

1 **The differing impact of air stagnation on summer ozone across Europe**

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13 **Keywords:** Air quality, air pollution, ozone, stagnation, meteorology

14 **Abstract**

15
16 We have examined the joint impact of daily maximum temperature and air stagnation on
17 observed maximum daily 8-h running average (MDA8) and hourly near-surface ozone
18 (O₃) over eight regions of Europe for the summer period 1998–2015. The percentage of
19 stagnant area in a region is a better predictor of summer MDA8 O₃ in central/southern
20 Europe (correlation coefficient R=0.50–0.70) than in the north (R=0.06–0.39). The
21 correlations of MDA8 O₃ with temperature are higher than with stagnation for most
22 regions, and the impact of stagnation on those correlations is region-dependent. MDA8
23 O₃ mixing ratios consistently increase over central/southern Europe under stagnant
24 conditions, but this has not been found for some temperature ranges in the north. Under
25 non-stagnant situations and relatively high temperatures (20–25 °C), southerly advection
26 often brings aged air masses from more polluted areas to the receptor regions in the north.
27 Furthermore, the occurrence of stagnation tends to amplify the ozone diurnal cycles in
28 the center/south of the continent, yielding lower nighttime and higher daytime mixing
29 ratios than on non-stagnant days, again with a less clear effect in the north. Changes in
30 planetary boundary layer height, accumulation of ozone and precursors, and subsequent
31 photochemical production during stagnant days are the presumed underlying mechanisms
32 for the amplification of the diurnal cycles. The results from this study prove that the
33 effects of stagnation on summer ozone are regionally dependent across Europe.
34 Consequently, climate model projections of increases in stagnation should not directly be
35 translated into degraded air quality without a proper assessment of the regional impacts.

36 **1. Introduction**

37 Tropospheric ozone (O₃) is one of the main pollutants linked to health and
38 environmental impacts (Ashmore, 2005; REVIHAAP, 2013; Tai et al., 2014). It is
39 transported from the stratosphere, and produced by photochemical reactions between
40 primary precursors such as nitrogen oxides (NO_x) and volatile organic compounds
41 (VOCs). Observational studies have shown strong relationships of O₃ with
42 meteorological parameters. Understanding such relationships is often complicated by the
43 covariance of the meteorological variables as well as by the effect of short- and long-term
44 emission changes. Ambient air temperature is the most correlated variable with daily
45 ozone maxima in summer (Bloomfield et al., 1996; Xu et al., 1996; Kuebler et al., 2001;
46 Tarasova and Karpetchko, 2003; Ordóñez et al., 2005). The relationship between both
47 variables is driven by several well-known temperature-dependent mechanisms: the
48 thermal decomposition of the NO_x reservoir species peroxyacetyl nitrate (PAN,
49 CH₃COO₂NO₂) (Orlando et al., 1992), the temperature-dependent emissions of both
50 VOCs from vegetation and NO_x from soil (Fehsenfeld et al., 1992; Simpson, 1995), the
51 extra evaporation of anthropogenic VOCs at high temperatures (Vautard et al., 2005), or
52 the high stomatal resistance at elevated temperatures, which limits the dry deposition of
53 ozone to vegetation (Wesely, 1989). Nevertheless, the temperature – O₃ relationship is
54 complex and can only partially be explained by the mentioned processes. As an example,
55 Porter and Heald (2019) have estimated that the covariance of temperature with other
56 meteorological variables explains over 40% of the ozone-temperature correlation in the
57 United States (US) and up to 60% in Europe. There are a number of relevant non-
58 temperature meteorological processes that may contribute to this covariance. Among
59 others, air stagnation is a potential candidate as it is characterized by stable weather
60 conditions with poor ventilation, leading to the accumulation of pollutants close to the
61 surface (Jacob and Winner, 2009; Dawson et al., 2014; Fiore et al., 2015).

62 Enhanced ozone concentrations under stagnant conditions have been documented
63 for the US (Leibensperger et al., 2008; Schnell and Prather, 2017) and Europe (Garrido-
64 Perez et al., 2018). The relationship between a simplified air stagnation index (ASI)
65 defined by Horton et al. (2012) – based on absolute thresholds of daily precipitation, 10
66 m and 500 hPa wind speed –, temperature and O₃ has been closely examined by some
67 studies for the US. Sun et al. (2017) found that the probability of high O₃ days increases
68 with the co-occurrence of high temperature and stagnation, and that the occurrence of
69 successive stagnation days further enhances this likelihood. Schnell and Prather (2017)
70 showed that air stagnation and extreme levels of O₃ and temperature tend to co-occur in
71 large-scale, multiday and coherent structures. In Europe, Garrido-Perez et al. (2018)
72 reported significant changes in the frequency distribution of O₃ when stagnation occurs
73 as well as a strong build-up of the O₃ mixing ratios during widespread stagnant episodes.
74 These evidences are in line with the hypothesis of Horton et al. (2012), who projected 21st
75 century increases in stagnation frequency over several highly populated regions of the
76 globe and suggested a potential to enhance the accumulation of near-surface ozone.
77 However, the relationship between stagnation and summer ozone has been reported to be
78 weak over some regions of the US (Kerr and Waugh, 2018). That study noted the limited
79 skill of the ASI to explain air pollution events and indicated an erratic overlap between
80 stagnation and pollutant extremes in the US, especially in the Northeast. Consequently,
81 they advised against the use of this common index as a metric to examine pollution events
82 and recommended testing new indices considering other meteorological predictors such
83 as planetary boundary layer (PBL) height and temperature.

84 Building on the results by Kerr and Waugh (2018) for the US, the aims of this
85 study are to (i) evaluate whether the ASI is an adequate index of summer ozone pollution
86 over different regions of Europe and (ii) analyze the complementarity of this index with
87 temperature to understand summer ozone variability in the continent. To do so, we firstly
88 investigate the temporal co-variability of ozone, temperature and air stagnation. Then we
89 evaluate whether the relationship between O₃ and temperature differs under stagnant and
90 non-stagnant situations. Finally, we explore the diurnal cycle of O₃ in order to understand
91 the response of this pollutant to stagnation and the mechanisms involved.

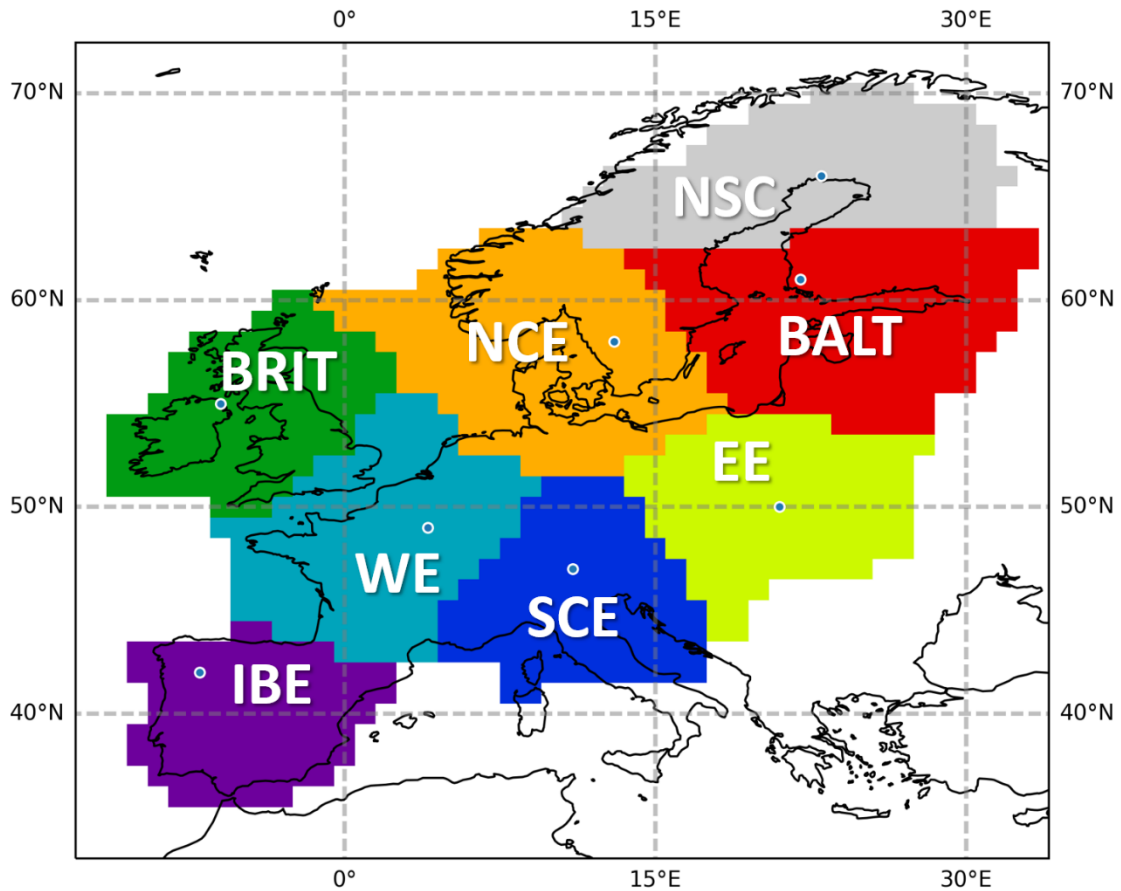
92 **2. Data and methods**

93 **2.1. Ozone and meteorological data**

94 We use interpolated datasets of observed maximum daily 8-h running average
95 near-surface ozone (MDA8 O₃) and hourly ozone volume mixing ratios over Europe at
96 1.0° × 1.0° resolution for the 18 summers (JJA) during the period 1998 – 2015. The
97 summer means of MDA8 O₃ from these datasets, excluding data over the Balkans (see
98 below), are illustrated in Supplementary Figure S1. The data have been generated by
99 merging non-traffic observations obtained from two databases – the European
100 Environment Agency’s air quality database (AirBase) and the European Monitoring and
101 Evaluation Programme (EMEP) (Tørseth et al., 2012) – with the objective mapping
102 algorithm developed by Schnell et al. (2014, 2015). These data have recently been used
103 by several studies (Otero et al., 2016, 2018; Carro-Calvo et al. 2017; Ordóñez et al., 2017;
104 Garrido-Perez et al., 2018) to understand the meteorological drivers of ozone in Europe.
105 In particular, Carro-Calvo et al. (2017) applied the k-means clustering technique to obtain
106 nine regions where O₃ shows coherent spatiotemporal patterns. The regions from that
107 study will be used here to examine the relationship among air stagnation, temperature and
108 O₃ in a consistent way. However, we have decided to exclude one of the regions (the
109 Balkans) from our analysis because of inhomogeneities in the O₃ time series reported by
110 that study. From west to east and from north to south, the regions analyzed here roughly
111 cover the British Isles (BRIT), northern-central Europe (NCE), northern Scandinavia
112 (NSC), the Baltic region (BALT), the Iberian Peninsula (IBE), western Europe (WE),
113 southern-central Europe (SCE) and eastern Europe (EE). This regionalization is displayed
114 on Figure 1.

