

1 **Consistency evaluation of tropospheric ozone from ozonesonde and**
2 **IAGOS aircraft observations: vertical distribution, ozonesonde types**
3 **and station-airport distance**

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16 **Abstract:** The vertical distribution of tropospheric O₃ from ozonesondes is compared with that from
17 In-service Aircraft for a Global Observing System (IAGOS) measurements at 23 pairs of sites
18 between about 30°S and 55°N, from 1995 to 2021. Profiles of tropospheric O₃ from IAGOS aircraft
19 are in generally good agreement with ozonesonde observations, for Electrochemical concentration
20 cells (ECC), Brewer-Mast, and Carbon-Iodine sensors, with average biases of 2.58 ppb, -0.28 ppb,
21 and 0.67 ppb, and correlation coefficients (R) of 0.72, 0.82, and 0.66, respectively. Agreement
22 between the aircraft and Indian-sonde observations is poor, with an average bias of 15.32 ppb and
23 R of 0.44. The O₃ concentration observed by ECC sondes is on average higher by 5-10% than that
24 observed by IAGOS aircraft, and the relative bias increases modestly with altitude. For other sonde
25 types, there are some seasonal and altitude variations in the relative bias with respect to IAGOS
26 measurements, but these appear to be caused by local differences. The distance between station and
27 airport within 4° has little effect on the comparison results. For the ECC ozonesonde, the overall
28 bias with respect to IAGOS measurements varies from 5.7 to 9.8 ppb, when the station pairs are
29 grouped by station-airport distances of <1° (latitude and longitude), 1-2°, and 2-4°. Correlations for

30 these groups are $R = 0.8, 0.9$ and 0.7 . These comparison results provide important information for
31 merging ozonesonde and IAGOS measurement datasets. They can also be used to evaluate the
32 relative biases of the different sonde types in the troposphere, using the aircraft as a transfer standard.

33 **Key words:** WOUDC; IAGOS; tropospheric O_3 ; vertical distribution; ozonesonde; aircraft

34

35 **1 Introduction**

36 Ozone (O_3) is a trace gas with small concentrations in the atmosphere (Ramanathan et al., 1985). It
37 is an important greenhouse gas in the upper troposphere. In the planetary boundary layer, it is a
38 major air pollutant (Lefohn et al., 2018; Monks et al., 2015). It can endanger human health, damage
39 ecosystems, and affect climate change (Fu and Tai, 2015; Lefohn et al., 2018; Percy et al., 2003).
40 Therefore, it is of importance to study the temporal and spatial variations in tropospheric O_3
41 including near-surface O_3 and mechanisms affecting the variations (Logan, 1985; Ma et al., 2020;
42 Sharma et al., 2017; Young et al., 2018).

43 A large number of studies have been carried out on the spatiotemporal distribution, formation
44 mechanisms, and transport characteristics of tropospheric O_3 (Li et al., 2020, 2021; Vingarzan, 2004;
45 Wang et al., 2017, 2023; Xu et al., 2021; Yu et al., 2021). However, due to the limitation of
46 observations, there are many unknowns on tropospheric O_3 , especially the vertical distribution of
47 tropospheric O_3 . Satellites provide an effective platform for measuring O_3 globally. Satellite O_3
48 instruments, including TES, GOME, GOME-2, SCIAMACHY, OMI, and TROPOMI, have been in
49 operation for decades (David et al., 2013; Ebojie et al., 2016; Hegarty et al., 2009; Hoogen et al.,
50 1999; Hubert et al., 2021; Miles et al., 2015). Although satellite observations can provide detailed
51 temporally- and horizontally-resolved maps of tropospheric O_3 columns, in general satellite data
52 lack vertical resolution. While tropospheric differential absorption lidar can also provide vertical
53 distribution information for tropospheric O_3 (Keckhut et al., 2004; Yang et al., 2023), there are very
54 few routinely operating stations.

55 The principal sources of vertically-resolved, trend-quality observations of tropospheric O_3 are
56 therefore balloon-borne ozonesondes, and IAGOS aircraft observations. The World Ozone and
57 Ultraviolet Radiation Data Centre (WOUDC) and the In-service Aircraft for a Global Observing
58 System database (IAGOS) house the data from these two observation programs with the longest

59 duration and the most global stations, which are the most widely used for tropospheric O₃ studies
60 (Gaudel et al., 2020; Liao et al., 2021; Tarasick et al., 2019; Wang et al., 2022; Zang et al., 2024).
61 These two datasets are used to study the distribution, variability and trends of tropospheric O₃, and
62 its sources and transport, as well as satellite and model validation (Hu et al., 2017; Gaudel et al.,
63 2018; 2020; Wang et al., 2022; Zhang et al., 2008). The first phase of the Tropospheric Ozone
64 Assessment Report (TOAR-I), initiated in 2014, utilized available surface, ozonesonde, aircraft, and
65 satellite observations to assess tropospheric O₃ trends from 1970 to 2014 (Schultz et al., 2017). Hu
66 et al. (2017) found that the largest bias in a chemical transport model, GEOS-Chem, with respect to
67 ozonesondes and IAGOS observations, is in high northern latitudes in winter-spring, where the
68 simulated O₃ is 10-20 ppb lower. Wang et al. (2022) examined observed tropospheric O₃ trends,
69 their attributions, and radiative impacts from 1995 to 2017, using aircraft observations from IAGOS,
70 ozonesondes, and a multi-decadal GEOS-Chem chemical model simulation, and found increases in
71 tropospheric O₃ (950-250 hPa) of 2.7 ± 1.7 ppbv per decade from IAGOS observations in the
72 Northern Hemisphere and at 19 of 27 global ozonesonde sites averaging 1.9 ± 1.7 ppbv per decade.
73 There are also a number of comparative studies on these two datasets (Zbinden et al., 2013; Staufer
74 et al., 2013, 2014; Tanimoto et al., 2015; Tarasick et al., 2019). Staufer et al. (2013, 2014) used
75 trajectory calculations to match air parcels sampled by both sondes and aircraft. Zbinden et al. (2013)
76 compared coincidences (± 24 hours) at three site pairs, while Tanimoto et al. (2015) examined
77 simultaneous observations (± 3 hours for sonde versus aircraft) at several site pairs less than 100 km
78 apart. In general, these studies show small (6% or less) negative biases of aircraft measurements
79 against ECC sondes. Tarasick et al. (2019) compared trajectory-mapped averages over 20°-70° N
80 of ozonesonde and MOZAIC/IAGOS profiles and concluded that over 1994-2012 ozonesonde
81 measurements were about $5 \pm 1\%$ higher in the lower troposphere and $8 \pm 1\%$ higher in the upper
82 troposphere.

83 As shown above, the global O₃ vertical distribution datasets observed by WOUDC and IAGOS have
84 been widely used in various studies. Still, a long-term and multi-site systematic comparison of these
85 two datasets is rare, especially for the observations in the past three decades. In this study, we
86 attempt to make the most comprehensive evaluation to date of the relative biases of IAGOS and
87 sonde profiles, using as many station pairs as possible. We identify 23 suitable pairs of sites in the

88 WOUDC and IAGOS datasets from 1995 to 2021, compare the average vertical distribution of
89 tropospheric O₃ shown by ozonesonde and aircraft measurements, and analyze their differences by
90 ozonesonde type and by station-airport distance.

91

92 **2 Data and methods**

93 **2.1 MOZAIC-IAGOS observations**

94 The MOZAIC (Measurements of OZone and water vapor on Airbus In-service airCraft) program,
95 initiated in 1994 and incorporated into the IAGOS (In-service Aircraft for a Global Observing
96 System; www.iagos.org) program since 2011, takes advantage of commercial aircraft to provide
97 worldwide in-situ measurements of several trace gases (e.g., O₃ and CO) and meteorological
98 variables (e.g., water vapor) throughout the troposphere and the lower stratosphere (Marenco et al.,
99 1998; Petzold et al., 2015; Nédélec et al., 2015). O₃ measurements are performed using a dual-beam
100 UV-absorption monitor (time resolution of 4 seconds) with an instrumental uncertainty of ± 2
101 ppbv+2% (Thouret et al., 1998; Blot et al., 2021). It should be noted that this is only the instrumental
102 uncertainty, and does not include sampling uncertainties (possible losses) caused by the inlet line
103 and the compressor before the UV-photometric measurements are made. Loss of O₃ on the inlet
104 pump was an issue in earlier aircraft O₃ sampling programs (Brunner et al., 2001; Dias-Lalcaca et
105 al., 1998; Schnadt Poberaj et al., 2007; Thouret et al., 2022), but Thouret et al. (1998) found it
106 negligible for MOZAIC/IAGOS.

107 More details on the new IAGOS instrumentation can be found in Nédélec et al. (2015). The
108 continuity of the dataset between the MOZAIC and IAGOS programs has been demonstrated based
109 on their 2-year overlap (2011-2012) (Nédélec et al., 2015). Blot et al. (2021) evaluated the internal
110 consistency of the O₃ measurements since 1994, which confirmed the instrumental uncertainty of
111 ± 2 ppb. Moreover, they found no bias drift amongst the different instrument units (six O₃ IAGOS-
112 MOZAIC instruments, nine IAGOS-Core Package1 and the two instruments used in the IAGOS-
113 CARIBIC aircraft).

114 **2.2 WOUDC ozonesonde observations**

115 The World Ozone and Ultraviolet Radiation Data Centre (WOUDC) is part of the Global
116 Atmosphere Watch (GAW) program of the World Meteorological Organization

117 (<https://woudc.org/data/explore.php>). The WOUDC is operated by Environment and Climate
118 Change Canada. WOUDC ozonesonde data have been evaluated in a number of WMO-sponsored
119 international field intercomparisons (Attmannspacher and Dütsch, 1970, 1981; Kerr et al, 1994) and
120 more recently in laboratory simulation chamber experiments using a standard reference photometer
121 (Smit et al., 2007, 2024; Thompson et al., 2019). In the global ozonesonde network, while different
122 ozonesonde types were common in the past, more than 95% of current sounding stations use
123 electrochemical concentration cells (ECC). ECC ozonesondes have a precision of 3-5% ($1-\sigma$) while
124 the precision of other sonde types is somewhat poorer, at about 5–10% for Brewer-Mast and the
125 Japanese KC (Carbon-Iodine) sonde, and somewhat larger for the Indian-sonde (Kerr et al., 1994;
126 Smit et al., 2007). Biases with respect to UV reference spectrometers have been estimated for ECC
127 sondes at 1-5% in the troposphere (Smit et al., 2021; Tarasick et al., 2019, 2021).

128 **2.3 Data processing**

129 The two datasets were first screened for airport-sonde station pairs within a latitude separation of
130 $<4^\circ$ and a longitude separation of $<4^\circ$. Many sonde stations have observational records that do not
131 overlap with the IAGOS period (1994-present). In addition, the IAGOS dataset has large gaps at
132 many airports, because the frequency of visits to airports by aircraft that take part in IAGOS depends
133 on commercial airlines' operating constraints. In total, 23 station pairs (Fig. 1) were identified with
134 a separation of less than 4° in both latitude and longitude, and coincident observations over at least
135 nine months. The majority of the 23 ozonesonde site records are ECC (17), while four are Indian-
136 sonde, one Brewer-Mast, and one Carbon-Iodine (the Japanese KC sonde). These stations were
137 divided into 3 groups according to the distance (D) between the ozonesonde station and the airport:
138 $D < 1^\circ$, $1^\circ < D < 2^\circ$, and $2^\circ < D < 4^\circ$. Specific information on the comparison stations is shown in Table
139 1.

