

2005–2017 ozone trends and potential benefits of local measures as deduced from air quality measurements in the north of the Barcelona Metropolitan Area

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Abstract

We analyzed 2005–2017 data sets on ozone (O_3) concentrations in an area (the Vic Plain) frequently affected by the atmospheric plume northward transport of Barcelona Metropolitan Area (BMA), the atmospheric basin of Spain recording the highest number of exceedances of the hourly O_3 information threshold ($180 \mu g m^{-3}$). We aimed at evaluating the potential benefits of implementing local-BMA short-term measures to abate emissions of precursors. To this end, we analyzed in detail spatial and time variations of concentration of O_3 and nitrogen oxides (NO and NO_2 , including OMI remote sensing data for the latter). Subsequently, a sensitivity analysis is done with the air quality (AQ) data to evaluate potential O_3 reductions in the North of the BMA on Sundays, compared with weekdays as a consequence of the reduction in regional emissions of precursors.

The results showed a generalized decreasing trend for regional background O_3 as well as the well-known increase of urban O_3 and higher urban NO decreasing slopes compared with those of NO_2 . The most intensive O_3 episodes in the Vic Plain are caused by (i) a relatively high regional background O_3 (due to a mix of continental, hemispheric–tropospheric and stratospheric contributions); (ii) intensive surface fumigation from mid-troposphere high O_3 upper layers arising from the concatenation of the vertical recirculation of air masses, but also by (iii) an important O_3 contribution from the northward transport/channeling of the pollution plume from the BMA. The high relevance of the local-daily O_3 contribution during the most intense pollution episodes is clearly supported by the O_3 (surface concentration) and NO_2 (OMI data) data analysis.

A maximum decrease potential (by applying short-term measures to abate emissions of O_3 precursors) of $49 \mu g O_3 m^{-3}$ (32%) of the average diurnal concentrations was determined. Structurally implemented measures, instead of episodically, could result in important additional O_3 decreases because not only the local O_3 coming from the BMA plume would be reduced but also the recirculated O_3 and thus the intensity of O_3 fumigation in the Plain. Therefore, it is highly probable that both structural and episodic measures to abate NO_x and volatile organic compounds (VOCs) emissions in the BMA would result in evident reductions of O_3 in the Vic Plain.

Keywords: tropospheric ozone, regional pollution, photochemistry, air quality trends.

1. Introduction

Tropospheric ozone (O_3) is a secondary atmospheric pollutant produced by the photooxidation of volatile organic compounds (VOCs) in the presence of nitrogen oxides ($NO_x = NO + NO_2$). Its generation is enhanced under high temperature and solar radiation (Monks et al., 2015 and references therein). Thus, O_3 maxima occur generally in the afternoon, with the highest levels typically registered in summer, when exceedances of regulatory thresholds are most frequent.

O_3 is one of the key air pollutants affecting human health and the environment (WHO, 2006, 2013a, 2013b; GBD, 2016; Fowler et al., 2009; IPCC, 2013). According to EEA (2018), in the period 2013–2015, more than 95% of the urban population in the EU-28 was exposed to O_3 levels exceeding the WHO guidelines set for the protection of the human health (maximum daily 8-h average concentration of $100 \mu g m^{-3}$).

44 On a global scale, approximately 90% of the tropospheric O₃ is produced photochemically within the
45 troposphere (Stevenson et al., 2006; Young et al., 2013), the remaining part being transported from the
46 stratosphere (McLinden et al., 2000; Olson et al., 2001). The main global sink of tropospheric O₃ is photolysis
47 in the presence of water vapor. Dry deposition, mainly by vegetation, is also an important sink in the
48 continental planetary boundary layer (PBL) (Jacob and Winner, 2009).

49 On a regional scale, O₃ levels vary substantially depending on the different chemical environments within the
50 troposphere. O₃ chemical destruction is largest where water vapor concentrations are high, mainly in the lower
51 troposphere, and in polluted areas where there is direct O₃ destruction by titration. Thus, the hourly, daily and
52 annual variations in O₃ levels at a given location are determined by several factors, including the geographical
53 characteristics, the predominant meteorological conditions and the proximity to large sources of O₃ precursors
54 (Logan, 1985).

55 Southern Europe, especially the Mediterranean basin, is the most exposed to O₃ pollution in Europe (EEA,
56 2018) due to the specific prevailing meteorological conditions during warm seasons, regional pollutant
57 emissions, high biogenic VOCs' (BVOCs) emissions in spring and summer and the vertical recirculation of air
58 masses due to the particular orographic features that help stagnation–recirculation episodes (Millán et al.,
59 2000; EC, 2002, 2004; Millán, 2009; Diéguez et al., 2009, 2014; Valverde et al., 2016). Periods with high O₃
60 concentrations often last for several days and can be detected simultaneously in several countries. Lelieveld
61 et al. (2002) reported that during summer, O₃ concentrations are 2.5–3 times higher than in the hemispheric
62 background troposphere. High O₃ levels are common in the area, not only at the surface but also throughout
63 the PBL (Millán et al., 1997; Gangoi et al., 2001; Kalabokas et al., 2007). Photochemical O₃ production is
64 favored due to frequent anticyclonic conditions with clear skies during summer, causing high insolation and
65 temperatures and low rainfall. Besides, the emissions from the sources located around the basin, which is
66 highly populated and industrialized, and the long-range transport of O₃ contribute to the high concentrations
67 (Millán et al., 2000; Lelieveld et al., 2002; Gerasopoulos, 2005; Safieddine et al., 2014).

68 In this context, the design of efficient O₃ abatement policies is difficult due to the following circumstances:

- 69 • The meteorology driving O₃ dynamics is highly influenced by the complex topography surrounding the
70 basin (see the above references for vertical recirculation of air masses and Mantilla et al., 1997;
71 Salvador et al., 1997; Jiménez and Baldasano, 2004; Stein et al., 2004).
- 72 • The complex nonlinear chemical reactions between NO_x and VOCs (Finlayson-Pitts and Pitts, 1993;
73 Pusede et al., 2015), in addition to the vast variety of the VOCs precursors involved and the
74 involvement of BVOCs in O₃ formation and destruction (Hewitt et al., 2011).
- 75 • The transboundary transport of air masses containing significant concentrations of O₃ and its
76 precursors, which contribute to increased O₃ levels, mainly background concentrations (UNECE, 2010).
- 77 • The contribution from stratospheric intrusions (Kalabokas et al., 2007).
- 78 • The fact that O₃ concentrations tend to be higher in rural areas (EEA, 2018), where local mitigation
79 plans are frequently inefficient, because the emission of precursors takes place mostly in distant urban
80 and industrial agglomerations.

81
82 Sicard et al. (2013) analyzed O₃ time trends during 2000–2010 in the Mediterranean and observed a slight
83 decrease of annual O₃ averages (–0.4% per year) at rural sites, and an increase at urban and suburban stations
84 (+0.6% and +0.4%, respectively). They attributed the reduction at rural sites to the abatement of NO_x and VOCs
85 emissions in the EU. Paradoxically, this led to an increase in O₃ at urban sites due to a reduction in the titration
86 by NO. Their results also suggested a tendency to converge at remote and urban sites. Paoletti et al. (2014)
87 also reported convergence in the EU and the US in the period 1990–2010 but found increasing annual averages
88 at both rural and urban sites, with a faster increase in urban areas. Querol et al. (2016) determined that O₃
89 levels in Spain remained constant at rural sites and increased at urban sites in the period 2000–2015. This was
90 suggested to be a result of the preferential reduction of NO versus NO₂, supported by the lack of a clear trend
91 in O_x (O₃ + NO₂). They also found that the target value was constantly exceeded in large areas of the Spanish
92 territory, while most of the exceedances of the information threshold took place in July, mainly downwind of

93 urban areas and industrial sites, and were highly influenced by summer heatwaves. The Vic Plain (located
94 north of Barcelona) was the area registering the most annual exceedances of the information threshold in
95 Spain, with an average of 15 exceedances per year per site.

96 In this study, we analyze NO, NO₂ and O₃ surface data around the Barcelona Metropolitan Area (BMA) and the
97 Vic Plain, as well as NO₂ satellite observations, in the period 2005–2017, with the aim of better understanding
98 the occurrence of high O₃ episodes in the area on a long-term basis. Previous studies in this region focused on
99 specific episodes, whereas we aim at assessing the spatial distribution, time trends and temporal patterns of
100 O₃ and its precursors, and the exceedances of the information threshold on a long time series. After better
101 understanding the 2005–2017 O₃ episodes, we aim to evaluate, as a first approximation using air quality
102 monitoring and OMI remote sensing data, the effect that episodic mitigation measures of O₃ precursors would
103 have in the O_x concentrations in the Vic Plain.

104 We recognize that the O₃ problem has to be studied with executable models with dispersion and
105 photochemical modules, which allow performing sensitivity analyses. It is also well recognized that there is a
106 complex O₃ phenomenology in the study area and that although models have greatly improved in the last 10
107 years, there are still problems in reproducing some of the processes in detail, such as the channeling of O₃
108 plumes in narrow valleys or the vertical recirculation patterns. Our study intends to obtain a sensitivity analysis
109 for O₃ concentrations using air quality data. Ongoing collaboration is being established with modelers to try to
110 validate model outputs with this experimental sensitivity analysis and then to implement a prediction system
111 for abating efficiently O₃ precursors to reduce O₃ concentrations, for which executable models are the solely
112 tool available.

113 **2. Methodology**

114

115 **2.1. The area of study**

116 The study is set in central Catalonia (Spain), in the north-eastern corner of the Iberian Peninsula (Figure 1).
117 Characterized by a Mediterranean climate, summers are hot and dry with clear skies. In the 21st century, heat
118 waves have occurred frequently in the area, often associated with high O₃ levels (Vautard et al., 2007; Guerova
119 et al., 2007; Querol et al., 2016; Guo et al., 2017).

120 The capital city, Barcelona, is located on the shoreline of the Mediterranean Sea. Two sets of mountain chains
121 lie parallel to the coastline (SW–NE orientation) and enclose the Pre-coastal Depression: the Coastal (250–500
122 m above sea level (a.s.l.)) and the Pre-Coastal (1000–1500 m a.s.l.) mountain ranges. The Vic Plain, situated
123 45–70 km North of Barcelona (500 m a.s.l.) is a 230 km² plateau that stretches along a S–N direction and is
124 surrounded by high mountains (over 1000 m a.s.l.). The complex topography of the area protects it from
125 Atlantic advections and continental air masses but also hinders the dispersion of pollutants (Baldasano et al.,
126 1994). The two main rivers in the area (Llobregat and Besòs) flow perpendicularly to the sea and frame the
127 city of Barcelona. Both rivers' valleys play an important role in the creation of air-flow patterns. The Congost
128 River is a tributary to the Besòs River and its valley connects the Vic Plain with the Pre-coastal Depression.

129 The BMA stretches across the Pre-Coastal and Coastal Depressions and is a densely populated (>1500 people
130 per km², MFom, 2017) and highly industrialized area with large emissions originating from road traffic, aircraft,
131 shipping, industries, biomass burning, power generation and livestock.

132 During summer, the coupling of daily upslope winds and sea breezes may cause the penetration of polluted
133 air masses up to 160 km inland, channeled from the BMA northward by the complex orography of the area.
134 These air masses are injected at high altitudes (2000–3000 m a.s.l.) by the Pyrenean mountain ranges. At night
135 time, the land breeze prevails, and winds flow toward the sea followed by subsidence sinking of the air mass,
136 which can be transported again by the sea breeze of the following day (Millán et al., 1997, 2000, 2002; Toll
137 and Baldasano, 2000; Gangoiti, 2001; Gonçalves et al., 2009; Millán, 2014; Valverde et al., 2016). Under
138 conditions of a lack of large-scale forcing and the development of a thermal low over the Iberian Peninsula
139 that forces the confluence of surface winds from coastal areas toward the central plateau, this vertical
140 recirculation of the air masses results in regional summer O₃ episodes in the Western Mediterranean. In

addition, there might be external O₃ contributions, such as hemispheric transport or stratospheric intrusions (Kalabokas et al., 2007, 2008, 2017; Querol et al., 2017, 2018).