115 We also use daily mean fields of sea level pressure (SLP); daily mean 10 m
116 (\hat{U}_{10m}), 850 hPa (\hat{U}_{850hPa}) and 500 hPa (\hat{U}_{500hPa}) wind; daily accumulated precipitation
117 (pr); daily maximum temperature (T_{max}), and PBL heights from the European Centre
118 for Medium-Range Weather Forecasts (ECMWF) ERA-Interim reanalysis (Dee et al.,
119 2011). We regridded the original 0.75° × 0.75° horizontal resolution data onto the 1.0° ×
120 1.0° grid of O₃ using bilinear interpolation. \hat{U}_{10m} , \hat{U}_{500hPa} and pr data are needed for the
121 calculation of the ASI (see Section 2.2), while SLP and \hat{U}_{850hPa} are employed for the
122 comparison of the atmospheric circulation patterns in the lower troposphere during
123 stagnant and non-stagnant days. Daily averages of SLP and wind speed have been
124 computed as the average of the four analysis values at 00, 06, 12 and 18 UTC for each
125 day. pr has been calculated by summing the 12-h accumulated precipitation forecast at
126 00 and 12 UTC. T_{max} has been computed as the maximum from all the temperature values
127 forecasted for each day, following the recommendations at
128 <https://confluence.ecmwf.int/pages/viewpage.action?pageId=58157862> (last access: 20
129 [September 2019](#)). The forecast values of PBL height have also been used at the standard

130 meteorological hours 00, 06, 12 and 18 UTC to better understand the diurnal cycles of
131 O₃.



132
133 **Figure 1.** Regionalization of summer MDA8 O₃. From west to east and from north to
134 south the regions roughly correspond to the British Isles (BRIT), northern-central Europe
135 (NCE), northern Scandinavia (NSC), the Baltic region (BALT), the Iberian Peninsula
136 (IBE), western Europe (WE), southern-central Europe (SCE) and eastern Europe (EE).
137 Blue circles represent the land grid cells where the correlations between the standardized
138 anomalies of the MDA8 O₃ time series and those of the respective centroid maximize (see
139 details in the main text).

140 2.2. Methods

141 Air stagnation is characterized by meteorological conditions that impede the
142 scavenging, horizontal dispersion and vertical escape of pollutants in the lower
143 troposphere (Horton et al., 2012, and references therein). Different air stagnation indices
144 have been proposed in the literature to identify the conditions conducive to air mass
145 stagnation (see e.g. Garrido-Perez et al., 2018, and references therein). Such indices are
146 usually defined by using predefined thresholds for daily average meteorological fields.

147 In this work we have used the simplified air stagnation index defined by Horton
148 et al. (2012). This ASI is an adaptation of another index originally developed for the US
149 that is currently being used by the US National Oceanic and Atmospheric Administration
150 (NOAA) National Climatic Data Center (NCDC) (Wang and Angell, 1999). With the
151 application of Horton's ASI, a reanalysis grid cell is considered as stagnant if three
152 conditions are simultaneously met on a given day: $\bar{U}_{10m} < 3.2 \text{ m s}^{-1}$, $\bar{U}_{500hPa} < 13.0 \text{ m s}^{-1}$
153 and $pr < 1 \text{ mm}$. This results in a high frequency of stagnation in some parts of

154 Mediterranean Europe (around 40% of the days), moderate frequency in central Europe
155 and Scandinavia (~25% of the days) and low over the British Isles (~15% of the days)
156 during the summer months of 1998–2015 (see Figure S2).

157 The index has recently been used to characterize the spatiotemporal variability
158 and impact of air stagnation on air quality in Europe (Garrido-Perez et al., 2018), but the
159 relationship between stagnation and summer ozone over different parts of the continent
160 might be sensitive to the ASI component threshold definition. To account for this, we
161 have repeated some of the analyses presented here by varying the three thresholds from
162 25 to 175% of their standard values. Overall, we have found relatively small changes both
163 in the ozone–stagnation correlations (Table 1) and the ozone dependence on stagnation
164 (Figure 3) in all regions when the three thresholds are within 75–125% of their standard
165 values (not shown). This indicates that the results presented in this study are not very
166 sensitive to the precise definition of stagnant conditions, as previously found by Kerr and
167 Waugh (2018) for the US. Finally, we would also like to note that more complex air
168 stagnation indices have recently been developed but they are specifically adapted to
169 particulate matter (e.g. Wang et al., 2016, 2018; Huang et al., 2018).

170 **3. Relationship between MDA8 O₃, temperature and stagnation**

171 Table 1 shows the Pearson correlation coefficients (R) between the daily time
172 series of average T_{max}, average MDA8 O₃ and the percentage of the area under stagnant
173 conditions for each region in summer. The significance of these correlations has been
174 determined through two-tailed t-tests as indicated in von Storch and Zwiers (1999), taking
175 into account that the data exhibit serial correlation, i.e. the daily values of these variables
176 are not independent of their own future and past values because of the effect of the
177 meteorological persistence (see e.g. Wilks, 2011). This implies that the effective number
178 of degrees of freedom (v_{eff}) is lower than that directly determined from the sample size,
179 increasing the p-value of the test and reducing the level of statistical significance.
180 Accordingly, we have calculated v_{eff} following Eq. 1 of Hu et al. (2017) before assessing
181 significance. This way, the three fields significantly correlate at the 95% confidence level
182 for all regions, with the exception of the stagnation – MDA8 O₃ correlation in NSC.
183 However, there is considerable spatial heterogeneity across Europe, with higher
184 correlations in the central/southern clusters (IBE, WE, SCE and EE) than in the northern
185 clusters (BRIT, NCE, NSC and BALT). The correlations are above 0.5 over the
186 central/southern regions, with the exception of T_{max} – MDA8 O₃ in IBE and T_{max} –
187 stagnation in EE, while they tend to be well below that value for the northern regions. As
188 expected, the correlations of MDA8 O₃ with T_{max} are higher than with the percentage of
189 stagnant area for most regions, although the latter can be considered as a good predictor
190 of ozone in central/southern Europe. On the other hand, T_{max} and stagnation present high
191 correlations in the southern clusters as well as in NCE, evidencing the strong covariability
192 between both variables. Hence, it is unclear whether the high correlations between O₃ and
193 stagnation in the southern regions reflect the ozone-temperature relationship or are due to
194 other processes.

195 **Table 1.** Pearson product-moment correlation coefficients (R) between the daily time
196 series of average T_{max}, average MDA8 O₃ and the percentage of stagnant area (AS) for
197 each region during summer (JJA) 1998 – 2015. The geographical locations of the regions
198 are displayed in Figure 1. All values are statistically significant at the 95% confidence
199 level, except where noted by asterisk (*).

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	BRIT	NCE	NSC	BALT	IBE	WE	SCE	EE
$T_{\max} - \text{MDA8 O}_3$	0.18	0.52	0.35	0.48	0.42	0.70	0.77	0.73
AS - MDA8 O₃	0.24	0.39	0.06*	0.27	0.56	0.62	0.70	0.50
$T_{\max} - \text{AS}$	0.23	0.47	0.24	0.33	0.51	0.53	0.58	0.44

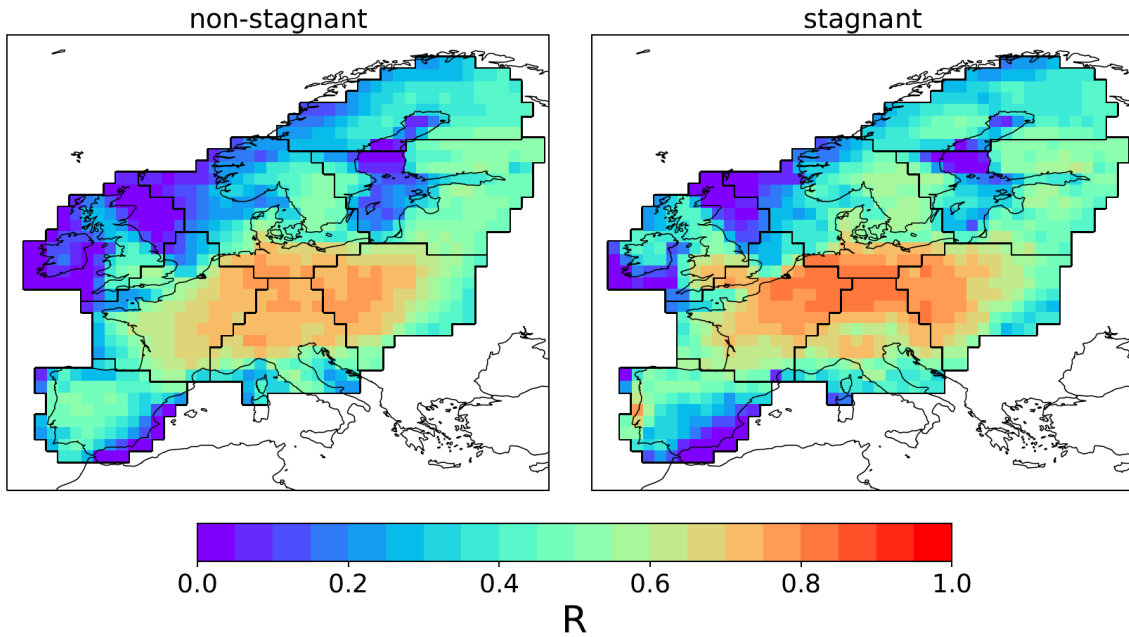
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203 To try to understand the influence of air stagnation in the $T_{\max} - \text{MDA8 O}_3$
 204 relationship, we have evaluated if this relationship changes with the occurrence of
 205 stagnation. For this purpose, we have first computed the correlation coefficients between
 206 the daily time series of T_{\max} and MDA8 O₃ for each grid cell, considering stagnant and
 207 non-stagnant days separately (Figure 2). The spatial distribution of the correlations
 208 displayed in the figure is consistent with that shown in Table 1, as the highest values are
 209 generally found for WE, SCE and EE. Nevertheless, Figure 2 presents a complex picture
 210 of the effect of stagnation on the $T_{\max} - \text{MDA8 O}_3$ relationship across Europe. On the one
 211 hand, stagnation seems to contribute to strengthening this relationship over large parts of
 212 the three regions that present high correlations. Regions characterized by relatively low
 213 ozone levels and stagnation frequency (e.g. the UK and NCE, see Figures S1 and S2) also
 214 present higher correlations when stagnation occurs. These results are in line with a
 215 previous study that has highlighted the contribution of stagnation to increasing the
 216 probability of occurrence of a high ozone day given a high-temperature day in the US
 217 (Sun et al., 2017). On the other hand, there are some parts of the south and east of the
 218 continent where these correlations do not increase with the occurrence of stagnation. The
 219 correlations even decrease over an area roughly covering the southeastern half of IBE,
 220 where stagnation is relatively common in summer (see Figure S2) and the day-to-day
 221 ozone variability – as measured by the relative standard deviation of summer MDA8 O₃
 222 – is lower than in the rest of the continent (not shown).