140 The observation times of the ozonesonde and aircraft are generally not the same. Ozonesondes are
141 typically launched once a week, although a few stations have more frequent launches. The aircraft
142 records generally contain more frequent observations, but observation times vary. For the selected
143 23 stations, we calculated the mean O_3 vertical profiles at 1km resolution (the first layer is from the
144 surface to 1 km above sea level) for each month during the observational period for the two datasets.
145 A minimum of four aircraft profiles were required to estimate a monthly mean profile; because

146 ozonesonde launches are typically only a few times per month, no minimum was required to
147 estimate a monthly mean profile. Only data with monthly means in both datasets were included for
148 further analysis. Comparisons between the two datasets were made by ozonesonde type and by
149 station-airport distance.

150

151 **3. Results and discussion**

152 **3.1 Comparison of the vertical profiles of tropospheric O₃ from four types of ozonesondes and** 153 **aircraft observations**

154 Previous intercomparisons of sondes launched on the same balloon (Attmannspacher and Dütsch,
155 1970, 1981; Beekmann et al, 1994, 1995; Deshler et al., 2008; Hilsenrath et al., 1986; Kerr et al,
156 1994; Smit et al., 2007) have shown that sondes of different types respond somewhat differently to
157 the same O₃ vertical profile; that is, they have relative biases, that vary with altitude. Fig. 2 therefore
158 compares the mean vertical profiles of tropospheric O₃ from ozonesonde and aircraft measurements,
159 separated by ozonesonde type. Both O₃ concentrations and absolute differences between
160 ozonesonde and aircraft increase with altitude, especially above 9 km. Average tropospheric O₃
161 profiles observed by ECC, Brewer-Mast, and Carbon-Iodine sondes are in good agreement with
162 aircraft measurements, with biases of 2.58 ppb, -0.28 ppb and 0.67 ppb, while the agreement with
163 the Indian-sonde is poorer, with a bias of 15.32 ppb. The Indian-sonde average also shows a linear
164 increase with altitude, while the aircraft measurements indicate an O₃ decrease with altitude above
165 8 km (Fig. 2b). This behavior is most clearly related to the comparisons of stations 2°-4° apart in
166 spring (Fig. S9).

167 These results are broadly consistent with those from JOSIE 1996 (Smit et al. 1996; Smit and Kley,
168 1998; Thompson et al., 2019), and with the northern hemisphere average result from Tarasick et al.
169 (2019). (Their Figure 20b; note that it is largely based on ECC sondes, and the scale is inverted
170 (IAGOS-sondes) from the sense we use here.)

171 Fig. 3 shows correlation plots of monthly mean O₃ at 1 km vertical intervals for months when both
172 IAGOS and ozonesonde data are available at the same location. While these monthly averages are
173 of data not necessarily coincident in time, Fig. 3 indicates that the data compare well on this
174 timescale, with correlation coefficients (R) of 0.71, 0.88 and 0.66, respectively (Fig. 3a-3c). The

175 agreement between the Indian-sonde and aircraft observations is poor, however, with an R of only
176 0.44 (Fig. 3d). The RMSE of O₃ observed with the four types of ozonesondes (ECC, Brewer-Mast,
177 Carbon-Iodine and Indian-sonde) and the aircraft is 15.99 ppb, 14.15 ppb, 16.26 ppb and 29.85 ppb,
178 respectively. After calculation, we obtained the slopes and offsets of ECC, Brewer-mast, Carbon-
179 iodine and Indian-sonde without forcing the fitted lines through zero, the slope is 0.71, 0.88, 0.56
180 and 0.74, respectively, and the offset is 18.94 ppb, 6.89ppb, 27.48ppb and 27.84ppb. When we force
181 the intercept to zero for the regressions, the slope is larger than the slope without forcing the fitted
182 lines through zero (fig. 3). In generally, when O₃ is zero both the ozonesondes and the aircraft will
183 measure zero. However, there is an offset in the fit of the two data sets due to potential causes for
184 systematic differences during the observation measurement process, e.g., high background current
185 in the sonde data.

186 Fig. 2 shows that the mean differences between ozonesonde and aircraft measurements vary
187 significantly with altitude. This can also be observed clearly from the relative differences (RD),
188 expressed as $(O_{3\text{-ozonesonde}} - O_{3\text{-aircraft}}) / O_{3\text{-aircraft}} \times 100\%$ (Fig. 4). O₃ concentrations from ECC
189 measurements are higher than those from aircraft measurements in all altitudes except at the surface.
190 Mean O₃ concentrations reported by Brewer-Mast sondes are lower than those from IAGOS below
191 7 km, but higher between 7 and 12 km. O₃ concentrations reported by Carbon-Iodine sondes are
192 higher than those observed from aircrafts below 2 km, but significantly lower above 8 km. In relative
193 terms, the bias between ECC sonde and aircraft measurements varies little with altitude, except near
194 the ground. The mean relative bias for Brewer-Mast measurements is at an absolute maximum of -
195 19 % near the ground, but increases slowly above 3 km, and is positive above 7 km, reaching more
196 than +10 % at 10-11 km. The relative bias for Carbon-Iodine measurements is about 8% below 2
197 km, becomes quite small from 2 - 8 km, and becomes large and negative above 8 km.

198 The Indian-sonde observations show much larger mean differences from the aircraft measurements.
199 Biases are everywhere positive, and as high as nearly 60% or 30 ppb, with much higher uncertainty
200 (standard errors) at each altitude as well (Fig. 2b, Fig. 4).

201 The region below 3 km has many local ozone sources and sinks (cities, airports, rural environment,
202 etc). In comparison, the region above 8 km is significantly influenced by stratosphere-troposphere
203 exchange, jet streams, and tropopause folds. Fig. S1 shows that the R between ozonesondes and

204 aircraft observations is higher near the ground (< 2 km) and at high altitudes (> 10 km). This shows
205 that although the influencing factors of O₃ near the ground and at high altitudes are more complex,
206 their long-term temporal variation characteristics are similar. The influences of cities, airports, rural
207 environment, stratosphere-troposphere exchange, jet streams, tropopause folds, etc., have a more
208 significant impact on the concentration of O₃ in the short term.

209 The correlation between four types of ozonesondes and aircraft observations also varies with altitude
210 (Fig. S1). From 0-8 km, the correlation between ECC and aircraft observations decreases with
211 altitude, with R being 0.71 at 0-1 km and reaching a minimum of 0.29 at 8-9 km; from 8-12 km, R
212 increases with altitude, reaching 0.49 at 11-12 km. The correlation between the other three
213 ozonesondes (Brewer-mast, Indian-sonde and Carbon-iodine) and the aircraft observations all vary
214 with altitude, with different inflection points. The number of stations for these three types of
215 ozonesondes is small (Table 1). Therefore, local variable influences on O₃ are more important, so
216 R varies more with altitude.

217 The bias and RMSE with respect to the aircraft observations of the four types of ozonesondes at 8-
218 12 km are higher than that at other altitudes. In contrast, the bias and RMSE values below 8 km are
219 smaller and vary less with altitude, consistent with the vertical distribution characteristics of O₃
220 concentration in Fig. 2. This is likely due to the higher concentration of O₃ and the typically larger
221 difference in spatial distance between ozonesonde and aircraft observations at 8-12km.

222 In addition, the bias and RMSE relative to the aircraft observations at different altitudes for ECC,
223 Carbon-iodine and Brewer-mast sondes are lower than those for the Indian-sonde, which is similar
224 to the results of the above analysis of O₃ concentration.

225 It should be noted that these comparisons only give an average relative bias between sondes and
226 IAGOS. The true value of the ozone profile remains unknown, as do the absolute biases of sondes
227 and IAGOS.

228

229 **3.2 Seasonal variations in relative biases between ozonesondes and IAGOS**

230 Fig. 5 compares mean profiles observed by ECC ozonesondes and IAGOS, separated by season.
231 There are modest seasonal differences in the relative bias profiles, with somewhat larger average
232 biases in winter and spring, but average biases are all positive (ECC sondes higher) and at all levels

233 the average seasonal biases are not statistically different.

234 The modest seasonal differences that are apparent in Fig. 5 and in Figs. S2-S4 are likely due to the
235 modest sample size (for ECC sondes) and small sample sizes (for other types). The actual
236 coincidence in time for profiles can range from less than one day to about 1-3 weeks, depending on
237 the number of ozonesonde and aircraft O₃ profiles collected within each month-bin. This means the
238 larger the atmospheric variability of O₃ is, the larger the real differences between ozonesonde and
239 aircraft O₃ can become, particularly when the number of profiles within a month-bin are small. In
240 addition, there are errors due to variations in the aircraft take-off and landing trajectories and the
241 balloon rise rate, the geographical location of the observation stations (and any associated
242 meteorological differences) and any systematic difference in standard observational times.

243 Table 2 indicates that in all four seasons ECC data correlate well with aircraft observations, with R
244 ranging from 0.71 to 0.76, but with larger average biases in winter and spring, as noted. It is not
245 clear if these seasonal average differences in bias are significant, as the uncertainty ranges on the
246 seasonal averages (lower plot of Fig. 5) overlap.

247 The vertical distribution of tropospheric O₃ observed by Brewer-Mast and IAGOS aircraft in the
248 four seasons is similar (Fig. S2). Differences are also similar, except above 7 km, where the
249 uncertainties are larger, and in general the uncertainty ranges on the seasonal difference averages
250 overlap. Since these comparisons come from only one station pair, some of the differences may be
251 attributable to local differences in topography and meteorology. Table 2 shows that correlations for
252 the ensemble of Brewer-Mast stations are higher than those for ECC stations. Like the ECC sondes,
253 average biases are all positive, but this is determined by the biases above 7 km (Fig. 4); unlike the
254 ECCs, biases are negative in the lowest 3 km.

255 The vertical distribution of tropospheric O₃ concentrations observed by Carbon-Iodine sondes and
256 IAGOS aircraft in the four seasons are similar, except in summer when the tropopause is high (Fig.
257 S3). The difference plots are fairly similar, except in the lowest 3 km, where differences become
258 quite large in summer. Like the previous comparison for Brewer-Mast sondes, these comparisons
259 come from only one station pair, and so the large differences in the boundary layer in summer are
260 likely due to local O₃ production sampled by the sonde but not the aircraft. Likely for this reason,
261 the consistency between Carbon-Iodine and aircraft observations is poor in summer, with R being

262 only 0.46 (Table 2). For the other three seasons it is fairly good.
263 The tropospheric O₃ observed by Indian-sondes displays a consistently high bias relative to IAGOS
264 in all seasons, and the seasonal difference plots are quite similar, except in the lowest 3 km in winter
265 (Fig. S4). This different behavior in winter is likely due to local ozone production sampled by the
266 aircraft but not the sonde. Temperature inversions are common in the winter in northern India and
267 trap local pollution. The very low values registered by the aircraft near the surface in summer also
268 suggest local effects, in this case titration by NO_x.
269 The tropospheric O₃ observed by Indian-sonde in the four seasons is 43.3-79.4 ppb, 31.4-80.2 ppb,
270 42.2-69.6 ppb and 51.5-87.5 ppb, and that observed by aircraft in the four seasons is 22.8-60.1 ppb,
271 14.8-47.1 ppb, 25.0-44.1 ppb and 35.6-53.3 ppb (Fig. S4). The tropospheric O₃ observed in Indian-
272 sonde in the four seasons increases with height almost linearly. The tropospheric O₃ observed by
273 aircraft first increases and then decreases with altitude in spring, summer and autumn, while in
274 winter, it first decreases and then increases with altitude. The tropospheric O₃ observed by the
275 Indian-sonde and the aircraft is quite different, and the RD in the four seasons is 6.3% to 47.5%,
276 22.6% to 52.9%, 26.4% to 40.6% and 5.13% to 39.13%. Table 2 indicates poor consistency between
277 Indian-sonde and aircraft observations in all four seasons, with R in winter only 0.18. The bias and
278 RMSE in winter are the largest, at 40.07 ppb and 64.99 ppb. The bias, R and RMSE in the other
279 three seasons are smaller, and the differences between them slight.

