

1 **Investigation of the summer 2018 European ozone air pollution episodes using**
2 **novel satellite data and modelling**

3 Richard J. Pope^{1,2}, Brian J. Kerridge^{3,4}, Martyn P. Chipperfield^{1,2}, Richard Siddans^{3,4}, Barry G. Latter^{3,4},
4 Lucy J. Ventress^{3,4}, Matilda A. Pimlott¹, Wuhu Feng^{1,5}, Edward Comyn-Platt⁶, Garry D. Hayman⁷,
5 Stephen R. Arnold¹ and Ailish M. Graham¹

6
7 *1: School of Earth and Environment, University of Leeds, Leeds, United Kingdom*

8
9 *2: National Centre for Earth Observation, University of Leeds, Leeds, United Kingdom*

10
11 *3: Remote Sensing Group, STFC Rutherford Appleton Laboratory, Chilton, United Kingdom*

12
13 *4: National Centre for Earth Observation, STFC Rutherford Appleton Laboratory, Chilton, United*
14 *Kingdom*

15
16 *5: National Centre for Atmospheric Science, University of Leeds, Leeds, United Kingdom*

17
18 *6: European Centre for Medium-Range Weather Forecasts, Reading, UK*

19
20 *7: Centre for Ecology and Hydrology, Wallingford, United Kingdom*
21

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23 Correspondence to: Richard J. Pope (r.j.pope@leeds.ac.uk)

24 **Abstract:**

25 In the summer of 2018, Europe experienced an intense heat wave which coincided with several
26 persistent large-scale ozone (O₃) pollution episodes. Novel satellite data of lower tropospheric
27 column O₃ from the Global Ozone Monitoring Experiment-2 (GOME-2) and Infrared Atmospheric
28 Sounding Interferometer (IASI) on the MetOp satellite showed substantial enhancements in 2018
29 relative to other years since 2012. Surface observations also showed ozone enhancements across
30 large regions of continental Europe in summer 2018 compared to 2017. Enhancements to surface
31 temperature and the O₃ precursor gases carbon monoxide and methanol in 2018 were co-retrieved
32 from MetOp observations by the same scheme. This analysis was supported by the TOMCAT
33 chemistry transport model (CTM) to investigate processes driving the observed O₃ enhancements.
34 Through several targeted sensitivity experiments we show that meteorological processes, and
35 emissions to a secondary order, were important for controlling the elevated O₃ concentrations at the
36 surface. However, mid-tropospheric (~500 hPa) O₃ enhancements were dominated by
37 meteorological processes. We find that contributions from stratospheric O₃ intrusions ranged
38 between 15 - 40%. Analysis of back trajectories indicates that the import of O₃-enriched air masses
39 into Europe originated over the North Atlantic substantially increasing O₃ in the 500 hPa layer during
40 summer 2018.

43 **1. Introduction**

44 Over the past two decades there have been several intense summer-time heatwaves over Europe
45 (e.g. 2003 over continental Europe (Scott et al., 2004), 2006 over north-western Europe (Rebetz et
46 al., 2008) and 2010 across eastern Europe and Russia (Matsueda et al., 2011)). With current and
47 future climate change, increasing average global surface temperature is expected to trigger more
48 frequent and intense heatwaves (Lhotka et al., 2017; Guerreiro et al., 2018). The summer-time 2018
49 heatwave across predominantly north-western and central Europe and Scandinavia generated
50 temperature anomalies of approximately 2.0-4.0 K (Li et al., 2020; Drouard et al., 2020). Dynamically,
51 it was caused by a combination of intense anticyclonic blocking systems, Rossby wave dynamics and
52 the positive phase of the summer-time North Atlantic Oscillation (NAO+) (Li et al., 2020; Liu et al.,
53 2020; Drouard et al., 2020). Environmentally, the summer 2018 heatwave caused severe drought
54 conditions with decreased precipitation and soil moisture content (Bastos et al., 2020; Dirmeyer et
55 al., 2020), while negatively impacting natural vegetation (e.g. decreased gross primary productivity
56 (Smith et al., 2020; Bastos et al., 2020)). From a human health perspective, the 2018 heatwave
57 caused 863 temperature related excess deaths in the UK (PHE, 2019).

58 As well as meteorological and vegetation responses, enhancements in atmospheric pollutants from
59 heatwaves can lead to a degradation in air quality (AQ) across Europe. Blocking systems (anticyclonic
60 conditions) have been shown to increase the level of air pollutions such as carbon monoxide (CO;
61 Thomas and Devasthale, 2014), nitrogen dioxide (NO₂; Pope et al., 2014) and particulate matter (i.e.
62 PM_{2.5}; Graham et al., 2020) to hazardous levels. Pope et al., (2016) focused on the 2006 UK
63 heatwave and detected enhancements in surface O₃ through the accumulation of pollutants (i.e.
64 atmospheric blocking) but also the higher temperatures yielding more active atmospheric chemistry
65 (i.e. ozone formation). Papanastasiou et al., (2015) found that Greek heatwave conditions (2001-
66 2010) typically yielded an increase in NO₂, PM_{2.5} and O₃ by 14-29%, 25-38% and 12%, respectively.
67 Rasilla et al., (2019) found that heatwaves in Madrid only moderately increased NO₂ and O₃ but
68 significantly increased PM₁₀ concentrations. However, they associated this with enhanced long-
69 range transport of African dust and then accumulation under heatwave conditions. García-Herrera et
70 al., (2010) provided a review of the 2003 European heatwave finding that the Alpine region had
71 substantially elevated surface ozone levels (peaking at 417 µg/m³ with 68% of sites from 23
72 countries reaching concentrations above 180 µg/m³) when compared with the previous 12 summers.
73 Biogenic volatile organic compound (BVOC) emissions from vegetation are known to increase under
74 drought conditions from temperature stress (e.g. in the 2003 European heatwave; Rennenberg et
75 al., 2006). Churkina et al., (2017) found that heatwave conditions (2006) in Berlin yielded an increase
76 in BVOC emissions which contributed up to 12% of the surface ozone formation. Heatwaves can also
77 trigger wildfires, which emit primary air pollutions and can form secondary gases such as surface
78 ozone on a regional and hemispheric scale (Honrath et al., 2004). Overall, elevated surface O₃ is
79 associated with adverse health impacts (Doherty et al., 2017; Heal et al., 2013; Jerrett et al., 2009)
80 with ailments such as asthma, reduced lung function and disease (WHO, 2021). It also has adverse
81 impacts on the natural biosphere (Sitch et al., 2007) and agriculture (Hollaway et al., 2012; van
82 Dingenen et al., 2009), in turn reducing deposition of surface ozone on vegetation. In this study, we
83 use surface and satellite observations of O₃, in combination with the well-evaluated TOMCAT global
84 chemical transport model (CTM), to investigate the impact of the summer 2018 heatwave on
85 European AQ and determine the key processes driving observed surface/tropospheric O₃

86 enhancements. We describe the observations and model we have used in Section 2. Section 3 and
87 Section 4 discusses our results and discussion/conclusions, respectively.

88 **2. Observations and Model**

89 **2.1. Satellite and Surface Observations**

90 We use satellite observations of lower tropospheric O₃ (i.e. sub-column O₃ (SCO₃) between the
91 surface and 450 hPa) from the Global Ozone Monitoring Experiment (GOME-2) and the Infrared
92 Atmospheric Sounding Interferometer (IASI) instruments on-board ESA's MetOp-A satellite, which
93 was launched in 2006 into a sun-synchronous polar orbit with equator crossing times of 9:30 (day)
94 and 21:30 (night). GOME-2 is a nadir-viewing spectrometer with spectral coverage in the ultraviolet-
95 visible (UV-Vis) of 240–790 nm (Riese et al., 2012) and a ground footprint of 40 km × 80 km in the
96 first part of the mission and 40 km x 40 km from 2013 (once Metop-B was commissioned). IASI is a
97 Michelson interferometer which observes the infrared spectral range 645 to 2760 cm⁻¹ with spectral
98 sampling of 0.25 cm⁻¹ (Illingworth et al., 2011). It measures simultaneously in four fields of view
99 (circular at nadir with a diameter of 12 km) which are scanned across track to sample a 2200 km-
100 wide swath (Clerbaux et al., 2009).

101 For GOME-2, the Rutherford Appleton Laboratory (RAL) scheme uses an optimal estimation
102 algorithm (Rodgers, 2000) to retrieve height-resolved ozone distributions spanning the stratosphere
103 and troposphere (Miles et al., 2015). The scheme applied to GOME-2 has been developed from that
104 used first for GOME-1 on-board ERS-2 (Munro et al., 1998; Forster et al., 2007). This is a multi-step
105 scheme in which profile information is first retrieved in the stratosphere by exploiting wavelength-
106 dependent absorption in the O₃ Hartley band (270-307nm) and is then extended into the
107 troposphere by exploiting temperature-dependent spectral structure in the O₃ Huggins bands (325-
108 335nm). For IASI, O₃ profiles are retrieved using an extended version of RAL's Infrared-Microwave-
109 Sounding (IMS) scheme, which is described in Pope et al., (2021), Palmer et al., (2022) and Pimlott et
110 al., (2022). The IMS core scheme was originally developed to retrieve temperature, water vapour
111 and stratospheric O₃ profiles along with surface spectral emissivity and cloud jointly from co-located
112 measurements by IASI, the Microwave Humidity Sounder (MHS) and the Advanced Microwave
113 Sounding Unit (AMSU-A) on MetOp (RAL Space, 2015). GOME-2 and IMS O₃ data were filtered for a
114 geometric cloud fraction less than 0.2, a solar zenith angle less than 80°, a cost function less than
115 200.0 and a convergence flag equal to 1.0. Examples of the vertical sensitivity to retrieving ozone
116 (i.e. averaging kernels) from GOME-2 and IMS are shown in **Supplementary Material (SM) 1**.

117 We also use surface O₃ observations from the European Monitoring and Evaluation Programme
118 (EMEP) network for May-August 2017 and 2018. The EMEP network contains >100 surface
119 measurement sites measuring information on a range of air pollutions (e.g. ozone, NO₂ and PM_{2.5}).
120 EMEP surface data can be used for multiple scientific applications such as trends analysis (Yan et al.,
121 2018) and atmospheric chemistry model evaluation (Schultz et al., 2017; Archibald et al., 2020) and
122 is hosted by the EBAS database infrastructure, developed by the Norwegian Institute for Air
123 Research. In total, we used 125 spatial collocated EMEP sites in both years across Europe. Here, data
124 at individual sites were selected where the corresponding data flag was set to 0.0.

