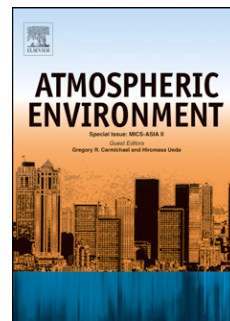


Accepted Manuscript

Title: Megacities as hot spots of air pollution in the East Mediterranean

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PII: S1352-2310(10)01016-2

DOI: [10.1016/j.atmosenv.2010.11.048](https://doi.org/10.1016/j.atmosenv.2010.11.048)

Reference: AEA 10063

To appear in: *Atmospheric Environment*

Received Date: 4 July 2010

Revised Date: 17 November 2010

Accepted Date: 30 November 2010

Please cite this article as: Kanakidou, M., Mihalopoulos, N., Kindap, T., Im, U., Vrekoussis, M., Gerasopoulos, E., Dermitzaki, E., Unal, A., Koçak, M., Markakis, K., Melas, D., Kouvarakis, G., Youssef, A.F., Richter, A., Hatzianastassiou, N., Hilboll, A., Ebojie, F., Wittrock, F., von Savigny, C., Burrows, J.P., Ladstaetter-Weissenmayer, A., Moubasher, H. Megacities as hot spots of air pollution in the East Mediterranean, *Atmospheric Environment* (2010), doi: 10.1016/j.atmosenv.2010.11.048

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1 Megacities as hot spots of air pollution in the East

2 Mediterranean

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29 **Abstract**

30 This paper provides a comprehensive overview of the actual knowledge on the
31 atmospheric pollution sources, transport, transformation and levels in the East
32 Mediterranean. It focuses both on the background atmosphere and on the similarities
33 and differences between the urban areas that exhibited important urbanization the past

34 years: the two megacities Istanbul, Cairo and the Athens extended area. Ground based
35 observations are combined with satellite data and atmospheric modeling. The overall
36 evaluation pointed out that long and regional range transport of natural and
37 anthropogenic pollution sources have about similar importance with local sources for
38 the background air pollution levels in the area.

39

40 *Keywords:* megacities, East Mediterranean, air pollution, transport, anthropogenic impact

41 **1. Introduction**

42 The increasing need of humans for facilities, security, health care and
43 employment have been the driving forces for increasing urbanization that gave birth
44 to the Megacities, urban agglomerations with more than 10 million of inhabitants
45 (<http://www.worldclimate.com>). This increasing urbanization not only affected the
46 neighboring landscape, air quality, regional climate and ecosystems in the megacities
47 but also downwind of these regions. During the last decades, the Mediterranean,
48 following the general trend, has experienced a rapid growth in urbanization, vehicle
49 use and industrialization as being reflected in pollutant emissions to the atmosphere.

50 The Eastern basin of the Mediterranean and the surrounding regions, include
51 two megacities: the Greater Cairo area (GCA) (>15 million, Egypt) at the south edge
52 of the basin and the Greater Istanbul Area (GIA) (>12 million inhabitants, Turkey) at
53 the North East edge, as well as several large urban centers like to its northern part the
54 Greater Athens area (GAA) (>4 million) in Greece (Table 1, Figures 1 and 2a) that
55 exhibited important urbanization the past years. The region covers rural (inland Greek
56 and Anatolian peninsulas), maritime (Crete and Cyprus islands) and desert (Anatolian
57 plateau, north Africa, Middle East) sites.

58 The Mediterranean located at the boundary between the tropical and mid-
59 latitudes, is subject to large (about 50%) changes in the total O₃ column (Ladstaetter-

60 Weissenmayer *et al.*, 2007), which have been attributed to changes in the location of
61 the sub-tropical front (Hudson *et al.*, 2003). It is also a crossroad of air masses coming
62 from Europe, Asia and Africa, where anthropogenic emissions, mainly from Europe,
63 Balkans and the Black Sea, meet with natural emissions from Saharan dust (e.g.
64 Kallos *et al.*, 1993, Kanakidou *et al.*, 2007), vegetation (e.g. Liakakou *et al.*, 2009)
65 and the sea (e.g. Kouvarakis *et al.*, 2002), as well as from biomass burning (e.g. Balis
66 *et al.*, 2003), which present a strong seasonal pattern. The transport of anthropogenic
67 pollutants from America also exerts a significant influence in the free troposphere
68 (Lelieveld *et al.*, 2002).

69 The typical Mediterranean climate is characterized by hot, dry summers and mild,
70 rainy winters. Evaporation is especially high in its eastern half basin, greatly
71 exceeding precipitation and river runoff in this region. This causes the sea water level
72 to decrease and salinity to increase eastward (Demirov and Pinardi, 2002). As a
73 consequence of its unique location and emissions, the Mediterranean is a climatically
74 sensitive region, often exposed to multiple stresses, such as a simultaneous water
75 shortage and air pollution exposure (IPCC, 2007) that is favored by the Mediterranean
76 climate and is likely to grow in the future due to the rapid urbanization.

77 Air pollution is one of the challenging environmental problems in the whole
78 East Mediterranean basin since both ozone and aerosol air quality limits are often
79 exceeded, in particular during summer. In contrast to Central and Northern Europe,
80 photochemical episodes can also occur during winter since at these latitudes solar
81 radiation is intensive year-around, driving photochemical reactions that favour air
82 pollution. The contribution of natural emissions to these exceedences seems
83 significant and remains to be determined. High ozone and aerosol concentrations are

84 harmful for human health and ecosystems, and they also cause agricultural crop loss
85 and climate change.

86 This paper summarizes the actual knowledge on the atmospheric pollution
87 sources, transport, transformation and levels in the Eastern Mediterranean. It first
88 outlines characteristics of the two megacities Istanbul and Cairo and the Athens
89 extended area, air transport patterns and meteorology. Then it discusses the
90 similarities and differences between these major pollution sources in the region and
91 compares them to the background atmosphere. Areas where further research is needed
92 to support mitigation strategy development are pointed out.

93 **2. The Megacities characteristics**

94 The studied urban areas are distributed over three continents: Europe, Asia and
95 Africa and present some common features as well as significant differences (Table 1).
96 Istanbul extends on two continents with the European part of the city being the oldest
97 one. It is separated from the Asian part by Bosphorus strait of 30-km length that
98 connects the Marmara Sea at the south with the Black Sea at the north.

99 The air circulation patterns at all three urban locations are affected by the
100 existence of hills: seven hills in GIA, the Mogattam hill to the east and the southeast
101 of GCA and the Parnes, Penteli and Hymettus mountains, all three over 1000 m,
102 surrounding mainly the North- and East boundaries of GAA. In Istanbul northeasterly
103 winds prevail during summer (Kindap, 2008) whereas southwesterly occur mainly
104 during winter (Koçak *et al.*, 2010). Istanbul is vulnerable to trans-boundary transport
105 of air pollutants from Europe, because of its location on the eastern end of the
106 continent in the zone of westerly synoptic air flow (Kindap *et al.*, 2006). Cairo
107 experiences two dominant wind sectors: the North sector and the South–West sector.
108 Although prevailing all year long, the north sector presents maximum occurrence

109 frequency in summer. The winter and spring seasons are significantly impacted by
110 south-western winds (Favez *et al.*, 2008a,b). Finally, in Athens, the prevailing wind
111 axis is north-east/ south-west and the ventilation takes place at northeasterly
112 directions (Melas *et al.*, 1995).

113 GIA and GAA are both subject to sea and land breeze local circulation
114 phenomena, favored during the weakening of the synoptic wind. During summer, the
115 southern part of GIA close to the Marmara Sea experiences such circulation patterns
116 that influence pollutants transport and accumulation in the boundary layer (Im *et al.*,
117 2006). The northern part of GIA is affected by the colder northern air masses and the
118 cooler Black Sea. In Athens sea/land breezes appear along the axis of the basin (NE to
119 SW) and anabatic/catabatic flows from the surrounding mountains. Under these
120 circumstances the ventilation of the basin is poor; the boundary layer is shallow and
121 the air pollution potential increases (Melas *et al.*, 1995 and references therein). The
122 sea breeze system from the Saronic Gulf, located to the south of GAA, sweeps
123 primary pollution from the city center, combined with O₃ titration, and favors
124 pollutant accumulation to the northern suburbs where significant episodes are
125 encountered. Air pollution episodes may occur in Athens during all seasons of the
126 year but most of these episodes are associated with the development of sea-breeze
127 (Kallos *et al.*, 1993).

128 2.1. Istanbul

129 The city of Istanbul (Table 1) is hosting almost 17% of Turkey's population.
130 Since the southern part of the GIA is the most urbanized, further growth will intensify
131 pressure on industrial and residential uses in the northern part of the metropolitan
132 region, where the natural protection areas and the watersheds are located (OECD,
133 2008). Average wind speed is highest in winter and lowest in summer with annual
134 average of about 2.7m/s. The humidity is high during all seasons (Ezber *et al.*, 2007).

135 The heating effect due to urbanization was found to produce two-cell structure during
136 summer, one on the European and one on the Asian side of the city. The cells extend
137 to about 600–800 m height in the atmosphere over the city and combine aloft (Ezber
138 *et al.*, 2007).

139

140 2.2. Cairo

141 Cairo (Al-Qāhirah), Egypt's capital (Table 1) situated south of the delta in the
142 Nile basin, is the largest rapidly expanding city in Egypt facing many environmental
143 problems. GCA's main populated area of about 200 km² is 4 km wide stretching 50
144 km along the banks of the Nile River. Outside GCA desert areas extend in the west
145 and east directions. Dust and sand storms frequently occur in spring and autumn
146 (Zakey and Omran, 1997). Hot desert cyclones known as the "Khamasin" depressions
147 pass over the desert during spring, always associated with strong hot and dry winds
148 often carrying dust and sand that increase particulate matter (PM) levels. During
149 winter the climate is generally cold, humid and rainy; while during the summer season
150 the predominant weather is hot and dry (Zakey *et al.*, 2008). The mean wintertime
151 wind is weaker than during summer, implying a lower ventilation of the area during
152 winter that could favor pollutant accumulation in the vicinity of the sources (Abu-
153 Allaban *et al.*, 2009). Robaa (2003) showed that rural and suburban parts of the city
154 have higher ventilation due to higher wind speeds than urban parts, which may lead to
155 higher pollutant levels in the urban regions of GCA. Cairo has a very poor dispersion
156 factor because of the advection patterns, its layout of tall buildings and narrow streets
157 and the lack of rain (Table 1). This results in a permanent haze over the city with PM
158 in the air reaching over three times the background levels.