280 **3.3 Dependence of relative biases on station-airport distances**

281 A major concern with comparing IAGOS and ozonesonde observations is that the stations and
282 airports are not generally co-located, and even where they are close, the flight paths taken by balloon
283 and aircraft are quite different. Fig. 6 compares the average vertical distribution of tropospheric O₃
284 observed at different station-airport distances by ECC sondes and IAGOS aircraft. Note that we
285 continue to separate sonde station data by type --- only ECC data are used here. Sonde-aircraft pairs
286 have been grouped by station-airport distance (Table 1). The differences in average bias vary only
287 very modestly between the different station-airport distance categories, and those differences are
288 not statistically different at the 95% confidence level (Fig. 6d). This, partially owing presumably to
289 the use of mean monthly averages, is encouraging, as this provides further evidence that the average
290 bias we have derived is an artifact strictly of instrument differences.

291 Table 3 indicates that the bias variation between ECC and aircraft observations at different station-
292 airport distances is small, ranging from 5.7 ppb to 9.8 ppb. Correlations for these groupings are also
293 fairly similar, at $R = 0.8, 0.9$ and 0.7 .

294 Compared with ECC sondes, the consistency between the Indian-sonde and aircraft observations is
295 poor at all station-airport distances, with much larger biases, and poor correlations, with $R = 0.2$ to
296 0.4 . Nevertheless, Fig. S5 shows that the profiles of average differences are quite similar for station-
297 airport distances $< 1^\circ$, and distances of 2° - 4° (Fig. S5c).

298 Fig. 7 and Figs. S6-S8 examine possible seasonal variation in the differences at different station-
299 airport distances, for ECC sondes. The mean differences for the different station-airport distance
300 categories are larger than for the annual averages (Fig. 6), but in general those differences are not
301 statistically different at the 95% confidence level (Figs. 7d and S6d-S8d).

302

303 **3.4 Comparison of ozonesonde relative biases under operational conditions using IAGOS** 304 **observations as a transfer standard**

305 The foregoing discussion demonstrates that, consistent with previous work, there is a fairly constant
306 relative bias between IAGOS and sondes, with considerable dependence on sonde type, as expected
307 from previous sonde intercomparisons like JOSIE 1996. Although uncertainties are sizeable due to
308 the relatively sparse nature of the available data, we find consistent differences at all sites, with little
309 dependence on season or on station-airport separation, and little regional dependence (not shown).
310 Notwithstanding this overall sonde-IAGOS bias, we can use these station-airport comparisons to
311 derive relative biases of the different sonde types in use in the global network.

312 This does not assume that the aircraft data are unbiased. The true value of the O_3 profile (or even its
313 average) remains unknown, as do the absolute biases of sondes and IAGOS. It does assume:

- 314 1. That the measurement errors are random and normally distributed;
- 315 2. That there is one, constant bias for each measurement type (that is, if, for example, the Indian
316 sonde has changed over the period of comparison, or the IAGOS instruments have different biases,
317 there would be additional error that is not included in our uncertainty estimate);
- 318 3. That the measurement biases are not dependent on the geographic location or other variability of
319 the O_3 profile. This does not assume that the average O_3 profile is the same, just that the instruments

320 respond in the same way.
321 With these assumptions we can use the results of Fig. 2 to estimate the relative biases of each sonde
322 type to each other. The uncertainty of the comparisons will be the quadratic sum of the uncertainties
323 of the two IAGOS-sonde comparisons. The results are shown in Table 4. This intercomparison of
324 the different sonde types has an important advantage: it compares ozonesonde relative biases under
325 operational conditions, as it compares the data that are actually in databases like the WOUDC. It
326 also fills a gap, as the last WMO international intercomparison involving all four sonde types was
327 JOSIE 1996. These results are broadly consistent with those from JOSIE 1996 (Smit and Kley, 1998;
328 their Table 8 and Fig. 11).

329 In fact, the types of ozonesonde have changed during long-term observations at some stations (e.g.
330 Uccle and Payerne). De Backer et al. (1998) showed that with the use of an appropriate correction
331 procedure, accounting for the loss of pump efficiency with decreasing pressure and temperature, it
332 is possible to reduce the mean difference between O₃ profiles obtained with both types of sondes
333 below 3%, which is statistically insignificant over nearly the whole operational altitude range (from
334 the ground to 32 km). Stübi et al. (2008) also found that the O₃ difference between the Brewer-Mast
335 and the ECC ozonesonde data shows good agreement between the two sonde types, and the profile
336 of the O₃ difference is limited to ±5% (±0.3 mPa) from the ground to 32 km. The results for Brewer-
337 Mast sondes in Table 4 should also be applicable to the older Payerne and Uccle records, and are
338 generally consistent with these results and with those for the older Canadian records (Tarasick et al.,
339 2002; 2016).

340 The results in Table 4 will be quite valuable for addressing the problem of relative biases when
341 merging ozonesonde data into global climatologies (e.g. McPeters et al., 2007; McPeters and Labow,
342 2012; Bodeker et al., 2013; Liu et al., 2013; Hassler et al., 2018;).

343 **4 Conclusions**

344 The vertical distribution of tropospheric O₃ observed by ozonesondes and IAGOS aircraft sensors
345 are compared at 23 pairs of sites between about 30°S and 55°N from 1995 to 2021. Overall, ECC,
346 Brewer-Mast, and Carbon-Iodine sondes agree reasonably well with aircraft observations, with
347 average biases of 2.58 ppb, -0.28 ppb, and 0.67 ppb, and correlation coefficients of 0.72, 0.82, and
348 0.66, respectively. The agreement between the aircraft and Indian-sonde observations is poor, with

349 an average bias of 15.32 ppb and R of 0.44. Ozonesondes and aircraft observations have smaller R
350 in the middle troposphere, but larger bias and RMSE in the upper troposphere. The bias and RMSE
351 relative to the aircraft observations at different altitudes for ECC, Carbon-iodine and Brewer-mast
352 sondes are lower than those for the Indian-sonde.

353 Notwithstanding this general agreement, all sonde types show significant average biases with
354 respect to IAGOS. The O₃ concentration observed by ECC sondes is on average higher by 5-10%
355 than that observed by IAGOS aircraft, and the relative bias increases modestly with altitude.
356 Seasonal variations in the relative bias are not in general statistically significant. The distance
357 between station and airport within 4° also has little effect on the comparison results. When the ECC
358 station pairs are grouped by station-airport distances of <1° (latitude and longitude), 1-2°, and 2-4°,
359 biases with respect to IAGOS measurements vary from 5.7 to 9.8 ppb, and correlations from 0.7 to
360 0.9.

361 Thus, the observed average relative bias between sondes and IAGOS found in this study, also noted
362 by previous authors (Zbinden et al., 2013; Staufer et al., 2013, 2014; Tanimoto et al., 2015; Tarasick
363 et al., 2019), is a robust result. Possible reasons for the difference include: side reactions that cause
364 sondes to produce excess iodine (Saltzman and Gilbert, 1959), and/or loss of O₃ on the inlet pump
365 that could cause IAGOS monitors to read low at pressures below 800 hPa. The latter was an issue
366 in earlier aircraft O₃ sampling programs (Schnadt Poberaj et al., 2007; Dias-Lalcaca et al., 1998;
367 Brunner et al., 2001), but Thouret et al. (1998) found it negligible for MOZAIC/IAGOS. A recent
368 intercomparison campaign at the World Calibration Centre for Ozone Sondes (WCCOS) in Julich
369 in June 2023 indicates that the pumps do not greatly influence the ozone IAGOS measurements
370 between 1000 and 200 hPa. The IAGOS-CORE O₃ measurements (Package 1 with pressurization
371 pumps) and IAGOS-CARIBIC O₃ measurements differ by less than 2%, and the WCCOS reference
372 UV photometer measurements are usually higher by 1-2% (to a maximum of 5%) compared to both
373 IAGOS instruments (Blot et al., 2021; Nédélec et al., 2015; Thouret et al., 2022). IAGOS-CARIBIC
374 does not have pressurization system, so that's why the good comparison between both IAGOS
375 systems means a lot.

376 However, as noted by Saltzman and Gilbert (1959), the differences in stoichiometry found at
377 different pH values imply that the chemistry of reaction of O₃ with KI is complex, involving

378 reactions that cause loss of iodine, as well as reactions other than the principal one that produce
379 additional iodine. Several authors have noted the existence of slow side reactions involving the
380 phosphate buffer, with a time constant of about 20 minutes, that may also increase the stoichiometry
381 from 1.0 (Tarasick et al., 2021, Smit et al., 2024). Furthermore, evaporation causes the concentration
382 of the sensing solution to increase, which can further enhance the stoichiometry, by concentrating
383 the phosphate buffer, and to a lesser degree, by increasing the concentration of the KI itself (Johnson
384 et al., 2002). These factors could contribute to the observed average relative bias between sondes
385 and IAGOS found in this study.

386 This result implies that care must be taken when merging ozonesonde and IAGOS measurement
387 datasets. While the aircraft and sonde measurements are often complementary, filling in important
388 spatial gaps that would otherwise exist if only one type were used, the records are not typically over
389 the same period, and so merging can introduce spurious jumps if relative biases are not taken into
390 account.

391 The importance of O₃ in the troposphere as an air pollutant and a greenhouse gas, and therefore of
392 accurate measurements of its temporal and spatial distribution implies that it will be important to
393 resolve the causes of this bias, and so further research involving more direct comparisons of IAGOS
394 instrumentation and ozonesondes, e.g. in the WCCOS chamber, are strongly recommended.

395 These results are also useful to evaluate the relative biases of the different sonde types in the
396 troposphere, using the aircraft as a transfer standard. This intercomparison of the different sonde
397 types has the advantage that it compares ozonesonde relative biases under operational conditions;
398 that is, the data that are actually in databases like the WOUDC. These results will be invaluable for
399 addressing relative biases when merging ozonesonde data into global climatologies (e.g. Bodeker
400 et al., 2013; Hassler et al., 2018; Liu et al., 2013; McPeters et al., 2007; McPeters and Labow, 2012).

401

402 **Competing interests.** The contact authors have declared that none of the authors has any competing
403 interests.

404 **Data availability.** The global ozone sounding data were acquired from the World Ozone and
405 Ultraviolet Radiation Data Center (<http://www.woudc.org>) operated by Environment Canada. The
406 IAGOS data are created with support from the European Commission, national agencies in Germany

407 (BMBF), France (MESR), and the UK (NERC), and the IAGOS member institutions
408 (<http://www.iagos.org/partners>).

409 **Author contributions.** **HW:** Data curation, Methodology, Validation, Visualization, Writing -
410 original draft preparation, Writing - review & editing, Funding acquisition. **LS:** Methodology,
411 Investigation, Writing - original draft. **DWT:** Data curation, Resources, Conceptualization,
412 Supervision, Writing - original draft preparation, Writing - review & editing. **JL:** Data curation,
413 Resources, Methodology, Conceptualization, Supervision, Writing - original draft preparation,
414 Writing - review & editing, Funding acquisition. **TZ:** Funding acquisition, Writing - review &
415 editing. **HGJS** and **RVM:** Writing - review & editing. **HGJS** and **RB:** Acquisition of IAGOS data,
416 Quality assessment IAGOS data.

417

418 **Acknowledgments.** We thank many whose dedication makes datasets used in this study possible.
419 The global O₃ sounding data were acquired from the World Ozone and Ultraviolet Radiation Data
420 Center (<http://www.woudc.org>) operated by Environment Canada, Toronto, Canada, under the
421 auspices of the World Meteorological Organization. Flight-based atmospheric chemical
422 measurements are from IAGOS. IAGOS is funded by the European Union projects IAGOS-DS and
423 IAGOS-ERI. The IAGOS data are created with support from the European Commission, national
424 agencies in Germany (BMBF), France (MESR), and the UK (NERC), and the IAGOS member
425 institutions (<http://www.iagos.org/partners>). The participating airlines (Lufthansa, Air France,
426 Austrian, China Airlines, Iberia, Cathay Pacific, Air Namibia, Sabena) supported IAGOS by
427 carrying the measurement equipment free of charge since 1994. We are also thankful to the Digital
428 Research Alliance of Canada at the University of Toronto for facilitating data analysis.

