Tropospheric Ozone Variability during the East Asian Summer Monsoon as Observed by Satellite (IASI), Aircraft (MOZAIC) and Ground Stations

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21 Abstract

Satellite measurements from the thermal Infrared Atmospheric Sounding Interferometer (IASI), 22 aircraft data from the MOZAIC/IAGOS project, as well as observations from ground based 23 stations, are used to assess the tropospheric ozone (O_3) variability during the East Asian Summer 24 Monsoon (EASM). Six years [2008-2013] of IASI data analysis reveals the ability of the 25 instrument to detect the onset and the progression of the monsoon seen by a decrease in the 26 tropospheric [0-6] km O₃ column due to the EASM, and to reproduce this decrease from one year 27 to the other. The year-to-year variability is found to be mainly dependent on meteorology. 28 Focusing on the period of May-August 2011, taken as an example year, IASI data show clear 29 30 inverse relationship between tropospheric [0-6] km O₃ on one hand and meteorological parameters such as cloud cover, relative humidity and wind speed, on the other hand. Aircraft 31 data from the MOZAIC/IAGOS project for the EASM of [2008-2013] are used to validate the 32 IASI data and to assess the effect of the monsoon on the vertical distribution of the tropospheric 33 O_3 at different locations. Results show good agreement with a correlation coefficient of 0.73 34 (12%) between the [0-6] km O₃ column derived from IASI and aircraft data. IASI captures very 35 36 well the inter-annual variation of tropospheric O₃ observed by the aircraft data over the studied domain. Analysis of vertical profiles of the aircraft data shows a decrease in the tropospheric O₃ that is more important in the free troposphere than in the boundary layer and at [10-20]^oN than elsewhere. Ground station data at different locations in India and China show a spatiotemporal dependence on meteorology during the monsoon, with decrease up to 22 ppbv in Hyderabad, and up to 5 ppbv in the North China Plain.

42 **1. Introduction**

Over South and East Asia, tropospheric ozone (O_3) concentrations have significantly increased 43 over the past few decades as a result of rapid urbanization (Cooper et al., 2014) with important 44 implications on regional and global air quality. South and East Asian countries are experiencing 45 increasing emissions of different pollutants, many of which are precursors of O₃ (Akimoto, 2003; 46 Ohara et al., 2007; Richter et al., 2005). For example, China showed an increase of NO₂ reaching 47 29% per year for the period 1996-2006 (van der A. et al., 2008), and about 50% over the 48 industrial areas of China over the period 1996-2004 (Richter et al., 2005), though recent NO₂ 49 observations from space are suggesting NO₂ decrease in 2013 and 2014 (Richter et al., 2015). An 50 increase in background O3 concentrations is also detected in Southern China during the last 51 decade (Wang et al. 2009). In Eastern China, a study by Xu et al., (2008), analyzing long term 52 53 trends at a background surface O₃ station in Linan, suggests enhanced O₃ variability linked to the 54 increase in NO_x (NO_x=NO₂+NO) concentrations. Over most India, increasing trends in tropospheric O₃ are consistent with the observed trends in emissions from NO_x and carbon 55 monoxide (CO) as well as coal and petroleum consumption (Lal et al., 2012). 56

The Asian monsoon circulation dominates the regional meteorology of southern and East Asia. In 57 summer, the East Asian Summer Monsoon (EASM) is characterized by torrential rain, strong 58 winds carrying clean air from the ocean over the heated tropical land and deep convection 59 60 processes forming cirrus clouds and further rain (Lawrence and Lelieveld, 2010). Surface observations have shown that the EASM is responsible for a decrease in surface O₃ at a rural site 61 near Beijing (Wang et al. 2008) and a coastal site near Hong Kong (Lam et al. 2001; Wang et al. 62 2009). Yang et al. (2014) found that the largest impacts of EASM on the decrease in surface O₃ 63 are found over central and western China, while Beijing (North East China) and Nanjing (East 64 China) experience a summertime O_3 maxima during June and July respectively (Ding et al., 65 2008; 2013). Ozonesondes measurements have also detected the effect of the EASM on lower 66 tropospheric O₃ (Chan et al., 1998; Zhou et al., 2013). Satellite measurements over South and 67 East Asia have been used to assess the daily variability of tropospheric O₃, notably from the 68 69 Infrared Atmospheric Sounding Interferometer (IASI) (Dufour et al., 2015). The effect of the EASM on the tropospheric O₃ was previously detected with a decrease in the O₃ partial column 70 using data from IASI over several Indian and Chinese cities (Dufour et al., 2010; Safieddine et 71 al., 2013). Using the Ozone Monitoring Instrument (OMI) and Microwave Limb Sounder (MLS) 72 measurements, together with a regional chemistry and transport model, Zhao et al. (2010) showed 73 that the air quality over southeastern China is affected by the EASM, leading to an influence 74 extending to central East China from June to July. At 300 hPa, the Tropospheric Emission 75