2.2. Air quality, meteorological and remote sensing data

We evaluated O₃ and NO_x AQ data together with meteorological variables and satellite observations of background NO₂.

The regional government of Catalonia (Generalitat de Catalunya, GC) has a monitoring network of stations that provides average hourly data of air pollutants (XVPCA, GC, 2017a, b). We selected a total of 25 stations (see Figure 2). To study the O₃ phenomenology in the Vic Plain, we selected the 8 stations marked in green, which met the following constraints: (i) location along the S–N axis (Barcelona–Vic Plain–Pre-Pyrenean Range); (ii) availability of O₃ measurements; (iii) availability of at least 9 years of data in the period 2005–2017, with at least 75% data coverage from April to September. The remaining selected stations (used only as reference ones for interpreting data from the main Vic-BMA axis stations) met the following criteria: (i) location across the Catalan territory, and (ii) availability of a minimum of 5 years of valid O₃ data in the period 2005–2017. We chose this period due to the poor data coverage of most of the AQ sites in the regional network of AQ monitoring stations before 2005.

In addition, we selected wind and temperature data from 5 meteorological stations from the Network of Automatic Meteorological Stations (XEMA, Meteocat, 2017) closely located to the previously selected AQ stations, as well as solar radiation data from two solar radiation sites from the Catalan Network of Solar Radiation Measurement Stations (ICAEN-UPC, 2018) located in the cities of Girona and Barcelona.

We also used daily tropospheric NO₂ column satellite measurements using the Ozone Monitoring Instrument (OMI) spectrometer aboard NASA's Earth Observing System (EOS) Aura satellite (see OMI, 2012; Krotkov and Veefkind, 2016). The measurements are suitable for all atmospheric conditions and for sky conditions where cloud fraction is less than 30% binned and averaged into 0.25° × 0.25° global grids.

2.3. Data analysis

2.3.1. O_x calculations

We calculated O_x concentrations to better interpret O₃ dynamics. Kley and Gleiss (1994) proposed the concept of O_x to improve the spatial and temporal variability analysis by decreasing the effect of titration of O₃ by NO with the subsequent consumption of O₃ in areas where NO concentrations are high. Concentrations were transformed to ppb units using the conversion factors at 20 °C and 1 atm (DEFRA, 2014).

O_x concentrations were only calculated if there were at least 6 simultaneous hourly recordings of O₃ and NO₂ from 12:00 to 19:00 h, June–August, in the period 2005–2017. The stations used for these calculations were those located along the S–N axis (Barcelona–Vic Plain–Pre-Pyrenean Range).

2.3.2. Variability of concentrations across the air quality monitoring network

To study the variability of concentrations of NO, NO₂, O₃ and O_x across the air quality monitoring network we calculated June–August averages (months recording the highest concentrations of O₃ in the area) from hourly concentrations provided by all the selected AQ sites. For each of them, we calculated daily averages and daytime high averages (12:00 to 19:00 h).

2.3.3. Time trends

By means of the Mann–Kendall method, we analyzed time trends for NO, NO₂ and O₃ for the period 2005–2017. In addition, we used the Theil–Sen statistical estimator (Theil, 1950; Sen, 1968) implemented in the R package Openair (Carslaw and Ropkins, 2012) to obtain the regression parameters of the trends (slope, uncertainty and p-value) estimated via bootstrap resampling. We examined the annual time trends of seasonal averages (April–September) for each pollutant. Data used for these calculations were selected according to the recommendations in EMEP-CCC (2016): the stations considered have at least 10 years of data (75% of the total period considered, 2005–2017) and at least 75% of the data is available within each season. In addition,

187 we analyzed annual time trends of tropospheric NO₂ measured by satellite along the S–N axis and of
188 greenhouse gases (GHGs) emitted in Catalonia and the average number of vehicles entering the city of
189 Barcelona.

190 **2.3.4. Assessment of O₃ objectives according to air quality standards**

191 We identified the maximum daily 8-hour average concentrations by examining 8-h running averages using
192 hourly data in the period 2005–2017. Each 8-h average was assigned to the day on which it ended (i.e., the
193 first average of one day starts at 17:00 h on the previous day), as determined by EC (2008).

194 To assess the time trends and patterns of the Exceedances of Hourly Information Thresholds (EHITs)
195 established by EC (2008) (hourly mean of O₃ concentration greater than 180 µg m⁻³), we used all the data,
196 independently of the percentage of data availability.

197 **2.3.5. Tropospheric NO₂ column**

198 We analyzed daily average Tropospheric Column NO₂ measurements from 2005 to 2017 aiming at two
199 different goals. On the one hand, to quantify the tropospheric NO₂ in the area along the S–N axis and obtain
200 annual time trends and monthly/weekly patterns. On the other hand, to assess qualitatively the tropospheric
201 NO₂ across a regional scale (Western Mediterranean Europe) in two different scenarios, by means of visually
202 finding patterns that might provide a better understanding of O₃ dynamics in our area of study. The scenarios
203 were: days with the maximum 8-h O₃ average above the 75th percentile at the Vic Plain stations, and days
204 with the maximum below the 25th percentile. See selected regions for retrieval of NO₂ satellite measurements
205 in Figure S1.

206 **2.3.6. Time conventions**

207 When expressing average concentrations, the times shown indicate the start time of the average. For example,
208 12:00–19:00 h averages take into account data registered from 12:00 h to 19:59 h. All times are expressed as
209 local time (UTC + 1 hour during winter and UTC + 2 hours during summer) and the 24-hour time clock
210 convention is used.

211 **3. Results and discussion**

212

213 **3.1. Variability of concentration of pollutants across the air quality monitoring network**

214 We analyzed the mean NO, NO₂, O₃ and O_x concentrations (June to August) in the study area in the period
215 2005–2017.

216 As expected, the highest NO and NO₂ concentrations are registered in urban/suburban (U/SU) traffic sites in
217 and around Barcelona (MON, GRA, MNR and CTL, 7–10 µg NO m⁻³ and CTL and MON 30–36 µg NO₂ m⁻³). Also,
218 as expected, the remote high-altitude rural background (RB) sites (MSY and MSC) register the lowest NO (<1
219 µg m⁻³) and NO₂ (2–4 µg m⁻³) concentrations, see Figure S2.

220 The lowest June–August average O₃ concentrations (45–60 µg m⁻³) are recorded in the same U/SU traffic sites
221 (MON, GRA, MNR and CTL) where titration by NO is notable, while the highest ones (>85 µg m⁻³) are recorded
222 at the RB sites, MSC being the station recording the highest June–August O₃ levels (102 µg m⁻³). These spatial
223 patterns are significantly different when we consider the 8-h daily averages of O₃ concentrations for June–
224 August 12:00–19:00 h (Figure 3a). Thus, these concentrations are repeatedly high (85–115 µg m⁻³) in the whole
225 area of study. The highest O₃ concentrations (>107 µg m⁻³) were recorded at the four sites located downwind
226 of BMA along the S–N corridor (MSY, TON, VIC and MAN), and downwind of Tarragona (PON, RB station).
227 Figure 3a also shows a positive O₃ gradient along the S–N axis (O₃ levels increase farther north) following the
228 BMA plume transport and probably an increase of the mixing layer height (MLH). The higher O₃ production
229 and/or fumigation in the northern areas are further supported by the parallel northward increasing O_x gradient
230 (δO_x Figure 3b). Time series show that in 85% of the valid data in June–August (849 out of 1001 days in 2005–
231 2017) this positive gradient is evident between CTL and TON (δO_{xTON-CTL} > 0). The average O_x increase between

232 CTL in Barcelona and TON is 15 ppb. Taking into account the low NO₂ concentrations registered at this station,
233 this is equivalent to approximately 29 µg m⁻³ of O₃ (+30% O_x in TON compared with CTL).

234 Thus, TON at the Vic Plain records the highest 12:00–19:00 h, June–August O_x and O₃ concentrations in the
235 study area. The MNR site also exhibits very high O_x levels (Figure 3b) but these are mainly caused by primary
236 NO₂ associated with traffic emissions.

237 3.2. Time patterns

238

239 3.2.1. Annual trends

240 Figure 4 shows the results of the trend analysis of NO, NO₂, O₃ and O_x averages (April to September, the O₃
241 season according to the European AQ Directive) by means of the Mann–Kendall test.

242 NO_x levels exhibit a generalized and progressive decrease during the time period across Catalonia. In
243 particular, NO₂ tended to decrease along the S–N axis during the period (U/SU sites CTL, MON and MAN
244 registered –1.6, –2.0 and –1.3% year⁻¹, respectively, with statistical significance in all cases). A similar trend
245 was found for NO in these stations, with higher negative slopes (–2.2, –4.3 and –1.1% year⁻¹, the latter without
246 statistical significance).

247 The annual averages of tropospheric NO₂ across the S–N axis decreased by 35% from 2005 to 2017 (–3.4%
248 year⁻¹ with statistical significance). The marked drop of NO₂ from 2007–2008 can be attributed to the
249 reduction of emissions associated with the financial crisis starting in 2008. The time trends of average traffic
250 (number of vehicles) entering Barcelona City on working days from 2005 to 2016 (Ajuntament de Barcelona,
251 2010, 2017) and the GHGs emitted in Catalonia attributed to industry and power generation sectors calculated
252 from the Emissions Inventories published by the Regional Government of Catalonia from 2005 to 2016 (GC,
253 2017c) (Figure 5a) support this hypothesis. We found both decreasing trends to be statistically significant but
254 the GHG emissions decreasing trend is significantly higher (–3.8% year⁻¹) than the traffic (–1.2% year⁻¹), which
255 suggests that the crisis had a more severe effect on industry and power generation than on road traffic. This
256 is also supported by a larger decrease of GHG emissions and OMI-NO₂ from 2005–2007 (precrisis) to 2008
257 (start of the crisis) than BMA traffic counting and urban NO_x levels (without a 2007–2008 steep change and a
258 more progressive decrease, Figure 5b). Thus, in the BMA, the financial crisis caused a more progressive
259 decrease (without a 2007–2008 steep change) of the circulating vehicles and therefore its associated
260 emissions.

261 April–September O₃ and O_x mean concentration trends are shown in Figure 4. The data show that seven out
262 of the eight RB sites registered slight decreases in O₃ concentrations during the period (BdC, AGU and STP; –
263 1.6% year⁻¹, –1.1% year⁻¹ and –1.4% year⁻¹, respectively, in all cases with statistical significance) while in BEG,
264 PON, LSE and GAN the trends were not significant (not shown). As in several regions of Spain and Europe
265 (Sicard et al., 2013; Paoletti et al., 2014; EEA, 2016; Querol et al., 2016; EMEP, 2016), the opposite trends are
266 found for U/SU sites, with increases in O₃ concentrations during the period at some stations (CTL, MON, MAN,
267 MAT, MNR and ALC; +0.4 to +3.2% year⁻¹ all with statistical significance). When considering O_x, the increasing
268 trends in U/SU sites are neutralized in some cases (CTL, MON, MAN, MAT and ALC). This, and the higher NO
269 decreasing slopes compared with those of NO₂, support the hypothesis that the U/SU O₃ increasing trends are
270 probably caused by less O₃ titration (due to decrements in NO levels) instead of a higher O₃ generation. The
271 marked decrease of the vehicle diesel emissions of NO/NO₂ time trends (Carslaw et al., 2016) might have
272 caused this differential NO and NO₂ trends, although other causes cannot be discarded.

