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XCO<sub>2</sub> in an emission hot-spot region: the COCCON Paris campaign 2015





during a field campaign in spring of 2015. Here, we describe and analyze the variations of
XCO<sub>2</sub> observed at different sites and how they changed over time. We find that observations
upwind and downwind of the city centre differ significantly in their XCO<sub>2</sub> concentrations,
while the overall variability of the daily cycle is similar, i.e., increasing during night-time with
a strong decrease (typically 2-3 ppm) during the afternoon.
An atmospheric transport model framework (CHIMERE-CAMS) was used to simulate XCO<sub>2</sub>

44 and predict the same behaviour seen in the observations, which supports key findings, e.g. 45 that even in a densely populated region like Paris (over 12 Million people), biospheric uptake of CO<sub>2</sub> can be of major influence on daily XCO<sub>2</sub> variations. Despite a general offset between 46 47 modelled and observed XCO<sub>2</sub>, the model correctly predicts the impact of the meteorological 48 parameters (e.g. wind direction and speed) on the concentration gradients between different 49 stations. Looking at the local gradients of XCO<sub>2</sub> for upwind and downwind station pairs, 50 which is less sensitive to changes in XCO<sub>2</sub> regional background conditions, we find the 51 model-data agreement significantly better. Our modelling framework indicates that the local 52 XCO<sub>2</sub> gradient between the stations is dominated by the fossil fuel CO<sub>2</sub> signal of the Paris 53 metropolitan area. This highlights the usefulness of XCO<sub>2</sub> observations to help optimise 54 future urban GHG emission estimates.

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### 56 1 Introduction

Atmospheric background concentrations of CO<sub>2</sub> measured since 1958 in Mauna Loa, USA, 57 58 have passed the symbolic milestone of 400 ppm (monthly mean) as of 2013 [Jones 2013]. 59 Properly quantifying fossil fuel CO<sub>2</sub> emissions (FFCO<sub>2</sub>) can contribute to define effective 60 climate mitigation strategies. Focussing our attention on cities is a critical part of this 61 endeavour as emissions from urban areas are currently estimated to represent from 53 % to 62 87 % of global FFCO<sub>2</sub>, depending on the accounting method considered, and are predicted to increase further [IPCC-WG3 2014, IEA 2008, Dhakal 2009]. As stated in the IPCC 5th 63 64 assessment report, "current and future urbanisations trends are significantly different from 65 the past" and "no single factor explains variations in per-capita emissions across cities and 66 there are significant differences in per capita greenhouse gas (GHG) emissions between cities within a single country" [IPCC-WG3 2014]. The large uncertainty of the global 67 68 contribution of urban areas to CO<sub>2</sub> emissions today and in the future is why a new 69 generation of city-scale observing and modelling systems are needed.

In recent years, more and more atmospheric networks have emerged that observe GHG concentrations using the atmosphere as a large-scale integrator, for example in Paris (France), Indianapolis (USA), and Toronto (Canada). The air measured at in-situ groundbased stations is considered to be representative of surface CO<sub>2</sub> fluxes of a larger surrounding area (1 km<sup>2</sup>-10000 km<sup>2</sup>) [e.g., Bréon et al. 2015, Xueref-Remy et al, 2018,





75 Turnbull et al. 2015, Vogel et al. 2013]. If CO2 measurements are performed both up-wind 76 and downwind of a city, the concentration gradient between the two locations is influenced 77 by the local net emission strength between both sites and atmospheric mixing [Xueref-Remy 78 et al, 2018, Bréon et al. 2015, Turnbull et al. 2015]. To derive quantitative flux estimates, 79 measured concentration data are typically assimilated into numerical atmospheric transport 80 models which calculate the impact of atmospheric mixing on concentration gradients for a 81 given flux space-time distribution. Such a data assimilation framework implemented for Paris 82 with three atmospheric CO<sub>2</sub> measurement sites [Xueref-Remy et al, 2018] previously allowed deriving quantitative estimates of monthly emissions and their uncertainties over one year 83 84 [Staufer et al. 2016].