159

160 2.3. Athens agglomeration

161 The GAA gathers about 40% of Greece's total population in a basin on the
162 west coast of the Attica peninsula. During the warmer part of the year, the mean wind
163 pattern in the atmospheric boundary layer is a persistent northeasterly flow of
164 relatively high constancy. GAA is also exposed to the summer monsoon circulation of
165 the Eastern Mediterranean. Etesians, a system of semi persistent summer northerly
166 winds, favor good ventilation of the basin prohibiting pollution episodes.

167

168 2.4. Outflow of pollution

169 Trajectories at approximately 700m height have been used to define air
170 pollution transport patterns from Istanbul, Cairo and Athens, in a regional scale. They
171 are based on 30-year (1961-1990) reanalysis data (NCEP/NCAR), available for every
172 six hours at a 2.5° resolution (Kindap *et al.*, 2009). The computed probability depends
173 on the grid size and increases with the trajectories length, with very small changes for
174 trajectories longer than 8 days (Kindap *et al.*, 2009). Figure 1 depicts the probability
175 of air masses originating from GIA, GCA and GAA to reach various locations in the
176 East Mediterranean, demonstrating the regional importance of air pollution from these
177 megacities. Istanbul pollution is exported mainly in the North East- South West
178 direction (Koçak *et al.*, 2010) whereas Cairo outflow is mainly affecting the south-
179 southwest locations and the Arabian Peninsula. Similarly, Athens plume is transported
180 mainly towards South East over the East Mediterranean Sea. These results are in good
181 agreement with the global modeling study by Lawrence *et al.* (2007).

182 3. Emission sources of air pollutants

183 All three cities experience heavy pollution from the transportation sector with
184 more than 2 million of cars in Athens and Istanbul and more than 1 million in Cairo,

185 of variable age and technical characteristics with the older ones in Cairo. A large
186 fraction of their country's industrial activities is also located in their vicinity.

187 The emissions inventories available for the entire East Mediterranean have
188 relatively coarse resolution (e.g. EMEP in 50 km resolution, Vestreng *et al.*, 2006,
189 and global inventories down to 1°x1° Granier *et al.*, 2005). The new EDGAR v4
190 inventory now becoming available, is making significant improvement increasing the
191 resolution to 0.1°x0.1° (<http://edgar.jrc.ec.europa.eu/>). However for large urban
192 agglomerations such as GIA, GAA and GCA higher resolution detailed emission
193 inventories would greatly improve our understanding of air pollution levels in the
194 area. Such inventories of anthropogenic sources have been developed by Markakis *et*
195 *al.* (2009; 2001a,b), in high spatial (2x2 km²) and temporal resolutions for the GIA
196 (reference year 2007) and for the GAA (reference year 2003), but appropriate
197 information is still missing for Cairo (Table 2). Weekend emissions are lower than
198 week days and diurnal profile fits with the rush hours due to the highest contribution
199 of traffic emissions (Markakis *et al.*, 2009). Application of the Markakis *et al.* (2009)
200 inventory has significantly improved the simulations of PM₁₀ levels (Im *et al.*, 2010)
201 in GIA.

202 Table 2 shows the annual sectoral distribution of pollutants. Industrial
203 activities are important sources of PM and responsible for almost 30 % of the SO₂
204 emissions. On-road traffic is the major contributor to CO, NO_x and non methane
205 volatile organic compounds (NMVOCs) in Istanbul and Athens. Residential
206 combustion and cargo shipping are significant pollution contributors in GIA and
207 GAA. Similar conclusions are reached for Istanbul by Koçak *et al.* (2010), based on
208 Positive Matrix Factorization (PMF) analysis of aerosol chemical characterization
209 observations (Theodosi *et al.*, 2010) from an urban background site in Istanbul.
210 Almost 20% of PM emissions in GAA originate from non-exhaust sources, including

211 tire, break wear and road abrasion. The central heating operations do not account for
212 more than a few percent in the annual totals (with the exception of SO_2 ~ 15%
213 contribution), but in the winter months they make a significant contribution.

214 Cairo shows different emissions fingerprint: Residential Combustion and
215 Industries being the major emitters of CO and NO_x whereas NMVOC emissions are
216 mostly from solvents use seconded by road transport. A significant portion of NO_x
217 (~50%) and SO_2 (~71%) originates from industrial activities. On-road traffic is also
218 an important source for CO (35%), NMVOC (37%) and $\text{PM}_{2.5}$ (36%). Anthropogenic
219 $\text{PM}_{2.5}$ in GCA originates mainly (54%) from residential combustion and open
220 burnings. Open fire burnings is a common practice and a major contributor to air
221 pollution in Egypt, as also seen on aerosol optical depth (AOD) seasonality derived
222 from satellite data with peaks in fall (Hatzianastassiou *et al.*, 2009).

223 To limit air pollution, measures were taken in all three urban centres around
224 1990-1995 with different level of implementation success.

225

226 3.1. Istanbul

227 Between 1980 and 1990 the consumption ratio of coal to fuel-oil increased
228 from 0.68 (in 1980) to 3.09 (in 1990; Tayanc, 2000). There has been the use of higher
229 quality coal and a shift from coal to natural gas for domestic heating purposes starting
230 from early 90s, leading to a decrease in the concentrations of primary pollutants such
231 as sulfur oxides (SO_x) and an increase in secondary pollutants such as secondary
232 aerosols and ozone (Tayanç, 2000). From the beginning of 1998 liquefied petroleum
233 gas (LPG) has been widely used in traffic. Low quality solid and liquid fuels with
234 high sulfur content, natural gas and LPG are the most commonly used fuel types in
235 the industrial activities that comprise 37% textile, 30% metal, 21% chemical, 5% food
236 and 7% other industries (Istanbul Chamber of Industry reports cited by Im *et al.*,

237 2006). Under these dense and various industrial activities, the region experiences very
238 complex air quality conditions.

239

240 3.2. Cairo

241 About 52% of the industries and 40% of the electricity production in Egypt are
242 located in the GCA (Nasralla, 2001). Cairo has many unregistered lead and copper
243 smelters which heavily pollute the city. GCA accommodates 50% of Egypt's road
244 transport fleet, 60% of which is over 10 years old, lacking modern emission cutting
245 features like catalytic converters (Mowafi and Atalla, 2005). The information
246 regarding the amounts of pollutants released in the atmosphere of Cairo is very
247 limited (El Mowafi and Atalla, 2005; Gurjar *et al.*, 2008; Table 2). Source
248 apportionment analysis based on simultaneous observations of several non methane
249 hydrocarbons (NMHC), including aromatics, and of aerosol components, including
250 metals, (Abu-Allaban *et al.*, 2002, 2007, 2009), pointed to mobile and industrial
251 emissions (lead smelting and LPG, considering that industrial processes may be
252 fueled by LPG) as the major source of NMHC during both summer and winter.

253 In 1995, the first environmental acts were introduced and the situation has
254 seen some improvement, with 36 air monitoring stations and emissions control on
255 cars. 20,000 buses have also been commissioned to the city to improve congestion
256 levels. In 2003, Egypt initiated an enforced vehicle emission-testing program in
257 Greater Cairo. The limits of CO, hydrocarbons and opacity for the vehicles have been
258 significantly reduced in 1995. However, the publicized information indicated an
259 overall failure rate of about 10% (El Mowafi and Atalla, 2005).

260

262 The massive number of registered vehicles in circulation, growing at a rate of
263 7% yearly, is allegedly the major cause of air pollution related problems in the area,
264 taking into account the large proportion of non-catalytic (0.8 million) or powered by
265 old technology diesel engines vehicles (0.2 million). Athens experiences very severe
266 congestion phenomena with the average speed not exceeding 12 km/h during rush
267 hours. Although the use of natural gas for domestic heating purposes has increased
268 lately, combustion of fuel oil is still primarily used for central heating. The large
269 industrial complexes are located in the Thriassion plain, several kilometres to the west
270 of the GAA. They are separated from the Athens basin by mount Aigaleo (up to 450
271 m) that acts as a physical barrier preventing most of the exchange of air pollutants
272 between the industrialized area and the city (Melas *et al.*, 1998).

273 4. Air pollution in the East Mediterranean

274 Enhanced levels of pollution (Figure 2) and increasing trends over the last
275 decade are seen by satellites over East Mediterranean and over the Middle East and
276 Cairo (Lelieveld *et al.*, 2008; Vrekoussis *et al.*, 2009). Background tropospheric O₃
277 levels in the area are high, particularly in spring and summer, depending on the
278 meteorological conditions since they are controlled by large-scale, long-range
279 transport and photochemical formation (Gerasopoulos *et al.*, 2005). Background PM
280 levels are also high due to a significant contribution of Sahara dust aerosol (Querol *et al.*,
281 2009) but also transported pollution (Mihalopoulos *et al.*, 2007). In the urban
282 atmosphere due to the high levels of primary pollutants, like PM and NO_x, maintained
283 by the anthropogenic emissions, O₃ titration by reaction with NO is leading to very
284 low O₃ levels over city centers, whereas NO_x and PM remain high. Primary pollutants
285 decrease downwind where O₃ and secondary aerosols build up photochemically. In

286 the urban regions, the temporal variability of primary gaseous pollutants reflects the
287 high emissions during winter time and the faster photochemical destruction during
288 summer time. Figure 2b depicts the tropospheric NO₂ columns as observed by
289 SCIAMACHY (SCanning Imaging Absorption SpectroMeter for Atmospheric
290 CHartographY) for the period 2005-2006 and highlights the local pollution sources all
291 around the Mediterranean. SCIAMACHY observations of NO₂ tropospheric column
292 over the region (Figure 2b) indicate high tropospheric columns of NO₂ over urban
293 sites around the Mediterranean with those over both the GIA and the GCA increasing
294 over the last years (Vrekoussis *et al.*, 2009). This distribution nicely contrasts to the
295 O₃ distribution shown in Figure 2a, that presents the largest enhancement actually
296 over the water, covering the whole East Mediterranean basin, which acts as receptor
297 of the surrounding pollution. Total columns of CO over the region range between 1.5
298 and 3×10^{18} molecules.cm⁻² maximizing in late winter/early spring (high emissions)
299 and minimizing in late summer/early fall (high photochemical destruction) (MOPITT:
300 Measurements of Pollution in The Troposphere; [ftp://14ftl01.larc.nasa.gov/MOPITT/
301 MOP03M.003/](ftp://14ftl01.larc.nasa.gov/MOPITT/MOP03M.003/)).