Spectrometer (TES) observations over India showed enhanced O₃ abundances during June and
 July followed by a decrease in August (Worden et al., 2009).

The main objective of this study is to document the effect of the monsoon on the regional and 78 79 vertical distribution of tropospheric O₃ in East Asia during the summer using different observation datasets and relating them to one another. After this introduction, section 2 will look 80 at six years of tropospheric [0-6] km O₃ data and on a case study of the monsoon of 2011. We 81 study the relationship between the tropospheric [0-6] km O₃ and CO total columns from IASI and 82 different meteorological parameters from the ECMWF (European Centre for Medium-Range 83 Weather Forecasts) Reanalysis (ERA) (winds, cloud cover and relative humidity) (Dee et al., 84 2011). Section 3 will look at a dense set of vertical O₃ airborne profiles (363 in total) used to 85 validate the IASI-O3 columns as well assess how well IASI captures the inter-annual variability 86 of the EASM as observed by the aircraft data. Section 4 will look at a ground stations dataset at 87 five locations (Hyderabad, Udaipur, Jabalpur, Pearl River Delta (PRD) and North China Plain 88 (NCP)) during the EASM of 2011. Conclusions are given in Section 5. 89

90 **2.** Tropospheric O₃ from IASI During the EASM

91 **2.1 The IASI instrument**

92 The IASI instruments launched onboard the MetOp platforms in October 2006 (IASI-A) and September 2012 (IASI-B) are nadir looking Fourier transform spectrometers that probe the 93 Earth's atmosphere in the thermal infrared spectral range between 645 and 2760 cm⁻¹, with a 94 spectral resolution of 0.5 cm⁻¹ (apodized) and 0.25 cm⁻¹ spectral sampling. In this study, and to 95 have a consistent O₃ product over the period 2008-2013, only IASI-A data have been used. The 96 IASI footprint is a matrix of 2 x 2 pixels, each with 12 km diameter at nadir. IASI monitors the 97 atmospheric composition at any location two times per day (the satellite's ground track is at about 98 99 09:30 local time in the morning and 21:30 in the evening). Each IASI measures many of the chemical components that play a key role in the climate system and in several aspects of 100 atmospheric pollution. Global distributions of O₃ vertical profiles are retrieved in near real time 101 using a dedicated radiative transfer and retrieval software for the IASI O₃ product, the Fast 102 Optimal Retrievals on Layers for IASI (FORLI-O₃) (Hurtmans et al., 2012). Data are selected 103 using a filter for scenes with no or low cloud coverage (below 13%), and by rejecting all 104 observations with root mean square (RMS) of the spectral fit residual larger than 3.5×10^{-8} 105 W/cm².sr.cm⁻¹. Details about the chemical components that can be measured by IASI can be 106 found in Clerbaux et al. (2009); Coheur et al. (2009) and Clarisse et al. (2011). IASI tropospheric 107 108 O₃ retrieved from FORLI has been extensively validated against ozonesondes during the period 2008-2014 and results show that IASI underestimates tropospheric O₃ by 12 to 14% in the mid-109 latitudes and tropics (Boynard et al., 2016). 110

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115 **2.2 Tropospheric O₃ during the EASM for 2008-2013**