223 From these results, no clear conclusions can be drawn on the contribution of
 224 stagnation to the ozone-temperature relationship. Actually, these correlations may depend
 225 on a number of factors whose effects are difficult to disentangle. First, the range of
 226 variability of both daily temperature and ozone differs considerably across Europe.
 227 Second, the ozone-temperature relationship tends to be linear only within specific
 228 temperature ranges (e.g. Bloomer et al., 2009), high enough to favor photochemical
 229 production but below some high thresholds that may lead to ozone suppression at some
 230 locations (Steiner et al., 2010; Shen et al., 2016; Meehl. et al., 2018). Finally, the
 231 strengthening of the O₃- T_{\max} correlations under stagnant conditions found for some
 232 regions might partly occur because elevated temperatures often coincide with stagnant
 233 days. Henceforth, to remove as much as possible the influence of the different factors that
 234 limit our understanding of the impact of stagnation on ozone, we will perform composite
 235 analyses within specific temperature bins separately for each region.

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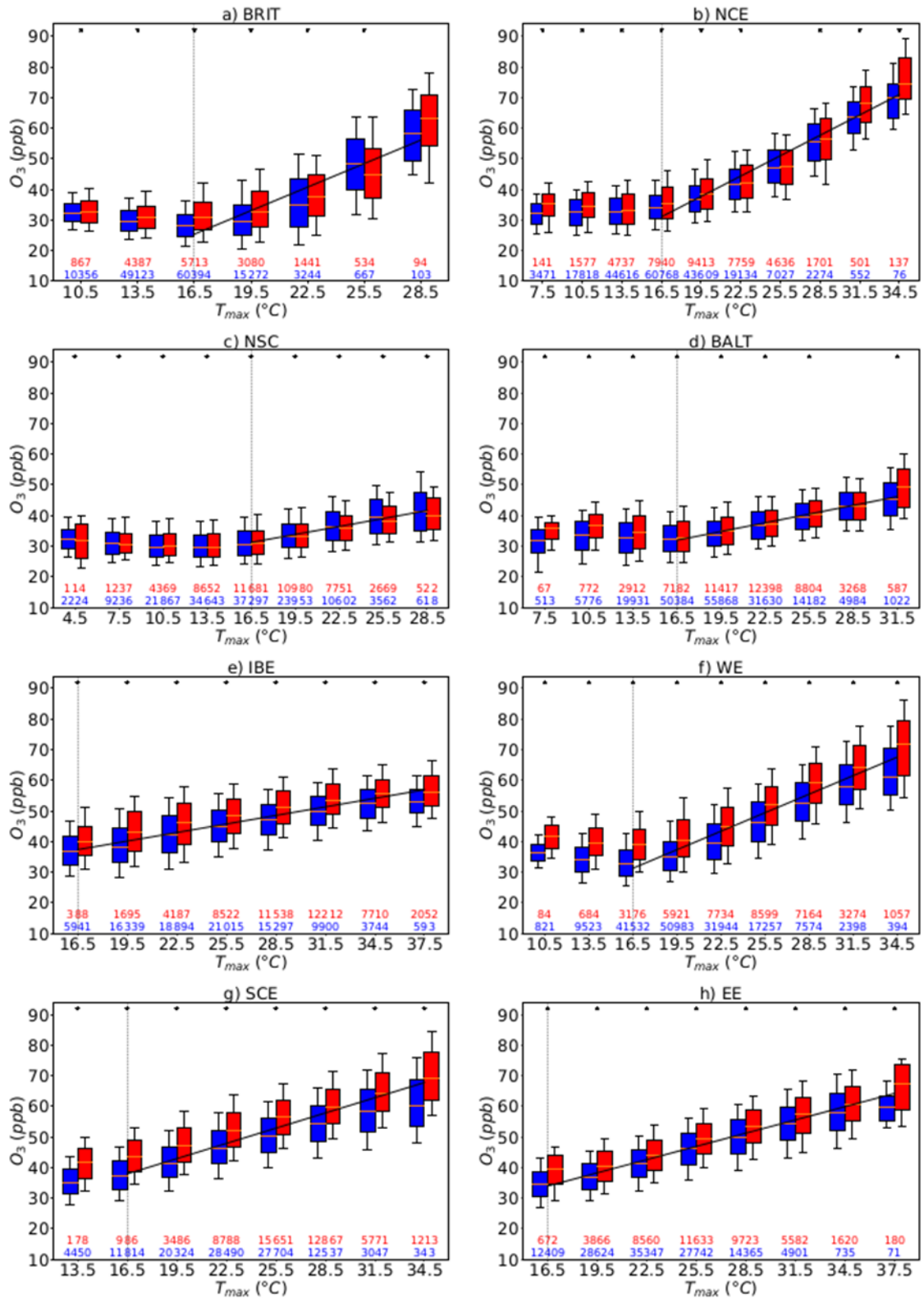
239 **Figure 2.** Pearson product-moment correlation coefficient (R) between the 1998 – 2015
 240 summer (JJA) daily time series of T_{\max} and MDA8 O_3 for days without (left) and with
 241 (right) stagnation in each grid cell.

242 Figure 3 shows boxplots of MDA8 O_3 within 3 °C temperature bins for days with
 243 (red) and without (blue) stagnation, considering every grid cell within each region.
 244 Overall, the relationship of ozone with temperature starts to be linear above ~16 °C, with
 245 coefficients of determination (R^2) above 0.93 for all regions. The ozone distributions
 246 under stagnant and non-stagnant conditions are significantly different at the 95%
 247 confidence level for most temperature bins and regions, as determined through the two-
 248 sample Kolmogorov-Smirnov test (e.g. Wilks, 2011) and indicated by stars on top of the
 249 pairs of boxes. In central/southern Europe, the presence of stagnant conditions causes
 250 consistently higher MDA8 O_3 regardless of the temperature bin (around 4–6 ppb on
 251 average considering the mean values of each bin), but this is not always the case for the
 252 northern regions. For instance, ozone is significantly reduced on stagnant days within 24–
 253 27 °C in BRIT or above ~21 °C in NSC, and there is a minor impact of stagnation on
 254 ozone within 21–30 °C in BALT. Furthermore, stagnation does not seem to exert a
 255 significant impact within 27–30 °C in that region and 24–27 °C in NCE.

256 To understand these results we have compared the typical circulation patterns in
 257 the lower troposphere for stagnant and non-stagnant days in all regions. As stagnant
 258 conditions in the reanalysis are determined on a grid cell by grid cell basis, we have first
 259 identified a representative location for each region and then produced composite maps of
 260 daily SLP and 850 hPa wind, considering stagnant and non-stagnant days at each location.
 261 For the selection of such locations we have looked for the land grid cells where the
 262 correlation between the daily time series of standardized anomalies of summer MDA8 O_3
 263 and that of the respective centroid (i.e. time series of the average standardized anomalies
 264 within the cluster) maximize. The choice of standardized anomalies instead of absolute
 265 values of MDA8 O_3 is justified by the fact that the former were used for the identification
 266 of the regions used here (Carro-Calvo et al., 2017). The eight representative locations are
 267 displayed as circles in Figure 1. Then we have composited SLP and 850 hPa wind data
 268 for stagnant and non-stagnant days at each location with T_{\max} within 20–25 °C. This
 269 temperature interval has been chosen because it is typical of fair weather conditions
 270 favorable for ozone production in northern Europe. Moreover, around 22–44% of the

271 summer days lay within this temperature range at 7 out of the 8 locations, with the
 272 exception of only ~3% of the days for the grid cell in BRIT. Consequently, this
 273 temperature interval also guarantees a representative enough number of occurrences to
 274 draw solid conclusions for most locations.