302 Mean satellite observations of short lived trace gases (NO₂, CHOCHO, HCHO
303 and O₃) and AOD over the region during the recent years are summarized in Figure 3.
304 High tropospheric columns of NO₂, HCHO, CHOCHO are observed over urban
305 locations (GIA, GCA, GAA) and low levels over the background receptor site of
306 Finokalia. The progressive reduction of tropospheric columns of NO₂ from Istanbul to
307 Athens and then to Cairo can be noticed together with a similar trend in CHOCHO
308 and HCHO, used as proxy for NMVOC levels. Remarkably, CHOCHO peaks over
309 GCA pointing to a higher NMVOC/NO_x ratio than over GIA and GAA. This
310 indicates higher O₃ formation potential of NO_x in GCA due to high NMVOC
311 loadings, in agreement with ground-based observations (Abu-Allaban *et al.*, 2009).

312 The HCHO/CHOCHO ratio appears different over GCA than over GIA and GAA,
313 indicating a different NMVOC speciation in this region, most probably strongly
314 marked by biomass burning emissions. Tropospheric O₃ columns indicate the elevated
315 O₃ background towards the south that maximizes over the Finokalia receptor site.
316 However, they minimize over GCA that is closer to tropics and thus affected by a
317 much lower total O₃ column (~18 DU lower than over Finokalia, based on TOMS /
318 OMI 2005-2008 data in 0.25°x0.25° grid; <http://gdata2.sci.gsfc.nasa.gov>).

319

320 *4.1. Ozone and its precursors*

321 Table 3 recapitulates the available measurements of ozone in the Eastern
322 Mediterranean at urban and regional background locations. A clear North to South
323 increasing gradient is evident. In particular, surface O₃ increases when moving from
324 rural background sites of Istanbul to Athens and then to Cairo, indicating significant
325 contribution from long-range transport sources in air masses that age in the region.
326 Ozone measurements along the Aegean Sea (NE Mediterranean, Kourtidis *et al.*,
327 2002; Kouvarakis *et al.*, 2000) confirmed that transport from the European continent
328 is the main mechanism controlling ozone levels in the region, especially in summer
329 (or spring depending on the prevailing air transport patterns), when ozone presents a
330 maximum of about 60±10 ppbv (Gerasopoulos *et al.*, 2005).

331 Kalabokas *et al.* (2007) analyzing aircraft data found that during summer in
332 the middle troposphere of the eastern basin, O₃ was only 5–10% higher than over
333 Central Europe and high tropospheric ozone values were mainly confined in the low
334 troposphere. Gerasopoulos *et al.* (2006b) analyzing 7 years of surface O₃ observations
335 at Finokalia, found that the entrainment of O₃ rich air masses from the free
336 troposphere (4–6% of the observed ozone levels) maximizes during summer, when
337 the chemical production of O₃ is also enhanced by photochemistry and long-range

338 transport. This summertime high regional source term of O₃ is almost balanced by the
339 enhanced O₃ destruction via deposition and chemistry. Below a brief presentation of
340 the ozone measurements at the various cities is presented.

341

342 4.1.1. *Istanbul*

343 Im *et al.* (2008) reported O₃ observations at two different urban locations
344 within GIA located at both its European (8±7 ppbv) and the Asian (11±8 ppbv) parts
345 from 2001 to 2005. The highest ozone levels were observed during sunny and warm
346 summer days (maximum temperatures >25 °C) with southwesterly surface winds.
347 Recent observations of ozone levels in semi-urban and rural stations in the GIA
348 during the period 2007-2009 (Im *et al.*, 2009), provide insight to the background
349 levels of ozone in the extended area. They show higher ozone levels than the urban
350 stations, reaching 30-35 ppbv on average, for high ozone seasons.

351

352 4.1.2. *Cairo*

353 Ozone in the southwestern Cairo area has been observed to exhibit a seasonal
354 and diurnal cycle with levels reaching 70 ppbv in summer (Egyptian Environmental
355 Affairs Agency, <http://www.eeaa.gov.eg/eimp/news8.html>). Year-long, mean levels
356 often exceed the Egyptian and European Union air quality standards of 60 ppbv for
357 daytime (8-h) O₃ mixing ratios. Khoder (2009) reported a year (Dec 2004-Nov 2005)
358 of observations of ground level O₃, nitrogen dioxide (NO₂) and nitric oxide (NO)
359 concentrations at Giza in the GCA with daytime mean O₃ values of 91 ppbv during
360 summer (Table 3). Air masses reaching Cairo during summer originate from the
361 Aegean and the Cretan Seas. Thus, considering the Finokalia regional background
362 values (60 ppbv), the observed mean value of 91 ppbv in Cairo indicates that despite
363 O₃ titration from the local NO_x emissions, significant photochemical O₃ production

364 occurs. This is additionally supported by high VOC levels (Abu-Allaban *et al.*, 2009)
365 in the GCA, in agreement with the satellite observations shown in Figure 3. Maxima
366 in O₃ levels occur in summer due to local photochemical production and long range
367 transport whereas the highest levels of NO_x are found in winter. The diurnal cycles of
368 O₃ revealed a uni-modal mid-day peak year-around. The diurnal variations in NO_x
369 concentrations during the winter and summer showed two daily peaks linked to traffic
370 density.

371

372 4.1.3. Athens

373 Kalabokas *et al.* (1999a,b) analyses of 11-year observations from the Greek
374 Ministry of Environment air pollution network in Athens since 1987, show a
375 significant downward trend for almost all primary pollutants in all stations.
376 Comparison between the 3-year periods 1988-1990 and 1995-1997 gave the highest
377 reduction in the center of GAA of 52%, 34%, 26% and 20% for SO₂, CO, NO_x and
378 black smoke, respectively. The concentrations of the secondary gaseous pollutants
379 remained essentially at the same levels since 1990, even though different
380 characteristics (e.g. in ozone trends) may be observed for different site types
381 (Hatzianastassiou *et al.*, 2007). Observations of O₃ prior to 2000 (Kalabokas and
382 Repapis, 2004) at three stations in the GAA and the surroundings were found to
383 exhibit characteristic seasonal variation of rural ozone concentrations, with lowest
384 winter afternoon values at about 25 ppbv in December–January and average summer
385 afternoon values at about 60 ppbv in July–August. These values are comparable to
386 observations at Finokalia (Gerasopoulos *et al.*, 2005; 2006b) and indicate significant
387 contribution from long range transport sources rather than local photochemistry.

388 The increased regional background in Athens is also supported by the CO-
389 NO_x molar ratios in GAA (Figure 4, derived from Table 3) that are between 20 and

390 30, whereas in GIA are lower ranging from 9.8 (Sarachane) to 12.6 (Kadikoy) close to
391 those in Mexico City (11) and higher than for Tokyo (8.5) and US cities (6.7 in 2003)
392 (Parrish *et al.*, 2009). Both in GIA and GAA, CO-to-NO_x molar ratios are lower than
393 the mean ratio of 41 observed in Beijing that has been attributed to significant
394 regional contribution to CO levels in that megacity (Parrish *et al.*, 2009). Ratios
395 higher than 50 are derived from the observations by Elminir *et al.* (2005) for a GCA
396 residential site and point to the different CO sources characteristics (like older cars,
397 domestic combustion and open fires) in GCA than in the other megacities. These
398 ratios are however much lower than those of about 100 to more than 300 observed
399 during summertime at Finokalia where long range transport is the dominant source for
400 CO.

401

402 4.2. Airborne particulate matter

403 The Mediterranean is one of the areas with the highest AOD in the world, also
404 seen from space (Hatzianastassiou *et al.*, 2009), which presents high temporal
405 variability due to the short lifetime of PM in the troposphere (of the order of a week).
406 Two-year (2005-2006) mean observations of AOD at 443nm over the area from
407 MISR (Multiangle Imaging Spectro Radiometer) and of the aerosol small mode
408 fraction derived from MODIS (Moderate Resolution Imaging Spectroradiometer,
409 using the Giovanni daily data of NASA GES DISC), are depicted in Figures 2c and
410 2d. Although the annual mean AOD distribution is marked by the Sahara dust
411 contribution, relatively high levels of AODs are also seen over the Aegean and the
412 Black Sea. In addition, Figure 2d indicates the existence of significant fraction (about
413 0.5 to 0.6) of fine particles in the region that are commonly associated with pollution
414 sources. Synergistic analysis of MODIS AOD and aerosol index TOMS data, used as
415 proxy for absorbing dust aerosol, enabled a first evaluation of the local anthropogenic

416 contribution to the AOD over the GAA and GCA at 15-30% and 25-50%,
417 respectively, during summer (Hatzianastassiou *et al.*, 2009).

418 Ground- based observations over the area show high concentrations of aerosols, in
419 both PM₁₀ and PM_{2.5} fractions (Querol *et al.*, 2009), with PM_{2.5}/PM₁₀ ratios around
420 0.5 (Table 3), in agreement with the satellite observations in Figure 2d. In the Eastern
421 Mediterranean, PM₁₀ has a similar seasonal behavior as PM_{2.5}, with maxima in spring
422 and fall in the eastern basin due to African dust transport. This is also seen by lidar
423 (Papayannis *et al.*, 2008), sun photometer (Fotiadi *et al.*, 2006) networks and satellite
424 based-sensors (Papayannis *et al.*, 2005; Kalivitis *et al.*, 2007). PM₁ behaves
425 differently showing small maxima during summer and is mainly dominated by
426 pollution components (Gerasopoulos *et al.*, 2007; Koçak *et al.*, 2008).