116 To look at the inter-seasonal and inter-annual variation of tropospheric O_3 during the monsoon, we show in Fig. 1 the monthly distribution (May-September) of the IASI [0-6] km O₃ column 117 over the period 2008-2013. Only IASI daytime observations are used, since better thermal 118 contrast, and hence better sensitivity to the lower troposphere, is usually obtained during the day 119 (Clerbaux et al., 2009). Earlier studies have shown that the information content in the satellite 120 measurement varies, and is generally maximal in the mid to upper troposphere, and lower at the 121 surface (Barret et al., 2011; Dufour et al., 2012; Safieddine et al., 2014). The [0-6] km O₃ column 122 is used to eliminate any possible stratospheric intrusions and previous studies have shown that 123 with the [0-6] km column, at least one piece of information is available for the IASI retrieval 124 (Barret et al., 2011; Dufour et al., 2012). Limitations in the sensitivity provide sources of error 125 126 that can influence our conclusions about the observed O₃ distribution, especially close to the surface. Another source of error emerge from the fact that the IASI observations used here are for 127 scenes (pixels) with no or low cloud contamination, and therefore reflect mostly the state of the 128 atmosphere before or after the rain/high cloud episodes that the EASM will generate. 129



Figure 1. Monthly averaged daytime tropospheric [0-6] km O₃ column from IASI over the EASM region and period (May-September) for the years 2008-2013. The decrease due to the monsoon is

- region and period (May-September) for the years 2008-2013.more prominent to the south of the domain.
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135 The monthly average tropospheric O_3 columns from IASI shown here have different observation count at each grid point, which may correspond to the average of one to more than 200 136 observations. Despite these limitations, this is the best-known dataset of remote infrared retrieved 137 O₃ columns covering the entire monsoon region, and we assume that the average effect of the 138 monsoon on the tropospheric O₃ column from one month to the other can be reflected in one or 139 more observations. The seasonal variation of O₃ as detected by IASI is highly dependent on 140 141 photochemical activity and is generally higher in summer and lower in winter (Safieddine et al., 2013). However, within the EASM region, IASI shows in Fig. 1 that this typical seasonality is 142 broken and on average, the [0-6] km O₃ columns are lower in June-July-August (JJA) in 143 comparison with May. The Asian monsoon onset date is around the mid-May and June 144 (Parthasarathy et al., 1994; Wang et al., 2009; Yang and Lau, 1998). Figure 1 shows that the 145 largest decrease is recorded in Southern India, where clean air masses from the Pacific starting 146 typically in May (generating the Indian Summer Monsoon, a subsystem of the EASM) will be 147 responsible of a decrease reaching 15-20 Dobson Unit (DU) in JJA. With the march of the 148 monsoon northeastward, the decrease becomes most prominent in July and August at higher 149 150 latitudes. To understand the year-to-year variability in tropospheric ozone, we looked at possible 151 stratospheric intrusions into the [0-6] km O₃ column. Potential vorticity (PV) and the water vapor mixing ratio (Qvap) measurements were used as markers of transport from the upper 152 troposphere-lower stratosphere (UTLS) to the troposphere, and extracted at different altitudes. 153 154 We used data from the ERA-interim reanalysis (more info about this dataset is provided in section 2.3), and it was seen that downward transport from the UTLS can have an effect at 155 altitude > 10 km and latitudes $> 30^{\circ}$ N (results not shown here). The analysis therefore suggests 156 that the [0-6] km O₃ monthly averaged columns studied here are not affected by transport from 157 the stratosphere. We also looked at the year-to-year variations associated with changes in 158 159 regional emissions to study their effect on local production of ozone. We use emission estimates of NO_x and non-methane hydrocarbons (NMVOCs), which are O₃ precursors, from EDGAR-160 HTAP project (http://edgar.jrc.ec.europa.eu/htap v2/index.php) available for only 2008 and 161 2010. An increase in both NOx and NMVOCS is observed between 2008 and 2010 over the 162 studied domain and months, except for Japan (see supplementary materials Fig. S1). Note that 163 strict controls on vehicles and industries emissions were implemented for the summer Olympic 164 and Paralympic Games in Beijing between July and September 2008 (United Nations 165 Environment Programme, 2009). Traffic was reduced by 22% during the Olympics (Wang and 166 167 Xie., 2009) and restrictions were applied on polluting industries in Beijing and surrounding 168 provinces (Li et al., 2009). While these standards did not include surface O₃ limitations, O₃ values were expected to be indirectly affected. When comparing 2008 to the rest of the years, O₃ 169 distributions do not show a significant decrease in Beijing and/or the surrounding regions (in fact 170 171 August 2010 and 2012 have lower O₃ values over Beijing than those recorded in 2008), a result