125 **2.2. Modelling & Sensitivity Experiments**

126 In this study the TOMCAT CTM (Chipperfield, 2006) is forced by European Centre for Medium-Range
127 Weather Forecasts (ECMWF) ERA-Interim reanalysis meteorology (Dee et al., 2011) and is run at a

128 horizontal resolution of $2.8^\circ \times 2.8^\circ$. The model has with 31 vertical levels from the surface to 10 hPa
129 with 5-7 (approximately 10) levels in the boundary layer (mid-troposphere), depending on latitude.
130 The model includes detailed tropospheric chemistry, including 229 gas-phase reactions and 82
131 advected tracers (Monks et al., 2017), and heterogeneous chemistry driven by size-resolved aerosol
132 from the GLOMAP module (Mann et al., 2010). Anthropogenic emissions used in this study come
133 from MACCity (Granier et al., 2011). The original dataset in Granier et al., (2011) derived emissions
134 up to 2010. Therefore, the Representative Concentration Pathways 8.5 (RCP 8.5) were used by
135 Granier et al., (2011) to generate emissions for later years (e.g. 2017 and 2018 as used in this study).
136 Fire emissions are from the Global Fire Assimilation System (GFAS, Kaiser et al., 2012) for 2017 and
137 2018. Year-specific off-line biogenic volatile organic compounds (VOCs) emissions for acetone,
138 methanol, isoprene and monoterpenes were simulated by the Joint UK Land Environment Simulator
139 (JULES – Pacifico et al., 2011; Best et al., 2011; Clark et al., 2011). All other biogenic VOC emissions
140 are climatological values and provided by the Chemistry-Climate Model Initiative (CCMI)
141 (Morgenstern et al., 2017). The global budgets of the JULES VOC emissions are low in comparison to
142 the climatological CCMI emissions, so were scaled up on a regional basis, while retaining the 2017-
143 2018 step change related to the 2018 summer heat wave. The full details of JULES VOC emissions
144 scaling can be found in **SM4**. Lightning emissions of NO_x are coupled to convection in the model,
145 which is derived from the meteorological reanalyses. Therefore, they vary in space and time
146 according to the seasonality and spatial pattern of convective activity (Stockwell et al., 1999). The
147 model was run for 2017 and 2018 with output at 6-hourly intervals (i.e. 00, 06, 12 and 18 UTC). Here,
148 each year was run with its respective meteorology and emissions and given the labels
149 Met17_Emis17 (representing 2017) and Met18_Emis18 (representing 2018).

150 To explore the importance of emission and meteorological processes behind the elevated European
151 summer 2018 tropospheric O_3 levels, a 1-year model sensitivity experiment was performed using
152 2018 meteorology but 2017 emissions (i.e. Met18_Emis17). Therefore, the difference between
153 Met18_Emis17 and Met17_Emis17 highlights the impact of fixed emissions (i.e. 2017 emissions for
154 both years), while the Met18_Emis18 minus Met18_Emis17 highlights the impact of fixed
155 meteorology (i.e. 2018 meteorology for both years – including BVOC emissions). These are
156 compared with the control differences for 2018-2017 (Met18_Emis18- Met17_Emis17). From here
157 on in, we refer to the control differences, fixed emission differences and the fixed meteorology
158 differences as CTL_DIFF, FIXED_EMIS_DIFF and FIXED_MET_DIFF, respectively. TOMCAT also
159 includes a stratospheric O_3 tracer, a common approach to tag stratospheric O_3 (e.g. Roelofs et al.,
160 2003; Akritidis et al., 2019), which can be used to investigate the impact of stratospheric O_3 intrusion
161 into the troposphere. The tracer is set equal to the model-calculated O_3 in the stratosphere. The only
162 tropospheric source of O_{35} is transport from the stratosphere while its sinks are via photolysis,
163 reactions with HO_2 , OH and H_2O through $\text{O}(^1\text{D})$ produced from O_{35} and surface deposition (Monks et
164 al., 2017). The tracer does not have a fixed lifetime but the loss rate in the troposphere depends on
165 the modelled local OH, HO_2 , H_2O and photolysis. Any O_3 that gets into the stratosphere will be
166 labelled as stratospheric before it returns. This was used to investigate the impact of stratospheric
167 O_3 intrusion into the troposphere.

168 TOMCAT has been used in a number of previous studies to investigate air quality and tropospheric
169 composition (e.g. Richards et al., 2013; Emmons et al., 2015; Pope et al., 2018; Pope et al., 2020)
170 whose results give confidence in the model's ability to simulate European tropospheric O_3 in this
171 study. Overall, when compared with observations, TOMCAT has good spatial agreement with both

172 GOME-2 and IASI and can reasonably reproduce the 2018 SCO₃ enhancement in 2018 versus 2017
173 (SM 5). The model also has good agreement, both in magnitude and seasonality, with the EMEP
174 observed surface concentrations (SM 5). TOMCAT surface ozone was also compared with higher
175 resolution modelling (reanalysis) data from the Copernicus Atmosphere Monitoring Service (CAMS),
176 which showed good spatial agreement between the modelling data sets and in the simulated surface
177 ozone absolute values during the European summer 2018 pollution episode (SM 5).

178 2.3 ROTRAJ Back-trajectories

179 We use the Reading Offline Trajectory Model (ROTRAJ) to generate air mass back-trajectories
180 (Methven et al., 2003) to assess the import of tropospheric O₃ into Europe. ROTRAJ is a Lagrangian
181 atmospheric transport model driven by meteorology from the same ECMWF ERA-Interim reanalyses
182 (horizontal resolution of 1.0125°) as used by TOMCAT. Velocity fields at the Lagrangian particle
183 positions are determined by cubic Lagrange interpolation in the vertical, bilinear interpolation in the
184 horizontal and linear interpolation in time. This method accounts for large scale advection since the
185 winds are resolved but does not resolve small scale sub-grid turbulent transport. Kinematic back-
186 trajectories were released at 6-hourly intervals (i.e. at 00, 06, 12 and 18 UTC) from Paris and Berlin,
187 both central locations over Europe in the region of summer-time 2018 O₃ enhancements, between
188 the 1st May and 31st August for both 2017 and 2018. The trajectories were released at the surface
189 and at approximately 500 hPa and integrated for 10 days with 6-hourly output (i.e. 41 trajectory
190 points including the starting location) to investigate the origin of air masses arriving in these altitude
191 regions of enhanced summer-time O₃ in 2018. In total, ROTRAJ was therefore run 8 times (2 years ×
192 2 altitudes × 2 locations).

193 To quantify the import of tropospheric O₃ into Europe, for each trajectory, all the trajectory points
194 were co-located with corresponding TOMCAT O₃ mixing ratio values (i.e. the horizontal and vertical
195 grid box the trajectory point sits within and corresponding time stamp) and then the average O₃-
196 weighted back-trajectory (O₃-WBT) determined (i.e. back-trajectories with larger O₃WBT values
197 come from air masses with larger O₃ content). This follows a similar approach to Graham et al.,
198 (2020) and Stirling et al., (2020), though using a model chemical tracer and not emission inventories.

199 3. Results

200 3.1 Surface Temperature

201 Several studies (e.g. Li et al., 2020; Liu et al., 2020; Drouard et al., 2020) have documented the
202 intense heat wave across Europe in the summer of 2018. This is further shown in **Figure 1** which
203 compares surface temperature, co-retrieved with ozone and other variables from MetOp-A by the
204 IMS scheme, between 2017 and 2018. In May, higher temperatures occur across Scandinavia (5.0-
205 10.0 K), eastern Europe (3.0-7.0 K) and the UK (1.0-3.0 K), but temperatures are lower (-3.0 to -1.0 K)
206 across Iberia. In June, a similar spatial distribution occurs but the magnitude of the differences is
207 smaller. In July the largest temperature increases range from 6.0-8.0 K in Scandinavia to 2.0-6.0 K in
208 the UK/France. Iberia continued to experience temperatures lower by -2.0 to 0.0 K. In August, there
209 are near-zero differences over the UK, Iberia and most of Scandinavia but with increases of 1.0-3.0 K
210 over eastern Europe and Finland.

211 3.2 Satellite Ozone

212 We investigate the longer-term variability in tropospheric O₃ (i.e. SCO₃) to determine if 2017 is a
213 suitable comparator for the 2018 summer O₃ enhancements as it is for temperature. **Figure 2** shows

214 the 2012-18 SCO_3 average between May and August for a domain over the Atlantic and Europe and
215 the difference for the same season between specific years and the 2012-18 average. In 2012 and
216 2013, there are significant positive differences from the average between 1.0 DU and 5.0 DU over
217 much of the domain. Over continental Europe, the differences are smaller. Here, the significance of
218 differences between the year-specific and long-term averages are determined using the Wilcoxon
219 Rank test (Pirovano et al., 2012) at the 95% confidence level. In 2014 and 2015, there are negative
220 differences across Europe (-4.0 DU to -1.0 DU). In 2016, similar negative differences are primarily
221 across the north and south-east of the domain. In 2017, there are near-zero differences across the
222 Atlantic, UK and western Europe. Over eastern Europe and Mediterranean, there are significant
223 negative differences of between -2.0 DU and -1.0 DU. In 2018, across continental Europe there are
224 significant positive differences between 2.0 DU and 4.0 DU. As the 2017 differences are relatively
225 small in magnitude with a low proportion of significant pixels (i.e. Sig Pixels % = 32.7 is the lowest
226 across the 7 years), it is representative of average conditions for comparison with 2018. For 2018,
227 the summer SCO_3 enhancements across continental Europe are the largest for the years shown with
228 a coherent cluster of significant differences. This illustrates that the summer 2018 SCO_3
229 enhancements are a substantial deviation from the average conditions (which we represent as 2017
230 hereon) and that this is an intense O_3 event.

231 Investigation of SCO_3 retrieved from both GOME-2 (**Figure 3**) and the IMS scheme (**Figure 4**) show
232 consistent enhancements in summer 2018. In 2017, between May and August, GOME-2 typically
233 observed SCO_3 values between 20.0-30.0 DU across continental Europe. Peak SCO_3 values occurred
234 over the Mediterranean (30.0-38.0 DU); relatively high ozone is a typical feature of the
235 Mediterranean in summer (Richards et al., 2013). In 2018, the seasonality is consistent with 2017,
236 but the continental European SCO_3 values ranged between 25.0 DU and 35.0 DU. For the 2018-2017
237 difference, SCO_3 enhancements occur across continental Europe in all four months but peaked in
238 May and July between 3.0 DU and 8.0 DU, while typically 1.0-5.0 DU in June and August. The spatial
239 distribution of IMS-retrieved SCO_3 is similar to that of GOME-2 in 2017 and 2018, although the
240 absolute values tend to be systematically lower by 3.0-4.0 DU. However, despite this systematic
241 offset, the 2018-2017 differences are reasonably consistent with GOME-2. Across continental
242 Europe, IMS SCO_3 shows 2018 enhancements in all months investigated, but peaks in May and July,
243 like GOME-2, between 3.0 DU and 6.0 DU. The differences range from 1.0 DU to 3.0 DU in June and
244 are approximately 1.0 DU in August (though a peak enhancement of 3.0-5.0 DU occurs over the
245 Mediterranean). Spatial correlations between the GOME-2 and IASI difference (i.e. 2018-2017) maps
246 for the months investigated ranged between 0.21 and 0.47 (see **SM 5**).

247 The GOME-2 and IASI instruments observe UV-Vis and IR wavelengths, with peak vertical sensitivities
248 to tropospheric O_3 in the lower and mid/upper troposphere, respectively. Consistency in the 2018
249 enhancements in SCO_3 indicates that these extend over the bulk of the troposphere and increases
250 confidence in the detected enhancements for both sensors.

251 Investigation of several satellite-retrieved O_3 precursor gases (see **SM 2**) showed enhancements in
252 total column methanol (TCCH_3OH , **Figure S2**), especially linked to May and July temperature
253 enhancements (**Figure 1**), minor increases in tropospheric column NO_2 (TCNO_2 , **Figure S3**) in May and
254 July over central Europe and widespread enhancements (weakest in July and strongest in August) in
255 total column carbon monoxide (TCCO , **Figure S4**). Investigation of the GOME-2 and IASI total column
256 O_3 (TCO_3) differences between 2017 and 2018 (**Figures S5 & S6**) showed these to be in close
257 agreement. Some spatial structure is similar to that of the SCO_3 difference patterns (**Figures 3 and**

258 4), with correlations of approximately 0.5 between TCO₃ and SCO₃ for each instrument (see **SM 3**).
259 Given the complex relationship between tropospheric O₃, precursor gases, atmospheric chemistry
260 (e.g. NO_x or VOC-limited regimes), surface deposition and meteorological conditions (e.g.
261 atmospheric temperatures and transport), a detailed chemistry transport model is required to assess
262 the key processes leading to the observed SCO₃ enhancements over Europe.