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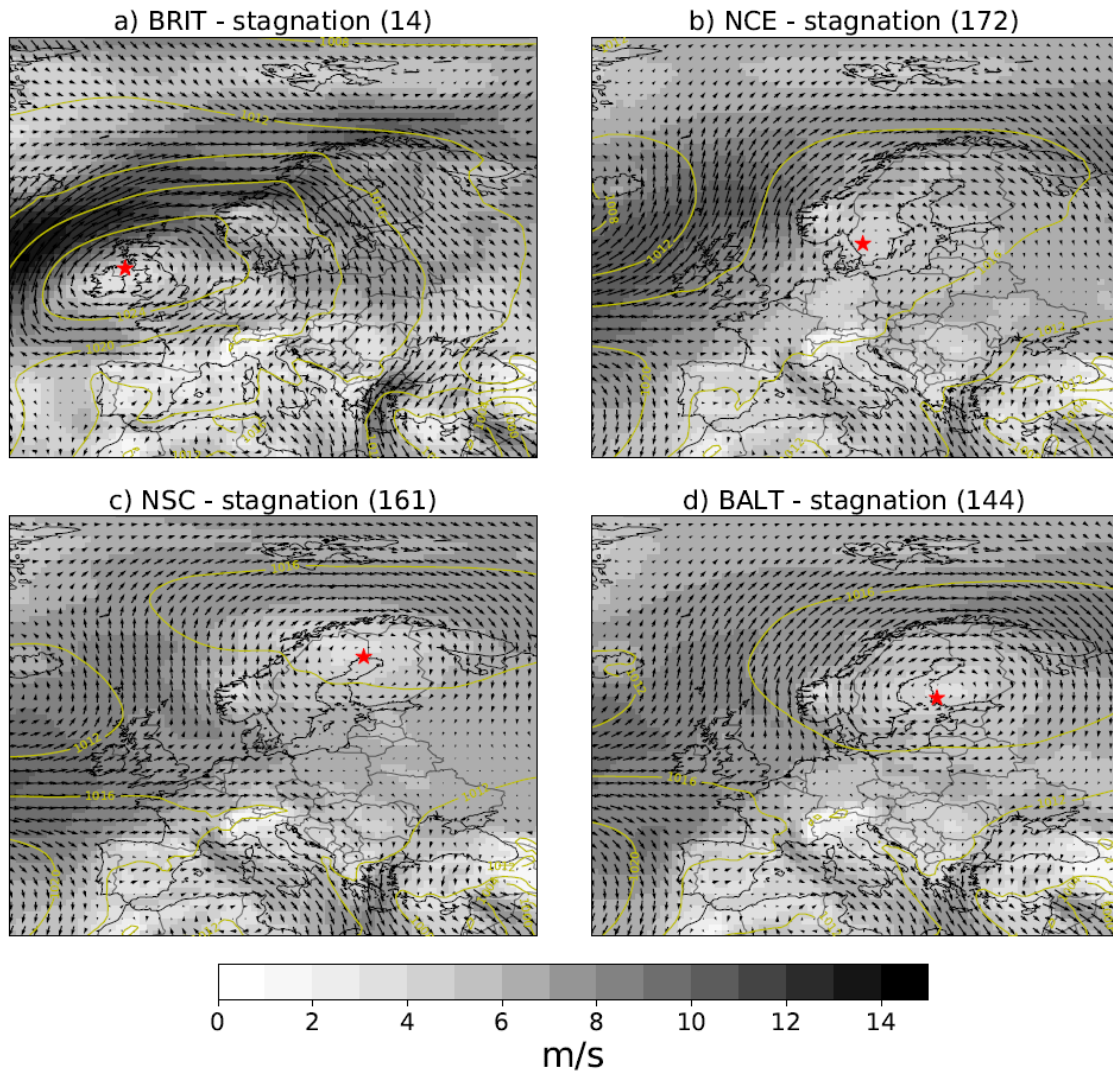
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277 **Figure 3.** Boxplots of 1998 – 2015 summer MDA8 O₃ in 3 °C bins of T_{max}, considering
278 all grid cells in each region. Blue (red) boxes represent days classified as non-stagnant
279 (stagnant). The boxes extend from the lower (Q1) to the upper (Q3) quartile values of the
280 data, with a horizontal line indicating the position of the median (Q2). The whiskers
281 extend from the boxes to show the range of the data between the 10th and 90th percentiles.
282 The x-axis tick labels represent the middle point for each temperature bin. Temperature
283 bins are only shown if they include at least 50 cases (number of cases = number of days
284 × number of sites) for both stagnant and non-stagnant conditions; as a result the range of
285 temperatures shown on the x-axis varies with the region. The numbers of stagnant (red)
286 and non-stagnant (blue) cases are indicated below the corresponding temperature bins.
287 Black stars are shown over each pair of boxes if their MDA8 O₃ distributions are
288 significantly different at the 95% confidence level (determined through the two-sample
289 Kolmogorov-Smirnov test). The vertical dotted lines indicate the temperature bins with
290 centers at 16.5°C. Linear regressions of the mean MDA8 O₃ within each temperature bin
291 (considering both stagnant and non-stagnant cases) on T_{max} (mid-points of the intervals)
292 have been drawn for the temperature bins ≥ 16.5°C.

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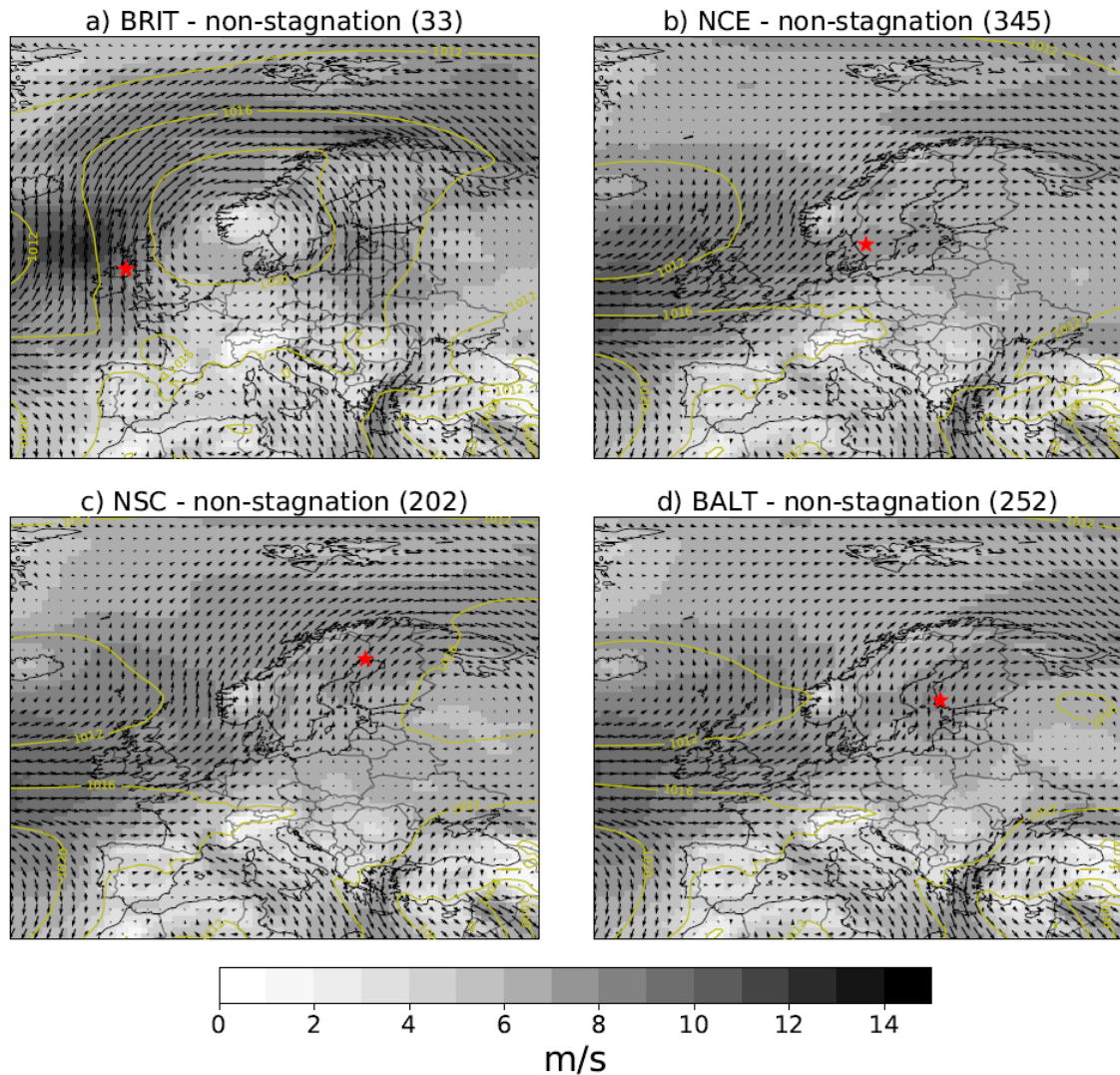
294 Figures 4 and 5 respectively illustrate the circulation patterns on stagnant and non-
295 stagnant days within the mentioned 20–25 °C temperature range in the northern locations
296 (BRIT, NCE, NSC and BALT). As expected, air stagnation situations are characterized
297 by anticyclonic circulation and weak wind in the lower troposphere over the affected
298 regions (Figure 4). The composite for NCE during non-stagnant days (Figure 5b) displays
299 westerly to southwesterly flow from the North Sea to the region. This situation enhances
300 the transport of relatively clean marine air masses to the receptor region, resulting in
301 ozone levels somewhat below those of stagnant days (Figure 3b). On the other hand,
302 BRIT, NSC and BALT are under the effect of southerly advection during non-stagnant
303 days (Figure 5, a, c, d) and, as shown above, do not display clear enhancements of MDA8
304 O₃ (Figure 3, a, c, d) despite the low wind speeds found on stagnant days (Figure 4, a, c,
305 d). The results for these three regions are in line with the findings by Carro-Calvo et al.
306 (2017), who reported that ozone extremes (exceedances of the summer 95th percentile) in
307 the north of the continent tend to be at least partly caused by the advection of polluted,
308 warm air masses from central/southern regions. This is consistent with the strong
309 meridional gradient of MDA8 O₃ in Europe (see Figure S1). In the receptor regions, wind
310 speeds are relatively high under such situations, which means that high ozone days often
311 concur with non-stagnant days. This explains the unclear impact of stagnation on MDA8
312 O₃ within some temperature bins in northern Europe. However, as anticipated before, the
313 results for BRIT should be interpreted with caution because of the reduced data sample
314 (only around 3% of the summer days with temperatures within 20–25 °C, less than one
315 third of them with stagnation). In the case of central-southern Europe, stagnant days are
316 associated with lower wind speeds than non-stagnant days for all regions, as well as with
317 a suppression of westerly advection over WE and EE (see Figures S3 and S4).