also detected in previous studies looking at O₃ surface measurements (Chou et al. 2011; Wang et 172 al. 2010). Studies suggested O₃ transport by winds from nearby polluted areas such as the North 173 174 China Plain (Wang et al., 2009, 2010), or O₃ formation in the boundary layer since the photostationary state of the nitrogen cycle was perturbed (Wang and Xie, 2009). Over the rest of 175 the domain, the increase in emissions of O₃ precursors, at least between 2008 and 2010, suggests 176 that tropospheric ozone values also increased between these two year during the different months, 177 178 but the IASI-O₃ columns do not show the same pattern. The analysis presented here suggests therefore that the meteorology associated with the EASM is the main driver of the O₃ regional 179 distribution and year-to-year variability. Over the different years, the seasonal variation is well 180 reproduced but the decrease in the tropospheric O₃ column will therefore change in magnitude 181 182 depending on the monsoon strength. For example, in June 2010, around 30°N and 120-130°E, the Western North Pacific region shows void area of IASI retrievals (with white pixels) due to large 183 cloud cover. The Western North Pacific Monsoon Index (WNPI, Wang and Fan, 1999) is the 184 highest in 2010 over this region in comparison with the rest of the years shown here. On the other 185 hand, 2011, is rather a typical monsoon season year, and will be used as a case study hereafter. 186

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188 **2.3 Case study of 2011**

In order to look at the O₃ response to change in meteorology during the monsoon, we show in 189 Fig. 2 the monsoon period [May-August] of 2011 taken as an example year of a typical monsoon. 190 191 Carbon monoxide (CO) total columns from IASI are also shown. CO is often used as an anthropogenic pollution and biomass-burning tracer (e.g. Edwards et al., 2004; McMillan et al., 192 2010). Note that the seasonal variation of CO is such that it is lower in summer, because of the 193 destruction of CO by the OH radical in the presence of sunlight. We consider different 194 195 meteorological parameters in order to highlight the relationship between change in meteorology and the [0-6] km O₃ column. These are: i) the total cloud cover that gives an insight on the 196 photochemical activity in the troposphere, ii) relative humidity, since increasing water vapor 197 increases O_3 loss as the production rate of the reactions $H_2O + O(^1D) \rightarrow 2OH$ increases (where 198 $O(^{1}D)$ is the product of the photo-dissociation of O_{3} in the presence of ultraviolet light), and iii) 199 horizontal and vertical wind fields. Horizontal wind speed and direction are used to assess 200 monsoon strength and possible transport. Vertical velocity is used to investigate possible 201 ascending motion of air masses from the boundary layer towards the free troposphere. All of the 202 meteorological parameters are extracted from the ECMWF re-analysis (Dee et al., 2011). The 203 204 data assimilation produces 4 analyses per day at 00, 06, 12 and 18 UTC at 37 pressure levels from 1000 to 1 hPa. Monthly means of total cloud cover, relative humidity at 850 hPa and u- and 205 v-components of horizontal wind directions at 850 hPa are extracted over a grid size of 0.75° x 206 0.75°. 207



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Figure 2. Monthly averaged tropospheric [0-6] km O₃ column from IASI, CO total columns from IASI, along with ECMWF total cloud cover, relative humidity, horizontal and vertical winds at 850 hPa for each of the months of May to August 2011. The black square and circle (upper right plot) are the regions the least affected by the EASM and discussed more in the text.