263 3.3 Surface Ozone

264 Increased temperatures during heat waves have been shown to enhance surface O₃ concentrations
265 (e.g. Jacob and Daniel, 2009; Vieno et al., 2010; Pyrgou et al., 2018). In the summer (May-June-July-
266 August, MJJA) of 2018, EMEP recorded larger O₃ mixing ratios across most of Europe in comparisons
267 to 2017 (**Figure 5a & b**). Over central Europe, surface O₃ mixing ratios ranged from approximately
268 45.0 ppbv to over 60.0 ppbv, while in 2017 it was 35.0 ppbv to 50.0 ppbv. Over the UK and north-
269 western Europe, surface O₃ mixing ratios ranged from 20.0 ppbv to 30.0 ppbv and then 25.0 ppbv to
270 35.0 ppbv in MJJA 2017 and 2018, respectively. In Scandinavia and eastern Europe, surface O₃ mixing
271 ratios ranged from 20.0 ppbv to 35.0 ppbv in MJJA 2017, while increasing to 25.0 ppbv to
272 approximately 40.0 ppbv in MJJA 2018. **Figure 5c** highlights these widespread enhancements where
273 domain-average surface O₃ mixing ratios are larger by typically 5.0-10.0 ppbv in May and from mid-
274 June to mid-August in 2018. **Figure 5d** shows that the domain median surface O₃ concentration
275 across MJJA was larger by 2.0-3.0 ppbv in 2018, but the 2018 extremes were greater with 75th and
276 95th percentiles of 45.0 ppbv and 55.0 ppbv in 2017 and 48.0 ppbv and 59.0 ppbv in 2018. Therefore,
277 surface observations of O₃ recorded widespread enhancements in MJJA 2018 compared to 2017
278 with peak site differences >10.0 ppbv. This is generally consistent with the 2018 layer-averaged
279 enhancements in the satellite-retrieved SCO₃ for regions where both datasets have spatial coverage.

280 3.4. Model Simulations

281 We use the TOMCAT model to investigate different factors potentially driving the observed
282 enhancements in tropospheric O₃. In comparisons with the observations (see **SM 5**) the model
283 reproduces the sign and spatial distribution of observed 2018-2017 differences reasonably well.
284 Although it has a tendency to underestimate the absolute magnitude, we are confident in the
285 model's ability to simulate the tropospheric O₃ enhancements relative to 2017.

286 At the surface (**Figure 6**), TOMCAT CTL_DIFF (i.e. Met18_Emis18 - Met17_Emis17) suggests that O₃ is
287 enhanced in May over Scandinavia (2.0- >5.0 ppbv), north-western Europe (0.0-2.0 ppbv), the Arctic
288 Ocean (>5.0 ppbv) and off the coast of Iberia (3.0-5.0 ppbv). However, negative values exist over
289 eastern Europe (-3.0 ppbv to -1.0 ppb) and the Atlantic west of Ireland (-3.0 ppbv to -1.0 ppb). In
290 June, the negative differences persist in eastern Europe (-3.0 ppbv to -1.0 ppb), but positive
291 differences are located over northern Scandinavia (1.0-2.0 ppbv) and the North Atlantic (2.0-4.0
292 ppbv). For July, CTL_DIFF shows the largest enhancements over continental Europe (i.e. Po Valley,
293 France, Benelux region and Iberia) and the UK (>5.0 ppbv). Negative differences of between -3.0
294 ppbv and -1.0 ppbv remain over eastern Europe. In August, the only clear differences are over Iberia
295 and the western Mediterranean, ranging between 3.0 ppbv and >5.0 ppbv. Overall, TOMCAT
296 simulates sub-regional surface O₃ enhancements over Europe, which are generally consistent with
297 EMEP observations apart from several sites over eastern Europe.

298 At 500 hPa, TOMCAT CTL_DIFF shows larger-scale O₃ enhancements in 2018 compared to 2017 (>5.0
299 ppbv) throughout May to August. In May and August, there are, however, a few negative differences

300 (-5.0 ppbv to -3.0 ppbv) over far eastern Europe. In June and July, the full domain is more or less
301 dominated by O₃ enhancements in 2018. In **Figures 3** and **4** (and **SM 5**), GOME-2 and IASI (and
302 TOMCAT with the instrument averaging kernels (AKs) applied to account for the vertical sensitivity of
303 the retrievals, see **SM 5** for more information) show SCO₃ enhancements during these months of
304 2018. Given the vertical extents and peak heights of their retrieval sensitivities and consistency in
305 spatial patterns (**Figs SM 9** and **11**) it is evident that the O₃ enhancements detected by GOME-2 and
306 IASI extend over the free troposphere. The model shows large-scale O₃ enhancements in the free
307 troposphere and similar patterns to GOME-2 and IASI when averaging kernels applied. So, the model
308 corroborates this finding from the satellite retrievals. Signals from EMEP and TOMCAT at the surface,
309 on the other hand, are more mixed across the domain.

310 The right-hand column of **Figure 6** shows the relative difference in the stratospheric O₃ contribution
311 to the 500 hPa O₃ layer (i.e. Strat % @ 500 hPa), from CTL_DIFF, between 2017 and 2018. Here, the
312 percentage of stratospheric O₃ contributing to the O₃ concentration at the 500 hPa is calculated for
313 2017 and 2018 and then the 2018-2017 difference determined. The largest enhancement to the 500
314 hPa layer was in July where the stratospheric O₃ contribution increased by 3.0% to >5.0% across
315 Europe. In June and August, the spatial patterns are similar with stratospheric O₃ contribution
316 enhancements of 3.0-5.0% across southern Europe, Scandinavia and the North Atlantic (above the
317 UK). In the North Atlantic, UK and northern Europe, there are near-zero changes in June and August.
318 In May, there are enhancements >5.0% across the northern region of the domain and northern
319 Africa, while smaller enhancements (1.0%-3.0%) over the UK and near-zero changes over eastern
320 Europe. This is partially supported by analysis of TCO₃ (see **SM 3**) where there are reasonable spatial
321 correlations (~0.5 to 0.6) between the SCO₃ 2017-2018 summer differences and the equivalent for
322 TCO₃. Therefore, these results indicate a potentially enhanced contribution of stratospheric O₃ into
323 the mid-troposphere during the summer of 2018 across Europe.

324 To quantify the separate importance of precursor emissions and meteorology in governing the
325 summer 2018 O₃ enhancements we compare the sensitivity experiments with the control runs.
326 **Figure 7** (left column) shows the results for the fixed emissions differences (i.e. FIXED_EMIS_DIFF)
327 between years (i.e. Met18_Emis17 – Met17_Emis17). At the surface, the FIXED_EMIS_DIFF show
328 similar spatial patterns to that of CTL_DIFF (**Figure 6** – left column). The domain spatial difference
329 correlations between these simulations is greater than 0.96 for all months considered. However,
330 FIXED_EMIS_DIFF (**Figure 7** - left column) tends to be lower than CTL_DIFF (**Figure 6** – left column)
331 by approximately 0.0-2.9 ppbv (i.e. positive red regions are weaker and negative blue regions
332 stronger in intensity). Therefore, the Met18_Emis17 run struggles to reproduce the absolute surface
333 O₃ enhancements in the Met18_Emis18 run. When the fixed meteorology differences
334 (FIXED_MET_DIFF, i.e. Met18_Emis18 - Met18_Emis17, **Figure 8** - left column) are compared with
335 CTL_DIFF, the surface 2018-2017 differences are substantially different.

336 Surface FIXED_MET_DIFF ranges between 0.0 ppbv and 2.0 ppbv across the domain in May and June
337 and is more confined to continental Europe in July and August. This shows that TOMCAT simulates
338 lower 2018 summer-time O₃ when 2017 emissions are used and indicates that emissions do have
339 some role in controlling O₃ levels at the surface. However, as the spatial difference pattern for
340 FIXED_MET_DIFF (**Figure 8** – left column) is different to that of CTL_DIFF (**Figure 6** – left column),
341 spatial correlations between them range from -0.53 to 0.54 over the four months, it suggests that
342 meteorology is important in governing the spatial distribution of CTL_DIFF. This is supported by the
343 fact that FIXED_MET_DIFF - CTL_DIFF (**Figure 8** left column – **Figure 6** left column) yields absolute

344 domain variations between 0.0 ppbv and 12.2 ppbv. Therefore, the two sensitivity experiments
345 suggest meteorology and emissions both play important roles in controlling surface O₃ during the
346 summer of 2018, but meteorology predominantly governs the spatial pattern and absolute
347 magnitude of the O₃ enhancements.

348 At 500 hPa, comparison of FIXED_EMIS_DIFF and CTL_DIFF show very consistent spatial patterns
349 across the four months with correlations all above 0.98. In terms of the absolute differences
350 between FIXED_EMIS_DIFF and CTL_DIFF (i.e. **Figure 7** centre column – **Figure 6** centre column) it
351 peaks at approximately 2.8 ppbv. For FIXED_MET_DIFF, the spatial correlation with CTL_DIFF, as for
352 the surface, is variable with values between -0.38 and 0.43. The absolute differences between
353 FIXED_MET_DIFF and CTL_DIFF (i.e. **Figure 8** centre column – **Figure 6** centre column) ranges from
354 0.0 ppbv to 14.8 ppbv. Therefore, emissions have a secondary role in controlling the O₃ while
355 meteorology is by far the dominant factor. For Strat % @ 500 hPa, the spatial correlations between
356 CTL_DIFF and FIXED_EMIS_DIFF are above 0.95 for all months and the absolute differences between
357 them (i.e. **Figure 7** right column - **Figure 6** right column) are near-zero. Comparison of
358 FIXED_MET_DIFF and TC_CTL shows spatial difference correlations ranging between -0.33 and 0.71
359 and absolute differences (i.e. **Figure 8** right column - **Figure 6** right column) peaking at 12.9%.
360 Therefore, as expected, meteorological processes are dominating the influence of the stratospheric
361 O₃ contribution (i.e. through stratosphere-troposphere exchanges) to the 500 hPa layer during the
362 summer 2018 O₃ enhancements over Europe.