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319

320 **Figure 4.** Composites of SLP and 850 hPa wind speed on days with stagnant conditions
 321 and T_{\max} within 20 – 25 °C, at the representative locations (red stars) of BRIT, NCE
 322 and BALT, during the summer months from 1998 to 2015. The yellow contour lines, grey
 323 shading and arrows respectively represent SLP (hPa), and module and vectors of 850 hPa
 324 wind speed (m/s). The numbers in brackets represent the number of days considered in
 325 each composite.



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327 **Figure 5.** As Figure 4 but for non-stagnant days.

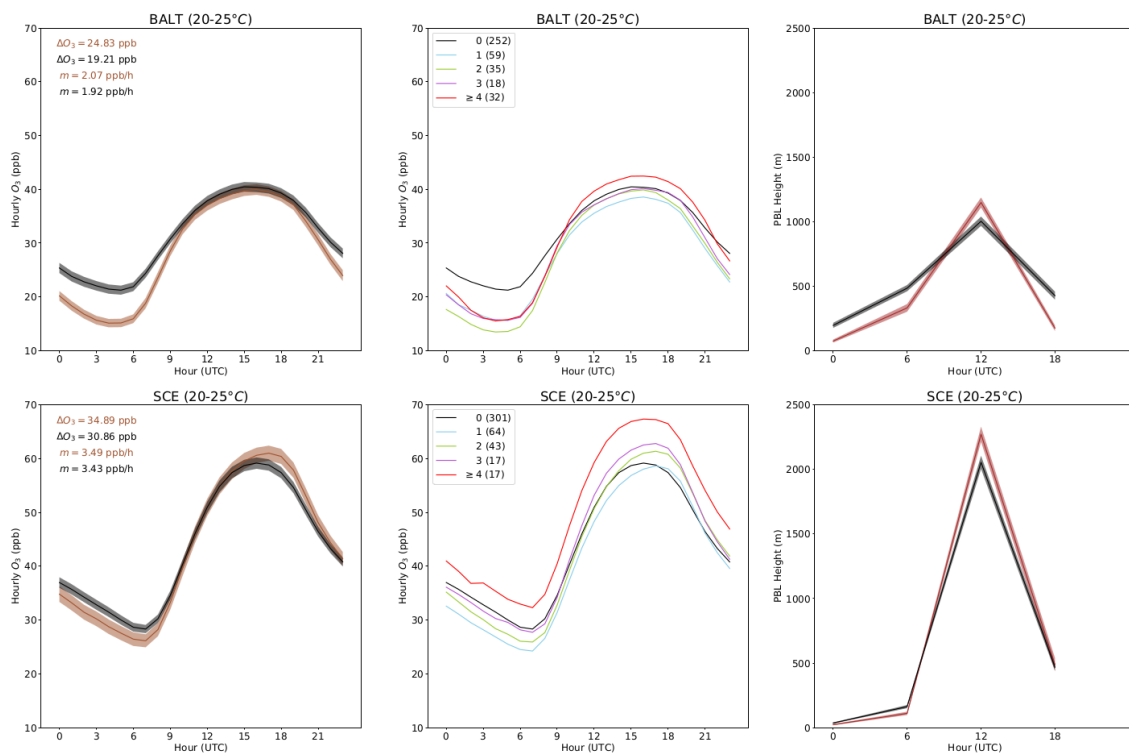
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329 **4. Impact of stagnation on the O₃ diurnal cycle**

330 To better understand the ozone response to stagnation, we have compared the
 331 diurnal cycles of this pollutant in each region for stagnant and non-stagnant days. Our
 332 analyses are inspired by those of Pyrgou et al. (2018), who examined the effect of
 333 heatwave conditions on the diurnal cycle of ozone on a Mediterranean island. As in that
 334 study, the main limitation of our approach is that the evaluation of diurnal cycles is done
 335 based on average or accumulated values of meteorological variables (wind speed and
 336 precipitation in our particular case) during the whole day. We have also examined the
 337 impact of the persistence of stagnation on these diurnal cycles to investigate if the day-
 338 to-day evolution of ozone is affected by stagnation. As before, for the sake of simplicity,
 339 these analyses have been carried out for a single land grid cell in each cluster (blue circles
 340 in Figure 1) considering days with T_{\max} within 20–25 °C. Results are reported here only
 341 for BALT and SCE as two representative regions of northern and central/southern
 342 Europe, respectively (Figure 6), and for the rest of clusters in the Supplementary Material
 343 (Figure S5 for the northern regions and Figure S6 for the central/southern regions).

344 It is well known that O₃ diurnal cycles are characterized by high concentrations
 345 during daylight hours and low values during the late night and early morning, as seen on
 346 the left and middle panels of Figures 6, S5 and S6. Both the entrainment of ozone-rich air
 347 from aloft during the development of the well-mixed PBL and photochemical production
 348 lead to the build-up of the ozone mixing ratios at daytime, with a daily maximum in the
 349 afternoon. On the other hand, O₃ mixing ratios decrease during the night because of the
 350 cease of production as well as the destruction by dry deposition to surfaces and titration
 351 by nitrogen monoxide (NO) in the shallow nocturnal PBL (e.g. Petetin et al., 2016, and
 352 references therein). The daily ozone minimum often occurs at a time with low PBL and
 353 elevated NO_x emissions from traffic during the morning rush hour. The amplitude of the
 354 diurnal cycle (defined as the difference between the maximum and the minimum hourly
 355 O₃ mixing ratios) is geographically dependent due to differences in the emission patterns
 356 and climates of each region.

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358

359 **Figure 6.** Composites of the diurnal cycles of hourly O₃ and PBL height at two
 360 representative locations in BALT and SCE on stagnant and non-stagnant days with T_{max}
 361 within 20 – 25 °C. The times displayed on the x-axes are UTC while summer local time
 362 is UTC+3 for BALT and UTC+2 for SCE. Left panels: Average diurnal ozone profiles
 363 for the summer (JJA) 1998 – 2015 days with (brown) and without (black line) stagnation
 364 in BALT (top) and SCE (bottom). Shadings cover the 95% confidence interval. The
 365 amplitudes and increase rates of the stagnant (brown) and non-stagnant (black) diurnal
 366 cycles are displayed as ΔO_3 and m on the upper-left corners of the panels. Middle panels:
 367 Average diurnal ozone profiles for summer days based on the persistence of stagnation in
 368 BALT (top) and SCE (bottom). The black line represents non-stagnation, while the blue,
 369 green, purple and red lines represent one, two, three and four or more consecutive days
 370 with stagnation, respectively. The numbers in brackets indicate the total number of days
 371 considered for each case. Right panels: Evolution of PBL height (using data at 0, 6, 12
 372 and 18 UTC) at both locations for stagnant (brown) and non-stagnant (black) days.

373 The left panels of Figures 6, S5 and S6 illustrate the average diurnal cycles of the
374 hourly O₃ mixing ratios at the selected locations in the 8 regions, considering stagnant
375 (brown) and non-stagnant (black) days, with 95% uncertainty estimates. The confidence
376 intervals have been calculated using standard t-statistics (von Storch and Zwiers, 1999).
377 Generally, the diurnal amplitude is significantly larger on stagnant days, as it increases
378 by 2.5 to 10 ppb in 6 out of 8 selected grid cells (see ΔO_3 values at the upper left corners
379 of the panels). The only exceptions are the locations in BRIT and NSC, where the
380 amplitude hardly changes. In the former location, the number of days with T_{\max} within 20
381 – 25 °C is not high enough to draw solid conclusions (see Figure S5 a, b), whereas NSC
382 is by far the cluster with the lowest sensitivity of MDA8 O₃ to stagnation (Table 1).

383 The differences between stagnant and non-stagnant O₃ diurnal cycles are often
384 due to the above-average daytime increase and/or nighttime decrease of O₃ during
385 stagnant days, again with the exception of BRIT and NSC. The low nighttime ozone
386 mixing ratios on stagnant days may be caused by enhanced chemical destruction and dry
387 deposition under a stable nocturnal PBL. Indeed, the right panels of Figures 6, S5 and S6
388 prove that the average PBL height at 6 UTC is often lower than usual during stagnant
389 nights. That is particularly the case for the locations with the strongest ozone decrease
390 under stagnant conditions at that time of the day (grid cells in BALT, NCE, WE and EE).
391 This results in a reduced O₃ baseline at the beginning of the day for these four locations
392 and, to a lesser extent, for those in SCE and BRIT, while no clear effect is found for the
393 grid cells in NSC and IBE. Thus we have shown that at most locations the occurrence of
394 stagnation leads to below-average early morning PBL height and ozone on the same day.

395 By contrast, the stronger build-up of daytime ozone on stagnant days could be
396 related to the accumulation of primary pollutants and subsequent photochemical
397 production as well as to enhanced subsidence and accumulation of ozone under the stable
398 anticyclonic conditions that characterize those days. The afternoon ozone maxima tend
399 to be higher than average on stagnant days for the central/southern locations, in particular
400 for the grid cells in WE and EE, and to lesser extent for those in IBE and SCE. The most
401 noticeable effect of stagnation on daytime ozone for the grid cell in SCE is the shift of
402 the highest mixing ratios to later hours in the evening, consistent with a synchronous shift
403 in temperature under stagnant conditions at the same location (not shown). The average
404 afternoon ozone maxima are also higher on stagnant than on non-stagnant days for the
405 grid cell in NCE, while there is no clear impact or even a decrease for the other northern
406 locations in BALT, BRIT and NSC. This is consistent with the unclear impact of
407 stagnation on the MDA8 O₃ - temperature relationship (Figure 3, a, c, d) and the effect of
408 southerly advection during non-stagnant days (Figure 5, a, c, d) shown previously for
409 those regions.