273 3.2.2. Monthly and daily patterns

274 Figure 6a shows 2005–2017 monthly average hourly O₃ concentrations measured at sites along the S–N axis,
275 showing the occurrence of chronic-type episodes with repeated high O₃ concentrations (90–135 µg m⁻³) in the
276 afternoon of April–September days at the Vic Plain sites (TON, VIC, MAN) and the remote RB sites (MSY and
277 PAR).

278 Typically, at the remote RB stations, O₃ concentrations are high during the whole day throughout the year,
279 and daily O₃ variations are narrower than at the other stations, with high average levels even during October–
280 February (MSY: 50–70 and PAR: 50–80 µg m⁻³). During the night these mountain sites are less affected by NO
281 titration, leading to high daily O₃ average concentrations. However, in summer, midday–afternoon
282 concentrations are relatively lower than at the stations located in the S–N valley (TON, VIC, MAN).

283 Regarding monthly average daily O_x (Figure 6b), the profiles of RB sites TON and MSY are very similar to the
284 respective O₃ profiles. In the case of the BMA U/SU sites (CTL, MON, GRA), the nocturnal O_x concentrations
285 increase with respect to O₃ due to the addition of secondary NO₂ from titration. Midday–afternoon O_x levels
286 are much lower at the BMA U/SU stations than those in the S–N valley (MAN, TON), similarly to O₃ levels,
287 supporting the contribution of local-regional O₃ from the BMA plume and/or from the fumigation of high-
288 altitude reserve strata as MLH grows (Millán et al., 1997, 2000; Gangoi et al., 2001; Querol et al., 2017) as
289 well as production of new O₃.

290 3.2.3. Weekly patterns

291 Accordingly, Figure 7 shows the O₃ weekly patterns for these O₃ average concentrations. As expected, the
292 variation of intra-annual concentration values is pronounced in the Vic Plain sites (TON, VIC, MAN; 20–45 µg
293 m⁻³ in December–January versus 110–125 µg m⁻³ in July), due to the higher summer photochemistry, the more
294 frequent summer BMA plume transport (due to intense sea breezing) and fumigation from upper atmospheric
295 reservoirs across the S–N axis, and of the high O₃ titration in the populated valleys in winter. However, at the
296 remote mountain sites of MSY and PAR, the intra-annual variability is much reduced (70–80 µg m⁻³ in
297 December versus 100–120 µg m⁻³ in July) probably due to the reduced effect of NO titration at these higher
298 altitude sites, and the influence of high-altitude O₃ regional reservoirs.

299 During the year, CTL, MON and GRA (U/SU sites around BMA) register very similar weekly patterns of the 8-h
300 maxima, with a marked and typical high O₃ weekend effect, i.e., higher O₃ levels than during the week due to
301 lower NO concentrations. From April to September, CTL O₃ 8-h concentrations are lower than MON's and
302 GRA's (the latter located north of BMA following the sea breeze air mass transport), despite being very similar
303 from October to March (when sea breezes are weaker). An O₃ weekend effect is also clearly evident during
304 the winter months in the Vic Plain sites (TON, VIC, MAN) and MSY. However, from June to August, a marked
305 inverse weekend effect is clearly evident at these same sites, with higher O₃ levels during weekdays. This
306 points again to the clear influence of the emission of precursors from the BMA on the O₃ concentrations
307 recorded at these inland sites.

308 We carried out a trend analysis of NO, NO₂ and O₃ levels measured at AQ sites and background NO₂ from
309 remote sensing (OMI) for weekday (W) and weekend (WE) days independently. To this end we averaged the
310 concentrations for 3 sites in the BMA (CTL, MON and GRA) and 3 receptor sites at the Vic Plain (TON, VIC and
311 MAN), and considering WE to be Saturday, Sunday and Monday for the Vic AQ sites data (adding Mondays to
312 account for the “clean Sunday effect”) and Saturday and Sunday for the BMA sites data.

313
314 We estimated time trends of W and WE concentrations separately by the Mann-Kendall method along the
315 study period. For O₃ (12:00 to 19:00 h) we found statistically significant increases in both the BMA and the Vic
316 Plain. Increases of O₃ in the BMA double the ones in the Vic Plain and trends of W and WE are very similar per
317 area (O₃ BMA W: +2.0 % year⁻¹, O₃ BMA WE: +2.2 % year⁻¹, O₃ Vic Plain W: +0.8 % year⁻¹, O₃ Vic Plain WE: +1.0
318 % year⁻¹). As seen before, both NO and NO₂ levels (daily averages) in the BMA decrease in a statistically
319 significant way, where NO decrements are larger than NO₂. We found that the decrease of W NO levels is
320 higher than the WE ones (NO BMA W: -3.4 % year⁻¹, NO BMA WE: -2.7 % year⁻¹) because emissions are higher
321 during W days and these decreased along the period. Regarding NO₂, W and WE decreases remain similar (NO₂
322 BMA W: -1.9 % year⁻¹, NO₂ BMA WE: -1.7 % year⁻¹) but lower than NO in both cases thus reducing the O₃
323 titration effects and increasing O₃ levels both in WE and W days. Regarding NO₂-OMI levels, only W levels show
324 a statistically significant decreasing trend (-3.4 % year⁻¹) and not the WE levels.

325

We then assessed the variations of WE concentrations with respect to W's per year and plotted them by short tilted lines in Figure 8, where the left and right side of each tilted line represent W and WE concentrations respectively. These W to WE variations are then plotted in percentage by continuous lines (>0 depicts increase and <0 decrease W to WE). The upper plot shows O_3 data averaged from 12:00 to 19:00 h from the BMA and the Vic Plain, the middle plot daily averages of NO and NO_2 concentrations in BMA and the bottom plot, daily NO_2 -OMI levels along the S-N axis. The results evidence again a constant drop in W to WE NO_x levels in the BMA along the period (negative percentages in the middle plot), with the subsequent O_3 weekend effect in the BMA (positive percentages in the upper plot). In the Vic Plain sites, O_3 concentrations remain constantly high along the study period showing inverse weekend effect almost during the whole period (negative percentages in the plot, except for 2005 to 2007 and 2017). Using the Mann-Kendall test to estimate trends for the W to WE variations we found a clear statistically significant decreasing trend along the period (reduction of the difference between W to WE levels: from -38% in 2005 to -17% in 2017, Figure 8 bottom). We attribute this to the decrease of W- NO_x levels, described before for the annual averages.

Furthermore we found a pattern of nearly parallel O_3 W to WE variation cycles between the Vic Plain and the BMA sites (Figure 8, upper). Due to the inverse W to WE O_3 at Vic and BMA, this parallel trend means in fact that maximum W to WE variations in the Vic Plain and the BMA tend to follow a reverse behavior, i.e. maximum W to WE variations in the BMA tend to occur when W to WE variations in the Vic Plain are minimum (for example 2007, 2010, 2014). NO_x W to WE variations tend to follow a similar behavior than O_3 W to WE variations in the Vic Plain sites (mostly from 2008 to 2016) where years with high W to WE variations of NO_x in the BMA tend to correspond to years with maximum O_3 W to WE variations in the Vic Plain (2009 and 2015). This behavior is probably associated to differences on air mass circulation patterns along the period (such as higher or lower breeze development). Those years with lower breeze development, the transport of the BMA plume is weaker; then NO_x would tend to accumulate at the BMA (low W to WE NO_x variation) which would generate more O_3 thus W to WE variation would be higher in the BMA and lower in the Vic Plain. As opposed, years with stronger breeze development and thus increased transport of the BMA plume, W to WE variations of NO_x in the BMA are higher, W to WE variations of O_3 in the BMA are lower (less O_3 is generated during WE) and higher W to WE O_3 variations are recorded in the Vic Plain sites.

3.3. Peak O_3 concentrations patterns along the S-N axis

July is the month of the year when most of the annual exceedances of the O_3 EHITs are recorded in Spain (Querol et al., 2016), including our area of study. Figure 9 shows the average O_3 and O_x July hourly concentrations along the S-N axis during 2005–2017. A progressive time-shift and a marked positive northward gradient of O_3 and O_x maxima are shown, pointing again to the gradual increase of O_3 and O_x due to the plume transport, new O_3 formation and fumigation from upper reservoirs as MLH grows.

Figure 10a shows the 2005–2017 trends of the EHITs from the European AQ Directive ($>180 \mu g m^{-3} h^{-1}$ mean; EC, 2008) registered at the selected sites in the S-N valley, as well as the average temperatures measured during July at early afternoon near Vic (at Gurb meteorological site), the background NO_2 measured by OMI (June to August) and the average solar radiation measured in Girona and Barcelona (June to August). In 2005, 2006, 2010, 2013, 2015 and 2017, the highest EHITs at almost all the sites were recorded. Temperature and insolation seem to have a major role in the occurrence of EHITs in 2006, 2010, 2015 and 2017. The effect of heat waves on O_3 episodes is widely known (Solberg et al., 2008; Meehl et al., 2018; Pyrgou et al., 2018; among others). However, because the emissions of precursors have clearly decreased ($\sim 30\%$ decrease on June to August OMI- NO_2 levels across the S-N axis from 2005 to 2017; $-2.7\% year^{-1}$ with statistical significance) the number of EHITs recorded in the warmest years has probably decreased with respect to a scenario where emissions would have been maintained. In any case, some years (for example 2009 and 2016) seem to be out of line for temperature and insolation being the driving forces, and other major causes also have to be relevant, with further research needed to interpret fully interannual trends. Otero et al. (2016) found that temperature is not the main driver of O_3 in the South-western Mediterranean, as it is in Central Europe, but the O_3 levels recorded the day before (a statistical proxy for the occurrence of Millán et al. (1997)'s vertical

recirculation of air masses). Again, the Vic Plain sites (TON, VIC, MAN) recorded most (75%) of the EHITs reported by the AQ monitoring stations in Catalonia (25%, 34% and 16%, respectively). The higher urban pattern of MAN, as shown by the higher NO concentrations, with respect to TON, might account for both the lower exceedances and the different interannual patterns.

Figure 10b shows that most EHITs occurred in June and July (30% and 57%, respectively), with much less frequency in May, August and September (6%, 8% and <1%, respectively). Although temperatures are higher in August than in June, the latter registers significantly more EHITs, probably due to both the stronger solar radiation and the higher concentrations of precursors (such as NO₂, see OMI-NO₂ and solar radiation in Figure 10b).

Figure 10c shows that EHITs occurred mainly between Tuesday and Friday (average of 19% of occurrences per day). On weekends and Mondays, EHITs were clearly lower (average of 9% of occurrences per day) than during the rest of the week, probably due to: (i) the lower emissions of anthropogenic O₃ precursors (such as NO_x, see OMI-NO₂) during weekends and (ii) to the effect of the lower Sunday emissions in the case of the lower exceedances recorded during Mondays. During weekends and in August, OMI-NO₂ along the S–N axis is relatively lower (–29% weekday average and –43% in August with respect to March) following the emissions patterns associated with industrial and traffic activity that drop during vacations and weekends (Figure 10). NO_x data from AQ monitoring sites follow similar patterns (not shown here).

Figure 10d shows that the frequency of occurrence of the EHITs at MSY (45 km north of Barcelona) is lower and earlier (maxima at 14:00 h) than at Vic Plain sites (TON, VIC, MAN). The EHITs occurred mostly at 15:00, 16:00, 16:00 and 19:00 h at TON, VIC, MAN and PAR (53, 63, 72 and 105 km north of Barcelona), respectively. PAR registered not only much later EHITs, but a much lower number than TON-VIC-MAN sites, again confirming the progressive O₃ maxima time-shift northward of Barcelona.