363 To investigate the importance of stratospheric-troposphere exchange to the middle troposphere
364 enhancement (i.e. as shown in the TOMCAT 500 hPa layer and the satellite SCO₃ data), **Figures 9** and
365 **10** show TOMCAT control run zonal 2018-2017 difference cross-sections (for the domain longitudes)
366 of O₃ profiles and the stratospheric O₃ contribution to each pressure layer. In May and June, in the
367 lower troposphere (approximately surface to 800 hPa), there are negative (-3.0% to 0.0%) and
368 positive (0.0% to 3.0%) differences between 30-50°N and 50-70°N, respectively. During June, there
369 are positive differences (0.0% to 5.0%) across most latitudes and in August, the opposite occurs to
370 that of May/June. In the mid-troposphere (800-300 hPa), positive differences occur in most months
371 (0.0-5.0% in May, 0.0-7.0% in June, >10% in July and 5.0-10.0% in August), though in May and August
372 negative differences (-5.0% to 0.0%) exist around 40°N and 55°N. This is consistent with the 500 hPa
373 O₃ differences in **Figure 6** (centre panels). In the upper troposphere – lower stratosphere (UTLS,
374 approximately 300-100 hPa) there are limbs of positive O₃ differences (i.e. >10%, 5.0-10.0 ppbv)
375 propagating into the mid-troposphere (30-40°N in May, 30-50°N in June, 40-50°N in July and 30-40°N
376 & 60-70°N in August), suggestive of stratospheric intrusion into the mid-troposphere. Using the
377 stratospheric O₃ tracer in TOMCAT, **Figure 10** shows the enhanced proportion of O₃ originating from
378 the stratosphere in the summer of 2018. Interestingly, for all months (apart from May between 30-
379 45°N), there are enhanced contributions of stratospheric O₃ (15.0% to >50.0%) in the lower-mid
380 troposphere (i.e. below 500 hPa). In absolute terms, this is only a minor contribution typically <1.0
381 ppbv below 800 hPa. Between 800-400 hPa, this increases to 1.0-5.0 ppbv (remains relatively
382 consistent in percentage terms) in most months and latitude bands. In the UTLS, it increases to 5.0-
383 10.0% enhancements in stratospheric O₃ contributions, which is consistent with its proximity to the
384 stratosphere. In comparison between **Figures 9** and **10**, where there are enhancements in the
385 stratospheric O₃ contribution but negative differences in O₃ (e.g. in June in the lower troposphere
386 between 50°N and 55°N) which is suggestive of different processes influencing the O₃ concentrations
387 (e.g. descent of relatively small stratospheric O₃ contributions but advection of tropospheric O₃ away

388 from the region). Overall though, in the mid-troposphere, where there are larger enhancements in
389 O₃, there are similar responses in the stratospheric O₃ contribution. For June, the mid-troposphere
390 O₃ enhancement is approximately 5.0-7.0 ppbv with a signal of 1.0-2.0 ppbv in the stratospheric
391 tracer. Therefore, in the more extreme cases, the stratospheric O₃ contribution is approximately
392 15.0-40.0% to the mid-tropospheric O₃ enhancements in summer 2018 over Europe. However, a
393 separate study would be required to undertake a detailed assessment of the meteorological
394 processes controlling the enhanced stratospheric intrusion of ozone in the summer of 2018 and how
395 it compares to other years (how does it compare with years other than 2017).

396 The two remaining factors, linked to meteorological processes (as suggested above), which may
397 affect the O₃ enhancements in 2018 are increased summer temperatures (e.g. through enhanced
398 kinetic rates), and the import of tropospheric O₃ from upwind (e.g. North America from the
399 prevailing winds). **Figure 11** shows the 2017-2018 zonal temperature differences (i.e. same as **Figure**
400 **9** but for temperature) with the correlation between the 2017 and 2018 temperature and O₃
401 differences overplotted. Qualitatively, the zonal differences in O₃ and temperature have some
402 similarities. There are positive differences (temperature differences of 0.0-1.0%) between 50-60°N at
403 the surface and 400 hPa in May and June. Then in July, collocated positive differences (peaking at
404 2.0% or 3.0 K) exist between 50-70°N from the surface to 300 hPa. In August, there is no clear
405 relationship between temperature and O₃ enhancements. In all months (to a lesser extent in
406 August), in the UTLS, there are spatial agreements with positive differences between approximately
407 30-45°N and negative differences between 50/55-70°N. In terms of correlations (i.e. temporal
408 correlation in each grid box using the TOMCAT 6-hourly time series), the spatial agreement is
409 relatively weak. In all months, most of domain has relatively small values ranging between -0.5 to
410 0.5. There are only a few locations with strong correlations (i.e. > 0.5), which are in the UTLS or in
411 the lower-mid troposphere between 50-70°N (June & August) and 45-55°N in July near the surface.
412 Overall, the relationship between increased temperatures and enhanced kinetic rates yielding more
413 ozone formation is non-linear, so it is unsurprising that the direct comparisons of temperature and
414 ozone 2018-2017 differences above shows no clear pattern. Therefore, future work could include a
415 further sensitivity experiment running TOMCAT for 2018, but with 2017 temperatures used in the
416 chemistry routines to quantify the role of temperature in the summer 2018 O₃ enhancements.

417 To investigate the potential advection of tropospheric O₃-rich air masses into Europe we have used
418 ROTRAJ back-trajectories to determine the O₃WBTs (i.e. an indicator of air mass O₃ content). As
419 shown in **SM 6**, there is large variability in the O₃WBT values and spatial distribution (i.e. **Figures S13**
420 **and 14**), so they have to be gridded onto the TOMCAT horizontal resolution (see **Figures S15 and 16**).
421 While this approach does not directly account for the frequency of trajectory points in each grid box,
422 **Figures S13 and S14** show there is widespread coverage across the North Atlantic. This results in
423 >500 trajectory points near the receptor sites (i.e. Paris and Berlin), ~100 trajectory points around
424 the edge of Europe and 25-50 trajectory points in the North Atlantic (not shown here). Overall, this
425 spatial distribution is relatively consistent and does not change substantially between years (typically
426 10%), thus this approach is suitable in this study. **Figure 12** shows the differences (2018-2017)
427 between the gridded O₃WBTs where the back-trajectories have been released at the surface from
428 Paris (**Figure 12a**), at the surface from Berlin (**Figure 12b**), at approximately 500 hPa from Paris
429 (**Figure 12c**) and at approximately 500 hPa from Berlin (**Figure 12d**). We selected Paris and Berlin as
430 they are situated in central Europe where the summer 2018 O₃ enhancements have been observed
431 while the surface and 500 hPa are the altitudes of primary focus in the modelling work.

432 At the surface, Paris and Berlin show consistent patterns. Over the North Atlantic (i.e. origin of the
433 prevailing winds into Europe), there are typically negative O_3 WBT values between -5.0 ppbv and -1.0
434 ppbv suggesting that advection of O_3 into Europe during the summer (i.e. May-August) was
435 predominantly larger in 2017 and did not strongly contribute to the 2018 observed surface O_3
436 enhancements. Advection of O_3 -rich air in 2018 did originate from Scandinavia into continental
437 Europe, though the number of trajectories is relatively low (see **Figure S13**). As both locations show
438 similar relationships, it provides confidence in this methodology. At 500 hPa, the 50-60°N spatial
439 pattern is less defined with values typically between -5.0 and 5.0 ppbv for both locations. However,
440 in the southern North Atlantic (30-50°N) there are positive differences of approximately 3.0-10.0
441 ppbv for both release locations. Note that as free-tropospheric winds tend to have larger horizontal
442 velocities, the back-trajectories generally start from further away closer to North America. Again,
443 given the broad similarity in differences between both release locations, it provides confidence in
444 this approach. Overall, our results indicate a larger transport of O_3 to the surface of continental
445 Europe in 2017, while at approximately 500 hPa the import of O_3 into Europe is larger in 2018. Here,
446 the positive differences originate from the southern North Atlantic (i.e. a larger range of locations,
447 absolute values and homogeneous signal than the mixed differences between 50-60°N).

448 One potentially important factor is dry deposition of O_3 to the land surface. Due to the heatwave,
449 stress on the biosphere and the associated die back of vegetation could potentially reduce the
450 efficiency of O_3 deposition decreasing the O_3 sink (i.e. O_3 is more likely to deposit onto land covered
451 by vegetation than bare soil). Investigation of the normalised difference vegetation index (NDVI),
452 from the IMS scheme, between the summers of 2017 and 2018 did not highlight any spatially
453 coherent changes (not shown here). As a result, there is no obvious large-scale spatial vegetation die
454 back in 2018 due to the heatwave and thus the impact this would have on ozone deposition in
455 TOMCAT. Therefore, we ran two further experiments where the bare soil fraction for each grid box
456 over Europe was increased and decreased by 25% in summer 2018. This was to investigate the
457 sensitivity of surface ozone deposition to changes in the land surface. For the increase in bare soil
458 fraction there was a moderate systematic increase in European summer ozone by 0.0-1.5 ppbv (i.e.
459 less ozone deposition). When the bare soil fraction was decreased by 25%, this yielded a small
460 decrease in surface ozone by approximately 0.5 ppbv. Overall, a sizable level of vegetation die back
461 would be required for decreased ozone dry deposition to substantially contribute to the summer
462 2018 surface ozone enhancements.

463 **4. Discussion and Conclusions**

464 The summer of 2018 produced an intense heatwave across most of Europe with a substantial impact
465 on tropospheric temperatures, droughts, stress on vegetation and human mortality. Observations of
466 surface temperature, precursor gases and total column O_3 (TCO_3) experienced enhancements in
467 2018 relative to 2017. In this paper, we have demonstrated a strong enhancement in surface and
468 tropospheric O_3 during the heatwave between May and August 2018. The EMEP surface data
469 suggest an average European enhancement, relative to 2017, peaking at approximately 10.0 ppbv in
470 July and August. Investigation of lower tropospheric O_3 (i.e. surface-450 hPa sub-column O_3 – SCO_3)
471 from the GOME-2 and IASI instruments also showed enhancements, peaking at 5.0-10.0 DU, relative
472 to 2017. Analysis of the long-term GOME-2 SCO_3 record indicates 2017 to be a suitably
473 neutral/average reference year and the enhancement in 2018 to be anomalously large. Our
474 comparisons were therefore made between the summers of 2017 and 2018.

475 Consistency between the UV (GOME-2) and IR (IASI) sounders was important to our analysis because
476 their vertical sensitivities peak in the lower and mid-upper troposphere, respectively. The similar
477 patterns of SCO₃ enhancement detected by the two sounders therefore indicate that these extend
478 over the bulk of the troposphere, supportive of surface/lower tropospheric ozone enhancements.
479 This consistency also provides confidence that the complementary vertical sensitivities of GOME-2
480 and IASI ozone retrievals could be exploited in further investigation of tropospheric ozone in the
481 future (e.g. long-term trends from multiple platforms/retrieval schemes have shown large-scale
482 inconsistencies in other studies e.g. Gaudel et al., (2018)).

483 Tropospheric O₃ behaviour is complex and the summer 2018 enhancements over Europe could
484 potentially have been caused by various factors: atmospheric chemistry, meteorology (e.g.
485 temperature, advection of O₃-rich air masses), anthropogenic and natural precursor emissions, dry
486 deposition and stratospheric intrusion. To investigate the interactions between these processes,
487 potentially leading to the summer 2018 O₃ enhancements, we used the well-evaluated TOMCAT 3D
488 CTM. Evaluation of the model in this study showed that it could accurately capture the spatial
489 pattern, temporal evolution and sign (i.e. positive 2018-2017 O₃ differences) of the O₃
490 enhancements and that, although it underestimated the observed enhancements, TOMCAT is an
491 adequate tool to investigate them.