427 In the background coarse mode aerosol (PM_{1.3-10}) dust and ionic components
428 contribute about 40% and 50%, respectively and organics about 10% (Koulouri *et al.*,
429 2008a). Mineral dust transport events are found to contribute about 8-12 $\mu\text{g m}^{-3}$ to the
430 background PM₁₀ annual mean levels in the East Mediterranean, whereas an
431 additional 5-10 $\mu\text{g m}^{-3}$ is attributed to transported anthropogenic regional sources and
432 sea-spray loads (Querol *et al.*, 2009). Re-suspension of dust is likewise a significant
433 and highly uncertain component of aerosols in the cities. Recent aerosol mass
434 spectrometer measurements of ultra fine aerosols on Crete Island during late spring
435 (Hildebrandt *et al.*, 2010), revealed highly oxidized background organic aerosol
436 throughout the campaign, regardless of the source region. These observations of aged
437 particles in air masses that circulated and were photochemically processed over the
438 extended region, support the role of the East Mediterranean basin as the ‘pressure-
439 cooker’ of transported air pollution. Compared to the colder Central and North
440 Europe, the high temperatures in the Mediterranean impose a low thermal stability of

441 ammonium nitrate in summer and favor the formation of nitric acid rather than
442 ammonium nitrate in the area (Querol *et al.*, 2009; Mihalopoulos *et al.*, 1997).

443 High sulfate background loadings in the East Mediterranean are mostly
444 attributed to the long-range transport of SO₂ (Zerefos *et al.*, 2000). In addition,
445 significant interactions exist in the Mediterranean between natural and anthropogenic
446 components in the atmosphere, both in the gas and aerosol phases. Observations and
447 modeling have shown that on a mean yearly basis, marine biogenic emissions
448 contribute up to 20% to the total sulphate production (Kouvarakis and Mihalopoulos,
449 2002). They also demonstrate that the reaction of dimethyl sulfide of marine origin
450 with nitrate radicals, which are mainly of anthropogenic origin, is responsible for
451 about 17% of the total HNO₃ production plus particulate nitrate formation
452 (Vrekoussis *et al.*, 2006). The deposition of these species is of great environmental
453 significance since it provides nutrients to the ocean. During summer in the eastern
454 Mediterranean, sulphate on fine particles is produced via gas phase reactions whereas
455 almost 90% of the supermicron nss-sulphate is formed via heterogeneous pathways,
456 coating natural aerosols (Mihalopoulos *et al.*, 2007).

457

458 4.2.1. Istanbul

459 Hourly PM₁₀ levels are monitored by the metropolitan Municipality of
460 Istanbul at the urban network stations of GIA since late 90's. GIA experiences high
461 and variable levels of PM₁₀ and PM_{2.5} particles (Table 3). Ozdemir *et al.* (2009)
462 reported average PM₁₀ levels of about 66 µg·m⁻³ observed at 10 Istanbul municipality
463 stations during the last 10 years with values ranging from 47 µg·m⁻³ to 115 µg·m⁻³.

464 A significant fraction of studied PM₁₀ episodes has been attributed to regional
465 transport of African dust and anthropogenic emissions. Kindap *et al.* (2006)
466 calculated that almost 50% of the wintertime PM₁₀ episodes in 2002 are associated

467 with air masses coming from Eastern Europe. Karaca and Camci (2010) attributed
468 about half of the studied high PM₁₀ levels in Istanbul in 2008 to distant source
469 contributions. On the other hand, Im et al. (2010) studied the effect of local emissions
470 on a 5-day PM episode in January 2008 using the high resolution emission inventory
471 of Markakis et al. (2009) and attributed 90% of the elevated PM₁₀ levels to local
472 anthropogenic emissions, combined with very low persisting vertical mixing. This is
473 in agreement with Koçak *et al.* (2010), who evaluated the contribution of the
474 anthropogenic sources to PM₁₀ levels at about 90%, in an independent analysis of the
475 same episode.

476 Recently, more than one year of aerosol observations at the background
477 Bõgaziçi University sampling station in Bosphorus strait coast, provided the first
478 complete chemical characterization measurements in GIA (Theodosi *et al.*, 2010).
479 They measured 9 different water-soluble ions, water soluble organic carbon (WSOC),
480 organic and elemental carbon (OC, EC) and several trace metals, between November
481 2007 and June 2009. Trace elements related to human activities obtained peak values
482 during winter due to domestic heating, whereas natural origin elements peaked during
483 the spring period due to dust transport from Northern Africa. During winter, OC was
484 found to be mostly primary and strongly linked to fuel oil combustion and traffic, as
485 EC. Both OC and EC concentrations increased during winter due to domestic heating.
486 The mean OC/EC ratio was about 2, lower than those in Athens and Finokalia, but
487 close to those observed in GCA (Table 3), indicating an overall dominance of primary
488 pollution. The higher WSOC to OC ratio observed during summer was mostly
489 attributed to the presence of secondary, oxidised and more soluble organics. Source
490 apportionment PMF analysis of these long term observations indicates that
491 approximately 80 % of the PM₁₀ in Istanbul is anthropogenic in origin (Koçak *et al.*,
492 2010). Secondary aerosols maximize during summer and are mainly due to long-range

493 transport sources that account for 20% of the PM₁₀ mass over the studied 1.5-years
494 period. Adding the contributions of crustal and sea salt (10.2 and 7.5 % of the
495 observed mass, respectively), regional sources can explain at least 38% of PM mass,
496 in line with the earlier mentioned studies.

497

498 4.2.2. Cairo

499 There have been a number of studies that evaluated the long-term surface
500 aerosol observations in Cairo (Abu-Allaban *et al.*, 2002; 2007) along with chemical
501 composition (Favez *et al.*, 2008a,b). These studies showed that the area is
502 characterized by elevated levels of surface PM, with annual averages around 100 µg
503 m⁻³ and above (Table 3). Favez *et al.* (2008a,b) reported more than 2 years (Jan.
504 2003- May 2006) of weekly observations of bulk aerosols at two GCA urban sites
505 (Table 3), along with their chemical characterization with respect to selected ionic
506 species and carbonaceous aerosols (sum of EC and OC). Dust aerosols displayed high
507 background levels (50 µg m⁻³) all year long, maximizing during the dust storm periods
508 (Favez *et al.*, 2008a). About 40% of Ca²⁺ on these dust aerosols was found to be
509 associated with ions of anthropogenic origin like SO₄⁼, NO₃⁻ and/or Cl⁻, pointing out
510 human driven processes that alter the chemical characteristics of dust and thus its
511 climatic impact on a regional scale. High concentration levels of non-sea-salt Cl⁻ (up
512 to 15 µg m⁻³ on a monthly basis), likely of industrial origin, were observed in autumn
513 and winter. During autumn, biomass burning aerosols originating from rice straw
514 burning in the Nile Delta, known as the “Black Cloud” event, have been estimated to
515 account for 12%, 35% and 50% of Cairo EC, water insoluble organic carbon (WIOC)
516 and WSOC mass concentrations, respectively.

517 Overall, non-dust aerosols were equally distributed between carbonaceous
518 aerosols and ions, and their concentrations were about 100 µg m⁻³ in autumn and

519 winter, and $60 \mu\text{g m}^{-3}$ in spring and summer. Remarkably, relatively low WSOC/OC
520 ratios (about 1/3) were obtained all the year-long. Favez *et al.* (2008b) further
521 investigated the carbonaceous content in the sub micron fraction of aerosols at an
522 urban site in GCA in spring 2005. They found well-marked diurnal patterns for the
523 WSOC/EC and WIOC/EC ratios, with minima during the traffic-influenced morning
524 period and maxima during the intense photochemical periods, suggesting significant
525 formation of both WSOC and WIOC during the afternoon. Applying the EC-tracer
526 method, they evaluated that freshly-formed secondary OC accounts for more than
527 50% of OC concentrations measured during the early afternoon period. This fresh
528 SOC was calculated to be mainly (~60%) composed of WIOC species. The latter
529 (unexpected) result has been tentatively attributed to low ambient relative humidity
530 and high anthropogenic volatile organic compounds in Cairo (Favez *et al.*, 2008b).

531

532 4.2.3. Athens

533 Grivas *et al.* (2008) analysed PM_{10} concentration data collected by the Greek air
534 quality monitoring network at 8 sites over the GAA, for the period of 2001-2004.
535 Daily concentrations, averaged over the whole study period, ranged between 32.3 and
536 $60.9 \mu\text{g m}^{-3}$ and the four-year average concentration of PM_{10} at five sites exceeded the
537 annual limit value of $40 \mu\text{g m}^{-3}$, while most of the sites surpassed the allowed
538 percentage of exceedances of the daily limit value ($50 \mu\text{g m}^{-3}$). The urban sites were
539 mainly affected by primary, combustion-related processes and especially vehicular
540 traffic, as deduced from the examination of the diurnal distribution of particulate
541 levels and by factor analysis. On the contrary, suburban background sites were subject
542 to particle transport from more polluted neighbouring areas and secondary particle
543 formation through gaseous precursors, both processes supported by favourable
544 meteorological conditions. The association of the PM_{10} levels with backward

545 trajectories indicated that a notable part of area-wide episodic events could be
546 attributed to trans-boundary transport of particles (Querol *et al.*, 2009b).

547 **5. Air pollution and impacts.**

548

549 *5.1. Climate*

550 In the Mediterranean, aerosols reduce the solar radiation absorption by the sea
551 by about 10%, alter the heating profile of the lower troposphere and exert a cooling
552 effect five times higher than the warming induced by the greenhouses gases
553 (Lelieveld *et al.*, 2002; Vrekoussis *et al.*, 2005). As a consequence, evaporation and
554 moisture transport, in particular towards North Africa and the Middle East, are
555 reduced. Satellite observation analysis (Rosenfeld, 2000) supported that aerosols
556 caused important perturbations to cloud microstructure and convection, probably
557 decreasing precipitation. Querol *et al.* (2009) analysis of available aerosol data in the
558 Mediterranean pointed out three very important climate relevant features of the
559 aerosols in the area: the increasing gradient of dust from the west towards the east; the
560 change of hygroscopic behavior of mineral aerosols (dust) via nitration and
561 sulphation; and the abundance of highly hygroscopic aerosols during high insolation
562 (low cloud formation) periods. Radiative forcing by aerosols also influences the
563 energy budget of the Mediterranean and the Black Sea, however the consequences of
564 this are still poorly understood. A changing energy budget and anomalous winds are
565 expected to influence the ocean circulation (Tragou and Lascaratos, 2003). Therefore,
566 aerosols may affect several components of the eastern Mediterranean atmosphere-
567 ocean system including the regional water cycle. These aerosol-generated effects are
568 already substantial today, even though sulphate from Europe has actually decreased in
569 the past two decades (Smith *et al.*, 2010) through the abatement of acidification.
570

572 During summer the persistent northerly winds carry large pollution loads from
573 Europe that can deposit onto the Mediterranean sea, for instance, nitrate and
574 phosphorus containing aerosols, which affect the water quality and could contribute to
575 eutrophication (Kouvarakis *et al.*, 2001; Markaki *et al.*, 2003). In addition, O₃ levels
576 in the regions downwind pollution sources are also often exceeding phytotoxicity
577 levels (Kourtidis *et al.*, 2002).