The results in Figure 11 clearly show that during non-EHIT days, the daily O₃ patterns are governed by the morning–midday concentration growth driven to fumigation and photochemical production, while on EHIT days there is a later abrupt increase, with maxima being delayed as we increase the distance from Barcelona along the S–N axis. This maximal second increase of O₃ is clearly attributable to the influence of the transport of the plume of the BMA (horizontal transport), as the secondary NO₂ peak at 15:00 h (Figure 11 left bottom), and the wind patterns (see Figure S3) seem to support. The differences in the late hourly O₃ concentration increases in EHIT versus non-EHIT days are even more evident when calculating hourly O₃ slopes (hourly increments or decrements of concentrations), Figure 11 (right). The first increment (fumigation and photochemistry) makes O₃ levels scale up to 120 µg m^{–3} during EHIT episodes and to nearly 100 µg m^{–3} during non-EHIT days. In EHIT days, the later peak (transport from BMA and causing most of the 180 µg m^{–3} exceedances) in the O₃ slope occurs again between 14:00 h and 20:00 h, depending on the distance to BMA, but this feature is not observed on non-EHIT days.

3.4. Relevance of local/regional pollution plumes in high O₃ episodes in NE Spain

Figure 12 depicts the basic atmospheric dynamics in the study area during a typical summer day, when the atmospheric conditions are dominated by mesoscale circulations. According to the previous references, indicated in Figure 12 with enclosed numbering (coinciding with the numbering below) the following O₃ contributions to surface concentrations in the study area can be differentiated:

- a. Vertical recirculation of O₃-rich air masses, which create reservoir layers of aged pollutants.
- b. Vertical fumigation of O₃ from the above reservoirs and the following sources aloft if the MLH growth is large enough:
 - b.1. Regional external O₃ layers (from other regions of southern Europe, such as southern France, Italy, Portugal and Tarragona).
 - b.2. High free tropospheric O₃ background due to hemispheric long-range transport.
 - b.3. High free tropospheric O₃ background due to stratospheric intrusions.

- c. Horizontal transport of O₃. Diurnal BMA plume northward transported and channeled into the Besòs–Congost valleys.
- d. Local production of O₃ from precursors.

During summer, the intense land heating due to strong solar radiation begins early in the morning. The associated convective activity produces morning fumigation processes (b in Figure 12) that bring down O₃ from the reservoir layers aloft, creating sharp increases in O₃ concentrations in the morning (see Figure 11 and S3). The breeze transports air masses from the sea inland and creates a compensatory subsidence of aged pollutants (including O₃) previously retained in reservoir and external layers and high free troposphere background aloft (Millán et al., 1997, 2000; Gangoiti et al., 2001). This subsided O₃ then affects the marine boundary layer and reaches the city the following day with the sea breeze, producing nearly constant O₃ concentrations in the city during the day (Figure S3 and Figure 9). As the breeze develops, coastal emissions and their photochemical products are transported inland, generating the BMA plume (c in Figure 12) that, in addition to the daily generated O₃, also contains recirculated O₃ from the marine air masses. Furthermore, during the transport to the Vic Plain, new O₃ is produced (d in Figure 12) by the intense solar radiation and the O₃ precursors emitted along the way (e.g., BVOCs from vegetation, NO_x from industrial and urban areas, highways).

This new O₃ gets mixed with the BMA plume and channeled northward to the S–N valleys until it reaches the Vic Plain and the southern slopes of the Pre-Pyrenees. As the BMA plume (loaded with O₃ and precursors) travels northward, a second increase in O₃ concentrations can be observed in the daily cycles of O₃ at these sites, (see Figure 11 and S3). This was described as the second O₃ peak by Millán et al. (2000).

The marked MLH increase in the Vic Plain compared with BMA (Soriano et al., 2001; Querol et al., 2017) may produce a preferential and intensive top-down O₃ transport (b in Figure 12) from upper O₃ layers (a, b.1, b.2 and b.3 in Figure 12), contributing to high O₃ surface concentrations. During the sea/mountain breezes' development, some air masses are injected upward to the N and NW return flows (controlled by the synoptic circulations dominated by the high-pressure system over the Azores) aloft helped by the orography (e.g., southern slopes of mountains) and again transported back to the coastal areas where, at late evening/night it can accumulate at certain altitudes in stably stratified layers.

Later, at night, land breezes returning to the coastal areas develop. Depending on the orography, these drainage flows of colder air traveling to the coastal areas can accumulate on the surface or keep flowing to the sea. The transported O₃ is consumed along the course of the drainage flows by deposition and titration. Next day, the cycle starts anew, producing almost closed loops enhancing O₃ concentrations throughout the days in the area. When the loop is active for several days, multiple O₃ EHITs occur over the Vic Plain.

The main complexity of this system arises from the fact that all these vertical/horizontal, local/regional/hemispheric/stratospheric contributions are mixed and all contribute to surface O₃ concentrations with different proportions that may largely vary with time and space across the study area. However, for the most intense O₃ episodes, the local-regional contribution might be very relevant to cause EHITs in the region. Furthermore, the intensity and frequency of O₃ episodes are partially driven by the occurrence of heat waves in summer and spring (Vautard et al., 2007; Geroova et al., 2007; Querol et al., 2016; Guo et al., 2017). If local and regional emissions of precursors are high, the intensity of the episodes will also be high. Thus, even though heat wave occurrences increase the severity of O₃ episodes, an effort to reduce precursors should be undertaken to decrease their intensity.

The generation of the O₃ episodes in 2005–2017 for the S–N corridor BMA–Vic Plain–Pre-Pyrenees occurs in atmospheric scenarios described in detail by Millán et al. (1997, 2000, 2002), Gangoiti et al. (2001), Kalabokas et al. (2007, 2008, 2017), Millán (2014) and Querol et al. (2018) for other regions of the Mediterranean basin, including Spain, or described in the same area for specific episodes (Toll and Baldasano, 2000; Gonçalves et al., 2009; Valverde et al., 2016; Querol et al., 2017). However, results from our study evidence a higher role of

the local-regional emissions on the occurrence of O₃ EHITs. Thus, our results demonstrate an increase in the EHITs northward from Barcelona to around 70 km and a decrease from there to 100 km from Barcelona following the same direction. There is also a higher frequency of occurrence of these in July (and June) and from Tuesday to Friday and a time-shift of the frequency of occurrence of EHITs from 45 to 100 km. The mountain site of MSY (located at 700 m a.s.l.) registered many fewer EHITs than the sites in the valleys (TON-VIC-MAN, 460–600 m a.s.l.) during the period, showing the key role of the valley channeling of the high O₃ and precursors BMA plume in July (when sea breeze and insolation are more intense). Furthermore, at the Vic Plain, we detected an inverse O₃ weekend effect, suggesting that local/regional anthropogenic emissions of precursors play a key role in increasing the number of EHITs on working days, with a Friday/Sunday rate of 5 for VIC for 2005–2017. Despite this clear influence of the BMA plume on EHITs' occurrence, Querol et al. (2017) demonstrated that at high atmospheric altitudes (2000–3000 m a.s.l.) high O₃ concentrations are recorded, in many cases reaching 150 µg m⁻³ due to the frequent occurrence of reservoir strata. As also described above, the higher growth of the MLH in TON-VIC-MAN as compared with the coastal area accounts also for higher top-down O₃ contributions. On the other side, close to the Pyrenees (PAR station), large forested and more humid areas give rise to a thinner MLH, hindering O₃ fumigation too. Furthermore, in these more distant northern regions O₃ consumption by ozonolysis of BVOCs might prevail over production due to weaker solar radiation during the later afternoon.

Figure 13 shows the distribution of average background OMI-NO₂ levels across the Western Mediterranean Basin in two different scenarios: when the O₃ levels in the Vic Plain are low (left) or high (right). To this end, we averaged the values from VIC and TON (in the Vic Plain) from all the maximum daily 8-h mean O₃ concentrations calculated for all the days in July within 2005–2017, and we calculated the 25th (93 out of 370 days, 105 µg m⁻³) and 75th (93 days, 139.5 µg m⁻³) percentiles of all the data (P25 and P75, respectively). For both scenarios, NO₂ concentrations are highest around large urban and industrial areas, including Madrid, Porto, Lisbon, Barcelona, Valencia, Paris, Frankfurt, Marseille and especially the Po Valley. The shipping routes toward the Gibraltar Strait and around the Mediterranean can be observed, as well as important highways such as those connecting Barcelona to France and Lyon to Marseille. As expected, the mountain regions (the Pyrenees and the Alps) are the areas with lower NO₂. Regional levels of background OMI-NO₂ in the P75 scenario are markedly higher with hotspots intensified and spanning over broader areas. Over Spain, new hotspots (marked in yellow), such as the coal-fired power plants in Asturias (a), ceramic industries in Castelló (c) and the coal-fired power plant in Andorra, Teruel (b), appear; in the latter case, with the pollution plume being channeled along the Ebro Valley with a NW transport. Furthermore, it is important to highlight that the maxima background NO₂ along the eastern coastline in Spain, including the BMA, tend to exhibit some north-northwest displacement, when compared with the P25 scenario, thus pointing to the relevance of the local emissions in causing inland O₃ episodes.

These qualitative results suggest in general less synoptic forcing in Western Europe in the P75 scenario; hence, in these conditions NO₂ is accumulated across the region and especially around its sources. In the east coast of the Iberian Peninsula, mesoscale circulations tend to dominate, hence the northwest displacement (taking the coastal regions as a reference) of the background NO₂. The bottom part of Figure 13 zooms our study area and shows the maximum daily 8-h mean O₃ concentrations in all the selected AQ sites averaged for both scenarios. As shown in the P75 scenario, NO₂ is significantly intensified across Catalonia, especially north of the BMA spreading to the Vic Plain. Comparing O₃ in both scenarios, in the P75 the O₃ levels are much higher (mostly >105 µg m⁻³), across the region except the urban sites in Barcelona (due to NO titration), reaching up to 154 µg m⁻³ in the Vic Plain.

Conversely, in the P25 scenario, background NO₂ concentrations are lower, the BMA NO₂ spot is significantly smaller and spreads along the coastline rather than being displaced to the north-northwest. In this case, synoptic flows seem to weaken sea breeze circulations and vertical recirculation, thus reducing the amount of background NO₂ and the inland transport from the coast. In these conditions, O₃ levels are markedly lower across the territory, the RB PON site (downwind of the city/industrial area of Tarragona) being the one recording the maximum daily 8-h mean O₃ concentration (99 µg m⁻³).

3.5. Sensitivity analysis for O_x using air quality monitoring data

We demonstrated above that the lower anthropogenic emissions of O₃ precursors in the BMA during weekends cause lower O₃ and O_x levels in the Vic Plain than during working days (inverse O₃ weekend effect). To apply a sensitivity analysis using air quality monitoring data for the O₃ levels in the Vic Plain if BMA's emissions were reduced, we compared weekend O₃ and O_x patterns with weekdays considering only data from June and July (August OMI-NO₂ levels are markedly lower, Figure 10b, therefore this month was not included).

Figure 14 shows the average O_x concentrations (12:00 to 19:00 h) in TON and MAN (both AQ sites in the Vic Plain) according to the day of the week for the period considered. Data in VIC cannot be used for O_x calculations due to the lack of NO₂ measurements. Despite the large variability in extreme values (i.e., maximum values with respect to minimum values, represented by whiskers), the interquartile range is quite constant on all the weekdays (between 13.6 to 17.3 ppb in TON and 12.7 to 19.1 in MAN). The average O_x decrease between the day with highest O_x levels (Wednesday in TON and Friday in MAN) and the day with the lowest O_x levels (Sunday in TON and Monday in MAN) is between 6.5 (TON) and 7.7 ppb (MAN), approximately 13 and 15 µg O₃ m⁻³, 10-12% decrease). The observed decrements on O_x levels downwind BMA due to the reduction in O₃ precursors' emissions in the BMA during weekends, can give us a first approximation of the effect that episodic mitigation measures could have on the O_x or O₃ levels in the Vic Plain. Thus, we considered feasible a scenario with a maximum potential of O_x reduction of 24.5 ppb (approximately 49 µg O₃ m⁻³, 32% decrease) when applying episodic mitigation measures (lasting 1-2 days equivalent to a weekend when, on average, NO and NO₂ are reduced 51 and 21%, respectively, compared with week days in the BMA monitoring sites). This was calculated as the difference between the P75 of O_x values observed on Wednesdays minus the P25 of O_x values on Sundays. Obviously, if these mitigation measures would be implemented structurally, instead of episodically, O_x and O₃ decreases would be probably larger because not only the local O₃ coming from the BMA plume would be reduced but also the recirculated O₃ and thus the intensity of O₃ fumigation in the Plain. Therefore, it is probable that both structural and episodic measures to abate VOCs and NO_x emissions in the BMA would result in evident reductions of O₃ in the Vic Plain, as evidenced by modeling tools by Valverde et al. (2016).