492 The results of several model simulations showed that the surface ozone enhancements (mainly in
493 north-western Europe) in the summer of 2018 were predominantly driven by meteorological
494 processes with emissions acting as a secondary factor. As the ROTRAJ back-trajectories suggest that
495 advection of summer-time O₃ was larger in 2017, the 2018 European O₃ enhancements at surface
496 level were likely from in-situ processes. The TOMCAT stratospheric O₃ tracer indicated a negligible
497 contribution of stratospheric O₃ to these surface enhancements. At 500 hPa, the enhancement in
498 tropospheric O₃ is much larger spatially across Europe and dominated by meteorological processes.
499 Intrusion of stratospheric O₃ into the mid-troposphere has a moderate influence on the
500 observed/modelled O₃ enhancements with contributions of up to 15.0-40.0%. Correlations between
501 TOMCAT temperature and O₃ enhancements show broad agreement at some latitudes (e.g. 50-70°N
502 in the lower-mid troposphere). However, this relationship is non-linear and difficult to quantify
503 without further simulations/model tracers, which was beyond the scope of this study. ROTRAJ back-
504 trajectories suggest that in 2018, relative to 2017, there is the advection of more O₃-rich airmasses
505 into the European mid-troposphere contributing to the summer 2018 O₃ enhancements at this
506 altitude. Therefore, in the summer of 2018 over Europe, in-situ meteorological processes appear to
507 be predominantly driving surface O₃ enhancements over Europe, while advection of tropospheric O₃-
508 rich air and stratospheric intrusion are driving the corresponding tropospheric O₃ enhancements

509 Overall, through our study focusing on the European summer 2018 air pollution episode, we have
510 demonstrated the use of novel satellite datasets and a modelling framework (i.e. targeted sensitivity
511 experiments and model tracers) suitable to investigate the air quality impacts from future European
512 heatwaves such as that which occurred in summer 2022.

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516 **Conflicting Interests**

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

518

519 **Date Availability**

520 The TOMCAT simulations are publicly available at

521 http://homepages.see.leeds.ac.uk/~earrjpo/european_summer_2018_o3/tomcat, while the RAL
522 Space satellite can be found at

523 http://homepages.see.leeds.ac.uk/~earrjpo/european_summer_2018_o3/satellite. The EMEP
524 surface O₃ data was obtained from <http://ebas-data.nilu.no/default.aspx>. The GOME-2 tropospheric
525 column NO₂ data was downloaded from EUMETSAT at https://acsaf.org/nrt_access.php. The
526 TOMCAT and RAL Space satellite data will be uploaded to the Zenodo open access portal
527 (<https://zenodo.org/>) if this manuscript is accepted for publication in ACP after the peer-review
528 process.

529 **Author Contributions**

530 RJP, MPC and BJK conceptualised and planned the research study. RJP performed the TOMCAT
531 model simulations with support from MPC and WF. The JULES BVOC emissions were provided by ECP
532 and GDH. RJP analysed the satellite data provided by RAL Space (BJK, RS, BGL and LJV) with support
533 from BJK, RS and BGL. RJP undertook the EMEP analysis. RJP ran ROTRAJ with technical support from
534 SRA and AMG. RJP prepared the manuscript with contributions from all co-authors.

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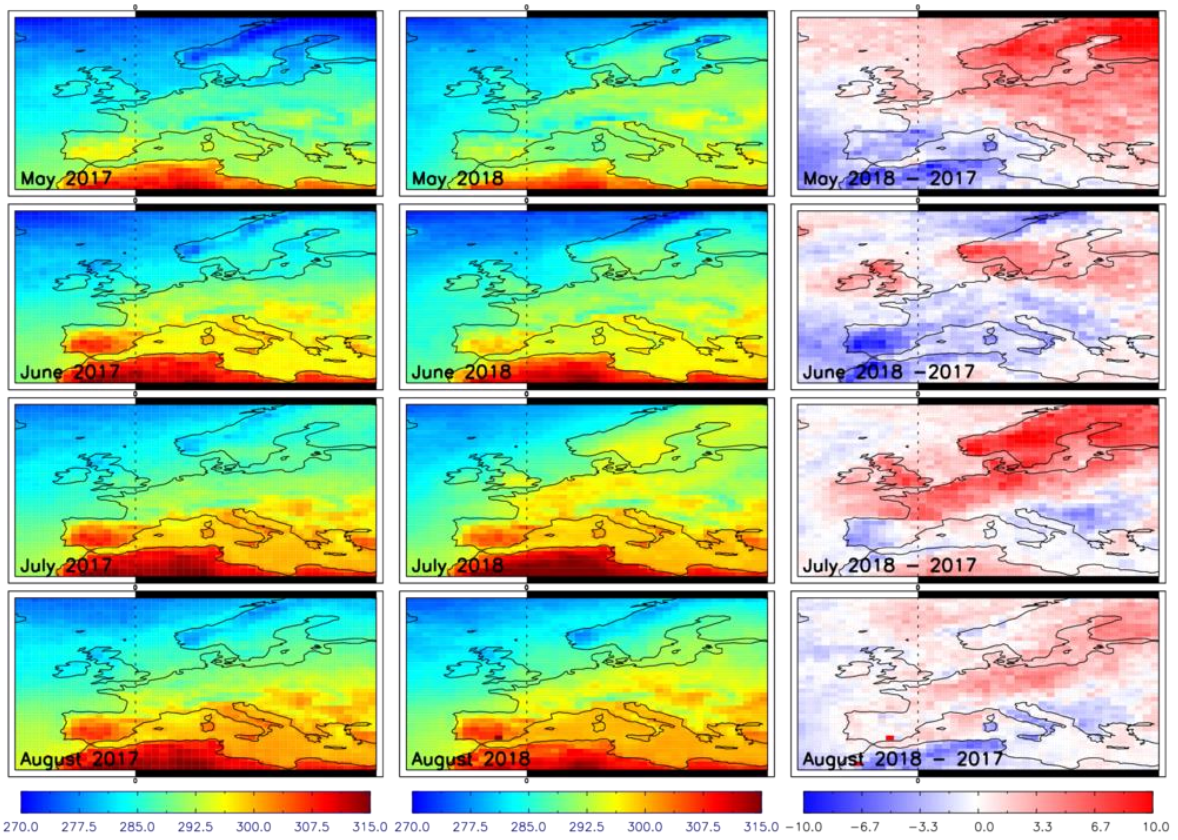
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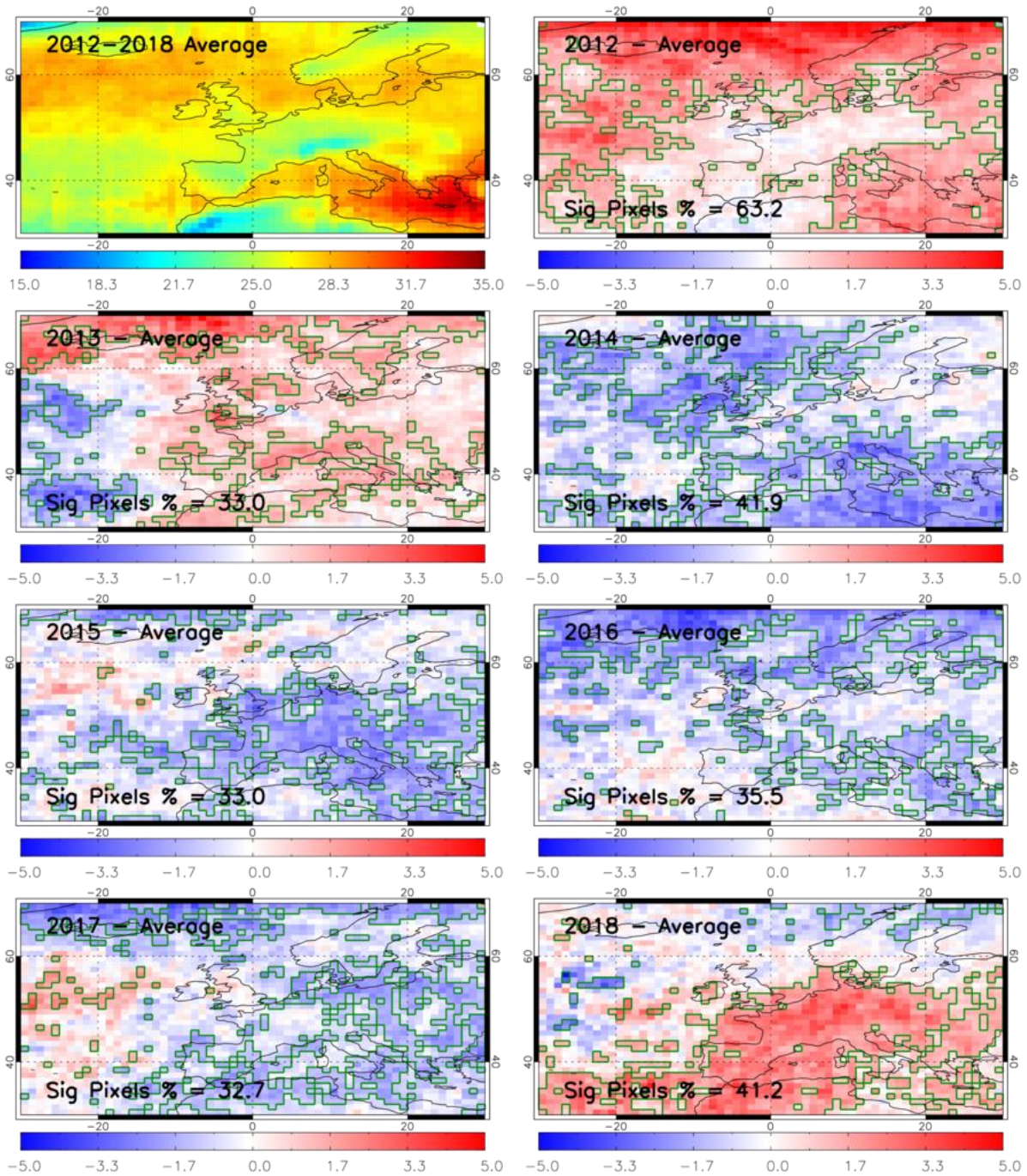
721 Figures:



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723 **Figure 1:** Surface temperature (K) over Europe for May to August in 2017 (left column), 2018 (centre
724 column) and 2018-2017 difference (right column) retrieved from MetOp-A IASI, MHS and AMSU by
725 the IMS scheme.

