selection of appropriate explanatory variables, J. Geophys. Res., 112,
 D11108, doi:10.1029/2006JD007694, 2007.
- 1159

- Miyazaki et al.: Decadal changes in global surface NOx emission from multi-constituent satellite
 data assimilation, Atmos. Chem. Phys., 17, 807–837, 2017.
- Moxim, W. J., and H. Levy II: A model analysis of the tropical South Atlantic Ocean tropospheric
 ozone maximum: The interaction of transport and chemistry, J. Geophys. Res., 105(D13), 17393–
 17415, d oi:10.1029/2000JD900175, 2000.
- 1166
- Neu, J.L., T. Flury, G. L. Manney, M. L. Santee, N. J. Livesey and J. Worden, Tropospheric ozone
 variations governed by changes in stratospheric circulation, Nat. Geosc., 7, 340–344,
 doi:10.1038/ngeo2138, 2014.
- 1170
- Oetjen, H., Payne, V.H., Kulawik, S.S., Eldering, A., Worden, J., Edwards, D.P., Francis, G.L.,
 Worden, H.M., Clerbaux, C., Hadji-Lazaro, J., Hurtmans, D. : Extending the satellite data record
 of tropospheric ozone profiles from Aura-TES to MetOp-IASI, Atmos. Meas. Tech., 7, 4223-4236,
 doi:10.5194/amt-7-4223-2014, 2014.
- 1175

Oltmans, S.J, A. S. Lefohn; D. Shadwick, J. M. Harris, H.-E. Scheel, I. Galbally, D. W. Tarasick,
B. J. Johnson, E.G. Brunke, H. Claude, G. Zeng, S. Nichol, F.J. Schmidlin, J. Davies, E. Cuevas,

1178 A. Redondas, H. Naoe, T. Nakano, T. Kawasato: Recent Tropospheric Ozone Changes - A Pattern 1179 Dominated by Slow or No Growth, Atmospheric Environment, 67, p. 331-351,

1180 10.1016/j.atmosenv.2012.10.057, 2013.

- 1181
- Oman, L. D., Douglass, A. R., Ziemke, J. R., Rodriguez, J. M., Waugh, D. W., and Nielsen, J. E.:
 The ozone response to ENSO in Aura satellite measurements and a chemistry–climate simulation,
 J. Geophys. Res.-Atmos., 118, 965–976, 2013.
- 1185

Parrington, M., P. I. Palmer, D. K. Henze, D. W. Tarasick, E. J. Hyer, R. C. Owen, D. Helmig, C.
Clerbaux, K. W. Bowman, M. N. Deeter, E. M. Barratt, P.-F. Coheur, D. Hurtmans, Z. Jiang, M.
George, and J. R. Worden: The influence of boreal biomass burning emissions on the distribution
of tropospheric ozone over north America and the north Atlantic during 2010, Atmos. Chem.
Phys., 12, 2077-2098, doi: 10.5194/acp-12-2077-2012, 2012.

- Parrish, D.D., Holloway, J.S., Trainer, M., Murphy, P.C., Forbes, G.L., and Fehsenfeld, F.C.:
 Export of North American Ozone Pollution to the North Atlantic Ocean, Science, 259, 1436–1439,
 1993.
- 1195

1199

1191

Parrish, D.D., K.S. Law, J. Staehelin, R. Derwent, O.R. Cooper, et al.: Long-term changes in lower
tropospheric baseline ozone concentrations at northern mid-latitudes. Atmos. Chem. Phys 12:
1198 11485–11504. doi:10.5194/acp-12-11485-2012, 2012.

Payne, V. H., J. L. Neu and H. M. Worden: Satellite observations for understanding the drivers ofvariability and trends in tropospheric ozone, J. Geophys. Res., in-press, 2017.