4. Conclusions

We analyzed 2005–2017 data sets on ozone (O₃) concentrations in an area frequently affected by the northward atmospheric plume transport of Barcelona Metropolitan Area (BMA) to the Vic Plain, the area of Spain recording the highest number of exceedances of the hourly O₃ information threshold (EHIT, 180 µg m⁻³). We aimed at evaluating the potential benefits of implementing local short-term measures to abate emissions of precursors. To this end, we analyzed in detail spatial and time (interannual, weekly, daily and hourly) variations of the concentration of O₃ and nitrogen oxides (including remote sensing data for the latter) in April–September and built a conceptual model for the occurrence of high O₃ episodes. Finally, a sensitivity analysis is done with the AQ data to evaluate potential O₃ reductions in the North of the BMA on Sundays, compared with weekdays, as a consequence of the reduction of emissions of precursors.

Results showed a generalized decrease trend for regional background O₃ ranging from –1.1 to –1.6% year⁻¹, as well as the well-known increase of urban O₃ (+0.4 to +3.2% year⁻¹) and higher urban NO decreasing slopes than those of NO₂ (–2.2 to –4.3 and –1.3 to –2.0% year⁻¹, respectively), that might account in part for the urban O₃ increase.

The most intensive O₃ episodes in the North of the BMA have O₃ contributions from relatively high regional background O₃ (due to a mix of continental, hemispheric–tropospheric and stratospheric contributions) as well as O₃ surface fumigation from the mid-troposphere high O₃ upper layers arising from the concatenation of the vertical recirculation of air masses (as a result of the interaction of a complex topography with intensive spring–summer sea and mountain breezes circulations (Millán et al., 1997, 2000; Gangoiti et al., 2001; Valverde et al., 2016; Querol et al., 2017). However, we noticed that for most EHIT days in the Vic Plain, the exceedance occurs when an additional contribution is added to the previous two: O₃ supply by the channeling of the BMA pollution plume along the S–N valley connecting BMA and Vic. Thus, despite the large external O₃

contributions, structural and short-time local measures to abate emissions of precursors might clearly influence spring–summer O₃ in the Vic Plain. This is supported by (i) the reduced hourly exceedances of the O₃ information threshold recorded on Sundays at the Vic AQ monitoring site (9 in 2005–2017) compared with those on Fridays (47), as well as by (ii) the occurrence of a typical and marked Sunday O₃ pattern at the BMA AQ monitoring sites and an also marked but opposite one in the sites of the Vic Plain; and (iii) marked increase of remote sensing OMI-NO₂ concentrations over the BMA and northern regions during days of the P75 diurnal O₃ concentrations compared with those of the P25.

Finally, we calculated the difference between the P75 of O_x diurnal concentrations recorded at two of the Vic Plain AQ monitoring stations for Wednesdays minus those of the P25 percentile of O_x for Sundays, equivalent to 1–2 days of emissions reductions in the BMA. A maximum decrease potential by applying short-term measures of 24.5 ppb (approximately 49 µg O₃ m⁻³, 32% decrease) of the diurnal concentrations was calculated. Obviously, structurally implemented measures, instead of episodic ones, would result probably in important additional O_x and O₃ abatements because not only the local O₃ coming from the BMA plume would be reduced but also the recirculated O₃, and thus the intensity of O₃ fumigation on the Plain. Therefore, it is highly probable that both structural and episodic measures to abate NO_x and VOCs emissions in the BMA would result in evident reductions of O₃ in the Vic Plain.

Author contributions

JM performed the data compilation, treatment and analysis with the aid of XQ, CC and ME. JM, CC, ME, JB, AA and XQ contributed to the discussion and interpretation of the results. JM and XQ wrote the manuscript. JM, CC, ME, JB, AA and XQ commented on the manuscript.

Competing interests

The authors declare that there is no conflict of interest.

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840 **FIGURE CAPTIONS**

841 Figure 1. Location and main topographic features of the area of study.

842 Figure 2. Location (left) and main characteristics (right) of the selected air quality monitoring sites (S–N axis:
843 green squares on the map and shaded gray on the table, rest of stations: white squares) and
844 meteorological/solar radiation stations (red circles) selected for this study. Types of air quality monitoring sites
845 are urban (traffic or background: UT, UB), suburban (traffic, industrial or background: SUT, SUI, SUB) and rural
846 (background or industrial: RB, RI). PLR (Palau Reial air quality monitoring site) and BCN (Barcelona)
847 meteorological and solar radiation sites are closely located.

848 Figure 3. Spatial variability of mean June–August O₃ (a) and O_x (b) concentrations from 12:00 to 19:00 h
849 observed in selected air quality monitoring sites. Data from Ciutadella (CTL), Palau Reial (PLR), Montcada
850 (MON), Granollers (GRA), Montseny (MSY), Tona (TON), Vic (VIC), Manlleu (MAN), Pardines (PAR), Montsec
851 (MSC), Begur (BEG), Bellver de Cerdanya (BdC), Berga (BER), Agullana (AGU), Santa Pau (STP), Mataró (MAT),
852 Manresa (MNR), Ponts (PON), Sort (SOR), Juneda (JUN), La Sénia (LSE), Constantí (CON), Gandesa (GAN),
853 Vilanova i la Geltrú (VGe) and Alcover (ALC) air quality monitoring stations.

854 Figure 4. Results of the time trend assessment carried out for annual season averages (April–September) of
855 NO (a), NO₂ (b), O₃ (c & d) and O_x (e) levels using the Theil–Sen statistical estimator shown graphically. Only
856 shown the trends with statistical significance. (d) Numerical results; the symbols shown for the p-values
857 related to how statistically significant the trend estimate is: $p < 0.001 = ***$ (highest statistical significance), p
858 $< 0.01 = **$ (mid), $p < 0.05 = *$ (moderate), $p < 0.1 = +$ (low). No symbol means lack of significant trend. Units
859 are $\mu\text{g m}^{-3}$. Shaded air quality monitoring sites belong to the S–N axis. Types of air quality monitoring sites are
860 urban (traffic or background: UT, UB), suburban (traffic, industrial or background: SUT, SUI, SUB) and rural
861 (background: RB). Data from AQ stations with at least 10 years of valid data within the period.

862 Figure 5. (a) Annual average traffic entering Barcelona City during weekdays (weekends not considered) during
863 2005–2016 versus GHG emissions (attributed to industry and power generation sectors) in Catalonia during
864 2005–2016. (b) Annual NO_x measured at CTL (Ciutadella) and MON (Montcada) air quality monitoring sites
865 versus annual OMI-NASA's measured background NO₂ during 2005–2017.

866 Figure 6. Monthly hourly average concentrations of O₃ (a) and O_x (b) along the S–N axis during 2005–2017.
867 Data from Ciutadella (CTL), Montcada (MON), Granollers (GRA), Montseny (MSY), Tona (TON), Vic (VIC),
868 Manlleu (MAN) and Pardines (PAR) air quality monitoring stations.

869 Figure 7. Monthly weekday average concentrations of O₃ concentrations calculated between 12:00 and 19:00
870 h along the S–N axis during 2005–2017. Data from Ciutadella (CTL), Montcada (MON), Granollers (GRA),
871 Montseny (MSY), Tona (TON), Vic (VIC), Manlleu (MAN) and Pardines (PAR) air quality monitoring stations.

872 Figure 8. Weekday (W) (Monday to Friday in the BMA and Tuesday to Friday in the Vic Plain) to Weekend (WE)
873 pollutant concentrations (O₃, NO and NO₂) measured at AQ sites and background NO₂ (remote sensing OMI)
874 for June to August, per year along the period 2005–2017. O₃ concentrations (top plot) are averaged from 12:00
875 to 19:00 h LT hourly concentrations, and NO and NO₂ concentrations are calculated from daily averages,
876 including OMI-NO₂. Each short line depicts the increasing or decreasing tendency of weekday concentrations
877 (left side of each short line) with respect to weekend levels (right side of the short line). Thus, a horizontal line
878 would represent same pollutant levels along the week (concentration in W = concentration in WE). We
879 consider BMA AQ sites: CTL, MON and GRA and Vic Plain AQ sites: TON and MAN. The continuous lines show
880 the percentage of variation of pollutant levels during weekends with respect to weekdays: increasing (>0) or
881 decreasing (<0) i.e. a quantification of the inclination of each short line.
882

883 Figure 9. (a) July O₃ and (b) O_x daily cycles plotted from mean hourly concentrations measured in air quality
884 monitoring sites located along the S–N axis during 2005–2017. The black arrows point to the O₃ and O_x maxima

time of the day. Data from Ciutadella (CTL), Montcada (MON), Granollers (GRA), Montseny (MSY), Tona (TON), Vic (VIC), Manlleu (MAN) and Pardines (PAR) air quality monitoring stations.

Figure 10. For the period 2005–2017, trends of the EHITs measured by air quality monitoring stations along the S–N axis (a) Annual trends of the EHITs, average temperatures measured in Vic (Gurb) (July during 13:00 to 16:00 h), background NO₂ measured by OMI-NASA (June to August) and average solar radiation measured at Girona and Barcelona (June to August). (b) Monthly patterns of the EHITs, average temperatures measured in Vic, background NO₂ measured by OMI and solar radiation measured at Girona and Barcelona. (c) Weekly patterns of the EHITs and background NO₂ measured by OMI. (d) Hourly patterns of the EHITs. Despite the incomplete data availability in MAN 2005, almost 20 EHITs were recorded. AQ data from Ciutadella (CTL), Montcada (MON), Granollers (GRA), Montseny (MSY), Tona (TON), Vic (VIC), Manlleu (MAN) and Pardines (PAR) monitoring stations.

Figure 11. Average hourly O₃ concentrations for all days with EHIT records and those without for Tona (TON), Vic (VIC), Manlleu (MAN) and Pardines (PAR) air quality monitoring stations, (left top) as well as for the NO₂ levels at TON (left bottom). Average hourly increments of O₃ concentrations for all days with and without EHIT records (right); in all cases for June–August 2005–2017.

Figure 12. Idealized two-dimensional section of O₃ circulations in the coastal region of Barcelona to the Pre-Pyrenees on a typical summer day (upper) and night (bottom). The gray shaded shape represents a topographic profile south to north direction, from the Mediterranean Sea to the south slopes of the Pre-Pyrenean Ranges (i.e., along the S–N axis). The colored dots and abbreviations depict the air quality monitoring stations located along the S–N axis: Ciutadella (CTL), Montcada (MON), Granollers (GRA), Montseny (MSY), Tona (TON), Vic (VIC), Manlleu (MAN) and Pardines (PAR). Modified and adapted to the S–N axis from Millán et al. (1997, 2000), Querol et al. (2017, 2018).

Figure 13. Daily average background NO₂ levels in Western Europe (top) and Catalonia (bottom), July 2005–2017 in two different scenarios. (Left) P25: days when the maximum daily 8-h mean O₃ concentrations in the Vic Plain are below the percentile 25 (<105 µg m⁻³) and (right) P75: same but concentrations being above the percentile 75 (>139.5 µg m⁻³).