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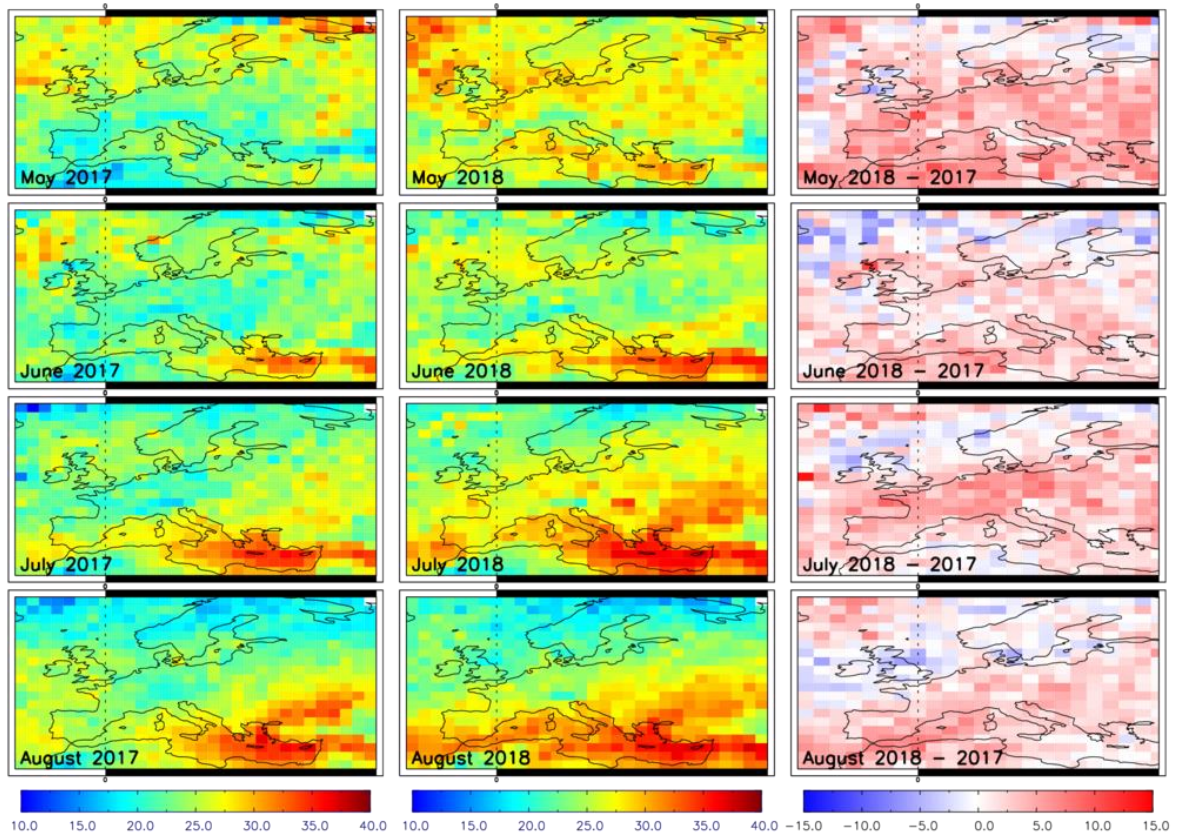
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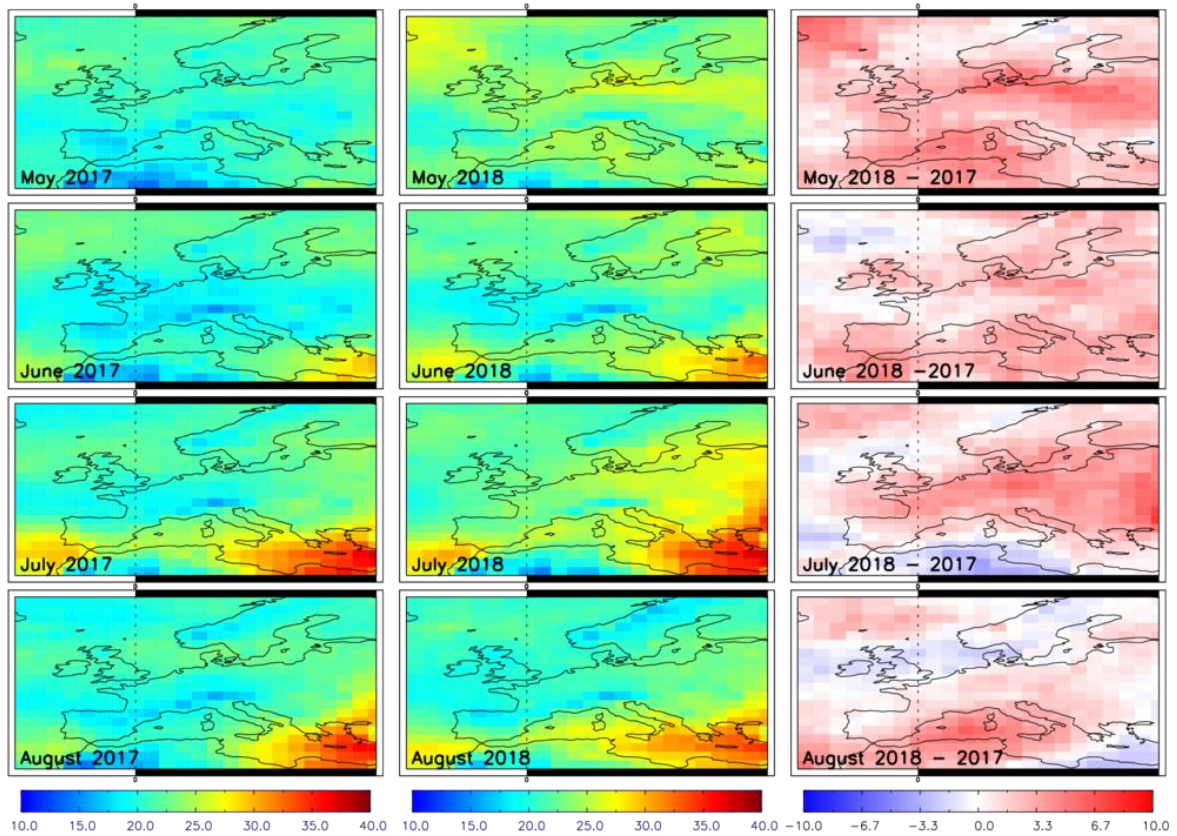
Figure 2: Sub-column ozone (SCO_3 , surface-450 hPa), in Dobson units (DU), retrieved from GOME-2 on Metop-A averaged across May to August between 2012 and 2018 (top left panel) and the corresponding difference from the 2012-18 mean for each year, respectively. The green-polygon-outlined regions show where the year-specific seasonal average is significantly different (95% confidence level based on the Wilcoxon Rank Test (WRT)) from the long-term (2012-2018) seasonal average. The “Sig Pixel %” label indicates the number of pixels in the domain with significant differences.



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737 **Figure 3:** SCO₃ (DU) from GOME-2 over Europe for May to August in (left column) 2017, (centre
 738 column) 2018 and (right column) 2018-2017 difference.

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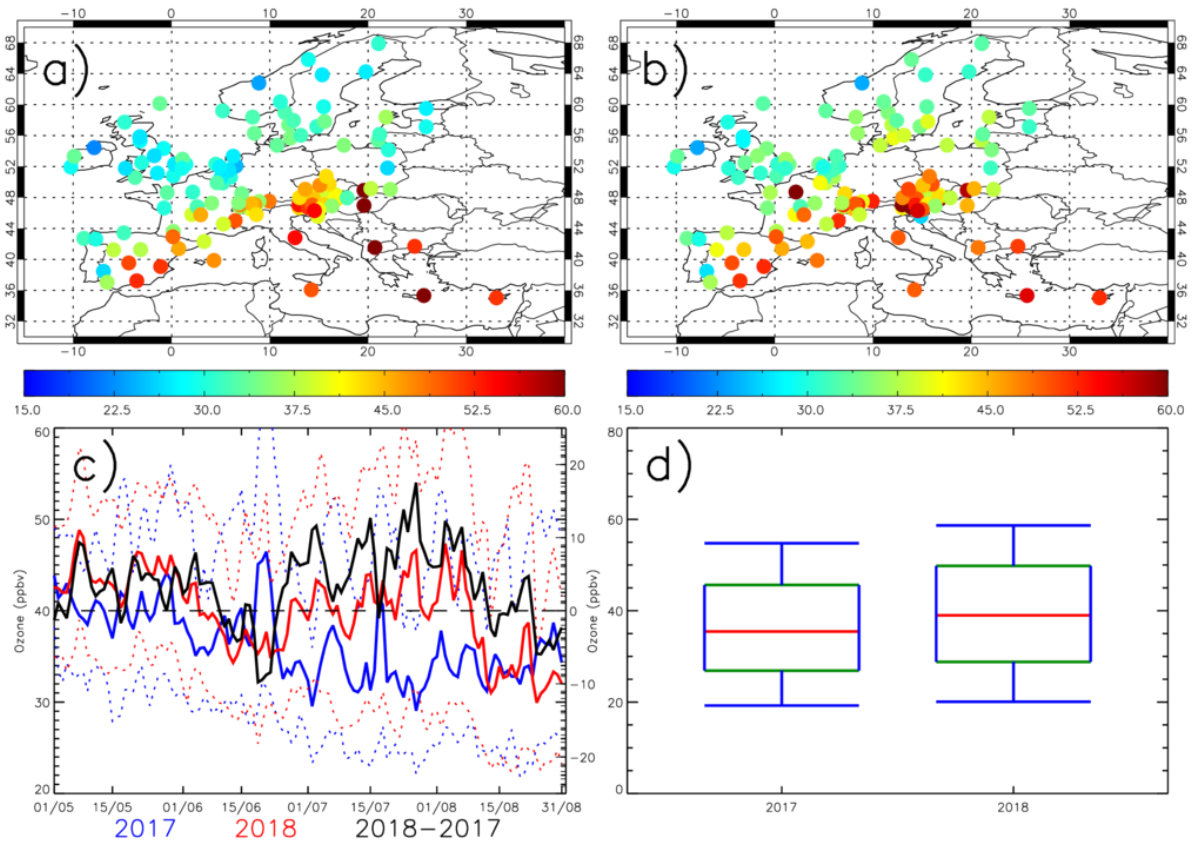
Figure 4: *SCO₃ (DU) for May to August in 2017 (left column), 2018 (centre column) and 2018-2017 difference (right column) over Europe retrieved from MetOp-A IASI, MHS and AMSU by the IMS scheme.*