429

430 **Financial support.** This study was supported by the Natural Science and Engineering Council of
431 Canada (Grant No. RGPIN-2020-05163), the National Key Research and Development Program of
432 China (Grant No., 2022YFC3701204), the National Natural Science Foundation of China
433 (42275196 and 41830965), the Natural Science Foundation of Jiangsu Province (BK20231300), and
434 Wuxi University Research Start-up Fund for Introduced Talents (2023r035).

435

436 **References**

- 437 Attmannspacher, A., and Dütsch, H.U., 1970. International ozone sonde intercomparison at the
438 Observatory Hohenpeissenberg, Ber. Dtsch. Wetterdienstes, 120, 1-85.
- 439 Attmannspacher, A., and Dütsch, H.U., 1981. Second international ozone sonde intercomparison at
440 the Observatory Hohenpeissenberg, Ber. Dtsch. Wetterdienstes, 157, 1-64.
- 441 Beekmann, M., Ancellet, G., Megie, G., Smit, H.G.J., Kley, D., 1994. Intercomparison campaign of
442 vertical ozone profiles including electrochemical sondes of ECC and Brewer-Mast type and a
443 ground based UV-differential absorption lidar. *Journal of Atmospheric Chemistry*, 19, 259-288.
444 19.
- 445 Beekmann, M., Ancellet, G., Martin, D., Abonnel, C., Duverneuil, G., Eideliman, F., Bessemoulin,
446 P., Fritz, N., Gizard, E., 1995. Intercomparison of tropospheric ozone profiles obtained by
447 electrochemical sondes, a ground based lidar and an airborne UV-photometer. *Atmospheric
448 Environment*, 29(9), 1027-1042.
- 449 Bernhard, G.H., Bais, A.F., Aucamp, P.J., Klekociuk, A.R., Liley, J.B., McKenzie, R.L., 2023.
450 Stratospheric ozone, UV radiation, and climate interactions. *Photochemical and
451 Photobiological Sciences*, 1-53.
- 452 Blot, R., Nedelec, P., Boulanger, D., Wolff, P., Sauvage, B., Cousin, J.-M., Athier, G., Zahn, A.,
453 Obersteiner, F., Scharffe, D., Petetin, H., Bennouna, Y., Clark, H., Thouret, V., 2021. Internal
454 consistency of the IAGOS ozone and carbon monoxide measurements for the last 25 years.
455 *Atmospheric Measurement Techniques*, 14, 3935-3951, [https://doi.org/10.5194/amt-14-3935-](https://doi.org/10.5194/amt-14-3935-2021)
456 2021.
- 457 Bodeker, G. E., Hassler, B., Young, P. J., Portmann, R. W., 2013. A vertically resolved, global, gap-
458 free ozone database for assessing or constraining global climate model simulations. *Earth
459 System Science Data*, 5, 31-43, <https://doi.org/10.5194/essd-5-31-2013>.
- 460 Brunner, D., J. Staehelin, D. Jeker, H., Wernli, U. Schumann, 2001. Nitrogen oxides and ozone in
461 the tropopause region of the Northern Hemisphere: Measurements from commercial aircraft in
462 1995/96 and 1997. *Journal of Geophysical Research: Atmospheres*, 106, 27673-27699.
- 463 Callis, L.B., Boughner, R.E., Natarajan, M., Lambeth, J.D., Baker, D.N., Blake, J.B., 1991. Ozone
464 depletion in the high latitude lower stratosphere: 1979-1990. *Journal of Geophysical Research*:

465 Atmospheres, 96(D2), 2921-2937.

466 David, L.M. and Nair, P.R., 2013. Tropospheric column O₃ and NO₂ over the Indian region observed
467 by Ozone Monitoring Instrument (OMI): Seasonal changes and long-term trends. *Atmospheric*
468 *Environment*, 65, 25-39.

469 De Backer, H., De Muer, D., De Sadelaeer, G., 1998. Comparison of ozone profiles obtained with
470 Brewer-Mast and Z-ECC sensors during simultaneous ascents. *Journal of Geophysical*
471 *Research: Atmospheres*, 103(D16), 19641-19648.

472 Deshler, T., Mercer, J.L., Smit, H.G., Stubi, R., Levrat, G., Johnson, B.J., Oltmans, S.J., Kivi, R.,
473 Thompson, A.M., Witte, J., Davies, J., 2008. Atmospheric comparison of electrochemical cell
474 ozonesondes from different manufacturers, and with different cathode solution strengths: The
475 Balloon Experiment on Standards for Ozonesondes. *Journal of Geophysical Research:*
476 *Atmospheres*, 113(D4), D04307, doi:10.1029/2007JD008975.

477 Dias-Lalcaca, P., Brunner, D., Imfeld, W., Moser, W., Staehelin, J., 1998. An automated system for
478 the measurement of nitrogen oxides and ozone concentrations from a passenger aircraft:
479 Instrumentation and first results of the NOXAR project. *Environmental Science and*
480 *Technology*, 32(20), 3228-3236.

481 Ebojie, F., Burrows, J.P., Gebhardt, C., Ladstätter-Weißmayer, A., Von Savigny, C., Rozanov, A.,
482 Weber, M., Bovensmann, H., 2016. Global tropospheric ozone variations from 2003 to 2011
483 as seen by SCIAMACHY. *Atmospheric Chemistry and Physics*, 16(2), 417-436.

484 Fu, Y. and Tai, A.P.K., 2015. Impact of climate and land cover changes on tropospheric ozone air
485 quality and public health in East Asia between 1980 and 2010. *Atmospheric Chemistry and*
486 *Physics*, 15(17), 10093-10106.

487 García, O.E., Sanromá, E., Schneider, M., Hase, F., León-Luis, S.F., Blumenstock, T., Sepúlveda,
488 E., Redondas, A., Carreño, V., Torres, C., Prats, N., 2022. Improved ozone monitoring by
489 ground-based FTIR spectrometry. *Atmospheric Measurement Techniques*, 15(8), 2557-2577.

490 García, O.E., Schneider, M., Sepúlveda, E., Hase, F., Blumenstock, T., Cuevas, E., Ramos, R., Gross,
491 J., Barthlott, S., Röhlings, A.N., Sanromá, E., 2021. Twenty years of ground-based NDACC
492 FTIR spectrometry at Izaña Observatory-overview and long-term comparison to other
493 techniques. *Atmospheric Chemistry and Physics*, 21(20), 15519-15554.

494 Gaudel, A., Cooper, O.R., Ancellet, G., Barret, B., Boynard, A., Burrows, J.P., Clerbaux, C., Coheur,
495 P.F., Cuesta, J., Cuevas, E., Doniki, S., 2018. Tropospheric Ozone Assessment Report: Present-
496 day distribution and trends of tropospheric ozone relevant to climate and global atmospheric
497 chemistry model evaluation. *Elementa: Science of the Anthropocene*, 6.

498 Gaudel, A., Cooper, O.R., Chang, K.L., Bourgeois, I., Ziemke, J.R., Strode, S.A., Oman, L.D.,
499 Sellitto, P., Nédélec, P., Blot, R., Thouret, V., 2020. Aircraft observations since the 1990s reveal
500 increases of tropospheric ozone at multiple locations across the Northern Hemisphere. *Science*
501 *Advances*, 6(34), eaba8272.

502 Gebhardt, C., Rozanov, A., Hommel, R., Weber, M., Bovensmann, H., Burrows, J.P., Degenstein,
503 D., Froidevaux, L., Thompson, A.M., 2014. Stratospheric ozone trends and variability as seen
504 by SCIAMACHY from 2002 to 2012. *Atmospheric Chemistry and Physics*, 14(2), 831-846.

505 Hassler, B., Kremser, S., Bodeker, G. E., Lewis, J., Nesbit, K., Davis, S. M., Chipperfield, M. P.,
506 Dhomse, S. S., Dameris, M., 2018, An updated version of a gap-free monthly mean zonal mean
507 ozone database. *Earth System Science Data*, 10, 1473-1490.

508 Hegarty, J., Mao, H., Talbot, R., 2009. Synoptic influences on springtime tropospheric O₃ and CO
509 over the North American export region observed by TES. *Atmospheric Chemistry and*
510 *Physics*, 9(11), 3755-3776.

511 Hilsenrath, E., Attmannspacher, W., Bass, A., Evans, W., Hagemeyer, R., Barnes, R., Komhyr, W.,
512 Mauersberger, K., Mentall, J., Proffitt, M., Robbins, D., 1986. Results from the balloon ozone
513 intercomparison campaign (BOIC). *Journal of Geophysical Research: Atmospheres*, 91(D12),
514 13137-13152.

515 Hoogen, R., Rozanov, V.V., Burrows, J.P., 1999. Ozone profiles from GOME satellite data:
516 Algorithm description and first validation. *Journal of Geophysical Research:*
517 *Atmospheres*, 104(D7), 8263-8280.

518 Hu, L., Jacob, D.J., Liu, X., Zhang, Y., Zhang, L., Kim, P.S., Sulprizio, M.P., Yantosca, R.M., 2017.
519 Global budget of tropospheric ozone: Evaluating recent model advances with satellite (OMI),
520 aircraft (IAGOS), and ozonesonde observations. *Atmospheric Environment*, 167, 323-334.

521 Hubert, D., Heue, K.P., Lambert, J.C., Verhoelst, T., Allaart, M., Compernelle, S., Cullis, P.D., Dehn,
522 A., Félix, C., Johnson, B.J., Keppens, A., 2021. TROPOMI tropospheric ozone column data:

523 geophysical assessment and comparison to ozonesondes, GOME-2B and OMI. *Atmospheric*
524 *Measurement Techniques*, 14(12), 7405-7433.

525 Johnson, B.J., Oltmans, S.J., Vömel, H., Smit, H.G.J., Deshler, T., Kroeger, C., 2002. ECC
526 Ozonesonde pump efficiency measurements and tests on the sensitivity to ozone of buffered
527 and unbuffered ECC sensor cathode solutions. *Journal of Geophysical Research: Atmospheres*,
528 107(4393), 10-1029. doi: 10.1029/2001JD000557.

529 Keckhut, P., McDermid, S., Swart, D., McGee, T., Godin-Beekmann, S., Adriani, A., Barnes, J.,
530 Baray, J.L., Bencherif, H., Claude, H., di Sarra, A.G., 2004. Review of ozone and temperature
531 lidar validations performed within the framework of the Network for the Detection of
532 Stratospheric Change. *Journal of Environmental Monitoring*, 6(9), 721-733.

533 Kerr, J.B., Fast, H., McElroy, C.T., Oltmans, S.J., Lathrop, J.A., Kyro, E., Paukkunen, A., Claude,
534 H., Köhler, U., Sreedharan, C.R., Takao, T., 1994. The 1991 WMO international ozonesonde
535 intercomparison at Vanscoy, Canada. *Atmosphere-Ocean*, 32(4), 685-716.

536 Lefohn, A.S., Malley, C.S., Smith, L., Wells, B., Hazucha, M., Simon, H., Naik, V., Mills, G.,
537 Schultz, M.G., Paoletti, E., De Marco, A., 2018. Tropospheric ozone assessment report: Global
538 ozone metrics for climate change, human health, and crop/ecosystem research. *Elementa:*
539 *Science of the Anthropocene*, 6.