Figure 14. Box plots of O_x measured in TON and MAN (12:00 to 19:00h) per weekday June and July 2005–2017 for those days with $\delta O_{x \text{ TON-CTL}} > 0$ (n = 545 for TON and n = 479 for MAN of valid data). Each box represents the central half of the data between the lower quartile (P25) and the upper quartile (P75). The lines across the box displays the median values. The whiskers that extend from the bottom and the top of the box represent the extent of the main body of data. The outliers are represented by black points.

FIGURES

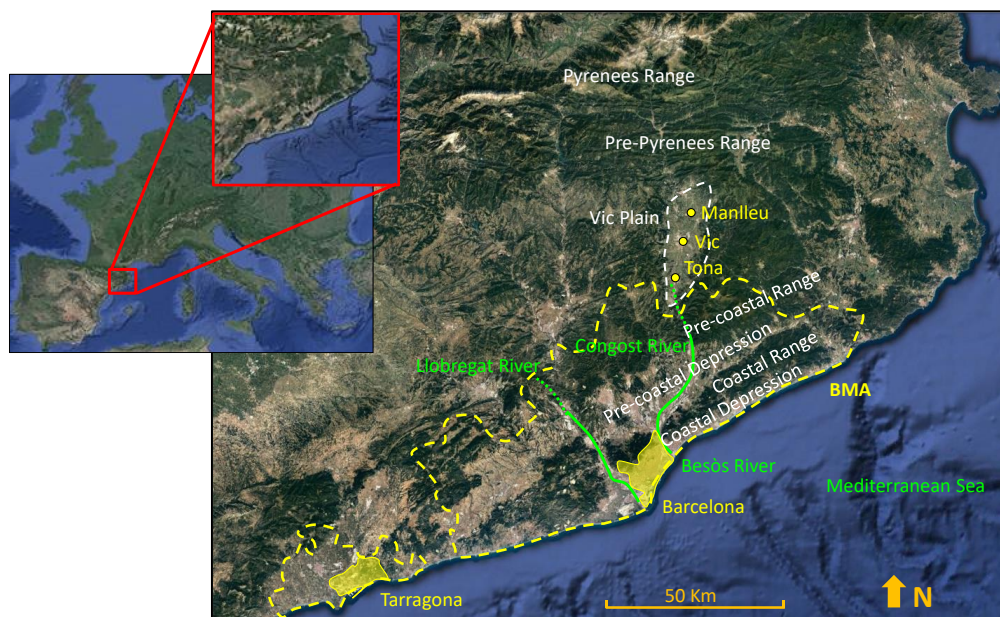


Figure 1

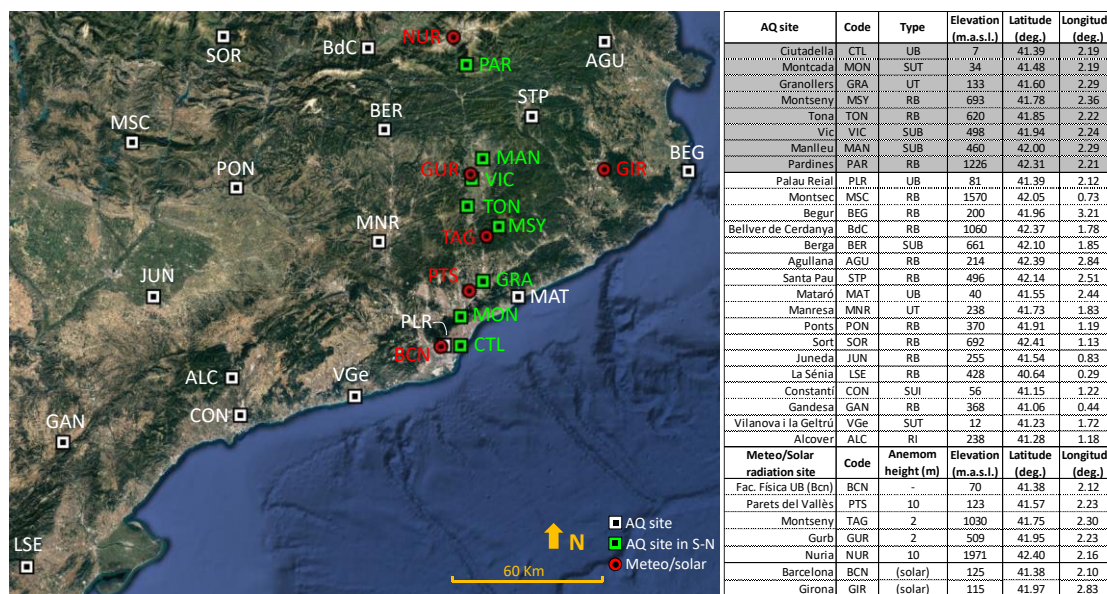
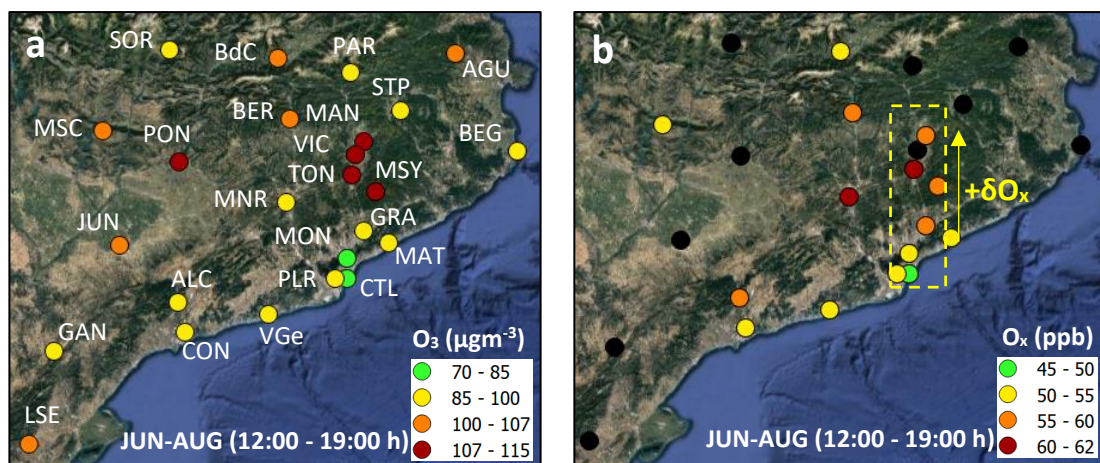
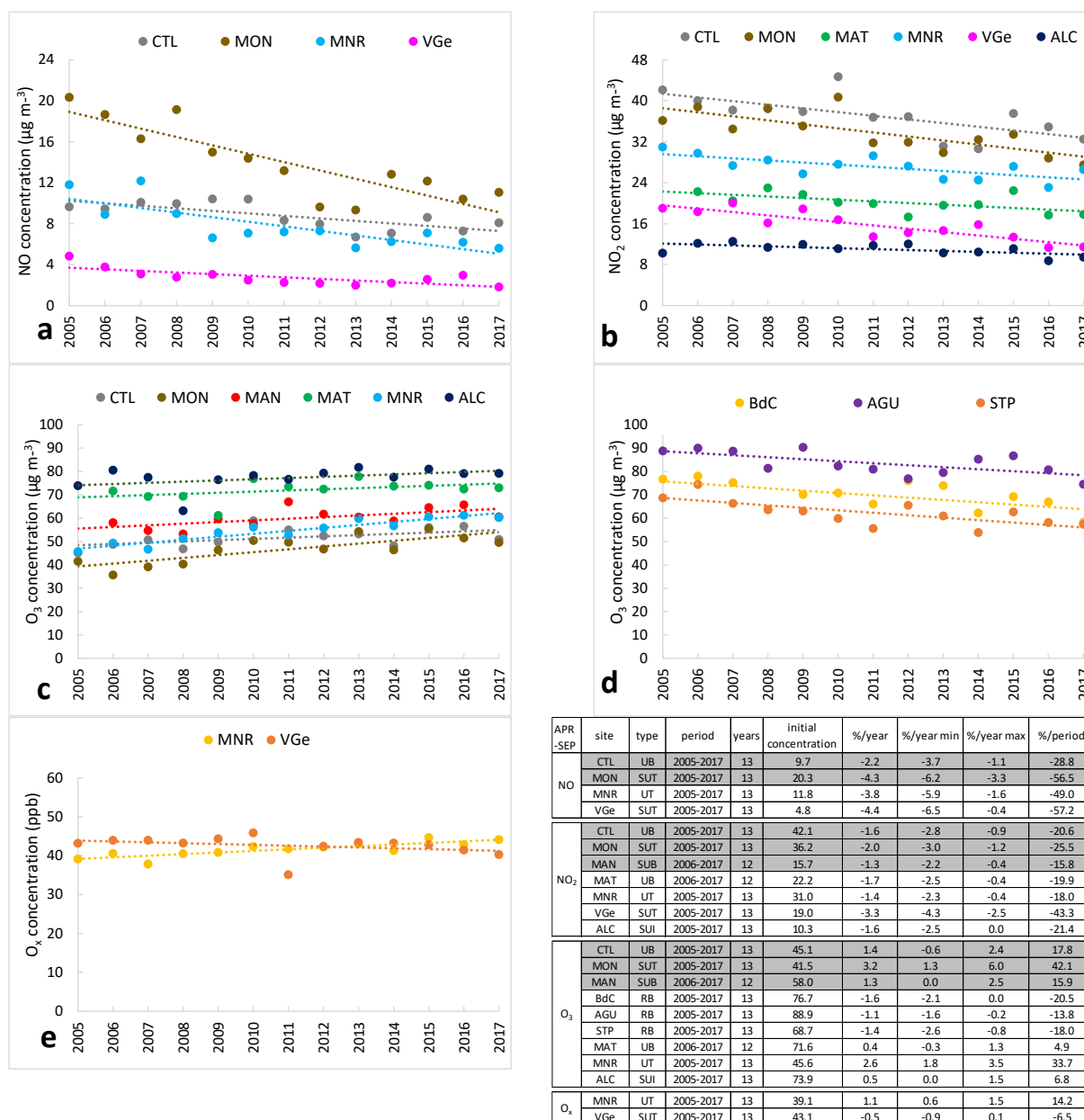