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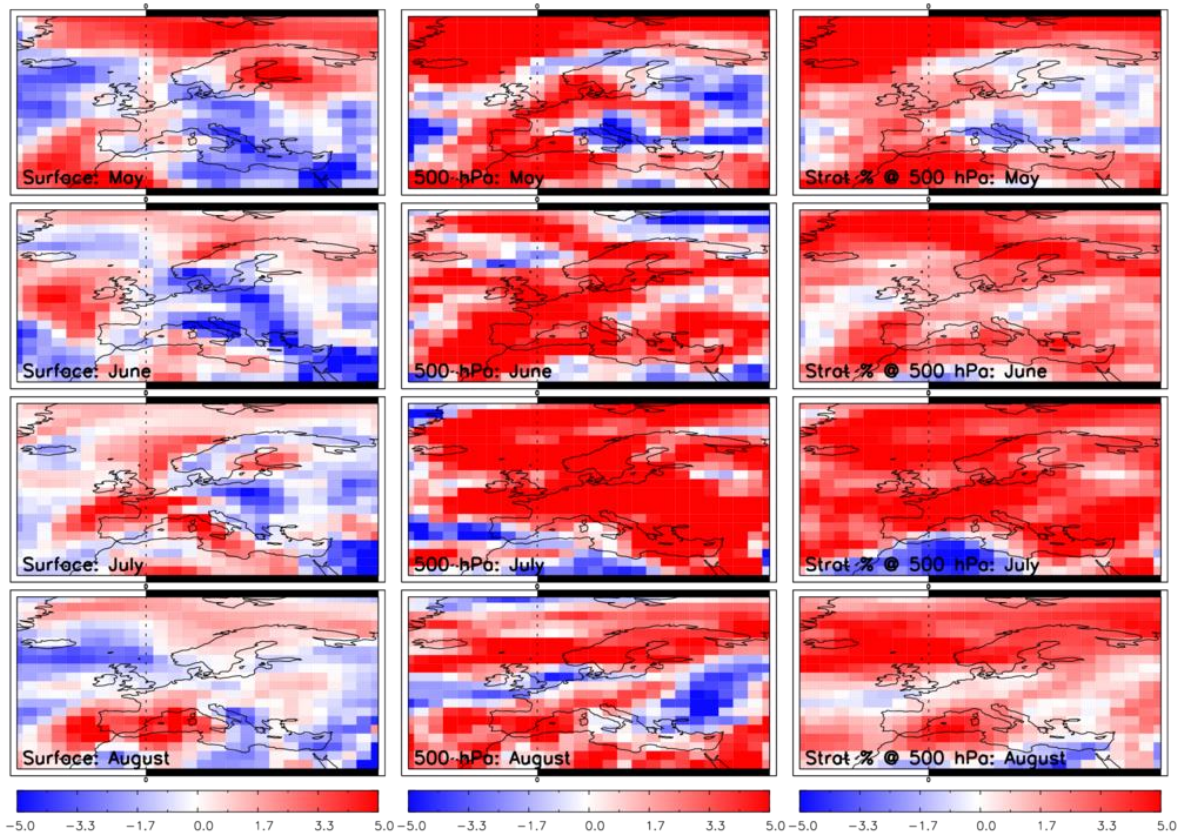
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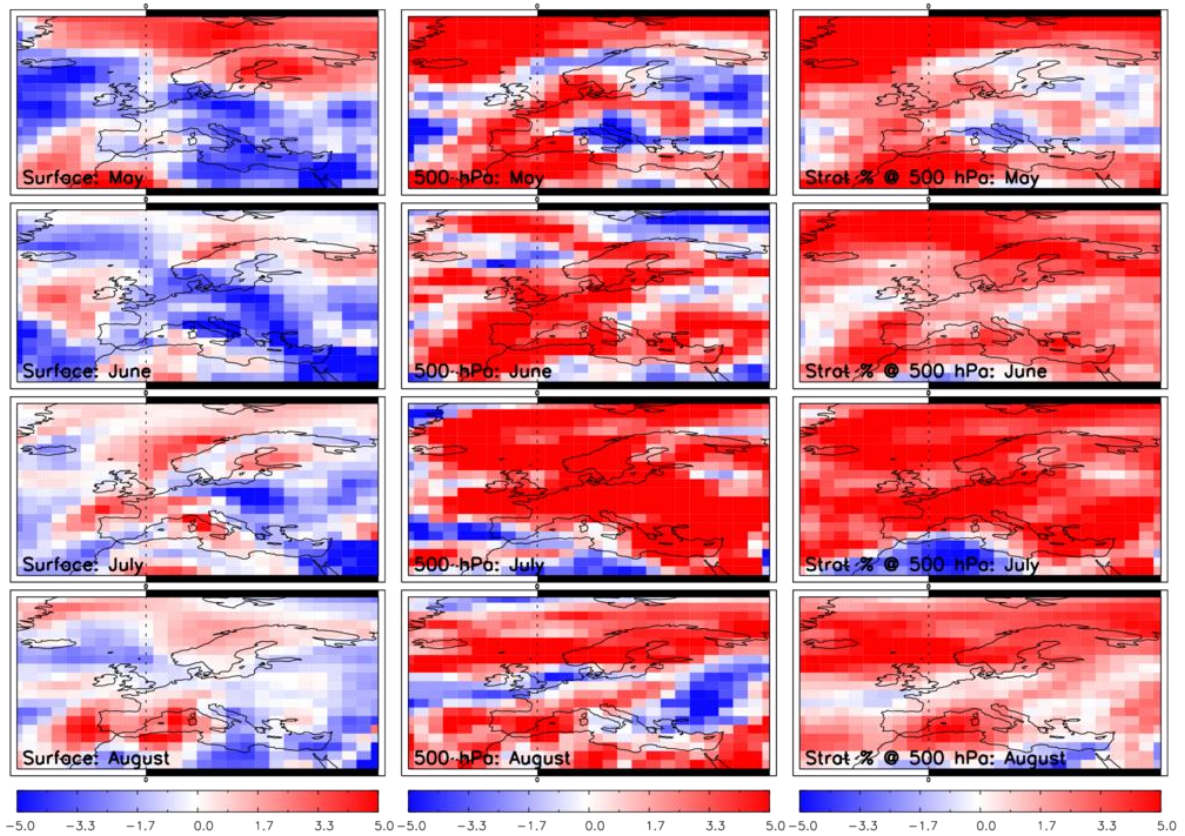
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Figure 5: European surface ozone (ppbv) for a) May-June-July-August (MJJA) 2017, b) MJJA 2018), c) regional mean time series (dotted lines show mean \pm standard deviation) for MJJA 2017 (blue), MJJA 2018 (red) and the 2018-2017 difference (black) and d) box-whisker plots for MJJA 2017 and 2018. In panel d) the median, 25th & 75th percentiles and 10th & 90th percentiles are shown by the red, green and blue lines, respectively.



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Figure 6: TOMCAT ozone (ppbv) 2018-2017 differences for May to August for the surface (left column), 500 hPa (centre column) and the stratospheric contribution (%) to the 500 hPa layer (right column).



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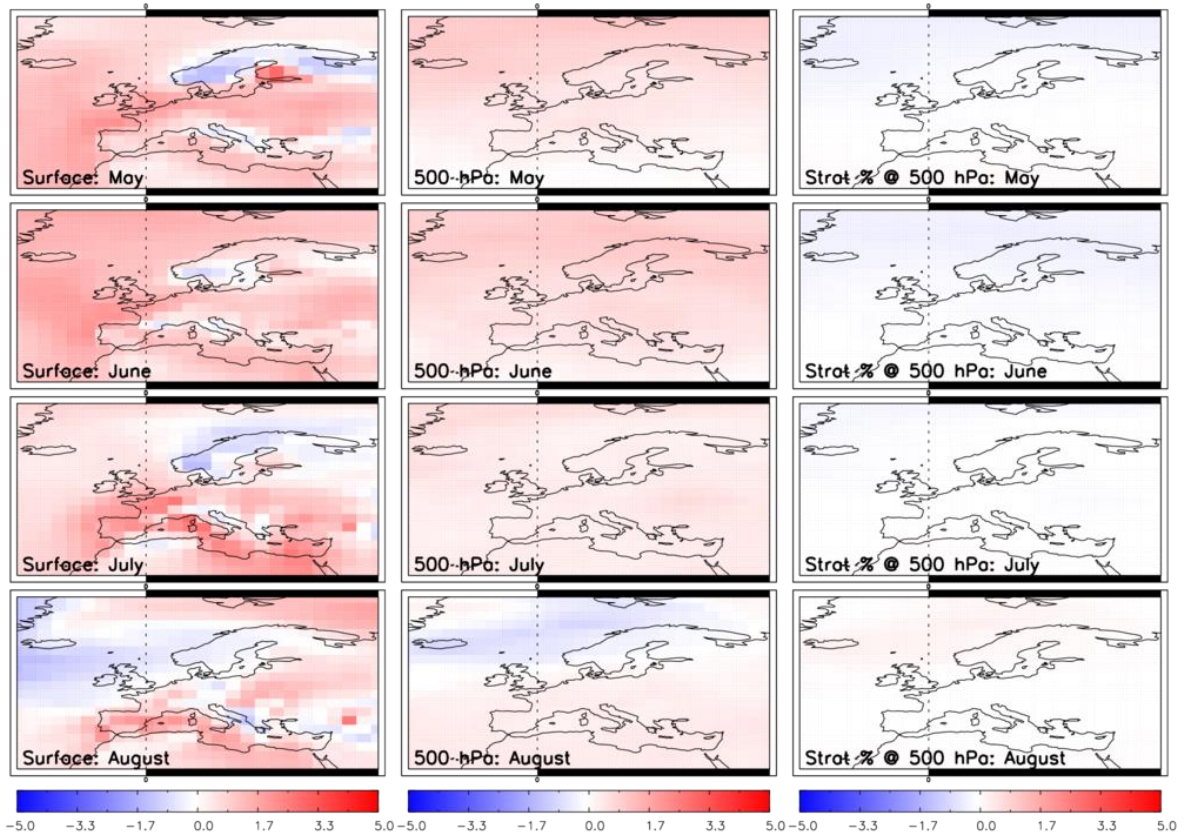
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Figure 7: TOMCAT ozone (ppbv) 2018-2017 differences for May to August for the fixed emissions simulation (Fixed_EMIS) for the surface (left column), 500 hPa (centre column) and the stratospheric contribution (%) to the 500 hPa layer (right column).



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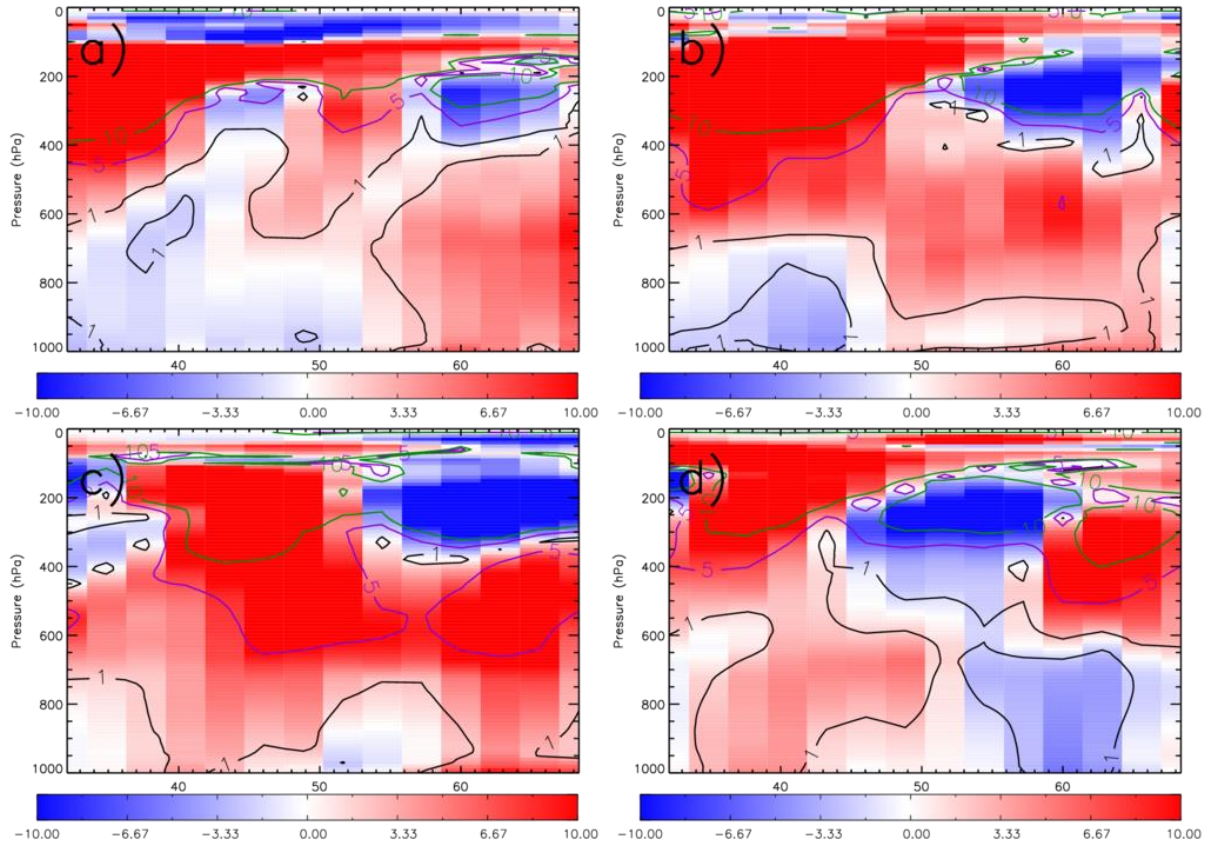
Figure 8: TOMCAT ozone (ppbv) 2018-2017 differences for May to August for the fixed meteorology simulation (Fixed_MET) for the surface (left column), 500 hPa (centre column) and the stratospheric contribution (%) to the 500 hPa layer (right column).