540 Li, K., Jacob, D.J., Liao, H., Qiu, Y., Shen, L., Zhai, S., Bates, K.H., Sulprizio, M.P., Song, S., Lu,
541 X., Zhang, Q., 2021. Ozone pollution in the North China Plain spreading into the late-winter
542 haze season. *Proceedings of the National Academy of Sciences*, 118(10), e2015797118.

543 Li, K., Jacob, D.J., Shen, L., Lu, X., De Smedt, I., Liao, H., 2020. Increases in surface ozone
544 pollution in China from 2013 to 2019: anthropogenic and meteorological
545 influences. *Atmospheric Chemistry and Physics*, 20(19), 11423-11433.

546 Liao, Z., Ling, Z., Gao, M., Sun, J., Zhao, W., Ma, P., Quan, J., Fan, S., 2021. Tropospheric ozone
547 variability over Hong Kong based on recent 20 years (2000-2019) ozonesonde
548 observation. *Journal of Geophysical Research: Atmospheres*, 126(3), e2020JD033054.

549 Liu, G., Liu, J., Tarasick, D.W., Fioletov, V.E., Jin, J.J., Moeini, O., Liu, X., Sioris, C.E., Osman,
550 M., 2013. A global tropospheric ozone climatology from trajectory-mapped ozone soundings.
551 *Atmospheric Chemistry and Physics*, 13(21), 10659-10675.

552 Logan, J.A., 1985. Tropospheric ozone: Seasonal behavior, trends, and anthropogenic
553 influence. *Journal of Geophysical Research: Atmospheres*, 90(D6), 10463-10482.

554 Ma, Y., Ma, B., Jiao, H., Zhang, Y., Xin, J., Yu, Z., 2020. An analysis of the effects of weather and
555 air pollution on tropospheric ozone using a generalized additive model in Western China:
556 Lanzhou, Gansu. *Atmospheric Environment*, 224, 117342.

557 McPeters, R.D., Labow, G.J., Logan, J.A., 2007. Ozone climatological profiles for satellite retrieval
558 algorithms. *Journal of Geophysical Research: Atmospheres*, 112(D5), D05308.
559 doi:10.1029/2005JD006823.

560 McPeters, R.D. and Labow, G.J., 2012. Climatology 2011: An MLS and sonde derived ozone
561 climatology for satellite retrieval algorithms. *Journal of Geophysical Research: Atmospheres*,
562 117(D10), D10303. doi:10.1029/2011JD017006.

563 Miles, G.M., Siddans, R., Kerridge, B.J., Latter, B.G., Richards, N.A.D., 2015. Tropospheric ozone
564 and ozone profiles retrieved from GOME-2 and their validation. *Atmospheric Measurement*
565 *Techniques*, 8(1), 385-398.

566 Meng, K., Zhao, T., Xu, X., Zhang, Z., Bai, Y., Hu, Y., Zhao, Y., Zhang, X., Xin, Y., 2022. Influence
567 of stratosphere-to-troposphere transport on summertime surface O₃ changes in North China
568 Plain in 2019. *Atmospheric Research*, 276, 106271.

569 Monks, P.S., Archibald, A.T., Colette, A., Cooper, O., Coyle, M., Derwent, R., Fowler, D., Granier,
570 C., Law, K.S., Mills, G.E., Stevenson, D.S., 2015. Tropospheric ozone and its precursors from
571 the urban to the global scale from air quality to short-lived climate forcer. *Atmospheric*
572 *Chemistry and Physics*, 15(15), 8889-8973.

573 Nédélec, P., Blot, R., Boulanger, D., Athier, G., Cousin, J.M., Gautron, B., Petzold, A., Volz-Thomas,
574 A., Thouret, V., 2015. Instrumentation on commercial aircraft for monitoring the atmospheric
575 composition on a global scale: the IAGOS system, technical overview of ozone and carbon
576 monoxide measurements. *Tellus B: Chemical and Physical Meteorology*, 67(1), 27791.
577 <https://doi.org/10.3402/tellusb.v67.27791@zelb20.2016.68.issue-s1>

578 Percy, K.E., Legge, A.H., Krupa, S.V., 2003. Tropospheric ozone: a continuing threat to global
579 forests? *Developments in Environmental Science*, 3, 85-118.

580 Perlwitz, J., Pawson, S., Fogt, R.L., Nielsen, J.E., Neff, W.D., 2008. Impact of stratospheric ozone

581 hole recovery on Antarctic climate. *Geophysical Research Letters*, 35(8).
582 <https://doi.org/10.1029/2008GL033317>.

583 Ramanathan, V., Cicerone, R.J., Singh, H.B., Kiehl, J.T., 1985. Trace gas trends and their potential
584 role in climate change. *Journal of Geophysical Research: Atmospheres*, 90(D3), 5547-5566.

585 Saltzman, B.E. and Gilbert, N., 1959. Iodometric microdetermination of organic oxidants and ozone.
586 Resolution of mixtures by kinetic colorimetry. *Analytical Chemistry*, 31(11), 1914-1920.

587 Schnadt Poberaj, C., Staehelin, J., Brunner, D., Thouret, V., Mohnen, V., 2007. A UT/LS ozone
588 climatology of the nineteen seventies deduced from the GASP aircraft measurement program.
589 *Atmospheric Chemistry and Physics*, 7(22), 5917-5936.

590 Schultz, M.G., Schröder, S., Lyapina, O., Cooper, O.R., Galbally, I., Petropavlovskikh, I., Von
591 Schneidmesser, E., Tanimoto, H., Elshorbany, Y., Naja, M., Seguel, R.J., 2017. Tropospheric
592 Ozone Assessment Report: Database and metrics data of global surface ozone
593 observations. *Elementa: Science of the Anthropocene*, 5.

594 Sharma, S., Sharma, P., Khare, M., 2017. Photo-chemical transport modelling of tropospheric ozone:
595 A review. *Atmospheric Environment*, 159, 34-54.

596 Smit, H.G.J., Sträter, W., Helten, M., Kley, D., Ciupa, D., Claude, H.J., Köhler, U., Hoegger, B.,
597 Levrat, G., Johnson, B., Oltmans, S.J., Kerr, J.B., Tarasick, D.W., Davies, J., Shitamichi, M.,
598 Srivastav, S.K., Vialle, C., 1996. JOSIE: The 1996 WMO international intercomparison of
599 ozonesondes under quasi-flight conditions in the environmental chamber at Jülich, *Proc.*
600 *Quadrennial Ozone Symposium 1996, l'Aquila, Italy*, edited by R. D. Bojkov and G. Visconti,
601 pp. 971-974, Parco Sci. e Tecnol. d'Abruzzo, Italy, 1996.

602 Smit, H.G., Straeter, W., Johnson, B.J., Oltmans, S.J., Davies, J., Tarasick, D.W., Hoegger, B., Stubi,
603 R., Schmidlin, F.J., Northam, T., Thompson, A.M., 2007. Assessment of the performance of
604 ECC-ozonesondes under quasi-flight conditions in the environmental simulation chamber:
605 Insights from the Juelich Ozone Sonde Intercomparison Experiment (JOSIE). *Journal of*
606 *Geophysical Research: Atmospheres*, 112(D19), D19306, doi:10.1029/2006JD007308.

607 Smit, H.G.J., 2014. Ozonesondes, in *Encyclopedia of Atmospheric Sciences, Second Edition*, edited
608 by G.R. North, J.A. Pyle, and F. Zhang, Vol 1, pp. 372–378, Academic Press, London.

609 Smit, H. G. J., Poyraz, D., Van Malderen, R., Thompson, A. M., Tarasick, D. W., Stauffer, R. M.,

610 Johnson, B. J., and Kollonige, D. E., 2024. New insights from the Jülich Ozone Sonde
611 Intercomparison Experiment: calibration functions traceable to one ozone reference instrument.
612 Atmospheric Measurement Techniques, 17, 73–112, <https://doi.org/10.5194/amt-17-73-2024>.

613 Smit, H.G.J., Thompson, A.M., and the ASOPOS 2.0 Panel, 2021. Ozonesonde Measurement
614 Principles and 1300 Best Operational Practices, WMO Global Atmosphere Watch Report
615 Series, No. 268, World Meteorological Organization, 1301 Geneva, [Available online at
616 https://library.wmo.int/doc_num.php?explnum_id=10884].

617 Solomon, S., 1999. Stratospheric ozone depletion: A review of concepts and history. Reviews of
618 Geophysics, 37(3), 275-316.

619 Staehelin, J., Harris, N.R., Appenzeller, C. and Eberhard, J., 2001. Ozone trends: A review. Reviews
620 of Geophysics, 39(2), 231-290.

621 Staufer, J., Staehelin, J., Stübi, R., Peter, T., Tummon, F., Thouret, V., 2013. Trajectory matching of
622 ozonesondes and MOZAIC measurements in the UTLS - Part 1: Method description and
623 application at Payerne, Switzerland. Atmospheric Measurement Techniques, 6, 3393-3406.
624 doi:10.5194/amt-6-3393-2013.

625 Staufer, J., Staehelin, J., Stübi, R., Peter, T., Tummon, F., Thouret, V., 2014. Trajectory matching of
626 ozonesondes and MOZAIC measurements in the UTLS - Part 2: Application to the global
627 ozonesonde network. Atmospheric Measurement Techniques, 7, 241-266. doi:10.5194/amt-7-
628 241-2014.

629 Stübi, R., Levrat, G., Hoegger, B., Viatte, P., Staehelin, J., Schmidlin, F.J., 2008. In-flight
630 comparison of Brewer-Mast and electrochemical concentration cell ozonesondes. Journal of
631 Geophysical Research: Atmospheres, 113(D13). <https://doi.org/10.1029/2007JD009091>.

632 Tanimoto, H., Zbinden, R.M., Thouret, V., Nédélec, P., 2015. Consistency of tropospheric ozone
633 observations made by different platforms and techniques in the global databases. Tellus B:
634 Chemical and Physical Meteorology, 67(1), 27073.

635 Tarasick, D. W., Davies, J., Anlauf, K., Watt, M., Steinbrecht, W., Claude, H. J., 2002. Laboratory
636 investigations of the response of Brewer-Mast ozonesondes to tropospheric ozone. Journal of
637 Geophysical Research: Atmospheres, 107(D16), ACH-14,
638 <https://doi.org/10.1029/2001JD001167>.

639 Tarasick, D. W., Davies, J., Smit, H. G., Oltmans, S. J., 2016. A re-evaluated Canadian ozonesonde
640 record: measurements of the vertical distribution of ozone over Canada from 1966 to 2013.
641 Atmospheric Measurement Techniques, 9(1), 195-214.

642 Tarasick, D.W., Galbally, I., Cooper, O.R., Schultz, M.G., Ancellet, G., LeBlanc, T., Wallington, T.J.,
643 Ziemke, J., Liu, X., Steinbacher, M., Stähelin, J., Vigouroux, C., Hannigan, J., García, O., Foret,
644 G., Zanis, P., Weatherhead, E., Petropavlovskikh, I., Worden, H., Neu, J.L., Osman, M., Liu, J.,
645 Lin, M., Granados-Muñoz, M., Thompson, A.M., Oltmans, S.J., Cuesta, J., Dufour, G., Thouret,
646 V., Hassler, B., Thompson, A.M., Trickl, T., 2019. TOAR- Observations: Tropospheric ozone
647 from 1877 to 2016, observed levels, trends and uncertainties. Elementa: Science of the
648 Anthropocene, 7(1), 39. DOI: <http://doi.org/10.1525/elementa.376>.

649 Tarasick, D.W., Smit, H.G., Thompson, A.M., Morris, G.A., Witte, J.C., Davies, J., Nakano, T., Van
650 Malderen, R., Stauffer, R.M., Johnson, B.J., Stübi, R., Oltmans, S.J., Vömel, H., 2021.
651 Improving ECC ozonesonde data quality: Assessment of current methods and outstanding
652 issues. Earth and Space Science, 8, e2019EA000914. <https://doi.org/10.1029/2019EA000914>.