During the EASM, the southwesterly monsoon flow brings warm, wet and clean air masses from 213 the Indian Ocean to South, Southeast and East Asia. The winds at 850 hPa in Fig. 2 show a 214 215 typical monsoon flow where in May and June, it mostly impacts South and Southeast Asia while the coastal region of East Asia is impacted by south-easterlies from the Pacific. In June, in 216 particular, the monsoon becomes stronger as the wind force and the cloud cover (and therefore 217 lower photochemical activity) increase over the regions $< 20^{\circ}$ N. A decrease in tropospheric O₃ 218 and total CO columns is recorded over India, and over all the countries around the Bay of Bengal 219 and South East China. In Southern India, negative (ascending winds, over the Arabian Sea) and 220 positive (descending winds, over land) vertical velocities are present from the surface up to 700 221 hPa (we show here an example at 850 hPa) suggesting exchanges of air masses vertically. Since 222 CO's chemistry does not depend on cloud cover, the decrease in CO between May and June 223 suggests that transport might be the main driver of decrease in pollutants. During July and 224 August, the monsoon reaches its maximal strength. Due to high cloud cover and strong horizontal 225 winds, the tropospheric O_3 columns show a large decrease. For latitudes $< 30^{\circ}N$, the drop in O_3 is 226 227 more notable, particularly over the Indo-Gangetic Plain where the decline in O_3 is driven by 228 decreasing photochemistry since CO values over this region do not follow the same trend.

229 Looking at specific regions, tropospheric [0-6] km O₃ columns in Korea show a decrease in particular in July for O₃ (and not for CO), which coincides with the high cloud cover and relative 230 231 humidity. On the other hand, and over North West of India and part of Pakistan, the low cloud cover and weak winds lead to the buildup of the high summer O₃ over this region. Enhanced IASI 232 233 CO columns over this same region suggest build-up of pollutants, and with little to no transport, the persistence of the [0-6] km O₃ values. Looking at the winds plots over the different months, 234 one can notice how the monsoon is stronger at the lower latitudes of the domain. Therefore the 235 areas of Beijing, Tianjin and the North China Plain (black squares in Fig. 2) are in general less 236 affected by the monsoon and they show much weaker O₃ decrease. High CO total columns, used 237 238 as pollution tracer, indicate the anthropogenic origin of the observed ozone enhancements. In 239 fact, the persistence of O₃ during the monsoon season in Beijing was previously documented using aircraft data from the MOZAIC program which suggested a summertime O₃ maximum 240 attributed to strong photochemical production (Ding et al., 2008). The other interesting region in 241 China that shows little or no change is the area between the Chongqing and Sichuan provinces 242 (and designated with a black circle in Fig. 2). This region does not exhibit any monsoon 243 characteristics with low cloud cover and weak winds. This region is also between two mountains, 244 making the persistence of O₃ and CO during summer favorable. The vertical velocity plots show 245 that the monsoonal convection responsible of uplift of pollutants from the boundary layer to the 246 free and upper troposphere, is more prominent for latitudes $< 30^{\circ}$ N, except for Southern India, in 247 248 accordance with previous studies (e.g. Randel et al., 2010; Fadnavis et al., 2013, 2015).

249 **3. Tropospheric O₃ from MOZAIC/IAGOS**

The Measurements of OZone and water vapor by in-service AIrbus airCraft (MOZAIC) program currently known as the European In-service Aircraft for a Global Observing System (IAGOS) program (Nedelec et al., 2015), has provided *in-situ* observations of ozone, water vapor, carbon monoxide and other trace gases made from multiple commercial aircraft since 1994 (Marenco et al., 1998; Thouret et al., 1998 <u>http://www.iagos.org</u>). In this study, we start by performing a validation analysis of the IASI data with 363 profiles from aircraft take-off and landing, we then check how well IASI captures the inter-annual variation of O_3 seen by the aircraft data and finally we look at the monthly averaged vertical profiles of tropospheric O_3 over our study domain.



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Figure 3. In red and scaled to number of observations: location of the MOZAIC/IAGOS flight
data at the different airports in our study domain and period. The "+" sign locations corresponds
to the ground stations used in Section 4 for the year 2011.

262 **3.1 Validation of IASI data**

The IASI-retrieved tropospheric O_3 product is not a real concentration profile, but an estimation of the true profile considering the ability of the instrument to discriminate different atmospheric layers. Therefore one cannot directly compare satellite-retrieved profiles with high-resolution *insitu* observations such as the aircraft data. Instead, each high-resolution O_3 profile measured by the aircraft needs to be convolved by the low-resolution IASI averaging kernel matrix with the *apriori* profile, following Rodgers (2000) formulation:

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$$x_{smoothed} = x_{a,IASI} + A_{IASI}(x_{aircraft} - x_{a,IASI})$$
 Eq.1