578 Furthermore, ageing of aerosols, such as coating of dust by pollution
579 compounds (Falkovich *et al.*, 2004) or chemical trapping of nitrogen on pollen
580 particles (Franze *et al.*, 2005), can be harmful for human health. Katsouyanni (1995)
581 points out that air pollution effects on health, partly determined by specific mixtures
582 of air pollutants, may be altered by other environmental, behavioural and social
583 patterns. She also points out that the health effects of the interactions between
584 pollutants and photochemical oxidants can be enhanced in the Mediterranean under
585 high temperatures and humidity patterns. She stresses that even if the health effects of
586 air pollution only slightly increase the risk to an individual, they are likely to be
587 important for public health because of the ubiquitous exposure of the population.

588 El Mowafi and Atalla (2005) cited that approximately 3% of the GCA
589 population is chronically exposed to PM₁₀ levels above 100 $\mu\text{g}\cdot\text{m}^{-3}$, compared to
590 48% exposed to 100-50 $\mu\text{g}\cdot\text{m}^{-3}$ and 49 % exposed to 50-5 $\mu\text{g}\cdot\text{m}^{-3}$ PM₁₀. Based on
591 ambient air pollutant concentrations Gurjar *et al.* (2008) have classified Cairo as a
592 megacity with extremely poor air quality, where measures for air pollution reduction
593 need to be taken urgently. It is estimated that 10,000 to 25,000 people a year in Cairo
594 die due to air pollution-related diseases. These findings indicate the significant
595 benefits that could be achieved by implementing the proper abatement measures to
596 improve air quality in Cairo.

598 Significant effort is recently paid on understanding atmospheric composition
599 change in the East Mediterranean due to human activities, supporting the role of the
600 basin as the ‘pressure-cooker’ of transported air pollution from distant anthropogenic
601 sources but also from surrounding urban centres. Air masses are mixed and aged in
602 the area under favourable meteorological conditions with high solar radiation.
603 Background O₃ observations show an increasing gradient towards the south that
604 partially compensates O₃ titration by NO_x in the urban sites. The increased regional
605 background contribution in Athens, Cairo and Finokalia compared to GIA are in line
606 with the observed CO/NO_x molar ratios. In GIA, CO/NO_x molar ratio is close to that
607 observed in Mexico City and Tokyo whereas in GCA is double or triple, indicating
608 significant regional contribution to CO levels. This ratio maximizes at the background
609 atmosphere ranging from about 100 to more than 300 observed during summertime at
610 Finokalia, where long range transport is the dominant source for CO. GCA
611 experiences also high levels of NMVOC that point to a high O₃ formation potential of
612 NO_x in this region. Satellite observations of HCHO and CHOCHO seem to indicate
613 different NMVOC speciation and sources over GCA than over GIA and GAA. Due to
614 the non linear dependence of O₃ on NO_x and NMVOC levels, control of NO_x
615 emissions is expected to lead to higher O₃ levels and thus O₃ exceedences in the cities.
616 Available information on NMVOC total amounts, reactivity and chemical speciation
617 is scarce, although the NMVOC/NO_x ratio and VOC reactivity is critical for the
618 build-up of air pollution. CO observations in rural areas are also limited, despite the
619 key role of CO in O₃ production. There is a clear need of such reliable and systematic
620 measurements of NMVOC, NO_x and CO in the region to support modelling of air
621 pollution and climate impacts.

622 PM, even in the urban regions, is also shown to have a significant contribution
623 by long range transport of African dust or distance anthropogenic pollution sources
624 over the region. Data analysis has shown that a significant number of PM
625 exceedences, registered in Istanbul and Athens as long range transport episodes, are
626 associated with regional pollution or natural dust transport. PMF analysis of ground
627 based aerosol chemistry observations indicates that local anthropogenic sources
628 account for about 60% of PM levels in GIA and an additional 20% of PM levels is
629 associated with transported anthropogenic pollution. Based on satellite derived AOD,
630 the local anthropogenic emissions in GAA and GCA have been estimated to
631 contribute by 15-30% and 25-50% to the total AOD, respectively. These estimates
632 need to be reconciled with ground based observations. On an annual mean basis, in
633 the East Mediterranean the background PM_{10} contains about $8-12 \mu g m^{-3}$ of
634 transported mineral dust and an additional $5-10 \mu g m^{-3}$ is attributed to transported
635 anthropogenic regional sources and to sea-spray loads. Dust transport increases
636 towards the east of the basin and dust aerosols are coated by pollution components
637 that modify their climate relevant properties. The climatic impact of this mixture
638 remains to be determined. The first limited number of available PM_1 data show that
639 their composition and variability is tightly linked to the anthropogenic sources in the
640 area. OC/EC observations help elucidating the ageing of pollution air masses and the
641 contribution of photochemistry versus primary sources. Further studies of PM_1 mass
642 and chemical characterisation will elucidate the sources and impact of PM pollution in
643 the area.

644

645 **Acknowledgements**

646 This work has been supported by the EU FP7 project CITYZEN. Analyses and
647 visualizations of TES, MISR and MODIS data used in this study were produced with
648 the Giovanni online data system, developed and maintained by the NASA Goddard

649 Earth Sciences Data and Information Services Center (GES DISC). We thank Drs. J.
 650 van Aardenne and U. Doering for the emission inventory developed for the
 651 Mediterranean in the frame of the EU FP6 project CIRCE. We also thank the
 652 reviewers for their useful comments. Scientific discussions within the IGAC SSC and
 653 the iCACGP are acknowledged.

654

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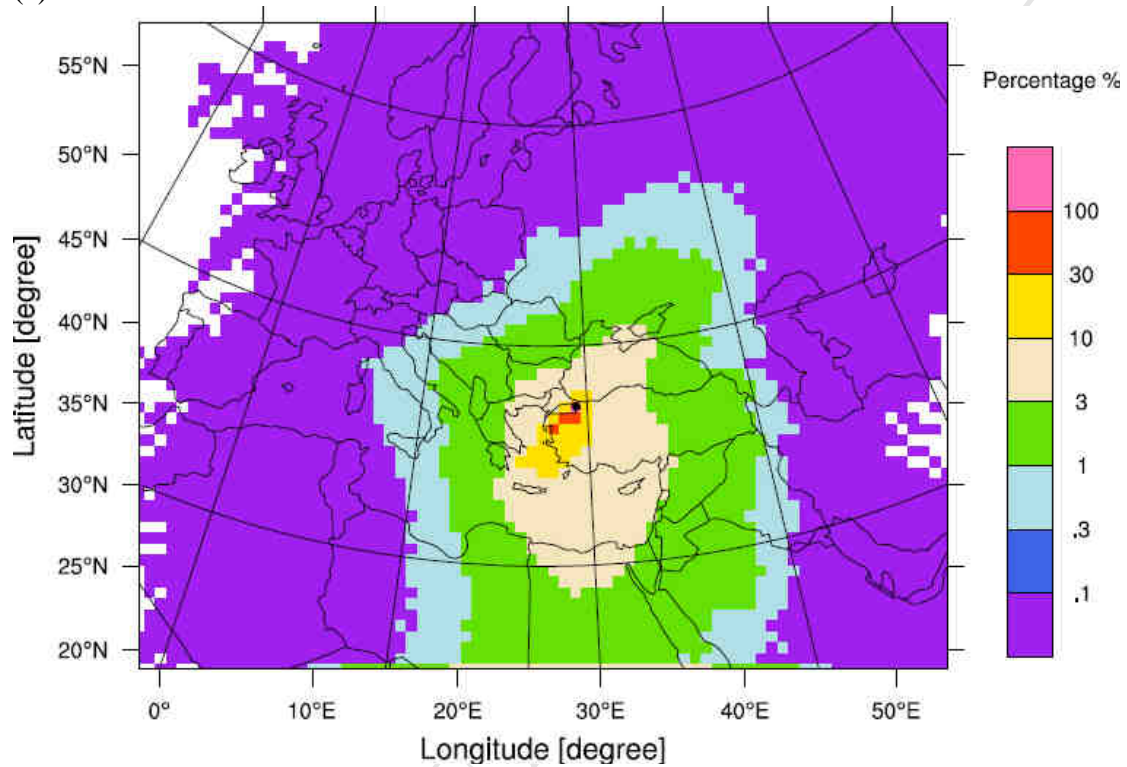
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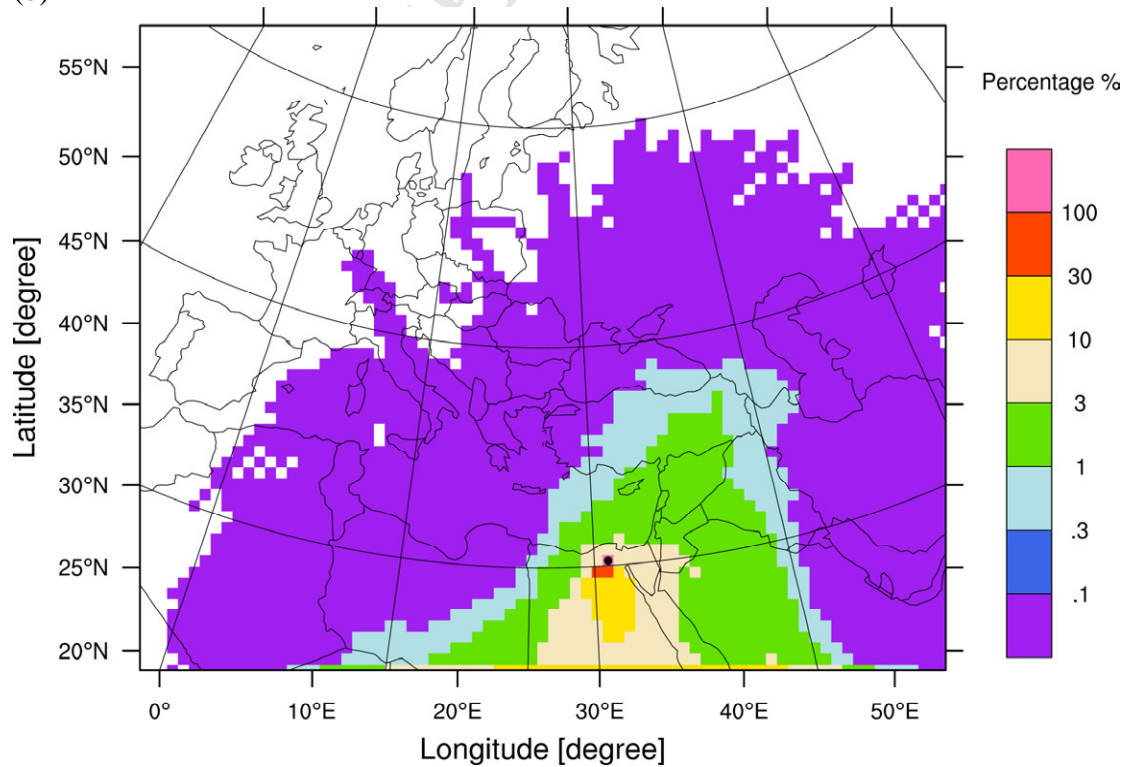
942 **Figures**