- Pommier, M., C. Clerbaux, K. S. Law, G. Ancellet, P. Bernath, P.-F. Coheur, J. Hadji- Lazaro, D.
 Hurtmans, P. Nedelec, J.-D. Paris, F. Ravetta, T. B. Ryerson, H. Schlager, and A. J.Weinheimer:
 Analysis of IASI tropospheric O₃ data over the arctic during POLARCAT campaigns in 2008,
 - 1206 Atmos. Chem. Phys., 12, 7371-7389, doi:doi:10.5194/acp-12-7371-2012, 2012. 1207
 - Rieder, H. E., Frossard, L., Ribatet, M., Staehelin, J., Maeder, J. A., Di Rocco, S., Davison, A. C.,
 Peter, T., Weihs, P., and Holawe, F.: On the relationship between total ozone and atmospheric
 dynamics and chemistry at mid-latitudes Part 2: The effects of the El Nino/Southern Oscillation,
 volcanic eruptions and contributions of atmospheric dynamics and chemistry to long-term total
 ozone changes, Atmos. Chem. Phys., 13, 165–179, 2013.
 - 1213

Rodgers, C. D.: Inverse Methods for Atmospheric Sounding: Theory and Practice, World
Scientific, Series on Atmospheric, Oceanic and Planetary Physics, 2, Hackensack, N. J., 2000.

- 1216
- 1217 Safieddine, S., Boynard, A., Coheur, P.-F., Hurtmans, D., Pfister, G., Quennehen, B., Thomas, J.
- 1218 L., Raut, J.-C., Law, K. S., Klimont, Z., Hadji-Lazaro, J., George, M., and Clerbaux, C.:
- 1219 Summertime tropospheric ozone assessment over the Mediterranean region using the thermal

infrared IASI/MetOp sounder and the WRF-Chem model, Atmos. Chem. Phys., 14, 10119–10131,
 doi:10.5194/acp-14-10119-2014, 2014.

1222

Saunois, M., Emmons, L., Lamarque, J.-F., Tilmes, S., Wespes, C., Thouret, V., and Schultz, M.:
Impact of sampling frequency in the analysis of tropospheric ozone observations, Atmos. Chem.
Phys., 12, 6757–6773, doi:10.5194/acp-12-6757-2012, 2012.

1226

1227 Sauvage, B., Thouret, V., Thompson, A. M., Witte, J. C., Cammas, J. P., Nedelec, P., and Athier, G.: Enhanced view of the "tropical Atlantic ozone paradox" and "zonal wave one" from the in situ 1228 MOZAIC and SHADOZ data. Geophys. Res.-Atmos., 111. D01301. 1229 J. doi:10.1029/2005jd006241, 2006. 1230

1231

Sauvage, B., Martin, R. V., van Donkelaar, A., and Ziemke, J. R.: Quantification of the factors
controlling tropical tropospheric ozone and the South Atlantic maximum, J. Geophys. Res.Atmos., 112, D11309, doi:10.1029/2006jd008008, 2007.

1235
1236 Scannell, C., D. Hurtmans, A. Boynard, J. Hadji-Lazaro, M. George, A. Delcloo, A. Tuinder, P.F.
1237 Coheur, and C. Clerbaux: Antarctic ozone hole as observed by IASI/MetOp for 2008-2010, Atmos.
1238 Meas. Tech., 5, 123-139, 2012.

1239

Simon, H., A. Reff, B. Wells, J. Xing, and N. Frank: Ozone Trends Across the United States over
a Period of Decreasing NOx and VOC Emissions, Environ. Sci. Technol., 49, 186–195,
dx.doi.org/10.1021/es504514z, 2015.

Stohl, A., S. Eckhardt, C. Forster, P. James, and N. Spichtinger: On the pathways and timescales
of intercontinental air pollution transport. J. Geophys. Res. 107 (D23), 4684,
doi:10.1029/2001JD001396, 2002.

1247

1243

Thompson, D. W. J. and J.M. Wallace: Annular modes in the extratropical circulation. Part I:month-to month variability, J. Climate, 13, 1000–1016, 2000.