410 We have also computed the daily ozone increase rate (defined as the daily
411 amplitude divided by the period of time between the maximum and the minimum O₃
412 mixing ratios) at each location under stagnant and non-stagnant days. These values are
413 displayed as m at the upper left corners of the left panels in Figures 6, S5 and S6. This
414 rate changes from 2.0–2.6 ppb h⁻¹ under non-stagnant conditions to 2.2–3.4 ppb h⁻¹ during
415 stagnant days at three of the central/southern locations, IBE, WE and EE (Figure S6). The
416 largest increases of this rate are found for WE and EE (above 0.7 ppb h⁻¹). However, the
417 increase at the grid cell in SCE is negligible (below 0.1 ppb h⁻¹), because of the mentioned
418 late afternoon ozone maxima at this location under stagnant conditions. The daily ozone
419 increase rate also rises at most of the northern locations under stagnant conditions, but
420 the typical values are lower than those shown above, ranging from 0.9 ppb h⁻¹ in NSC to
421 2.0–2.3 ppb h⁻¹ in BRIT. These results suggest higher photochemical production in the

422 central/southern than in the northern locations, in line with expectations, as well as an
423 enhancement of photochemical activity during stagnant days. Nevertheless, some of the
424 processes previously mentioned (e.g. decrease of the early morning ozone baseline
425 followed by enhanced vertical mixing and subsidence at daytime on stagnant days,
426 southerly advection to northern regions under non-stagnant conditions) may be relevant
427 to explain these differences.

428 Previous studies have considered four or more consecutive days with air
429 stagnation at a given location as an air stagnation episode (Wang and Angell, 1999; Huang
430 et al., 2017). Such episodes are not common in northern Europe (Garrido-Perez et al.,
431 2018), but it is still possible to examine changes in the diurnal evolution of ozone during
432 successive stagnation days. The middle panels of Figures 6, S5 and S6 display composites
433 of O₃ diurnal cycles for non-stagnant days (black) as well as for one (blue), two (green),
434 three (purple) and four or more (red) consecutive days with stagnation. Despite some
435 limitations due to the small number of consecutive stagnant days at some locations,
436 especially the grid cell in BRIT, the figures indicate that daytime ozone mixing ratios
437 tend to increase with the persistence of air stagnation. The only exceptions are that
438 location in BRIT and the one selected in NSC. Our results are in the same line as previous
439 analyses which found the number of successive stagnation days as a good indicator of the
440 build-up of MDA8 O₃ in most regions of the US (Sun et al., 2017). In addition, a number
441 of studies have related the persistence of air stagnation or the associated stable
442 anticyclonic conditions to the build-up of summer daily O₃ maximum and MDA8 O₃ in
443 some regions of Europe (Ordóñez et al., 2005, 2017; Garrido-Perez, 2018). This also
444 seems to be consistent with the well-known persistence of elevated MDA8 O₃ in Europe
445 (e.g. Otero et al., 2016). However, this increase does not necessarily apply to the whole
446 diurnal cycle. In fact, the first days with stagnation consistently show a lower daily O₃
447 minimum than non-stagnant days, but this minimum gradually increases with the
448 persistence of stagnation, as found during daylight hours, again with the exception of
449 BRIT and NSC. If air stagnation persists long enough, the early morning O₃ baseline can
450 recover and even exceed the typical levels found during non-stagnant days at the locations
451 in SCE and IBE. BALT, NCE, WE and EE present a similar build-up, but the early
452 morning O₃ baseline during the most persistent stagnation episodes remains below that
453 of non-stagnant days. Hence, it appears that the nighttime destruction of O₃ is enhanced
454 under the shallow PBL during the first stagnant days at most locations and that this
455 destruction is at least partially compensated by the accumulation of O₃ when consecutive
456 days with stagnation occur. The increase in the early morning baseline at some locations,
457 together with mixing of boundary layer air with ozone-rich air from the residual layer
458 after sunrise as well as daytime photochemical production, will favor the occurrence of
459 elevated MDA8 O₃ in the afternoon.

460 **5. Discussion and conclusions**

461 Climate change is expected to increase the risk of extreme air pollution events in
462 the future. Southern Europe has been identified as a climate change hot spot (Diffenbaugh
463 and Giorgi, 2012; de Sherbinin, 2014) and is projected to have multiple impacts. Among
464 them, climate models have projected strong increases in temperature (Giorgi and
465 Lionello, 2008; Barkhordarian et al., 2012) and atmospheric stagnation (Horton et al.,
466 2012, 2014) in this region. Both phenomena are closely linked to elevated ozone
467 concentrations and thus their exacerbation should make it harder to achieve air quality
468 goals for this pollutant in the future. However, a recent study has questioned the
469 relationship between summer air pollution and stagnation (Kerr and Waugh, 2018).

470 Keeping in mind that daily maximum temperature is the meteorological variable
471 that mostly controls the afternoon ozone concentrations, this study has investigated the
472 added value of a simple air stagnation index (Horton et al, 2012) to understand summer
473 ozone variability in Europe. Building on the regional classification of MDA8 O₃ by
474 Carro-Calvo et al. (2017), we have evaluated the joint effect of temperature and stagnation
475 on summer ozone in eight regions (BRIT, NCE, NSC, BALT, IBE, WE, SCE, EE),
476 finding considerable heterogeneity across the continent. The main findings are as follows:

- 477 • MDA8 O₃ mixing ratios consistently increase over central/southern Europe (IBE,
478 WE, SCE, EE) and, to a lesser extent, NCE under stagnant conditions, but this has not
479 been found for some temperature bins in three of the northern regions (BRIT, NSC,
480 BALT). Under non-stagnant conditions and T_{max} within 20–25 °C (typical
481 temperatures of fair weather conditions that allow photochemical production in
482 northern Europe), such regions are affected by southerly advection, which is a known
483 mechanism for the occurrence of ozone extremes in the north of the continent (Carro-
484 Calvo et al., 2017).
- 485 • The ozone diurnal cycle in the central/southern regions and NCE exhibits larger
486 amplitudes than usual when stagnation occurs, with low nighttime and high daytime
487 mixing ratios in most of these regions. Stagnant nights are associated with stable
488 shallow PBL and, presumably, enhanced dry deposition and chemical destruction of
489 ozone. After sunrise, with the development of the PBL, mixing with air from the
490 residual layer, accumulation of ozone and precursors, and photochemical production
491 seem to be the main mechanisms involved in the build-up of daytime ozone.
- 492 • Low nighttime ozone mixing ratios during stagnant days are also a feature of BALT
493 and BRIT, while afternoon ozone and consequently MDA8 O₃ are not clearly affected
494 by stagnation. In particular, for the 20–25 °C temperature range considered here,
495 afternoon ozone can even decrease under stagnant conditions in BRIT. However,
496 these results should be interpreted with caution considering the limited number of
497 occurrences observed and the fact that MDA8 O₃ in the same region increases on
498 stagnant days for other temperature ranges.
- 499 • NSC is the region with the weakest sensitivity of ozone to stagnation. This is firstly
500 evidenced from the poor correlation between the daily time series of average MDA8
501 O₃ and the percentage of stagnant area in the region (R=0.06). Moreover, the
502 occurrence of stagnation hardly affects the ozone diurnal cycles for T_{max} within 20–
503 25 °C and even yields decreases in the MDA8 O₃ mixing ratios for T_{max} above ~20
504 °C. Nonetheless, we note the results for this region are based on relative fewer surface
505 ozone observations as compared to those in central/southern regions (i.e. less than 30
506 sites north of 62.5° N).

507 In short, we have been able to identify regions with different responses of summer
508 ozone to the occurrence of air stagnation. It is remarkable that some of the
509 central/southern European regions where stagnation has a clear impact on ozone have
510 already undergone significant upward trends in stagnation over the last decades (Garrido-
511 Perez et al., 2018) and are also likely to experience increases in the future (Horton et al.,
512 2014). On the other hand, stagnation seems to exert a minor control on summer ozone
513 over most of northern Europe. Consequently, observations of air stagnation occurrence
514 and projections of increases in air stagnation should not directly be translated into
515 enhanced summer ozone pollution if the sensitivity of this pollutant to stagnation has not
516 been proved for a particular region.

517 Another relevant result of this paper is that the occurrence of air stagnation
518 amplifies the diurnal cycles of summer ozone over some regions. While simulations of
519 surface ozone by Chemical Transport Models (CTMs) are often evaluated against
520 observations of MDA8 O₃ or other relevant cumulative metrics (e.g. Lupaşcu and Butler,
521 2019), understanding the differences in the performance of a set of model simulations for
522 such metrics would require assessing the skill to reproduce the diurnal cycle of ozone. In
523 fact, current global and regional CTMs present limitations at night and under stable
524 conditions, partly because of the difficulty in resolving the stratified conditions near the
525 surface and the depletion of ozone through surface deposition (Travis and Jacob, 2019,
526 and references therein). Nighttime PBL dynamics and chemistry are crucial because they
527 will determine the early morning ozone baseline and also influence morning radical
528 chemistry, with important implications for the formation of daytime ozone (e.g. Brown
529 and Stutz, 2012). On the other hand, following the development of the PBL at daytime,
530 the entrainment of ozone-rich air from the upper levels to the ground has been proved to
531 be an important transport mechanism of summer ozone smog (e.g. Hu et al., 2018, and
532 references therein). Therefore, special attention should be given to quantifying the
533 contribution of individual processes such as horizontal and vertical advection, vertical
534 diffusion, dry deposition and chemistry to the ozone budget under different
535 meteorological conditions. This can be achieved through the use of probing techniques
536 like Integrated Process Rate analysis (IPR), which provides detailed mass balance
537 information for the aforementioned processes in some regional CTMs (e.g. Liu et al.,
538 2007; Gonçalves et al., 2009; Lyu et al., 2019). Our findings about the differing impacts
539 of air stagnation on the ozone diurnal cycles across Europe provide good observational
540 constrains for the evaluation of such processes under stagnant vs. non-stagnant
541 conditions. Furthermore, periods of persistent stagnant situations can be very appropriate
542 to test the ability of CTMs to reproduce PBL dynamics and the evolution of ozone
543 concentrations.