653 Thouret, V., Clark, H., Petzold, A., Nédélec, P., Zahn, A., 2022. IAGOS: Monitoring Atmospheric
654 Composition for Air Quality and Climate by Passenger Aircraft. In Handbook of Air Quality
655 and Climate Change (pp. 1-14). Singapore: Springer Nature Singapore.

656 Thouret, V., Marenco, A., Logan, J.A., Nédélec, P., Grouhel, C., 1998. Comparisons of ozone
657 measurements from the MOZAIC airborne program and the ozone sounding network at eight
658 locations. Journal of Geophysical Research: Atmospheres, 103(D19), 25695-25720. doi:
659 10.1029/98JD02243.

660 Thompson, A. M., Smit, H. G. J., Witte, J. C., Stauffer, R. M., Johnson, B. J., Morris, G., von der
661 Gathen, P., Van Malderen, R., Davies, J., Piters, A., Allaart, M., Posny, F., Kivi, R., Cullis, P.,
662 Hoang Anh, N. T., Corrales, E., Machinini, T., da Silva, F. R., Paiman, G., Thiong'o, K., Zainal,
663 Z., Brothers, G. B., Wolff, K. R., Nakano, T., Stübi, R., Romanens, G., Coetzee, G. J. R., Diaz,
664 J. A., Mitro, S., Mohamad, M., Ogino, S., 2019. Ozonesonde quality assurance: The JOSIE-
665 SHADOZ (2017) experience. Bulletin of the American Meteorological Society, 100(1), 155-
666 171.

667 Vingarzan, R., 2004. A review of surface ozone background levels and trends. Atmospheric

668 Environment, 38(21), 3431-3442.

669 Vömel, H., Smit, H. G. J., Tarasick, D., Johnson, B., Oltmans, S. J., Selkirk, H., Thompson, A. M.,
670 Stauffer, R. M., Witte, J. C., Davies, J., van Malderen, R., Morris, G. A., Nakano, T., Stübi, R.,
671 2020. A new method to correct the electrochemical concentration cell (ECC) ozonesonde time
672 response and its implications for “background current” and pump efficiency, *Atmospheric*
673 *Measurement Techniques*, 13, 5667-5680, <https://doi.org/10.5194/amt-13-5667-2020>.

674 Wang, H., Ke, Y., Tan, Y., Zhu, B., Zhao, T., Yin, Y., 2023. Observational evidence for the dual roles
675 of BC in the megacity of eastern China: Enhanced O₃ and decreased PM_{2.5}
676 pollution. *Chemosphere*, 327, 138548.

677 Wang, H., Lu, X., Jacob, D.J., Cooper, O.R., Chang, K.L., Li, K., Gao, M., Liu, Y., Sheng, B., Wu, K.,
678 Wu, T., 2022. Global tropospheric ozone trends, attributions, and radiative impacts in 1995-2017:
679 an integrated analysis using aircraft (IAGOS) observations, ozonesonde, and multi-decadal
680 chemical model simulations. *Atmospheric Chemistry and Physics*, 22(20), 13753-13782.

681 Wang, T., Xue, L., Brimblecombe, P., Lam, Y.F., Li, L., Zhang, L., 2017. Ozone pollution in China:
682 A review of concentrations, meteorological influences, chemical precursors, and
683 effects. *Science of the Total Environment*, 575, 1582-1596.

684 Williamson, C.E., Neale, P.J., Hylander, S., Rose, K.C., Figueroa, F.L., Robinson, S.A., Häder, D.P.,
685 Wängberg, S.Å., Worrest, R.C., 2019. The interactive effects of stratospheric ozone depletion,
686 UV radiation, and climate change on aquatic ecosystems. *Photochemical and Photobiological*
687 *Sciences*, 18(3), 717-746.

688 Xu, J., Huang, X., Wang, N., Li, Y., Ding, A., 2021. Understanding ozone pollution in the Yangtze
689 River Delta of eastern China from the perspective of diurnal cycles. *Science of the Total*
690 *Environment*, 752, 141928.

691 Yang, T., Li, H., Wang, H., Sun, Y., Chen, X., Wang, F., Xu, L., Wang, Z., 2023. Vertical aerosol
692 data assimilation technology and application based on satellite and ground lidar: A review and
693 outlook. *Journal of Environmental Sciences*, 123, 292-305.

694 Young, P.J., Naik, V., Fiore, A.M., Gaudel, A., Guo, J., Lin, M.Y., Neu, J.L., Parrish, D.D., Rieder,
695 H.E., Schnell, J.L., Tilmes, S., 2018. Tropospheric Ozone Assessment Report: Assessment of
696 global-scale model performance for global and regional ozone distributions, variability, and

697 trends. *Elementa: Science of the Anthropocene*, 6.

698 Yu, R., Lin, Y., Zou, J., Dan, Y., Cheng, C., 2021. Review on atmospheric Ozone pollution in China:
699 Formation, spatiotemporal distribution, precursors and affecting factors. *Atmosphere*, 12(12),
700 1675.

701 Zang, Z., Liu, J., Tarasick, D., Moeini, O., Bian, J., Zhang, J., Thompson, A. M., Van Malderen, R., Smit,
702 H. G. J., Stauffer, R. M., Johnson, B. J., Kollonige, D. E., 2024. An improved Trajectory-mapped
703 Ozonesonde dataset for the Stratosphere and Troposphere (TOST): update, validation and
704 applications, *EGUsphere* [preprint], <https://doi.org/10.5194/egusphere-2024-800>, 2024.

705 Zbinden, R.M., Thouret, V., Ricaud, P., Carminati, F., Cammas, J.P., Nédélec, P., 2013. Climatology of
706 pure tropospheric profiles and column contents of ozone and carbon monoxide using MOZAIC in
707 the mid-northern latitudes (24° N to 50° N) from 1994 to 2009. *Atmospheric Chemistry and Physics*,
708 13(24), 12363-12388, <https://doi.org/10.5194/acp-13-12363-2013>, 2013.

709 Zhang, L., Jacob, D.J., Boersma, K.F., Jaffe, D.A., Olson, J.R., Bowman, K.W., Worden, J.R., Thompson,
710 A.M., Avery, M.A., Cohen, R.C., Dibb, J.E., 2008. Transpacific transport of ozone pollution and the
711 effect of recent Asian emission increases on air quality in North America: an integrated analysis
712 using satellite, aircraft, ozonesonde, and surface observations. *Atmospheric Chemistry and*
713 *Physics*, 8(20), 6117-6136.

714 Zhao, K., Huang, J., Wu, Y., Yuan, Z., Wang, Y., Li, Y., Ma, X., Liu, X., Ma, W., Wang, Y., Zhang,
715 X., 2021. Impact of stratospheric intrusions on ozone enhancement in the lower troposphere
716 and implication to air quality in Hong Kong and other South China regions. *Journal of*
717 *Geophysical Research: Atmospheres*, 126(18), e2020JD033955.

Figures

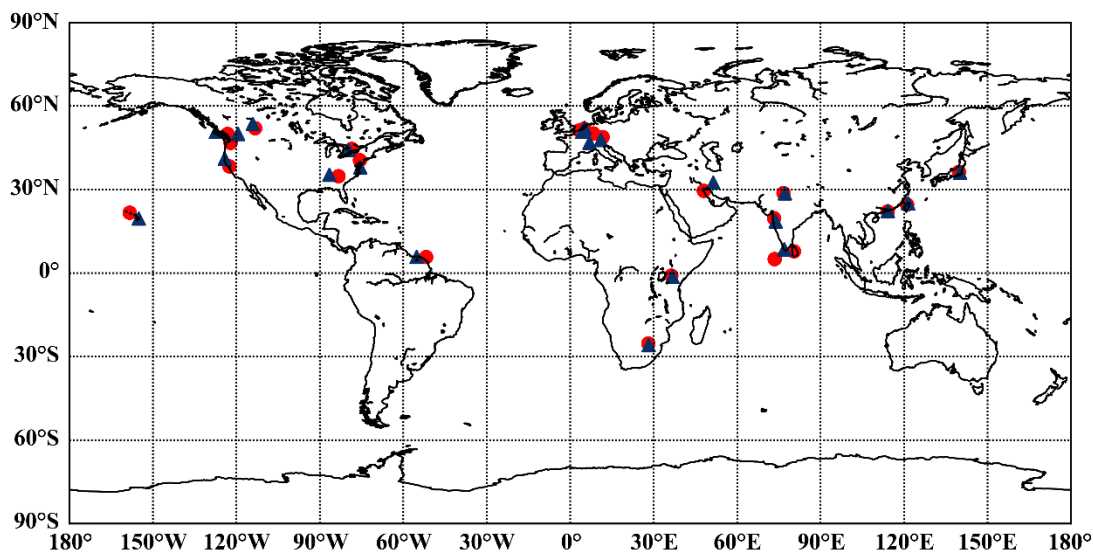


Figure 1. Map of 23 pairs of sites used in this study. Red circle markers are IAGOS sites, blue triangle markers are WOUDC sites.

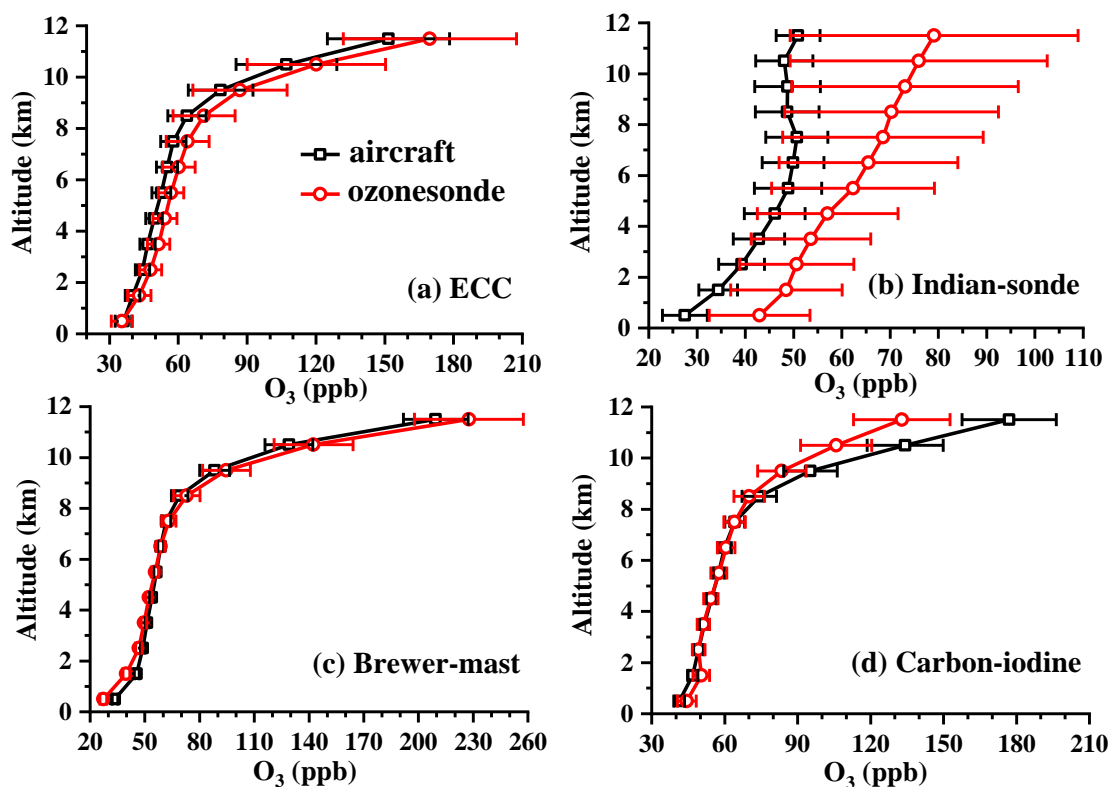


Figure 2. Comparison of the vertical profiles of tropospheric O_3 observed between aircraft measurements and four types of ozonesondes, ECC, Indian-sonde, Brewer-mast, and Carbon-iodine. The error bar length is 4 times the standard error (SE) of the mean (equivalent to 95% confidence limits on the averages).