Where x_{smoothed} is the smoothed profile that uses low-resolution measurement characteristics. A_{IASI} is the low-resolution averaging kernel matrix. x_{aircraft} is the high-resolution profile given by the aircraft, and $x_{a,\text{IASI}}$ is the low-resolution IASI *apriori* profile constructed from the McPeters/Labow/Logan climatology of O₃ vertical distribution, which combines long-term satellite limb measurements and measurements from ozonesondes (see McPeters, et al., 2007; Hurtmans et al. 2012). Before the smoothing, the validation profile has to cover the whole retrieval altitude range, which is from the ground up to 41 km. The aircraft data profiles were therefore completed by the same *apriori* used for IASI so that the matrix calculation of Eq. 1 is valid. We have also used a IASI spatial coincidence criterion of \pm 200 km around the position of the aircraft ascent or descent. In this section, both IASI day and nighttime observations are used for the validation of the aircraft data, and the temporal coincidence is \pm 10h.



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Figure 4. The [0–6] km O₃ columns retrieved from IASI correlation with 363 coincident MOZAIC/IAGOS profiles convolved with IASI averaging kernels for the period May–August of [2008-2013]. No aircraft data is available for 2010. Grey dots correspond to raw aircraft O₃ columns before smoothing.

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Figure 4 shows the correlation of [0–6] km O₃ column retrieved from spatio-temporal 288 coincidence of 363 IASI and MOZAIC/IAGOS smoothed profiles during May-August of [2008-289 2013] (except for 2010 where no aircraft profiles were available), and over the airports located in 290 the study domain (see Fig. 3 for the location). Over the five years, a good agreement between the 291 292 two datasets is found with correlation of 0.73 and absolute relative bias of 12%±9%. Analysis of 293 each year data leads to correlations ranging between 0.7 and 0.8 and bias ranging between 11 and 19%. Our results suggest a good ability of IASI to reproduce O_3 variability in the troposphere 294 over this region. Discrepancies arise from the spatial resolution of the IASI footprint resulting in 295 an observation averaged over tens of kilometers around the airport and therefore may include 296 other surface O₃ contributions. Moreover, the aircraft observation takes place at different times 297 during the day whereas IASI observation is at around 9:30 a.m. and 9:30 p.m. local time. With its 298 limited sampling time during the day and its lower sensitivity towards the surface, IASI 299 observation is not able to capture the diurnal variation of O₃ like an aircraft profile. Our results 300 301 show a declined correlation between IASI and aircraft products as compared to Barret et al. (2011) where they reported a correlation coefficient of 0.87 ($12\% \pm 6\%$). This could be due to the 302

different retrieval algorithm used: SOFRID (Barret et al., 2011) vs FORLI (Hurtmans et al., 303 2012). A discussion of the differences between the two algorithms can be found in Dufour et al. 304 305 (2012). Another source of difference may arise from the different season and time period studied (Barret et al. (2011) uses a 6-month profiles over the period July-December 2008). Over the 306 monsoon period, and in particular in May and June, the diurnal variability of tropospheric ozone 307 is much more pronounced and highly dependent on the local meteorology. Therefore, 308 discrepancies between IASI and the aircraft profile will carry larger discrepancies given the +/-309 10h coincidence criteria we used. Moreover, our study takes only the column from [0-6] km O₃ 310 column from IASI whereas the lower tropospheric column used by Barret et al. 2011 is based on 311 the column from the surface up to 250 hPa (~10 km) and IASI is known to have a better 312 313 sensitivity in the upper middle troposphere (Boynard et al., 2009; Safieddine et al., 2013).

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Figure 5 shows the time series of the columns plotted in Fig. 4 averaged over three latitude bands 315 (the vertical bars are the standard deviation around the mean). The IASI-O₃ product captures very 316 well the inter-annual variability of the EASM as observed by the aircraft data over the different 317 latitude bands. The decrease in the O₃ columns is most important at [10-20]^oN, and over the 318 319 different years, a result that can also be seen in Fig. 1. Less data is available at [20-30]°N (no colocated observations are recorded for 2008, 2009 and 2013). For 2011 for example, a decrease -320 321 though smaller in magnitude than the one in the [10-20]°N- is recorded both by IASI and the aircraft observations between May and June. Between July and August, a decrease is recorded in 322 323 2011, and an increase is recorded in 2012, and both events are well captured by IASI. At [30-40]^oN, the effect of the EASM on the tropospheric O₃ column, as Figs. 1 and 2 showed, is weak 324 or not clear. IASI captures very well the variability during both 2011 (the consistent slight 325 decrease between May to August 2011), and 2013 (various increasing/decreasing behavior). 326