544 **Acknowledgments**

545 JMGP was supported by a predoctoral research grant awarded by the Spanish
546 Ministerio de Educación, Cultura y Deporte (FPU16/01972) and CO by the Ramón y
547 Cajal Programme of the Spanish Ministerio de Economía y Competitividad [grant number
548 RYC-2014-15036]. JLS was funded by the Ubben Program for Climate and Carbon
549 Science. We also acknowledge support from STEADY (CGL2017-83198-R), project
550 funded by the Spanish Ministerio de Economía, Industria y Competitividad. ERA-Interim
551 reanalysis was provided by ECMWF. We are grateful to Leopoldo Carro-Calvo for
552 providing the regionalization of the gridded ozone dataset. The authors thank three
553 anonymous reviewers for their useful comments.

554 **References**

555 Ashmore, M.R., 2005. Assessing the future global impacts of ozone on vegetation.
556 *Plant, Cell Environ.* 28, 949–964. <https://doi.org/10.1111/j.1365-3040.2005.01341.x>

557 Barkhordarian, A., Bhend, J., von Storch, H., 2012. Consistency of observed near
558 surface temperature trends with climate change projections over the Mediterranean
559 region. *Clim. Dyn.* 38, 1695–1702. <https://doi.org/10.1007/s00382-011-1060-y>

560 Bloomer, B.J., Stehr, J.W., Piety, C.A., Salawitch, R.J., Dickerson, R.R., 2009,
561 Observed relationships of ozone air pollution with temperature and emissions. *Geophys.*
562 *Res. Lett.* 36, L09803. <https://doi.org/10.1029/2009GL037308>

563 Bloomfield, P., Royle, J.A., Steinberg, L.J., Yang, Q., 1996. Accounting for
564 meteorological effects in measuring urban ozone levels and trends. *Atmos. Environ.* 30,
565 3067–3077. [https://doi.org/10.1016/1352-2310\(95\)00347-9](https://doi.org/10.1016/1352-2310(95)00347-9)

566 Brown, S.S., Stutz, J., 2012. Nighttime radical observations and chemistry. *Chem.*
567 *Soc. Rev.* 41, 6405–6447. <https://doi.org/10.1039/c2cs35181a>

568 Carro-Calvo, L., Ordóñez, C., García-Herrera, R., Schnell, J.L., 2017. Spatial
569 clustering and meteorological drivers of summer ozone in Europe. *Atmos. Environ.* 167,
570 496–510. <https://doi.org/10.1016/j.atmosenv.2017.08.050>

571 Dawson, J.P., Bloomer, B.J., Winner, D.A., Weaver, C.P., 2014. Understanding
572 the meteorological drivers of U.S. particulate matter concentrations in a changing climate.
573 *Bull. Am. Meteorol. Soc.* 95, 521–532. <https://doi.org/10.1175/BAMS-D-12-00181.1>

574 de Sherbinin, A., 2014. Climate change hotspots mapping: What have we learned?
575 *Clim. Change* 123, 23–37. <https://doi.org/10.1007/s10584-013-0900-7>

576 Dee, D.P., Uppala, S.M., Simmons, A.J., Berrisford, P., Poli, P., Kobayashi, S.,
577 Andrae, U., Balmaseda, M.A., Balsamo, G., Bauer, P., Bechtold, P., Beljaars, A.C.M.,
578 van de Berg, L., Bidlot, J., Bormann, N., Delsol, C., Dragani, R., Fuentes, M., Geer, A.J.,
579 Haimberger, L., Healy, S.B., Hersbach, H., Hólm, E. V., Isaksen, I., Kållberg, P., Köhler,
580 M., Matricardi, M., McNally, A.P., Monge-Sanz, B.M., Morcrette, J.J., Park, B.K.,
581 Peubey, C., de Rosnay, P., Tavolato, C., Thépaut, J.N., Vitart, F., 2011. The ERA-Interim
582 reanalysis: Configuration and performance of the data assimilation system. *Q. J. R.*
583 *Meteorol. Soc.* 137, 553–597. <https://doi.org/10.1002/qj.828>

584 Diffenbaugh, N.S., Giorgi, F., 2012. Climate change hotspots in the CMIP5 global
585 climate model ensemble. *Clim. Change* 114, 813–822. <https://doi.org/10.1007/s10584-012-0570-x>

587 Fehsenfeld, F., Calvert, J., Fall, R., Goldan, P., Guenther, A.B., Hewitt, C.N.,
588 Lamb, B., Trainer, M., Westberg, H., Zimmerman, P., 1992. Emissions of volatile organic
589 compounds from vegetation and the implications for atmospheric chemistry. *Global*
590 *Biogeochem. Cycles* 6, 389–430. <https://doi.org/10.1029/92GB02125>

591 Fiore, A.M., Naik, V., Leibensperger, E.M., 2015. Air quality and climate
592 connections. *J. Air Waste Manag. Assoc.* 65, 645–685.
593 <https://doi.org/10.1080/10962247.2015.1040526>

594 Garrido-Perez, J.M., Ordóñez, C., García-Herrera, R., Barriopedro, D., 2018. Air
595 stagnation in Europe: Spatiotemporal variability and impact on air quality. *Sci. Total*
596 *Environ.* 645, 1238–1252. <https://doi.org/10.1016/j.scitotenv.2018.07.238>

597 Giorgi, F., Lionello, P., 2008. Climate change projections for the Mediterranean
598 region. *Glob. Planet. Change* 63, 90–104.
599 <https://doi.org/10.1016/j.gloplacha.2007.09.005>

600 Gonçalves, M., Jiménez-Guerrero, P., Baldasano, J.M., 2009. Contribution of
601 atmospheric processes affecting the dynamics of air pollution in South-Western Europe
602 during a typical summertime photochemical episode. *Atmos. Chem. Phys.* 9, 849–864.
603 <https://doi.org/10.5194/acp-9-849-2009>

604 Horton, D.E., Harshvardhan, Diffenbaugh, N.S., 2012. Response of air stagnation
605 frequency to anthropogenically enhanced radiative forcing. *Environ. Res. Lett.* 7, 044034.
606 <https://doi.org/10.1088/1748-9326/7/4/044034>

607 Horton, D.E., Skinner, C.B., Singh, D., Diffenbaugh, N.S., 2014. Occurrence and
608 persistence of future atmospheric stagnation events. *Nat. Clim. Chang.* 4, 698–703.
609 <https://doi.org/10.1038/nclimate2272>

610 Hu, J., Emile-Geay, J., Partin, J., 2017. Correlation-based interpretations of
611 paleoclimate data – where statistics meet past climates. *Earth Planet. Sci. Lett.* 459, 362–
612 371. <https://doi.org/10.1016/j.epsl.2016.11.048>

613 Hu, J., Li, Y., Zhao, T., Liu, J., Hu, X.M., Liu, D., Jiang, Y., Xu, J., Chang, L.,
614 2018. An important mechanism of regional O₃ transport for summer smog over the
615 Yangtze River Delta in eastern China. *Atmos. Chem. Phys.* 18, 16239–16251.
616 <https://doi.org/10.5194/acp-18-16239-2018>

617 Huang, Q., Cai, X., Song, Y., Zhu, T., 2017. Air stagnation in China (1985 –
618 2014): climatological mean features and trends. *Atmos. Chem. Phys.* 17, 7793–7805.
619 <https://doi.org/10.5194/acp-17-7793-2017>

620 Huang, Q., Cai, X., Wang, J., Song, Y., Zhu, T., 2018. Climatological study of the
621 Boundary-layer air Stagnation Index for China and its relationship with air pollution.
622 *Atmos. Chem. Phys.* 18, 7573–7593. <https://doi.org/10.5194/acp-18-7573-2018>

623 Jacob, D.J., Winner, D.A., 2009. Effect of climate change on air quality. *Atmos.*
624 *Environ.* 43, 51–63. <https://doi.org/10.1016/j.atmosenv.2008.09.051>

625 Kerr, G.H., Waugh, D.W., 2018. Connections between summer air pollution and
626 stagnation. *Environ. Res. Lett.* 13, 084001. <https://doi.org/10.1088/1748-9326/aad2e2>

627 Kuebler, J., Van Den Bergh, H., Russell, A.G., 2001. Long-term trends of primary
628 and secondary pollutant concentrations in Switzerland and their response to emission
629 controls and economic changes. *Atmos. Environ.* 35, 1351–1363.
630 [https://doi.org/10.1016/S1352-2310\(00\)00401-5](https://doi.org/10.1016/S1352-2310(00)00401-5)

631 Leibensperger, E.M., Mickley, L.J., Jacob, D.J., 2008. Sensitivity of US air quality
632 to mid-latitude cyclone frequency and implications of 1980–2006 climate change. *Atmos.*
633 *Chem. Phys.* 8, 7075–7086. <https://doi.org/10.5194/acp-8-7075-2008>

634 Liu, L., Andreani-Aksoyoglu, S., Keller, J., Ordóñez, C., Junkermann, W., Hak,
635 C., Braathen, G.O., Reimann, S., Astorga-Llorens, C., Schultz, M., Prévôt, A.S.H.,
636 Isaksen, I.S.A., 2007. A photochemical modeling study of ozone and formaldehyde
637 generation and budget in the Po basin. *J. Geophys. Res. Atmos.* 112, D22303.
638 <https://doi.org/10.1029/2006JD008172>

639 Lupaşcu, A., Butler, T., 2019. Source attribution of European surface O₃ using a
640 tagged O₃ mechanism. *Atmos. Chem. Phys. Discuss.* <https://doi.org/10.5194/acp-2019-225>

642 Lyu, X., Wang, N., Guo, H., Xue, L., Jiang, F., Zeren, Y., Cheng, H., Cai, Z., Han,
643 L., Zhou, Y., 2018. Causes of a continuous summertime O₃ pollution event in Jinan, a
644 central city in the North China Plain. *Atmos. Chem. Phys.* 19, 3025–3042.
645 <https://doi.org/10.5194/acp-19-3025-2019>