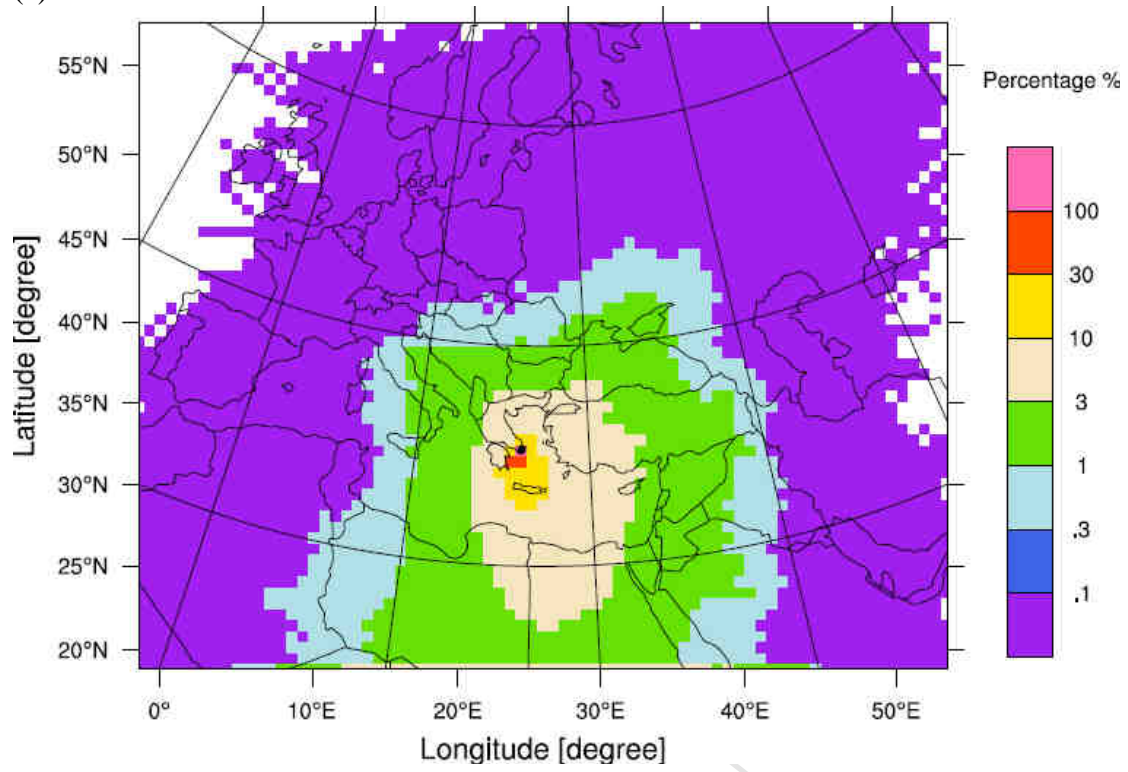
943 Figure 1: Map for the probability of arrival of trajectories starting from (a) Istanbul,
944 (b) Cairo, (c) Athens, over the 30 years period based on NCEP 6-hourly
945 meteorological data at 2.5° resolution, see text. Dot points indicate the city of
946 Istanbul, Cairo and Athens respectively.

947 (a)

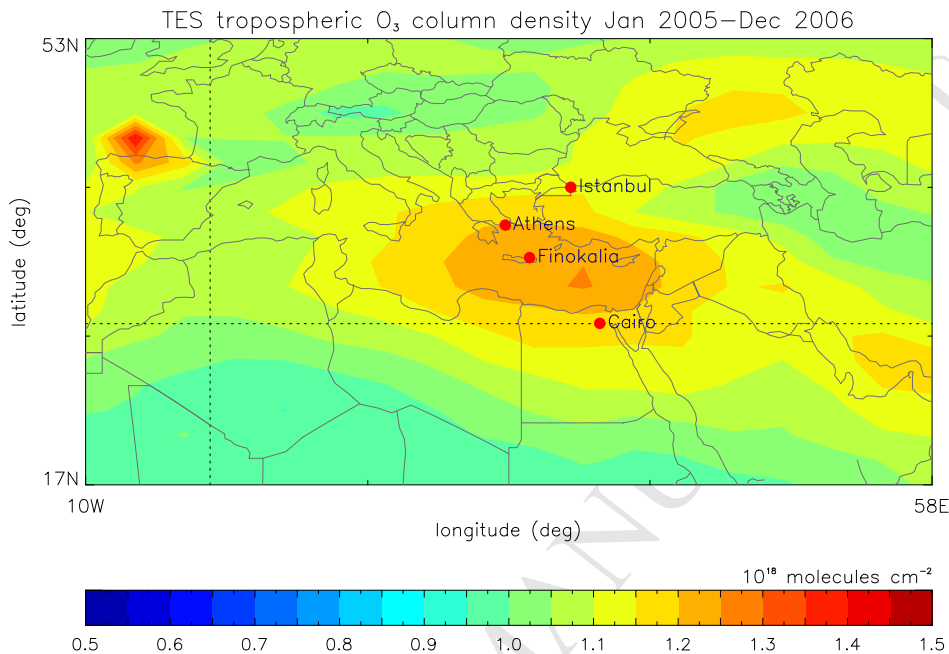
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(b)

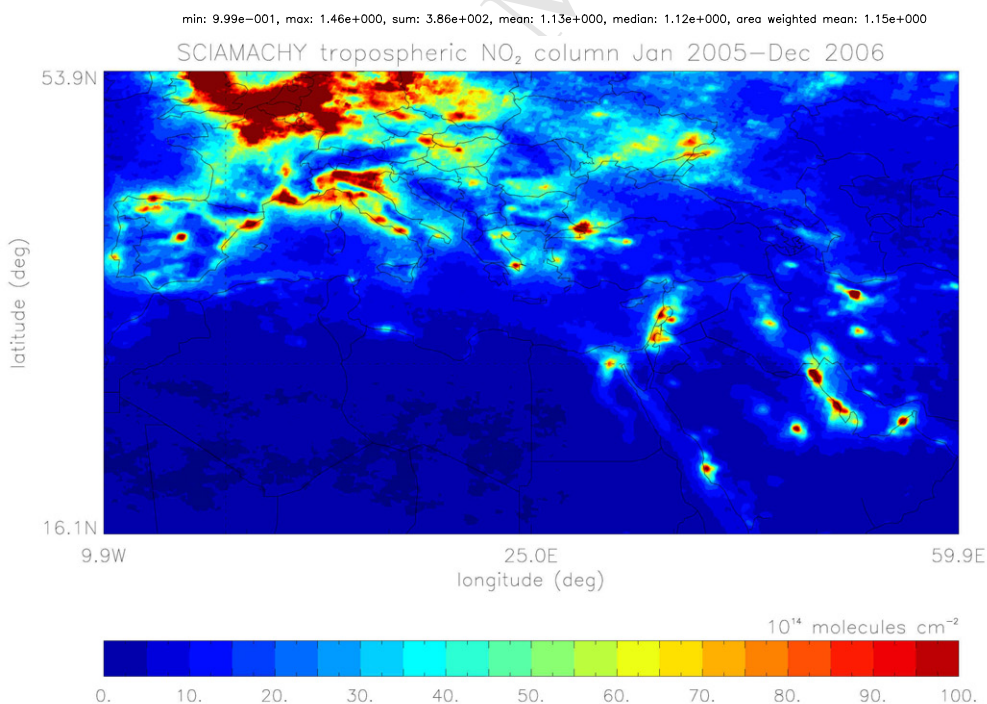
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955 Figure 2: (a) Tropospheric O₃ column as deduced from TES (Tropospheric Emission
 956 Spectrometer) satellite sensor gridded in 2°x4° lat x lon – The locations of Istanbul,
 957 Athens, Cairo and Finokalia are indicated; (b): Tropospheric NO₂ column from
 958 SCIAMACHY; (c) MISR aerosol optical thickness (AOT) at 443 nm in 0.5°x0.5° and
 959 (c) MODIS aerosol small mode fraction in 1°x1° resolution. Mean columns for the
 960 years 2005-2006. (a, c, d) have been derived from daily data using the Giovanni
 961 visualization tool of NASA (Acker and Leptouck, 2007).