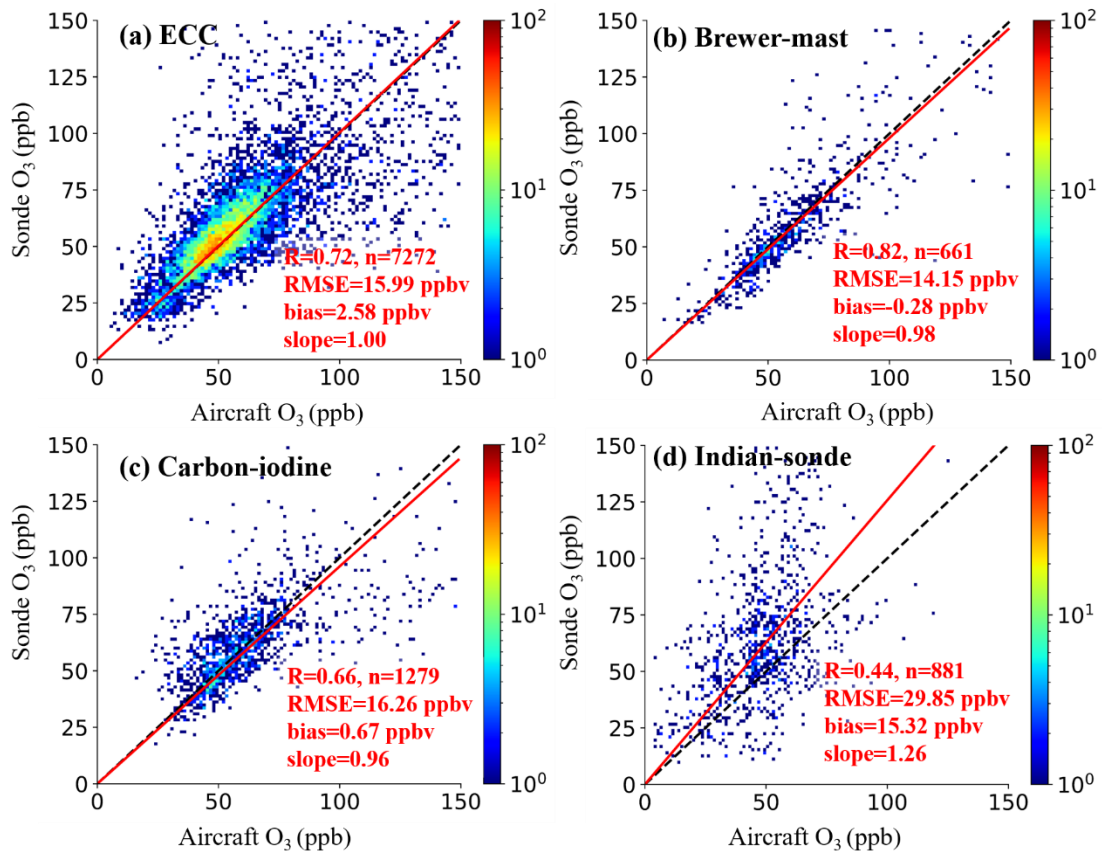


Figure 3. Correlation (R) of monthly mean ozone mixing ratios between ozonesonde and aircraft measurements. While IAGOS does measure in the lower stratosphere these values are usually far from the airport, so the sonde-aircraft distance will be large, we only plots data below 150 ppb. The black dashed line shows the 1:1 axis, the red line shows the linear fit (with the intercept set to 0), the color bar shows the data counts. Correlations are significant at the 99% level ($p < 0.01$). N denotes the number of data points, R is the correlation coefficient, Bias is the overall average difference in monthly mean values [Ozonesonde ozone – Aircraft ozone, in ppb], RMSE is the root mean square error, slope is the slope of the linear fit line. All data points are based on the monthly mean.

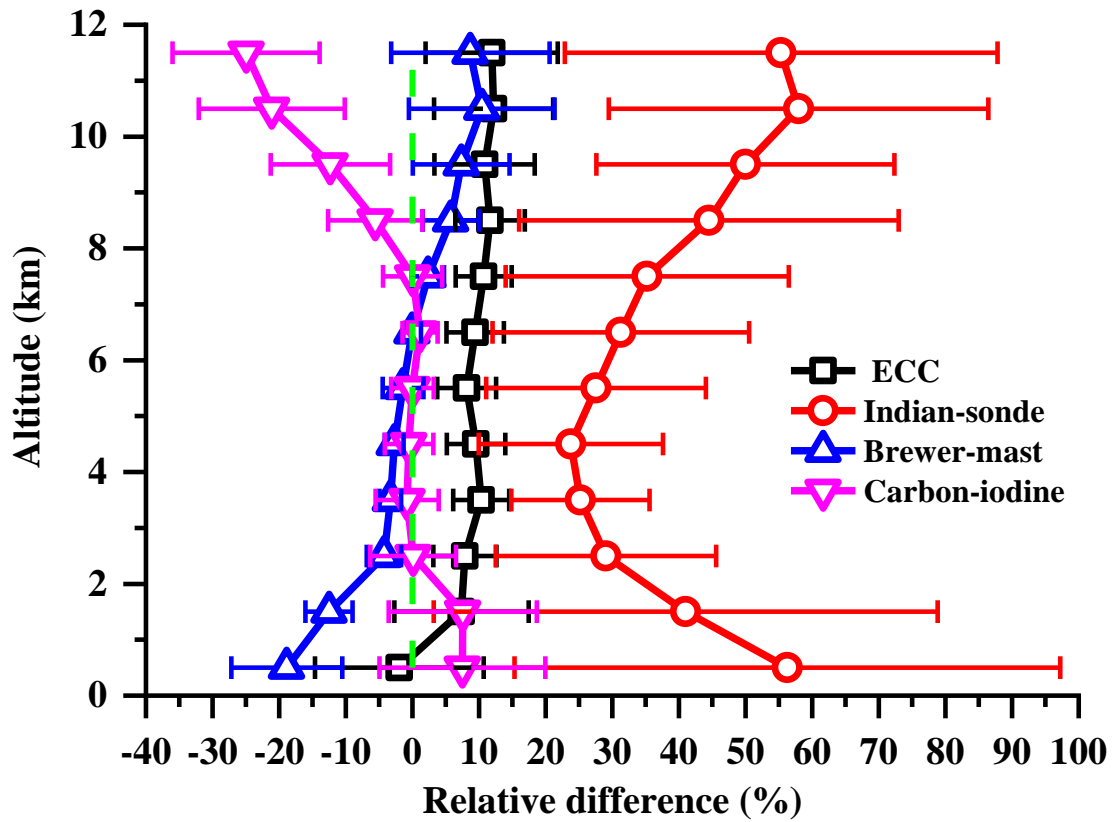


Figure 4. Mean relative difference (RD) between the ozonesonde O_3 and aircraft O_3 data. RD is calculated from $(O_3\text{-ozonesonde} - O_3\text{-aircraft}) / O_3\text{-aircraft} \times 100\%$. The green dashed line is the zero line.

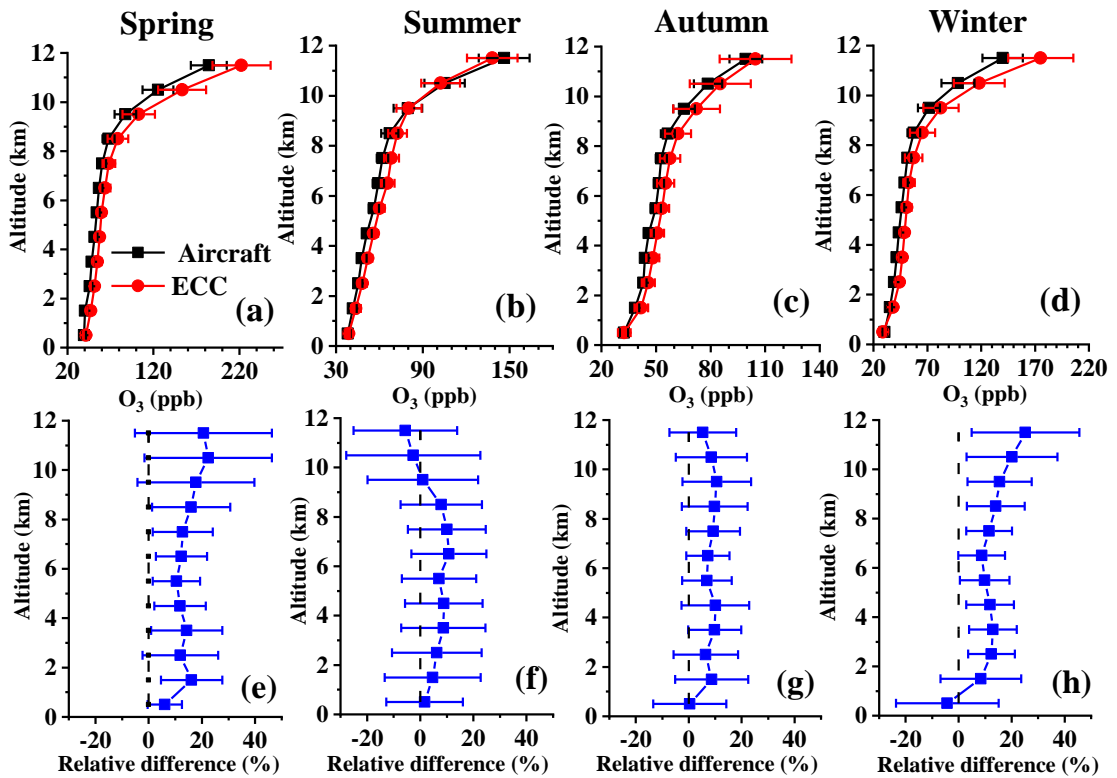


Figure 5. The mean difference in vertical profiles of the tropospheric O₃ between ECC ozonesonde and aircraft observations in four seasons (a-d) and their mean relative difference, the black dashed line is the zero line (e-h).