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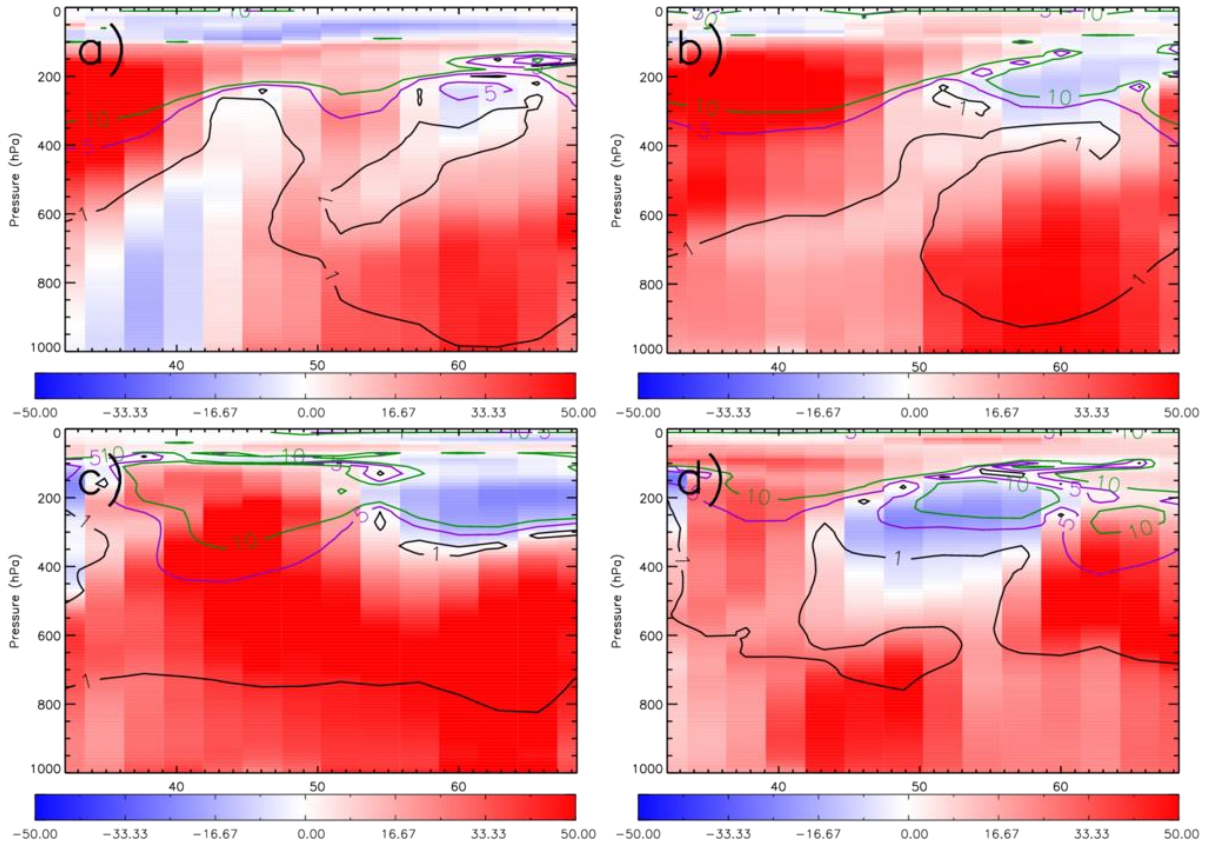
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772 **Figure 9:** TOMCAT ozone, zonally averaged between 20°W and 40°E, 2018-2017 percentage
 773 differences (absolute difference (ppbv) shown as solid lines) from the control simulation. Panels a)-d)
 774 represent the monthly averages for May, June, July and August.

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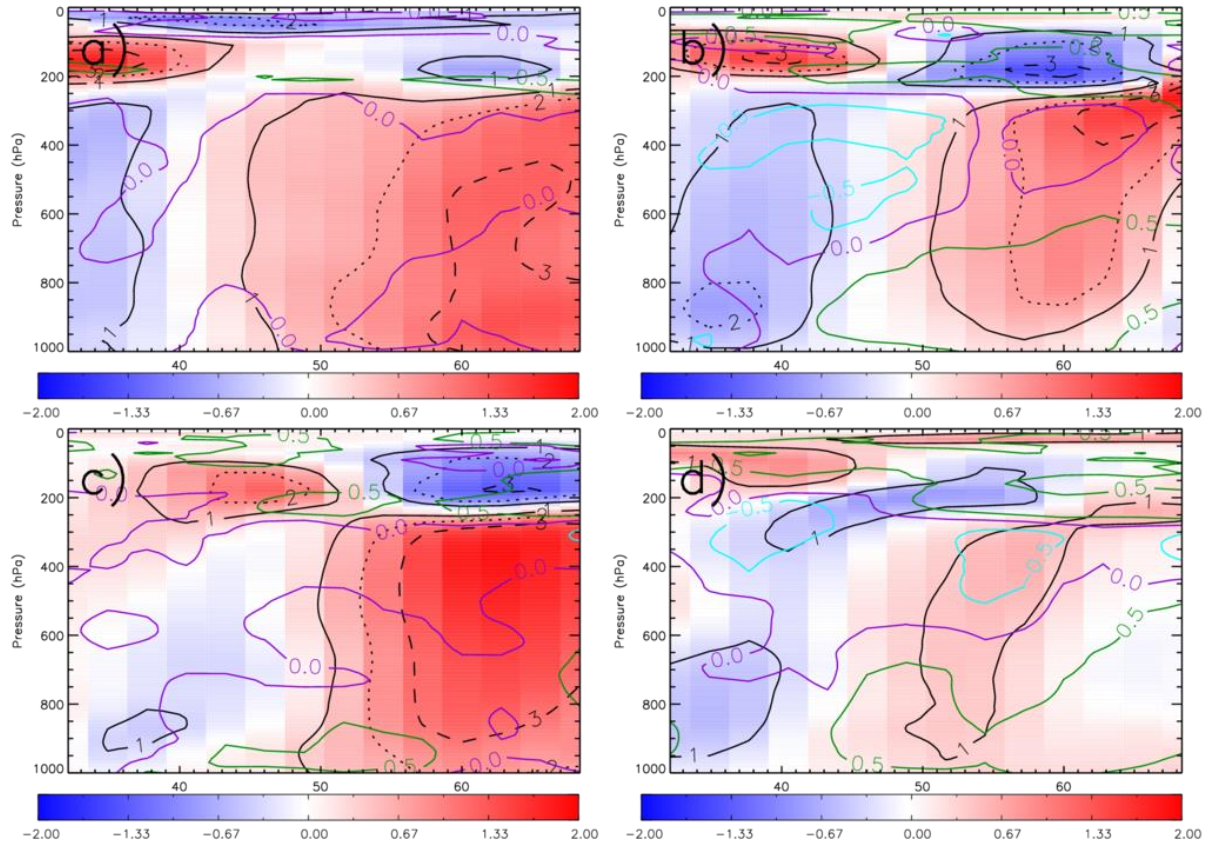
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777 **Figure 10:** TOMCAT stratospheric ozone tracer, zonally averaged between 20°W and 40°E, 2018-2017
 778 percentage differences (absolute difference (ppbv) shown as solid lines) from the control simulation.
 779 Panels a)-d) represent the monthly averages for May, June, July and August.

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784 **Figure 11:** TOMCAT temperature, zonally averaged between 20°W and 40°E, 2018-2017 percentage
 785 differences (absolute difference (K) shown by black solid, dotted and dashed lines) from the control
 786 simulation. Overplotted are contours of the temporal correlation (i.e. within each grid box) between
 787 the temperature and ozone 2018-2017 differences. Panels a)-d) represent the monthly averages for
 788 May, June, July and August.

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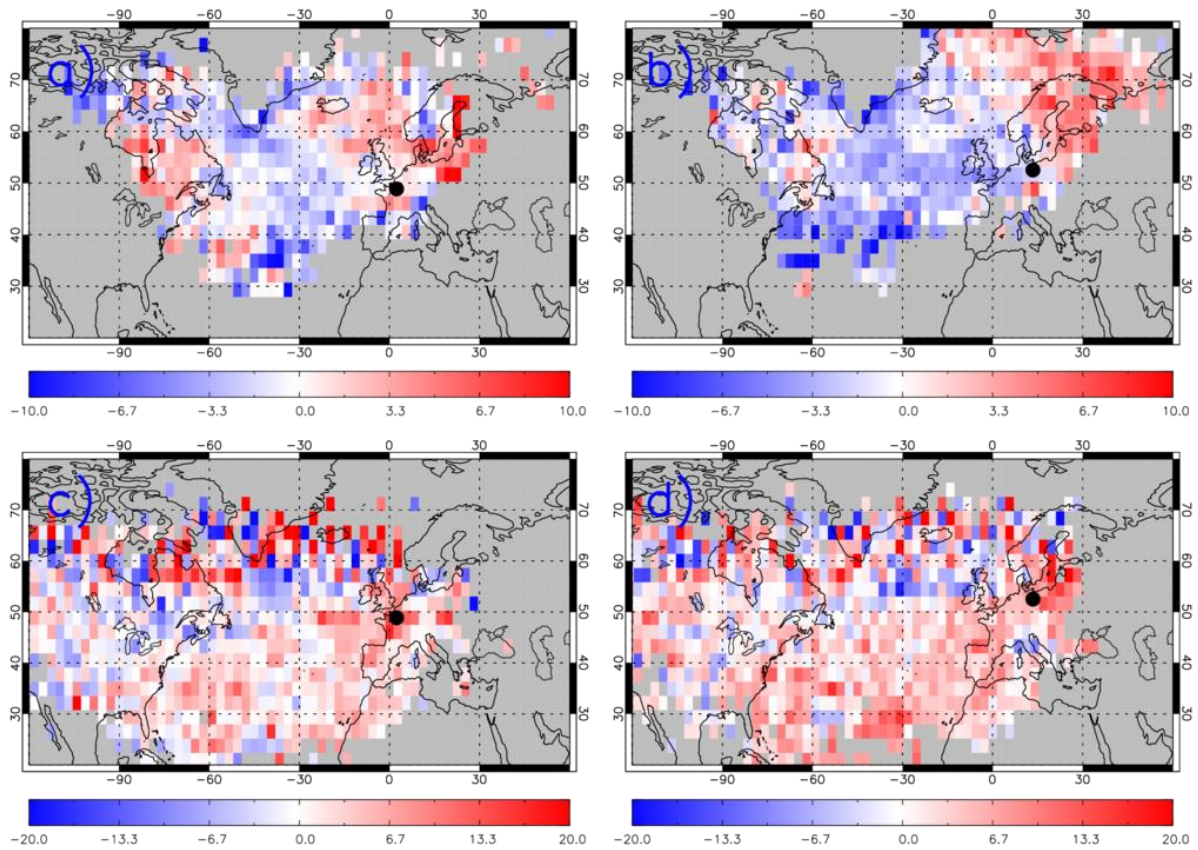
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Figure 12: The difference between May-August 2018 and May-August 2017 (i.e. 2018-2017) ROTRAJ back-trajectories (10 days), weighted by the average TOMCAT O₃ (ppbv) concentration along each trajectory path, gridded onto the TOMCAT horizontal resolution for a) Paris at the surface, b) Berlin at the surface, c) Paris at approximately 500 hPa and d) Berlin at approximately 500 hPa. The black circles represent the location of Paris or Berlin, where the trajectories were released from.