85 Space-borne measurements of the column-average dry air mole fraction of CO<sub>2</sub> (XCO<sub>2</sub>) are 86 increasingly considered for the monitoring of urban CO<sub>2</sub>. This potential was shown with 87 OCO-2 and GOSAT XCO<sub>2</sub> measurements, even though the spatial coverage and temporal 88 sampling frequency of these two instruments were not optimized for FFCO<sub>2</sub> [Kort et al., 89 2012, Janardanan et al. 2016, Schwandner et al. 2017], while other space-borne sensors 90 dedicated to FFCO<sub>2</sub> and with an imaging capability are in preparation [O'Brien et al, 2016, 91 Broquet et al. 2017]. Important challenges of satellite measurements are that they are not as 92 accurate as in-situ ones, having larger by systematic errors, while the XCO<sub>2</sub> gradients in the 93 column are typically 7-8 times smaller than in the boundary layer. Another difficulty of space-94 borne imagery with passive instruments is that they will only sample city XCO<sub>2</sub> plumes 95 during clear sky conditions at around mid-day.

96 The recent development of a robust portable ground-based FTIR (Fourier Transform 97 InfraRed) spectrometer as described in Gisi et al. [2012] and Hase et al. [2015] (EM27/SUN, 98 Bruker Optik, Germany) greatly facilitates the measurement of XCO<sub>2</sub> from the surface, with 99 better accuracy than from space and with the possibility of continuous daytime observation 100 during clear sky conditions. Typical compatibility (uncorrected bias) of the EM27/SUN 101 retrievals of the different instruments in a local network is better than 0.01 % (i.e. ` 0.04 ppm) 102 after a careful calibration procedure and a harmonized processing scheme for all 103 spectrometers [Frey et al. 2015]. The Collaborative Carbon Column Observing Network (COCCON) [Frey et al. 2018] intends to offer such a framework for operating the 104 105 EM27/SUN. This type of spectrometers therefore represents a remarkable opportunity to 106 document XCO<sub>2</sub> variability in cities as a direct way to estimate FFCO<sub>2</sub> [Hase et al. 2015] or 107 in preparation of satellite missions.

108 When future low-Earth-orbit operational satellites with imaging passive spectrometers of 109 suitable capabilities to invert FFCO<sub>2</sub> will sample different cities, this will likely be limited to 110 clear sky conditions and at a time of the day close to local noon. Increasing the density of 111 the COCCON network around cities will allow to evaluate those XCO<sub>2</sub> measurements and to





monitor XCO<sub>2</sub> during the early morning and afternoon periods, which will not be sampled
with satellites, except from geostationary orbit [Butz et al., 2015].

- 114 This study focuses on the measurements of XCO<sub>2</sub> from ground based EM27/SUN 115 spectrometers deployed within the Paris metropolitan area during a field campaign in the 116 spring of 2015, and modelling results. This campaign can be seen as a demonstration of the 117 COCCON network concept applied to the quantification of an urban FFCO<sub>2</sub> source. Several 118 spectrometers were operated by different research groups, while closely following the 119 common procedures suggested by Frey at al. [2015]. The paper is organised as follows. 120 After the instrumental and modelling setup descriptions of section 2, the observations of the 121 field campaign and the modelling results will be presented in section 3. Results are 122 discussed in section 4 together with the study conclusions.
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#### 124 2 Methods and materials