1250
1251 Thompson, A.M., J.C. Witte, H.G.J. Smit, S.J. Oltmans, B.J. Johnson, V.W.J H. Kirchhoff, and
1252 F.J. Schmidlin: Southern Hemisphere Additional Ozonesondes (SHADOZ) 1998-2004 tropical

1252 r.J. Schindini. Southern Henrisphere Additional Ozonesondes (SHADOZ) 1996-2004 tropical 1253 ozone climatology: 3. Instrumentation, station-to-station variability, and evaluation with simulated

- 1254 flight profiles, J. Geophys. Res., 112, D03304, doi:10.1029/2005JD007042, 2007.
- 1255 Ingin p

1256 Tarasick et al.: Tropospheric Ozone Assessment Report: Tropospheric ozone observations, Elem.1257 Sci. Anth., in-review, 2017.

1258

Thouret, V., J.-P. Cammas, B. Sauvage, G. Athier, R. Zbinden, R., P. Nédélec, P. Simon, and F.
Karcher: Tropopause referenced ozone climatology and inter-annual variability (1994–2003) from
the MOZAIC programme, Atmos. Chem. Phys., 6, 1033–1051, doi:10.5194/acp-6-1033-2006,
2006.

Tiao, G. C., G. C. Reinsel, D. Xu, J. H. Pedrick, X. Zhu, A. J. Miller, J. J. DeLuisi, C. L. Mateer,
and D. J. Wuebbles, Effects of autocorrelation and temporal sampling schemes on estimates of
trend and spatial correlation, J. GeophysR. es., 95, 20,507-20,517, 1990.

Tocquer, F., Barret, B., Mari, C., Le Flochmoen, E., Cammas, J. P., and Sauvage, B.: An upper tropospheric 'ozone river' from Africa to India during the 2008 Asian post-monsoon season, Tellus B, 67, 25350, doi:10.3402/tellusb.v67.25350, 2015.

1271

1267

Valks, P., N. Hao, S. Gimeno Garcia, D. Loyola, M. Dameris, P. Jöckel, and A. Delcloo: Tropical
tropospheric ozone column retrieval for GOME-2, Atmos. Meas. Tech., 7, 2513–2530,
doi:10.5194/amt-7-2513-2014, 2014.

1276 Van der A, R. J., et al.: Cleaning up the air: effectiveness of air quality policy for SO₂ and NO_x
1277 emissions in China, Atmos. Chem. Phys., 17, 1775–1789, 2017.

1278

1282

1275

1279 Verstraeten, W.W., J.L. Neu, J.E. Williams, K.W. Bowman, J.R. Worden and K.F. Boersma:
1280 Rapid increases in tropospheric ozone production and export from China, Nature Geosciences,
1281 doi: 10.1038/NGEO2493, 2015.

Voulgarakis, A., Telford, P. J., Aghedo, A. M., Braesicke, P., Faluvegi, G., Abraham, N. L.,
Bowman, K. W., Pyle, J. A., and Shindell, D. T.: Global multu-year O₃-CO correlation patterns
from models and TES satellite observations, Atmos. Chem. Phys., 11, 5819–5838,
doi:10.5194/acp-10-2491-2010, 2010.

1287

Voulgarakis, A., P. Hadjinicolaou, J. A. Pyle: Increases in global tropospheric ozone following an
El Nino event: examining stratospheric ozone variability as a potential driver, Atmos. Sci. Let. 12:
228–232, doi: 10.1002/asl.318, 2011.

1291

Weatherhead, E.C., G. C. Reinsel, G. C. Tiao, X.-L. Meng, D. Choi, W.-K. Cheang, T. Keller, J.
DeLuisi, D. J. Wuebbles, J. B. Kerr, A. J. Miller, S. J. Oltmans and J. E. Frederick: Factors affecting the detection of trends: Statistical considerations and applications to environmental data, J. Geophys. Res. Atmos., 103, 17149–17161, 1998.

Wespes, C., D. Hurtmans, L.K. Emmons, S. Safieddine, C. Clerbaux, D.P. Edwards, and P.-F.
Coheur: Ozone variability in the troposphere and the stratosphere from the first six years of IASI observations (2008-2013), Atmos. Chem. Phys., 16, 5721-5743, 2016.

Wespes, C., D. Hurtmans, C. Clerbaux, and P.-F. Coheur: O3 variability in the troposphere as
observed by IASI over 2008–2016 — Contribution of atmospheric chemistry and dynamics, J.