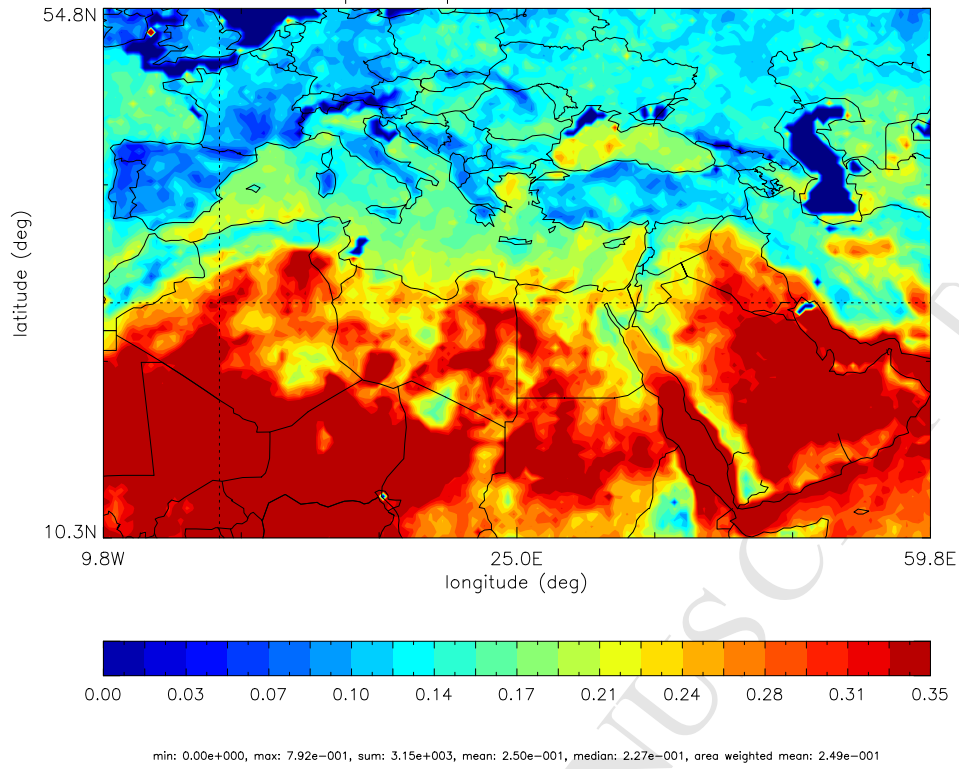
962 (a)



963 (b)

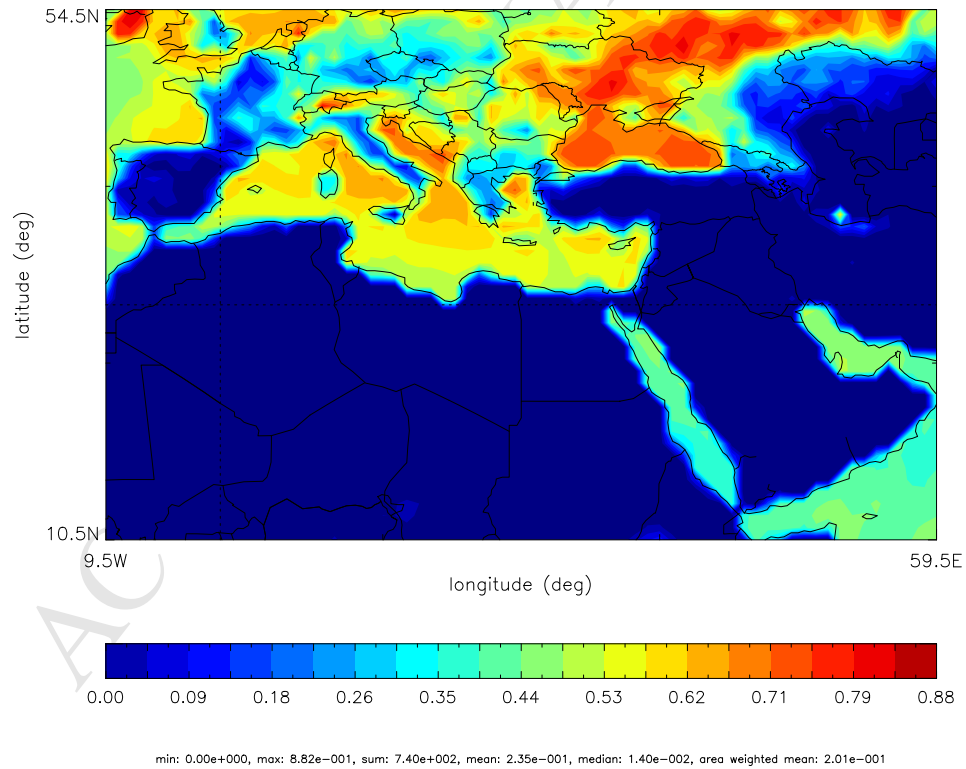
min: *****, max: 3.33e+002, sum: 2.78e+006, mean: 1.63e+001, median: 8.33e+000, area weighted mean: 1.44e+001

MISR aerosol optical depth at 443nm Jan 2005–Dec 2006



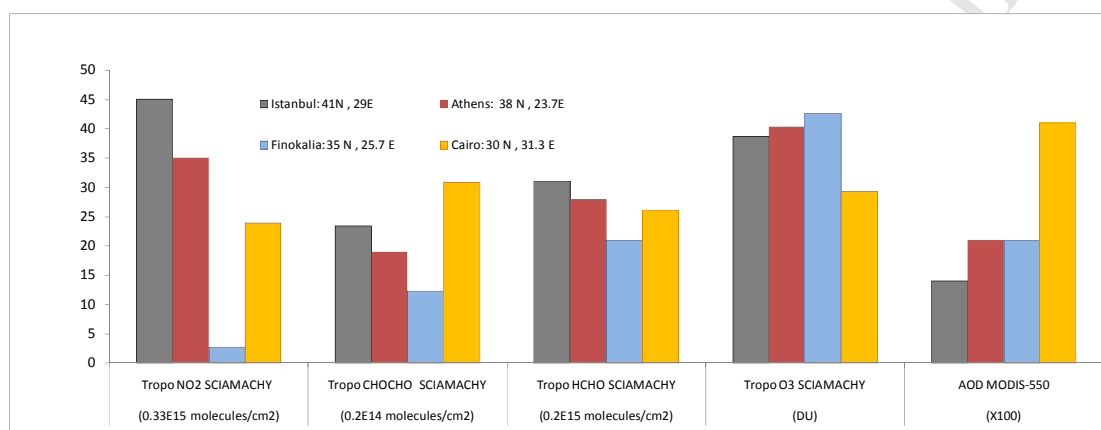
964 (c)

Fraction of fine aerosols MODIS Jan 2005–Dec 2006



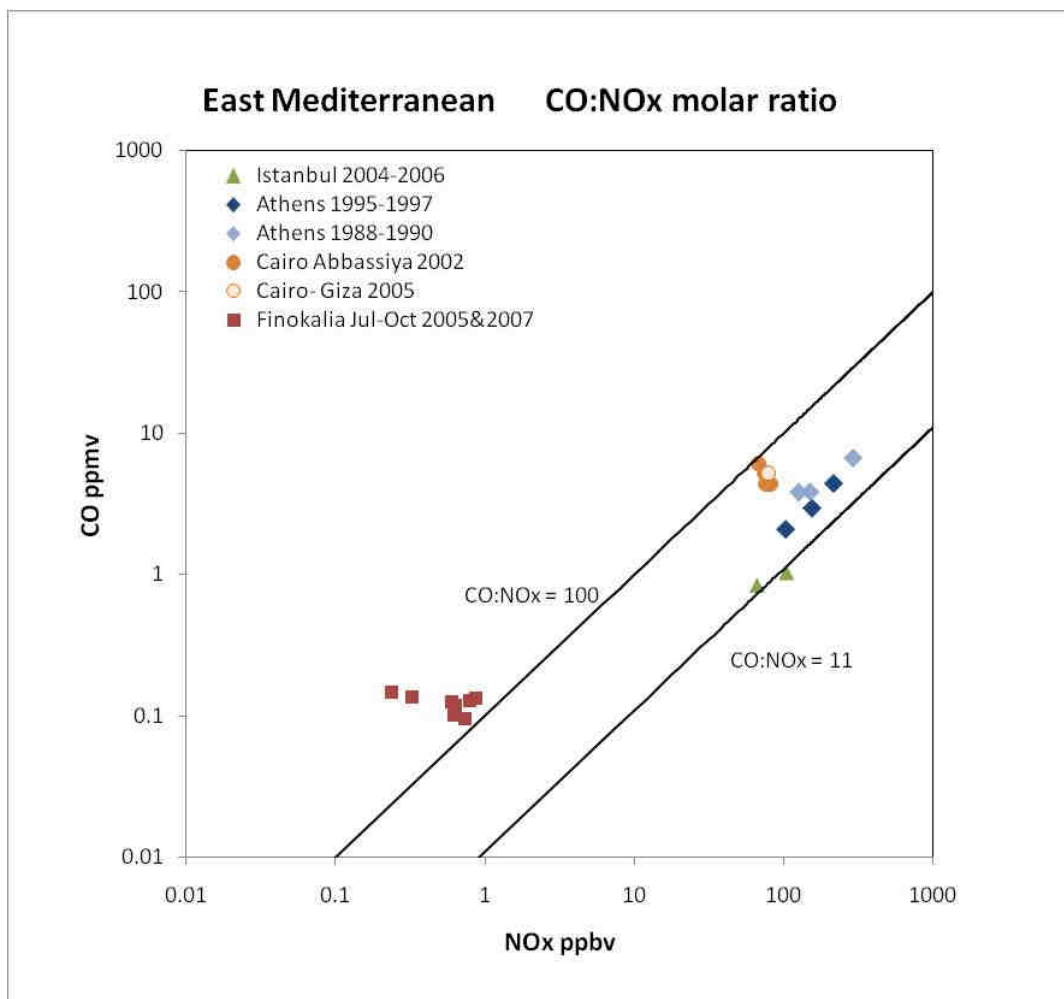
965 (d)

966 **Figure 3.** Satellite observations of air pollutants over GIA, GCA, GAA and Finokalia
 967 in the East Mediterranean. Mean over the period 2003-2009 from SCIAMACHY:
 968 Tropospheric columns of NO₂ in 10¹⁵ molecules cm⁻² and CHOCHO and HCHO in
 969 10¹⁴ molecules cm⁻² (multiplied by 3, 5 and 5 respectively) in a grid of 0.25°x0.25°
 970 covering the city. Mean tropospheric column of O₃ as deduced from SCIAMACHY
 971 (2003-2009) based on limb-nadir-matching and mean AOD at 550 nm from MODIS
 972 (2000-2008) in 1°x1° grid (multiplied by 100).
 973



974

975 **Figure 4.** Relationship between mean observations of CO (ppmv) and NO_x (ppbv)
976 levels in Istanbul, Athens, Cairo and Finokalia based on data reported in Table 3 and
977 references therein. Lines correspond to the CO:NO_x molar ratios of 11 and 100.