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Figure 5. Time series of [0-6] km available O₃ columns from co-located IASI (black) and aircraft (smoothed in red and raw in grey) data at the airports (Fig.3) averaged between [10-20]°N, [20-30] °N and [30 40] °N from May till August of 2008 to 2013. No aircraft data was recorded in 2010.

333 3.2 Monsoon effect on the vertical profiles

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We show in Fig. 6 the monthly averaged (with the horizontal bars as standard deviation) raw MOZAIC profiles during the EASM of 2011 taken as an example year.



Figure 6. Monthly averaged vertical profiles of tropospheric O₃ from MOZAIC during the period May–August 2011 at [10-20]°N, [20-30]°N and [30-40]°N. Error bars correspond to the standard deviation.

At [10-20]^oN, and from May to June, a clear decrease of around 20 ppby in O₃ volume mixing 340 ratios (VMR) is recorded at 1 and 2 km. At higher altitudes up to around 7 km, an important 341 decrease is also detected from May to June and then from June to both July and August. With a 342 small standard deviation around the O₃ profiles, the observations at [10-20]^oN, suggest consistent 343 averaged monthly behavior and shows a decrease in the O₃ profile at different altitudes from 0 to 344 7 km, which was also seen over the whole [0–6] km O₃ column from IASI in Figs. 1 and 2. At 345 [20-30]^oN, the O₃ VMR in the lower troposphere (1 to 5 km, no monsoon signature at the 346 surface) decreases from May to June of 10-20 ppbv but then increases back in July and/or 347 August. At [5–8] km, the different months averaged O₃ VMR becomes comparable. Looking at 348 this latitude band in Fig. 2, we can see how the decrease in the lower tropospheric O_3 in June can 349 be explained by the increase in the cloud cover and in particular an increase in the wind speed at 350 850 hPa coming from the west Pacific. The profiles located at [30-40]°N, except at the surface, 351 show a decrease in the O₃ VMR is detected from June to July and August (till 6 km). All three 352 353 averaged profiles show no monsoon signature at the surface, probably due to enhanced O₃ precursor emissions near the ascent and descent of the aircraft. 354

Ground-based measurements 355 4

The MOZAIC profiles illustrated in Fig. 6 have shown that the effect of the EASM on 356 surface/boundary layer O₃ largely depends on the location. In this section, we investigate ground 357 station data during the monsoon of 2011 over five different locations (see Fig. 3, "+" sign). We 358 distinguish between the Indian and Chinese stations by the sampling method. The three Indian 359 stations shown here are provided with monthly means only. The two Chinese sites are a 360 collection of stations and have hourly values: the data from the Pearl River Delta is the daily 361 average of 12 stations, and the data from the North China Plain site are daily average over 7 362 stations. Details on the location, type and sampling method of the Indian and Chinese stations as 363 well as discussion of each of the Chinese station data are provided in Supplementary Materials. 364 Figure 7 shows the surface O_3 VMR for the Indian ground stations in panel *a* and for the Chinese 365 366 ground stations in panel b.







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372 Figure 7. Ground station data during the EASM of 2011 over India: Hyderabad, Udaipur and Jabalpur in panel (a); and over China, Pearl River Delta (PRD), and North China Plain (NCP) in 373 panel (b). The shaded region in panel (b) corresponds to the standard deviation of the 24-hour 374 running average of the different stations in the PRD and NCP sites respectively. The location of 375 the stations is plotted in Fig. 3, and more information is provided in Supplementary Materials. 376