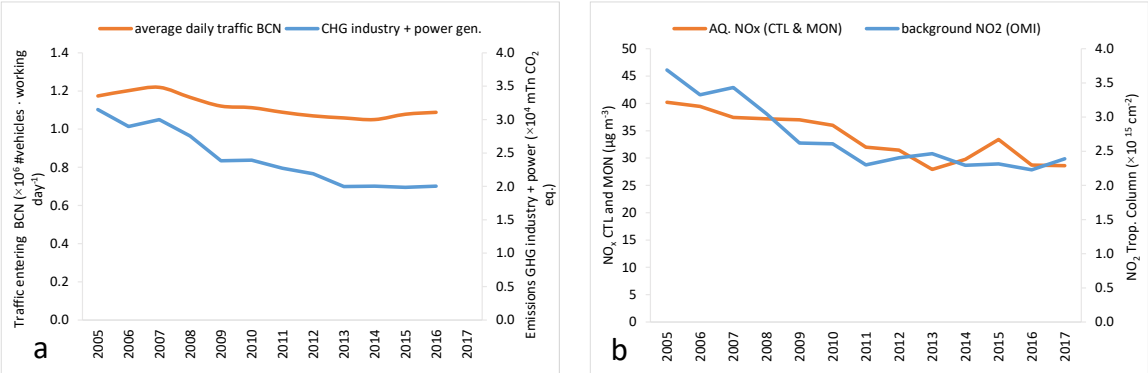
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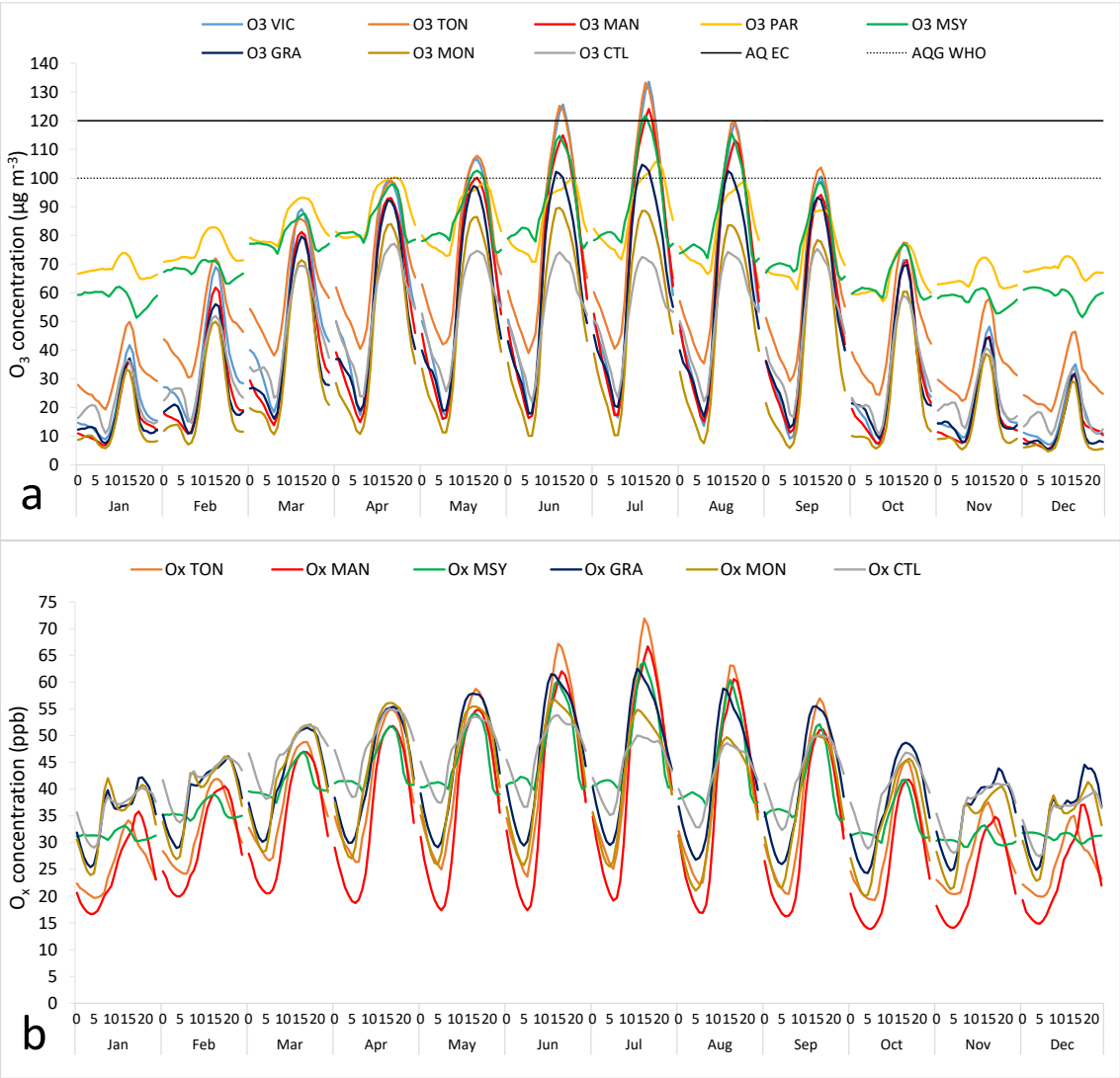
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948 Figure 4



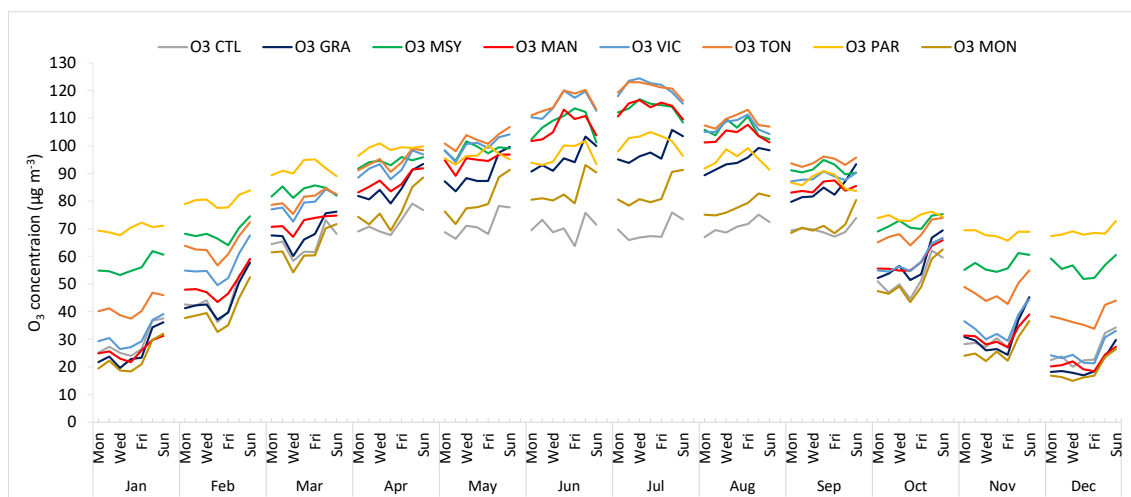
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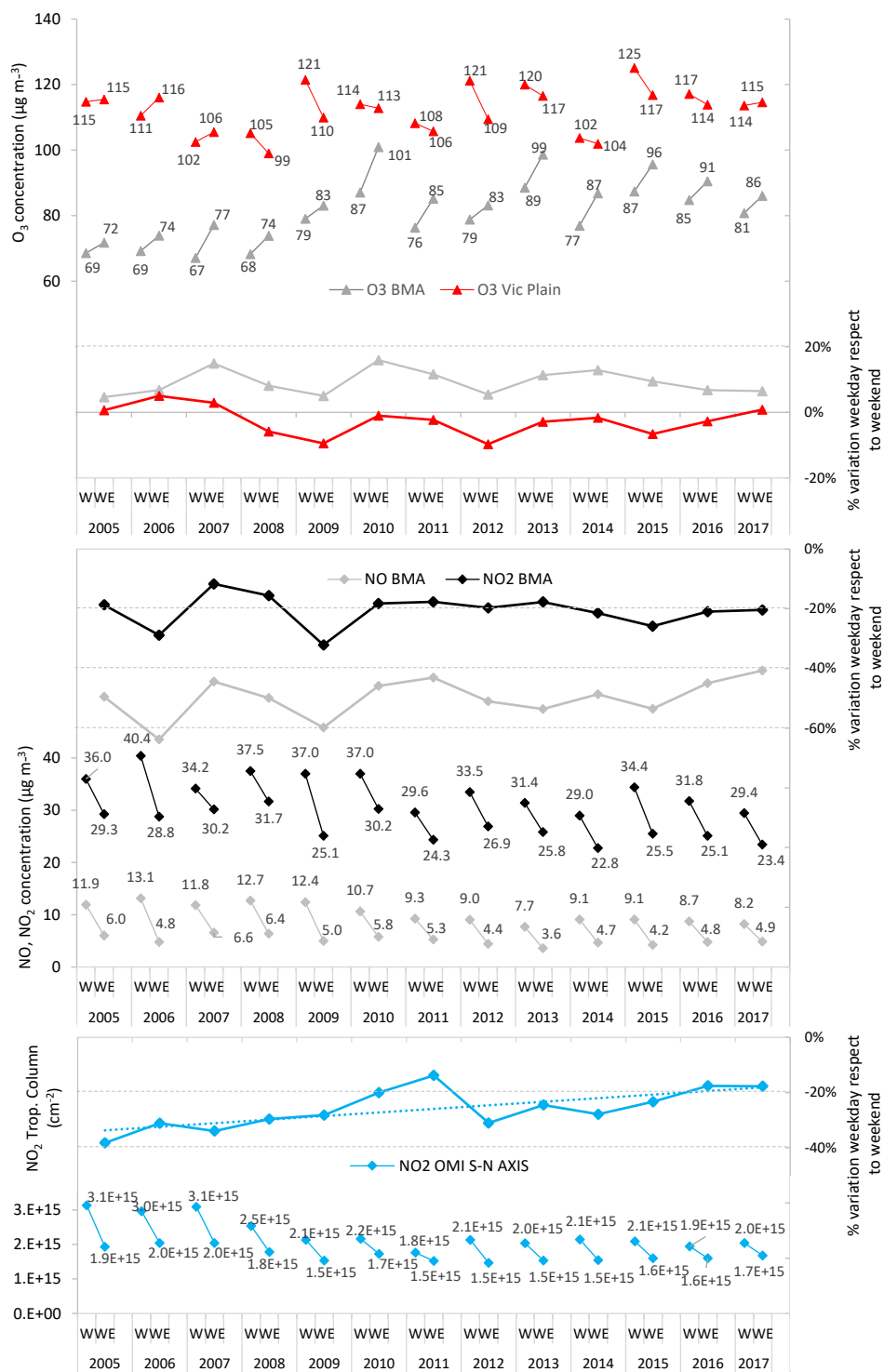
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952 Figure 6