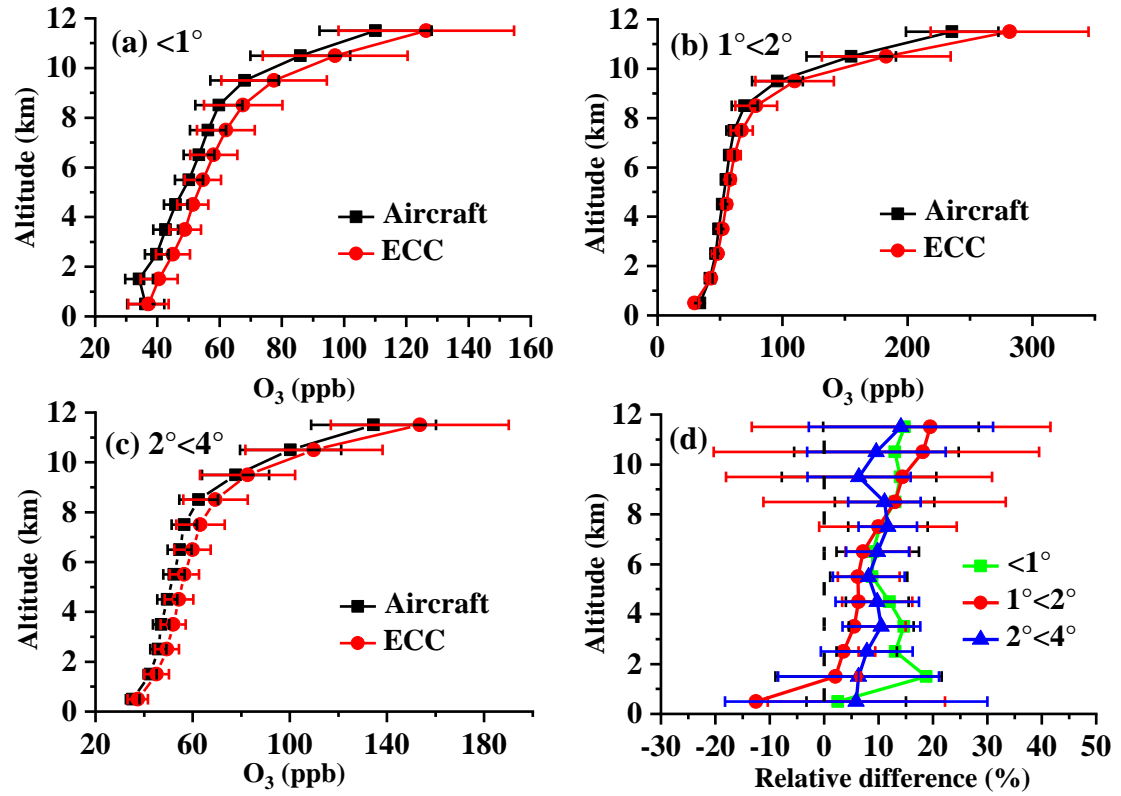


Figure 6. The annual mean vertical profiles of tropospheric O₃ between ECC ozonesonde and aircraft observations at station-pair distances (D) of $D < 1^\circ$ (a), $1^\circ < D < 2^\circ$ (b), and $2^\circ < D < 4^\circ$. The relative differences for the three categories are shown in (d), the black dashed line is the zero line.

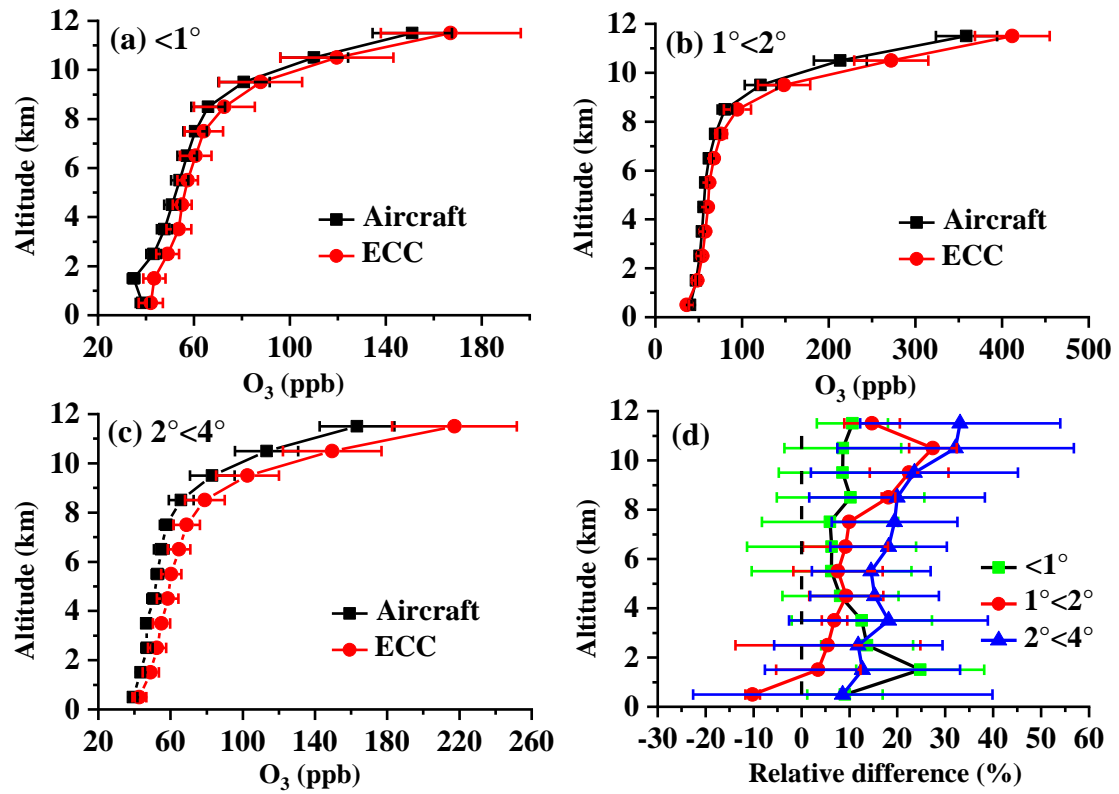


Figure 7. The seasonal mean vertical profiles of tropospheric O₃ in spring between ECC ozonesonde and aircraft observations at station-pair distances (D) of $D < 1^\circ$ (a), $1^\circ < D < 2^\circ$ (b), and $2^\circ < D < 4^\circ$. The relative differences for the three categories are shown in (d), the black dashed line is the zero line.

Tables

Table 1. Summary of the station information, including station's name, geolocation, the number of profiles, observational period, and the station-pair distance used in this study.

MOZAIC-IAGOS				WOUDC				Station- airport		No. valid data		observation	
Station name	Lon	Lat	No. profiles	Station name	Lon	Lat	No. profiles	Type	distance (km)	months	period	months	period
Toronto	-78.50	44.58	321	Egbert	-79.78	44.23	181	ECC	108.87	33	2004-2008		
Dusseldorf	4.96	51.82	412	De Bilt	5.18	52.10	333	ECC	34.59	63	1995-2013		
Munich	11.78	48.35	2136	Hohenpeissenberg	11.02	47.81	1032	Brewer-mast	82.42	67	1996-2006		
Johannesburg	28.07	-25.32	199	Irene	28.22	-25.91	135	ECC	67.30	26	1998-2003		
Nairobi	36.33	-0.94	114	Nairobi	36.75	-1.30	42	ECC	61.50	10	1997-1998		
Mumbai	73.26	19.05	122	Pune	73.85	18.53	56	Indian-sonde	84.85	35	1996-2003		
Delhi	76.65	28.73	342	New Delhi	77.18	28.63	88	Indian-sonde	52.88	50	1995-2016		
Hongkong	114.11	22.10	123	King's Park	114.17	22.31	115	ECC	24.15	25	2000-2005		
Taipei	121.08	24.59	2115	Taipei	121.48	25.02	58	ECC	62.58	31	2014-2018		
Tokyo	139.73	36.33	1342	Tateno (Tsukuba)	140.13	36.05	655	Carbon-iodine	47.52	116	1995-2006		

Calgary	-113.25	52.03	170	Edmonton	-114.10	53.55	112	ECC	178.41	17	2009-2011
Brussels	3.24	51.21	2412	Uccle	4.36	50.80	736	ECC	148.40	55	1997-2009
Honolulu	-158.33	21.66	169	Hilo (HI)	-155.07	19.58	107	ECC	410.56	16	2015-2017
Vancouver	-123.14	49.95	595	Kelowna	-127.38	50.69	594	ECC	312.01	68	2003-2015
San-Francisco	-122.50	38.30	34	Trinidad Head (CA)	-124.15	41.05	53	ECC	336.78	10	1999-2001
Portland	-122.06	46.76	385	Kelowna	-119.38	49.97	317	ECC	408.08	45	2003-2009
Atlanta	-83.28	34.78	34	Huntsville (AL)	-86.58	35.28	85	ECC	305.54	10	1999-2006
Washington	-75.59	40.52	610	Wallops Island (VA)	-75.46	37.94	616	ECC	287.09	80	1994-2014
Cayenne	-51.78	5.75	200	Paramaribo	-55.21	5.81	64	ECC	379.50	9	2002-2013
Frankfurt	8.30	50.16	12742	Payenne	6.94	46.81	2673	ECC	385.72	204	2002-2020
Kuwait-City	48.01	29.52	105	Esfahan	51.43	32.48	34	ECC	463.15	17	2001-2004
Male	73.51	5.00	76	Trivandrum	76.95	8.48	45	Indian-sonde	543.73	24	1997-2000
Colombo	80.41	7.79	31	Trivandrum	76.95	8.48	37	Indian-sonde	388.49	11	1998-2000

1 **Table 2.** Bias, correlation coefficient (R), and RMSE for four types of ozonesonde and aircraft
 2 observations in four seasons.

Type	Season	Bias (O_3 -ozonesonde - O_3 -aircraft) (ppb)	R	RMSE (ppb)
ECC	Spring	17.34	0.76	65.52
	Summer	1.96	0.76	40.15
	Autumn	1.75	0.71	34.47
	Winter	7.61	0.71	51.74
Brewer-mast	Spring	10.22	0.94	43.51
	Summer	2.99	0.83	48.79
	Autumn	6.53	0.79	29.40
	Winter	6.11	0.88	45.45
Carbon-iodine	Spring	-9.19	0.84	38.34
	Summer	3.83	0.46	29.31
	Autumn	2.33	0.68	15.10
	Winter	-16.68	0.88	44.72
Indian-sonde	Spring	19.64	0.44	44.30
	Summer	19.58	0.57	37.44
	Autumn	20.38	0.45	37.30
	Winter	40.07	0.18	64.99

3
 4 **Table 3.** Bias, correlation coefficient(R) and RMSE for ECC and Indian-sonde ozonesonde and
 5 aircraft observations at different station-airport distances.

Type	Station-pair distance	Bias (O_3 -ozonesonde - O_3 -aircraft) (ppb)	R	RMSE (ppb)
ECC	<1°	9.78	0.78	47.46
	1°-2°	8.91	0.90	40.73
	2°-4°	5.65	0.67	51.00
Indian-sonde	<1°	26.71	0.37	49.54
	2°-4°	15.35	0.24	30.86

6

7 **Table 4.** Comparison of the sondes of each type to IAGOS. (average \pm 2 times the standard error
 8 (SE)) Indian-sonde/ECC is (Indian-sonde/IAGOS)/(ECC/IAGOS), Brewer-mast/ECC is (Brewer-
 9 mast/IAGOS)/(ECC/IAGOS), Carbon-iodine/ECC is (Carbon-iodine /IAGOS)/(ECC/IAGOS)

Altitude(km)	Indian-sonde/ECC	Brewer-mast/ECC	Carbon-iodine/ECC	ECC/ IAGOS
0-1	1.59 \pm 1.74	0.83 \pm 0.96	1.10 \pm 1.36	0.98 \pm 1.28
1-2	1.31 \pm 1.83	0.81 \pm 0.90	1.00 \pm 1.05	1.07 \pm 1.58
2-3	1.20 \pm 1.62	0.89 \pm 0.97	0.93 \pm 0.85	1.08 \pm 1.54
3-4	1.14 \pm 1.57	0.88 \pm 0.94	0.90 \pm 0.87	1.10 \pm 1.48
4-5	1.13 \pm 1.61	0.89 \pm 1.02	0.91 \pm 0.99	1.10 \pm 1.44
5-6	1.18 \pm 1.76	0.91 \pm 1.05	0.92 \pm 1.04	1.08 \pm 1.37
6-7	1.20 \pm 1.89	0.91 \pm 1.00	0.92 \pm 0.82	1.09 \pm 1.54
7-8	1.22 \pm 1.92	0.92 \pm 0.94	0.90 \pm 0.64	1.11 \pm 1.69
8-9	1.29 \pm 2.09	0.95 \pm 0.99	0.85 \pm 0.55	1.12 \pm 1.61
9-10	1.35 \pm 2.35	0.97 \pm 1.09	0.79 \pm 0.62	1.11 \pm 1.46
10-11	1.41 \pm 3.26	0.98 \pm 1.21	0.70 \pm 0.68	1.12 \pm 1.37
11-12	1.39 \pm 4.61	0.97 \pm 1.19	0.67 \pm 0.72	1.12 \pm 1.42

10