646 Meehl, G.A., Tebaldi, C., Tilmes, S., Lamarque, J.F., Bates, S., Pendergrass, A.,
647 Lombardozzi, D., 2018. Future heat waves and surface ozone. *Environ. Res. Lett.* 13,
648 064004. <https://doi.org/10.1088/1748-9326/aabdc>

649 Ordóñez, C., Barriopedro, D., García-Herrera, R., Sousa, P.M., Schnell, J.L.,
650 2017. Regional responses of surface ozone in Europe to the location of high-latitude

651 blocks and subtropical ridges. *Atmos. Chem. Phys.* 17, 3111–3131.
652 <https://doi.org/10.5194/acp-17-3111-2017>

653 Ordóñez, C., Mathis, H., Furger, M., Henne, S., Hüglin, C., Staehelin, J., Prévôt,
654 A.S.H., 2005. Changes of daily surface ozone maxima in Switzerland in all seasons from
655 1992 to 2002 and discussion of summer 2003. *Atmos. Chem. Phys.* 5, 1187–1203.
656 <https://doi.org/10.5194/acp-5-1187-2005>

657 Orlando, J.J., Tyndall, G.S., Calvert, J.G., 1992. Thermal decomposition
658 pathways for peroxyacetyl nitrate (PAN): Implications for atmospheric methyl nitrate
659 levels. *Atmos. Environ. Part A, Gen. Top.* 26, 3111–3118. [https://doi.org/10.1016/0960-](https://doi.org/10.1016/0960-1686(92)90468-Z)
660 [1686\(92\)90468-Z](https://doi.org/10.1016/0960-1686(92)90468-Z)

661 Otero, N., Sillmann, J., Mar, K.A., Rust, H.W., Solberg, S., Andersson, C.,
662 Engardt, M., Bergström, R., Bessagnet, B., Colette, A., Couvidat, F., Cuvelier, C., Tsyro,
663 S., Fagerli, H., Schaap, M., Manders, A., Mircea, M., Briganti, G., Cappelletti, A., Adani,
664 M., D’Isidoro, M., Pay, M.T., Theobald, M., Vivanco, M.G., Wind, P., Ojha, N., Raffort,
665 V., Butler, T., 2018. A multi-model comparison of meteorological drivers of surface
666 ozone over Europe. *Atmos. Chem. Phys.* 18, 12269–12288. [https://doi.org/10.5194/acp-](https://doi.org/10.5194/acp-18-12269-2018)
667 [18-12269-2018](https://doi.org/10.5194/acp-18-12269-2018)

668 Otero, N., Sillmann, J., Schnell, J.L., Rust, H.W., Butler, T., 2016. Synoptic and
669 meteorological drivers of extreme ozone concentrations over Europe. *Environ. Res. Lett*
670 *11*, 024005. <https://doi.org/10.1088/1748-9326/11/2/024005>

671 Petetin, H., Thouret, V., Athier, G., Blot, R., Boulanger, D., Cousin, J.-M.,
672 Gaudel, A., Nédélec, P., Cooper, O., 2016. Diurnal cycle of ozone throughout the
673 troposphere over Frankfurt as measured by MOZAIC-IAGOS commercial aircraft. *Elem.*
674 *Sci. Anthr.* 4, 000129. <https://doi.org/10.12952/journal.elementa.000129>

675 Porter, W.C., Heald, C.L., 2019. The mechanisms and meteorological drivers of
676 the ozone-temperature relationship. *Atmos. Chem. Phys. Discuss.*
677 <https://doi.org/10.5194/acp-2019-140>

678 Pyrgou, A., Hadjinicolaou, H., Santamouris, M., 2018. Enhanced near-surface
679 ozone under heatwave conditions in a Mediterranean island. *Sci. Rep.* 8, 9191.
680 <https://doi.org/10.1038/s41598-018-27590-z>

681 REVIHAAP, 2013. Review of evidence on health aspects of air pollution -
682 REVIHAAP Project. Technical Report. World Health Organization Regional Office for
683 Europe. Bonn. Available at:
684 [http://www.euro.who.int/__data/assets/pdf_file/0004/193108/REVIHAAP-Final-](http://www.euro.who.int/__data/assets/pdf_file/0004/193108/REVIHAAP-Final-technical-report-final-version.pdf)
685 [technical-report-final-version.pdf](http://www.euro.who.int/__data/assets/pdf_file/0004/193108/REVIHAAP-Final-technical-report-final-version.pdf).

686 Schnell, J.L., Holmes, C.D., Jangam, A., Prather, M.J., 2014. Skill in forecasting
687 extreme ozone pollution episodes with a global atmospheric chemistry model. *Atmos.*
688 *Chem. Phys.* 14, 7721–7739. <https://doi.org/10.5194/acp-14-7721-2014>

689 Schnell, J.L., Prather, M.J., 2017. Co-occurrence of extremes in surface ozone,
690 particulate matter, and temperature over eastern North America. *Proc. Natl. Acad. Sci.*
691 *114*, 2854–2859. <https://doi.org/10.1073/pnas.1614453114>

692 Schnell, J.L., Prather, M.J., Josse, B., Naik, V., Horowitz, L.W., Cameron-Smith,
693 P., Bergmann, D., Zeng, G., Plummer, D.A., Sudo, K., Nagashima, T., Shindell, D.T.,
694 Faluvegi, G., Strode, S.A., 2015. Use of North American and European air quality

695 networks to evaluate global chemistry-climate modeling of surface ozone. *Atmos. Chem.*
696 *Phys.* 15, 10581–10596. <https://doi.org/10.5194/acp-15-10581-2015>

697 Shen, L., Mickley, L.J., Gilleland, E., 2016. Impact of increasing heat waves on
698 U.S. ozone episodes in the 2050s: Results from a multimodel analysis using extreme value
699 theory. *Geophys. Res. Lett.* 43, 4017–4025. <https://doi.org/10.1002/2016GL068432>

700 Simpson, D., 1995. Biogenic emissions in Europe: 2. Implications for ozone
701 control strategies. *J. Geophys. Res.* 100, 22891–22906.
702 <https://doi.org/10.1029/95jd01878>

703 Steiner, A.L., Davis, A.J., Sillman, S., Owen, R.C., Michalak, A.M., Fiore, A.M.,
704 2010. Observed suppression of ozone formation at extremely high temperatures due to
705 chemical and biophysical feedbacks. *Proc. Natl Acad. Sci.* 107, 19685–90. <https://doi.org/10.1073/pnas.1008336107>

707 Sun, W., Hess, P., Liu, C., 2017. The impact of meteorological persistence on the
708 distribution and extremes of ozone. *Geophys. Res. Lett.* 44, 1545–1553.
709 <https://doi.org/10.1002/2016GL071731>

710 Tai, A.P.K., Martin, M.V., Heald, C.L., 2014. Threat to future global food security
711 from climate change and ozone air pollution. *Nat. Clim. Chang.* 4, 817–821.
712 <https://doi.org/10.1038/nclimate2317>

713 Tarasova, O.A., Karpetchko, A.Y., 2003. Accounting for local meteorological
714 effects in the ozone time-series of Lovozero (Kola Peninsula). *Atmos. Chem. Phys.* 3,
715 941–949. <https://doi.org/10.5194/acp-3-941-2003>

716 Tørseth, K., Aas, W., Breivik, K., Fjæraa, A.M., Fiebig, M., Hjellbrekke, A.G.,
717 Lund Myhre, C., Solberg, S., Yttri, K.E., 2012. Introduction to the European Monitoring
718 and Evaluation Programme (EMEP) and observed atmospheric composition change
719 during 1972-2009. *Atmos. Chem. Phys.* 12, 5447–5481. <https://doi.org/10.5194/acp-12-5447-2012>

721 Travis, K.R., Jacob, D.J., 2019. Systematic bias in evaluating chemical transport
722 models with maximum daily 8 h average (MDA8) surface ozone for air quality
723 applications: a case study with GEOS-Chem v9.02. *Geosci. Model Dev.*
724 <https://doi.org/10.5194/gmd-12-3641-2019>

725 Vautard, R., Honoré, C., Beekmann, M., Rouil, L., 2005. Simulation of ozone
726 during the August 2003 heat wave and emission control scenarios. *Atmos. Environ.* 39,
727 2957–2967. <https://doi.org/10.1016/j.atmosenv.2005.01.039>

728 von Storch, H., Zwiers, F.W., 1999. *Statistical analysis in climate research*,
729 Cambridge University Press, Cambridge, UK.
730 <https://doi.org/10.1017/CBO9780511612336>

731 Wang, J.X.L., Angell, J.K., 1999. *Air Stagnation Climatology for the United*
732 *States*. NOAA/Air Resour. Lab. ATLAS 1.

733 Wang, X., Dickinson, R.R.E., Su, L., Zhou, C., Wang, K., 2018. PM_{2.5} pollution
734 in China and how it has been exacerbated by terrain and meteorological conditions. *Bull.*
735 *Am. Meteorol. Soc.* 99, 105–120. <https://doi.org/10.1175/BAMS-D-16-0301.1>

736 Wang, X., Wang, K., Su, L., 2016. Contribution of atmospheric diffusion
737 conditions to the recent improvement in air quality in China. *Sci. Rep.* 6, 36404.
738 <https://doi.org/10.1038/srep36404>

- 739 Wesely, M.L., 1989. Parametrization of surface resistances to gaseous dry
740 deposition in regional-scale numerical models. *Atmos. Environ.* 23, 1293–1304.
741 [https://doi.org/10.1016/0004-6981\(89\)90153-4](https://doi.org/10.1016/0004-6981(89)90153-4)
- 742 Wilks, D. S.: *Statistical Methods in the Atmospheric Sciences*, 3rd Edition,
743 Academic Press, Oxford, UK, 2011.
- 744 Xu, D., Yap, D., Taylor, P.A., 1996. Meteorologically adjusted ground level ozone
745 trends in Ontario. *Atmos. Environ.* 30, 1117–1124. [https://doi.org/10.1016/1352-](https://doi.org/10.1016/1352-2310(95)00331-2)
746 [2310\(95\)00331-2](https://doi.org/10.1016/1352-2310(95)00331-2)