- 1303 Geophys. Res. Atmos., 122, 2429–2451, doi:10.1002/2016JD025875, 2017.
- 1304

1300

Wilson, R. C., Z.L. Fleming, P.S. Monks, G. Clain, S.Henne, I.B. Konovalov, S. Szopa, and L.
Menut: Have primary emission reduction measures reduced ozone across Europe? An analysis of
European rural background ozone trends 1996–2005, Atmos. Chem. Phys., 12, 437–454,
doi:10.5194/acp-12-437-2012, 2012.

Worden, J., Jiang, Z., Jones, D.B.A., Alvarado, M., Bowman, K., Frankenberg, C., Kort, E.A.,
Kulawik, S., S., Lee, M., Liu, J., Payne, V., Wecht, K., Worden, H.: El Niño, the 2006 Indonesian
peat fires, and the distribution of atmospheric methane, Geophys. Res. Let., V40, 1–6,
doi:10.1002/grl.50937, 2013.

- 1314
- Zbinden, R. M., J.-P. Cammas, V. Thouret, P. Nédélec, F. Karcher, and P. Simon: Mid-latitude
 tropospheric ozone columns from the MOZAIC program: climatology and interannual variability,
 Atmos. Chem. Phys., 6, 1053–1073, doi:10.5194/acp-6-1053-2006, 2006.
- 1318
- 1319

Zeng, G., Morgenstern, O., Shiona, H., Thomas, A. J., Querel, R. R., and Nichol, S. E.: Attribution
of recent ozone changes in the Southern Hemisphere mid-latitudes using statistical analysis and
chemistry-climate model simulations, Atmos. Chem. Phys. 17, 10495-10513, 2017.

1323

Zhang, L., Jacob, D. J., Bowman, K. W., Logan, J. A., Turquety, S., Hudman, R. C., Li, Q., Beer,
R., Worden, H. M., Worden, J. R., Rinsland, C. P., Kulawik, S. S., Lampel, M. C., Shephard, M.
W., Fisher, B. M., Eldering, A., and Avery, M. A.: Ozone-CO correlations determined by the TES
satellite instrument in continental outflow regions, Geophys. Res. Lett., 33, L18804,
doi:10.1029/2006GL026399, 2006.

1329

Zhang, L., Jacob, D. J., Boersma, K. F., Jaffe, D. A., Olson, J. R., Bowman, K. W., Worden, J. R.,
Thompson, A. M., Avery, M. A., Cohen, R. C., Dibb, J. E., Flock, F. M., Fuelberg, H. E., Huey,
L. G., McMillan, W. W., Singh, H. B., and Weinheimer, A. J.: Transpacific transport of ozone
pollution and the effect of recent Asian emission increases on air quality in North America: an
integrated analysis using satellite, aircraft, ozonesonde, and surface observations, Atmos. Chem.
Phys., 8, 6117–6136, doi:10.5194/acp-8-6117-2008, 2008.

1336

1337 Zhang, L., Li, Q. B., Murray, L. T., Luo, M., Liu, H. and co-authors: A tropospheric ozone
1338 maximum over the equatorial Southern Indian Ocean. Atmos. Chem. Phys. 12, 4279-4296, 2012.
1339

1340 Zhang, Y., O. R. Cooper, A. Gaudel, A. M. Thompson, P. Nédélec, S.-Y. Ogino and J. J. West
1341 (2016), Tropospheric ozone change from 1980 to 2010 dominated by equatorward redistribution
1342 of emissions, *Nature Geoscience*, doi: 10.1038/NGEO2827, 2016.

1343

Zhao, B., Wang, S. X., Liu, H., Xu, J. Y., Fu, K., Klimont, Z., Hao, J. M., He, K. B., Cofala, J.,
and Amann, M.: NOx emissions in China: historical trends and future perspectives, Atmos. Chem.
Phys., 13, 9869-9897, doi:10.5194/acp-13-9869-2013, 2013.

1347

Ziemke, J. R., A.R. Douglass, L.D. Oman, S.E. Strahan, and B.N. Duncan: Tropospheric ozone
variability in the tropics from ENSO to MJO and shorter timescales, Atmos. Chem. Phys., 15,
8037–8049, 2015.