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979 Table 1 – Megacities and receptor location (Finokalia, Crete, Greece) characteristics (reference year 2009). General sources: Thomas
 980 Brinkhoff, 2009; <http://www.worldclimate.com/> (from data prior to 1990) * Extended region, in brackets highly populated area.

Characteristic	Istanbul	Cairo	Athens	Finokalia
Latitude, longitude	41.01°N, 28.97°E	30.03°N, 31.3°E	37.96°N, 23.71°E	35.33°N, 25.66°E
Continent	Europe-Asia	Africa	Europe	Europe
Surface (km ²)	6220	8815* (200)	3808* (450)	-
Population (Millions)	12.5	15.2	4.4	-
Ranked as Megacity	21 st	16 th	-	Background
population growth % over the last decade	45 29.6 (urban parts) 81 (rural parts)	16.4 (all Egypt) 18 (urban – Egypt)	6	-
Typical air temp. (°C)				2001-2009
winter average	8	15	10	11.6
summer average	28	28	26	24.2
Wind speed m/s	last 30 years	1995-2000	1984-2004 – Thissio	2001-2009
Annual mean	2.7	Urban; suburban; rural	3.3	5.8
Winter	3.0	2.2 3.7 3.1	3.4	5.8
summer	2.4	2.1 4.3 3.5	3.5	6.6
Mean precipitation (mm/yr)	800	25	400	350
Type of climate	Mediterranean (southern part) Cooler + wetter (northern part)	Sub tropical	Mediterranean Hot dry summer Wet mild winter	Mediterranean
Heat island (°C max surface temp. change)	1°	≤2.1°	exceeding 4° in 20% of studied cases	-
References	Ezber et al., 2007; Kindap, 2008	Zakey and Omran, 1997; Khoder, 2009; Zakey et al., 2008; Robaa, 2003	Kallos et al., 1993; Melas et al., 1995; Kassomenos & Katsoulis, 2006	Gerasopoulos et al., 2005 ; 2006; Vrekoussis et al., 2006

981 Table 2 – Anthropogenic emissions from Istanbul (reference 2007: Markakis *et al.*, 2009), Athens (reference year 2003; Markakis *et al.*,
 982 2010a,b) and Cairo (reference year 2005; van Aardenne *et al.*, 2009, Duering *et al.*, 2009; # Cairo inventory concerns PM2.5 emissions)
 983 greater areas.

	Residential Combustion %	Industry %	Fuel Extr./ Distribution %	Solvent Use %	Road Transport %	Off-road %	Maritime %	Waste %	Energy %	Total Ktons/yr
CO										
Istanbul	10.8	3.7	-	-	83.1	-	0.3	0.7	0.7	437
Athens	8.0	3.2	-	-	75.6	13.0	0.2	-	-	473
Cairo	28.8	31.2			35.5			2.2	2.4	285
NO _x										
Istanbul	2.1	2.4	-	-	79.4	2.8	9.5	-	3.2	305
Athens	3.1	22.4	-	-	51.0	17.8	3.1	-	2.6	78
Cairo	4.0	50.2			11.4	3.37		0.12	30.9	222
SO ₂										
Istanbul	14.7	23.2	2.3	-	2.3	4.1	17.6	-	35.6	91
Athens	14.9	29.1	8.4	-	3.2	7.2	11.3	-	25.9	31
Cairo	7.6	71.5			4.4					135
NMVOC										
Istanbul	2.6	0.5	-	29.8	44.8	0.4	0.6	20.4	0.2	77
Athens	3.2	2.1	2.0	13.8	70.6	5.7	0.5	-	2.1	93.2
Cairo	11	2.6		43.8	36.9			0.8		62.3
PM10 #										
Istanbul	7.1	64.9	0.1	-	17.4	3.9	3.1	1.7	1.8	61
Athens	18.0	62.7	-	-	13.0	0.8	1.9	3.6	-	21
Cairo #	53.4	4.3			35.9			4.4		6.4

984 Table 3 – Comparison of surface air pollution levels in Istanbul, Cairo, Athens and Finokalia -Crete (background site) in the East
 985 Mediterranean. PM₁₀ and PM_{2.5} are particles of diameter smaller than 10 and 2.5 microns, respectively.

Pollutant	Season/Date	Average	Location	Reference
O ₃ ppbv	1998-2008 2001-2005 2008-2009	<30 8 ± 7 11 ± 8 25.3 ± 16.8 19.9 ± 14.2	Istanbul * Saraçhane- Europe Kadikoy-Asia Buyukada Kandilli	Ozdemir et al. 2009 Im et al., 2009 Im et al., 2009 “
O ₃ ppbv	Winter 2005 Spring 2005 Summer 2005 Fall 2005 2002	Day / Dial 44 / 30 65 / 48 91 / 64 58 / 43 23.4	Cairo (Giza) Abbassiya	Khoder 2009 Elminir et al., 2005
O ₃ ppbv	1987–1996 winter (Dec.–Jan.) summer (Jul-Aug)	(12:00-18:00 LT) ~ 25 ~ 60	Athens	Kalabokas and Repapis, 2004
O ₃ ppbv	1997-2004 July-Aug. Dec	49 ± 11 58 ± 10 36 ± 7	Finokalia-Crete	Gerasopoulos et al., 2006b
NO ₂ ppbv	2001-2005	25± 18.9 (NO: 24± 46.3) 8.8±7.8 (NO: 2 ± 5.8)	Kadıköy Sarachane	Im et al., 2008

NO ₂ ppbv	Dec. 2004 - Nov. 2005 (hourly) Winter (hourly) Summer (hourly) 2002	60-150 80-200 (NO: 95-200) 60-130 (NO: 45-125) ~40	Cairo- Giza Cairo- Giza Cairo- Giza Abbassiva	Khoder, 2009 Khoder, 2009 Elminir et al., 2005
NO ₂ ppbv	1987-1997	57±5.3 (NO: 140.5±9.6) 18 ±4 (NO:31.9±18.0) 42.6±4.3 (NO:73.5±18.0)	Athens-Patission Maroussi Athinas	Kalabokas et al., 1999b
NO ₂ ppbv	June 2001 – Sept. 2003	0.35±0.31 (NO:0.033±0.020)	Finokalia-Crete	Vrekoussis et al., 2006
CO mg m ⁻³	2004-2006	1.181± 0.957 0.956±1.233	Sarachane Kadikoy	Im et al., 2008
CO mg m ⁻³	2002	~6 (4-10)	Cairo- Abbassiya	Elminir et al., 2005
CO mg m ⁻³	1987-1997	6.2± 1.2 1.9±0.6 3.8±0.5	Athens-Patission Maroussi Athinas	Kalabokas et al., 1999b
CO mg m ⁻³	July-Oct 2005 and Jul-Oct 2007	~ 0.143	Finokalia-Crete	Unpublished data
SO ₂ µg m ⁻³	1998-2008	~22 (7.5 - 45)	Istanbul *	Ozdemir et al., 2009
SO ₂ µg m ⁻³	Winter 1999-2000 Summer 2000	125±21.6 83±17.6	Cairo (Giza)	Khoder, 2002
SO ₂ µg m ⁻³	1995-1997	25±3 40±4	Athinas- Athens Patission-Athens	Kalabokas et al., 1999b
SO ₂ µg m ⁻³	1997-1999	2.7±0.9	Finokalia -Crete	Kouvarakis et al., 2002
PM ₁₀ µg m ⁻³	Jul 2002-Jul 2003 1998-2008 Nov 2007- Jun 2009	47.1 66 (47 – 115) 39.1	Istanbul * Background- Boğaziçi Univ.	Karaca et al., 2005 Ozdemir et al., 2009 Theodosi et al., 2010
PM ₁₀ (bulk aerosol)	2005: Win., Spr., Sum., Fall	215, 190, 115, 165	Cairo (Giza & El-Gomhoreya)	Favez et al., 2008

$\mu\text{g m}^{-3}$	2001-2002	170 \pm 25 140 \pm 40	Cairo (17 sites) Background -Cairo	Zakey et al., 2008 «
PM ₁₀ $\mu\text{g m}^{-3}$	Jun1999-May 2000	75.5 \pm 27.5	Athens	Chaloulakou et al., 2003
PM ₁₀ $\mu\text{g m}^{-3}$	2001-02 & 2004-05 2004-2006	28 \pm 30 32.5 \pm 27.7	Finokalia-Crete	Gerasopoulos et al, 2006a ; 2007; Koulouri et al., 2008
PM _{2.5} $\mu\text{g m}^{-3}$	Jul 2002-Jul 2003	20.8	Istanbul	Karaca et al., 2005
PM _{2.5} $\mu\text{g m}^{-3}$	2001-2002	85 \pm 12	Cairo (17 sites)	Zakey et al., 2008
PM _{2.5} $\mu\text{g m}^{-3}$	Jun1999-May 2000 2004-2006	40.2 \pm 16.7 23.7 \pm 10.7 29.3 \pm 10.4	Athens-Aristotelous Athens-Lykovrissi Athens-Goudi	Chaloulakou et al., 2003 Koulouri et al., 2008b “
PM _{2.5} $\mu\text{g m}^{-3}$	2004-2006	18.2 17.9 \pm 12.4	Finokalia-Crete	Gerasopoulos et al., 2007 Koulouri et al., 2008
OC/EC	Nov 2007- Jun 2009	1.98 (PM ₁₀)	Istanbul – urban Background	Theodosi et al., 2010
OC/EC	March- April 2005 2005	1.4 \pm 0.3 (morning) 2.9 \pm 0.5 (early afternoon) 2.5 - 5.0 Bulk aerosol	Cairo : El-Gomhoreya and Giza	Favez et al., 2008a « Favez et al., 2008b
OC/EC	June–July 2003	3.9 \pm 0.9 (PM _{2.5}) 24 \pm 17 (PM _{2.5-10})	Athens	Sillanpaa et al., 2006
OC/EC	July 2004-July 2006	4.0 (PM _{1.3}) 4.0 (PM _{1.3-10} non-dust cases)	Finokalia- Crete	Koulouri et al., 2008a

986 *10 municipality stations