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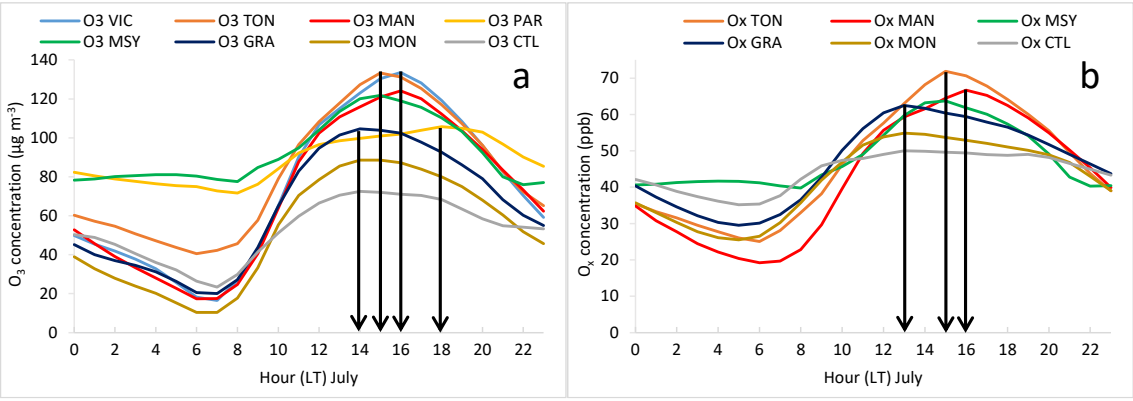
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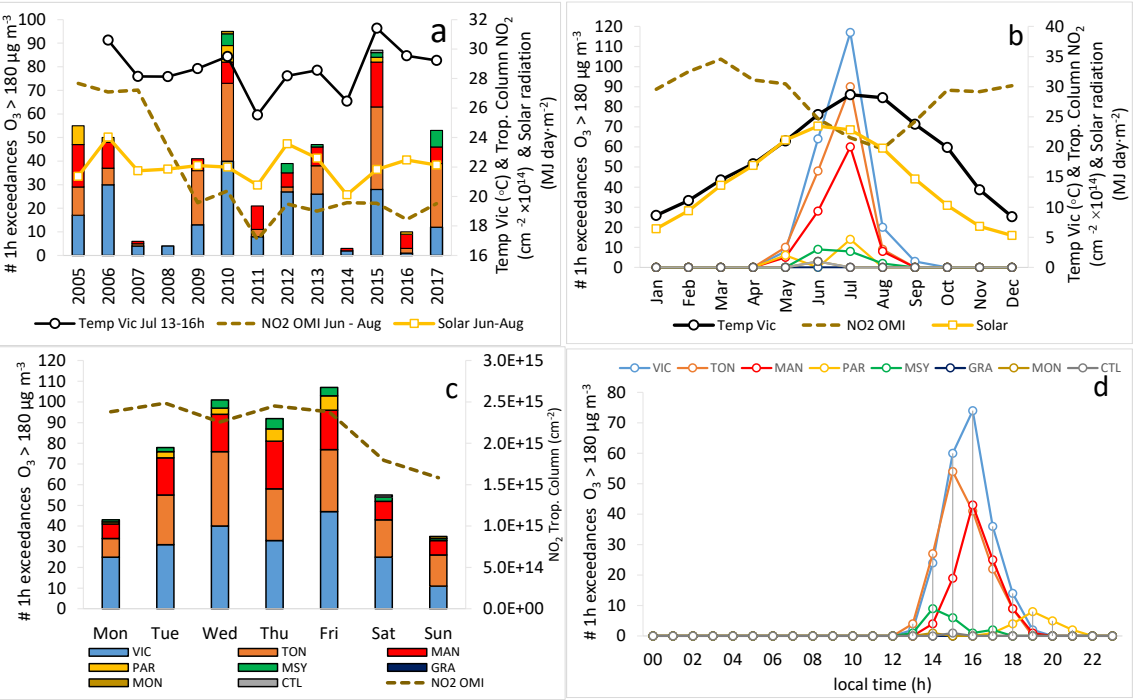
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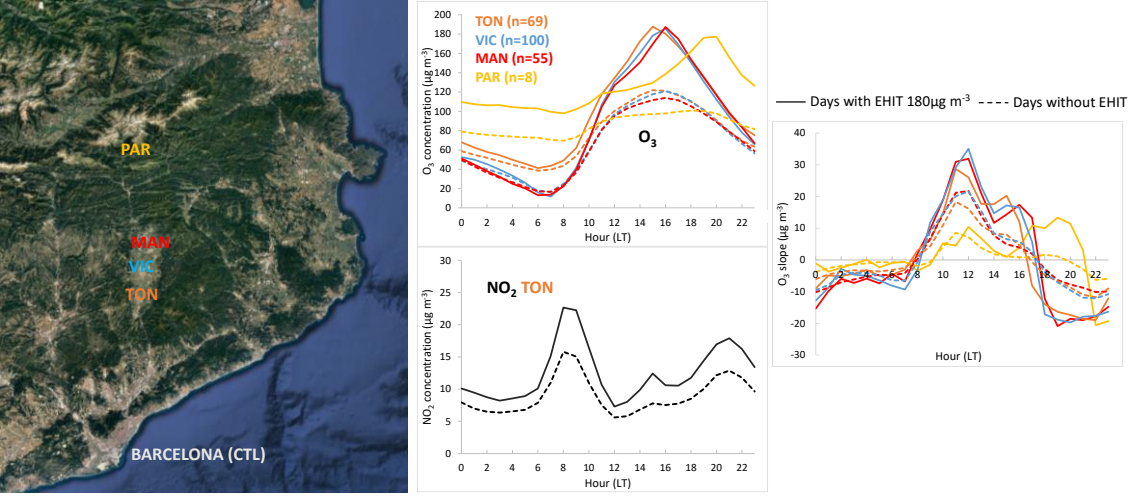
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958 Figure 9



959 Figure 10



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961 Figure 11

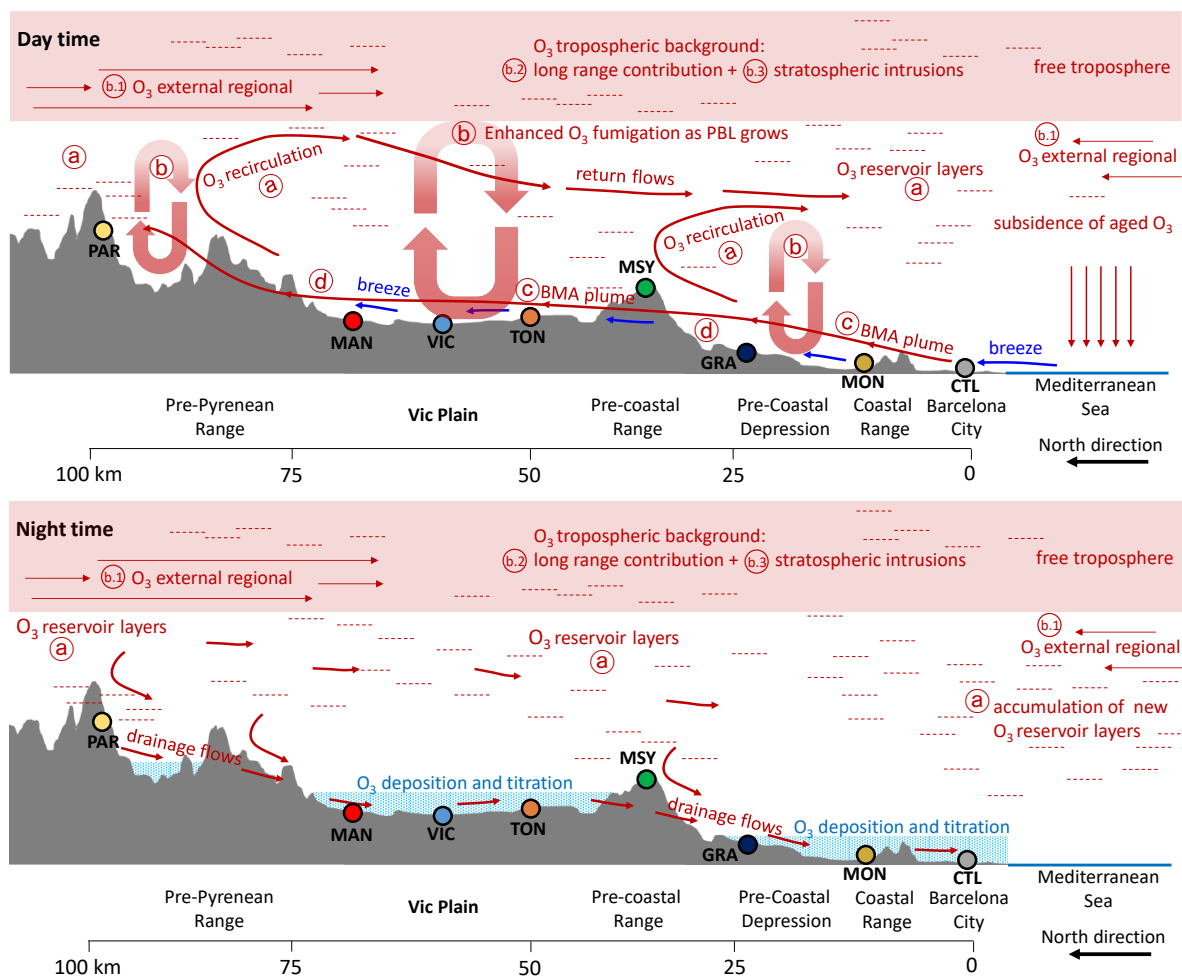


Figure 12

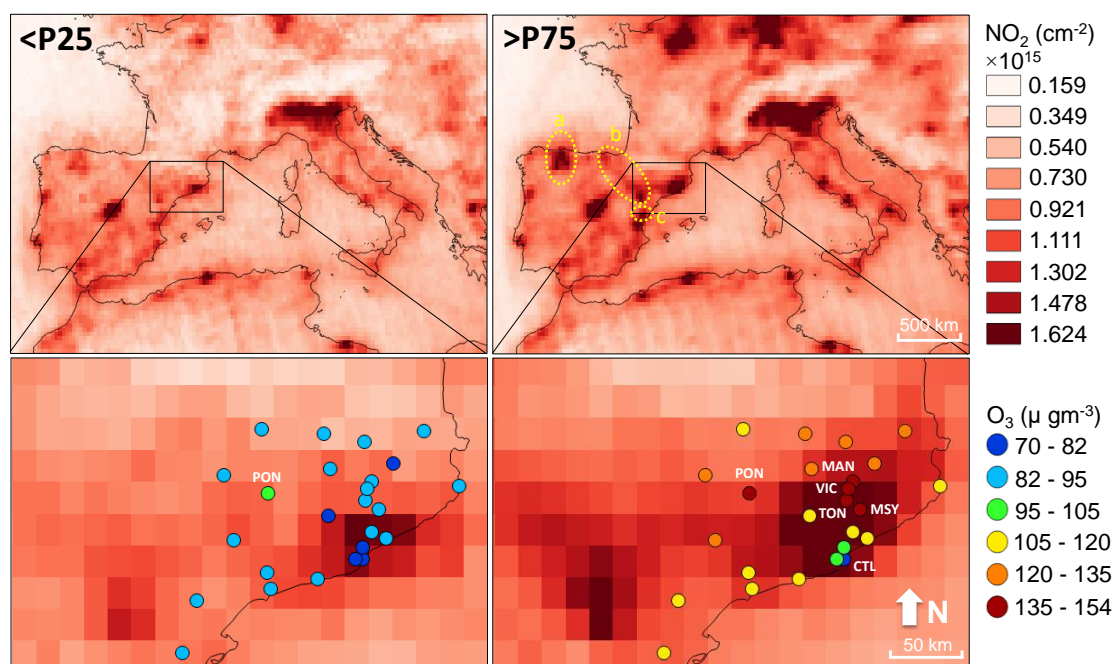
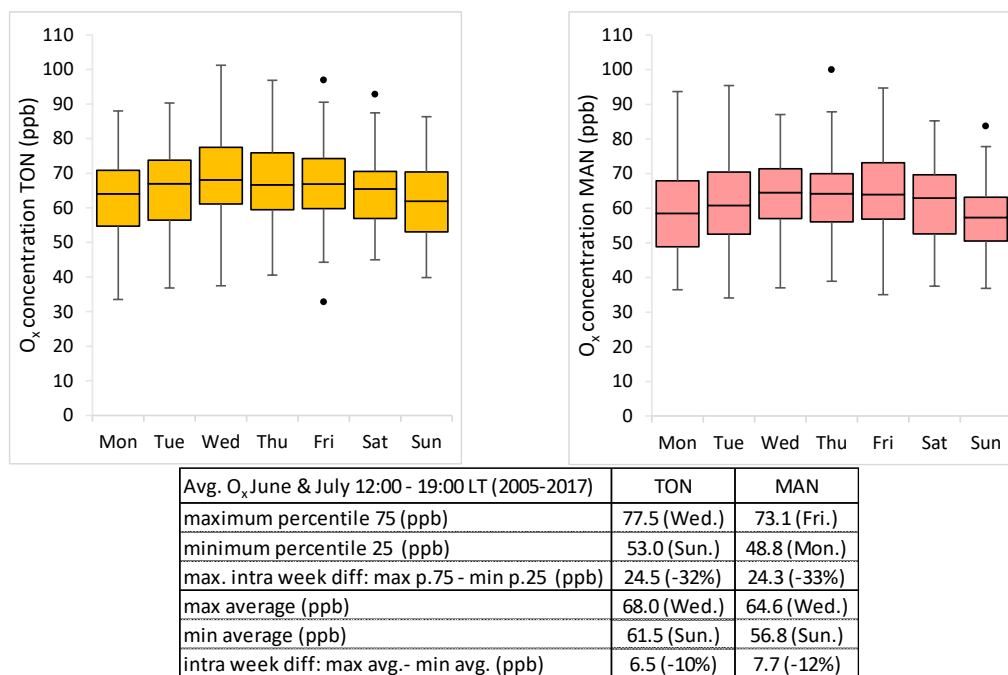


Figure 13



967 Figure 14

968