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Hyderabad shows continuous decreasing O₃ values from May to August, of total magnitude of 22 378 ppbv in accordance with Fig. 6 at the same latitude band. Jabalpur is located in a region where 379 the monsoon effect is strong as seen in Figs 1 and 2, and the O₃ at the surface behaves similarly, 380 with a decrease of 20 ppby. Udaipur lies in a region where the monsoon is milder, leading to a 381

small decrease in the summer-time O₃ values that decrease of 5 ppbv between May and July and 382 then increase back in August to 32 ppby. Panel b station data are the 24-h running average (and 383 384 the associated standard deviation in shaded blue) of 12 stations in the PRD region and 7 stations in the NCP region. Since we are interested in the regional EASM effect on O₃, we show the 385 average of the stations here (the station are between 25 to 300 km away). For details and more 386 timely resolved observations for each of the station, please check the supplementary material. 387 388 The ground observations of the PRD stations, represented also by the [20-30]°N MOZAIC profiles in Fig. 6, detect a decrease of > 35 ppbv from June to July coinciding with when the 389 northwesterly winds from the Pacific become stronger (see Fig. 2). The O₃ VMR increase slightly 390 afterwards during July and August due to the decrease in monsoon strength over this region (also 391 392 seen in Fig. 2). Panel b for the NCP stations shows a weak monthly decrease in O₃ concentrations 393 from June to July and August of 5–10 ppby. The IASI and meteorological data presented in Fig. 394 2, also suggest the same decreasing pattern in O_3 concentrations driven by the slight increase in cloud cover. 395

396 397

5. Conclusions

398 The East Asian Summer Monsoon plays an important role in changing the pollutants concentration as well as the weather and climate system over the monsoon regions leading to 399 400 effects on the global air quality and climate system (Rodwell and Hoskins, 2001). The study of the dynamics and variability of the East Asian monsoon provides useful information to analyze 401 402 the distribution and losses of pollutants such as tropospheric O_3 . The latter is shown to have a particular seasonal variation over south and East Asia due to the monsoon. This study shows that 403 the monsoon variability is recorded and well captured over the different years by the infrared 404 remote sensor IASI during [2008-2013]. The IASI [0-6] km O₃ columns decrease starting the 405 months of May/June of each year, and reach a minimum in July-August. This decrease is most 406 407 prominent in south Asia where the monsoon is stronger. In order to assess the monsoon 408 meteorological signature on tropospheric O_3 , we compare tropospheric O_3 to cloud cover, relative humidity and wind fields from the ERA-interim archive during the monsoon of 2011 taken as an 409 example year. We add CO total columns from IASI to check if the regions showing persistence of 410 ozone are stimulated by the presence of anthropogenic precursors. Over most of the domain, clear 411 inverse relationship is seen between the IASI [0-6] km tropospheric O₃ on one hand and cloud 412 cover and winds on the other hand. This explained by the fact that the high cloud cover that the 413 monsoon generates, accompanied with high relative humidity in the troposphere lead to a lower 414 production rate since the photochemical activity will be much lower and relative humidity is a 415 sink of O_3 in particular in the background troposphere. On the other hand, the winds are also 416 strong during the monsoon and O₃ during this period can be transported either vertically, or 417 418 horizontally to/from the Pacific and to the globe. Validation of the IASI-O₃ columns with aircraft data shows a good correlation (r=0.73 (12%)) between the [0-6] km columns from IASI to that 419 420 derived from 363 aircraft profiles. IASI is shown to reproduce very well the year-to-year variability in O_3 seen by the aircraft data. A monsoon signature is detected on the O_3 profiles over 421

the different latitude bands of our study domain that is in agreement with the IASI [0-6] km O_3 column spatial distribution. Ground station measurements of O_3 also show spatial dependence and are anti-correlated to the EASM strength. The stations in central and north India as well as in Pearl River Delta show a signature of the monsoon on ground O_3 that can be explained by the change in meteorology over these regions. In northwest India, and the North China plain, winds are weaker, and the cloud cover is smaller and thus the ground O_3 shows a smaller decrease during the EASM.

More generally, this study shows that in lack of good coverage of ground and aircraft measurements, satellite observations, such as the one provided in this work using IASI, provide valuable information that could help investigating the regional and global monsoon impact on the distribution of pollutants in the troposphere and in particular tropospheric O₃. With IASI-C to be launched in 2018 onboard MetOp-C and a new series of three IASI-NG instruments on MetOp Second Generation Satellites around 2022 (Crevoisier et al., 2014), long term IASI O₃ time series obtained at high spatial resolution will be available to characterize and detect the onset and the

436 progression of the monsoon.

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