#### 125 2.1 Description of study area and field campaign design

- During the COCCON field campaign (April 28th to May 13th, 2015) five portable FTIR spectrometers (EM27/SUN, Bruker Optik, Karlsruhe, Germany) were deployed in the Parisian region (administratively known as *Île-de-France*) and within the city of Paris. The campaign was conducted in early spring as the cloud cover is typically low in April and May and the time between sunrise and sunset is more than 14 hours.
- 131 The Paris metropolitan area houses over 12 million people, with about 2.2 million inhabiting the city of Paris. This urban region is the most densely populated in France with ~1000 132 inhabitants/km<sup>2</sup> and over 21000 inhabitants/km<sup>2</sup> for the city of Paris itself [INSEE 2016 -133 134 https://www.insee.fr/fr/statistiques]. The estimated CO2 emissions from the metropolitan 135 region are 39 Mt/year with on-road traffic emissions and residential and the tertiary (i.e. 136 commercial) sector as main sources (accounting for over 75 %), and minor contributions 137 from other sectors such as industrial sources and airports [AIRPARIF 2016]. It was crucial to understand the spatial distribution of these CO<sub>2</sub> sources to optimally deploy the COCCON 138 139 spectrometers. To this end a 1 km emission model for France by IER (Institut fuer 140 Energiewirtschaft und Rationelle Energieanwendung, University of Stuttgart, Germany) was used as a starting point [Latoska 2009]. This emission inventory is based on the available 141 142 activity data such as, e.g., traffic counts, housing statistics, or energy use, and the temporal disaggregation was implemented according to Vogel et al. [2013]. In brief, the total 143 144 emissions of the IER model were re-scaled to match the temporal factors for the different 145 emission sectors according to known national temporal emission profiles.
- To quantify the impact of urban emissions on XCO<sub>2</sub>, the FTIR instruments were deployed along the dominant wind directions in this region in spring, i.e., southwesterly [Staufer et al 2016], in order to maximize the likelihood to capture upwind and downwind air masses (see





Figure 1). The two southwesterly sites (GIF and RES) are located in a less densely populated area, where emissions are typically lower than in the city centre, where the station JUS is located. The data in Fig. 1 show that the densest FFCO2 emission area extends northwards and eastwards. The two Northwesterly sites (PIS and MIT) were placed downwind of this area. All instruments were operated manually and typically started for operations around 7-8 am local time from which they continuously observe XCO<sub>2</sub> until 5-6

### 156 2.2 Instrumentation, calibration, and data processing

The EM27/SUN is a portable FTIR spectrometer which has been described in detail in, e.g., 157 158 Gisi et al. [2012] and Frey et al. [2015]. Here, only a short overview is given. The centre 159 piece of the instrument is a Michelson interferometer which splits up the incoming solar 160 radiation into two beams. After inserting a path difference between the beams, the partial 161 beams are recombined. The modulated signal is detected by an InGaAs detector covering 162 the spectral domain from 5000 to 11000 cm<sup>-1</sup> and is called an interferogram. As the 163 EM27/SUN analyzes solar radiation, it can only operate in daylight sunny conditions. A Fourier transform of the interferogram generates the spectrum and a DC correction is 164 165 applied to remove the background signal and only keep the AC signal (see Keppel-Aleks et 166 al. [2007]). A numerical fitting procedure (PROFFIT code) [Schneider and Hase et al., 2009] 167 then retrieves column abundances of the concentrations of the observed gases from the spectrum. The single-channel EM27/SUN is able to measure total columns of O<sub>2</sub>, CO<sub>2</sub>, CH<sub>4</sub> 168 and H<sub>2</sub>O.The ratio over the observed O<sub>2</sub> column, assumed to be known and constant, 169 delivers the column-averaged trace gas concentrations of XCO2, XCH4 in µmol / mol dry air, 170 171 with a temporal resolution of one minute. XCO<sub>2</sub> is the dry air mole fraction of CO<sub>2</sub>, defined as 172  $XCO_2$  = Column[CO<sub>2</sub>] / Column[Dry Air]. Applying the ratio over the observed oxygen (O<sub>2</sub>) 173 column reduces the effect of various possible systematic errors; see Wunch et al. (2011).

174 In order to correctly quantify small differences in XCO<sub>2</sub> columns between Paris city upstream 175 and downstream locations, measurements were performed with the five FTIR instruments 176 side by side before and after the campaign, as we expect small calibration differences 177 between the different instruments due to slightly different alignment for each individual 178 spectrometer. These differences are constant over time and can be easily accounted for by 179 applying a calibration factor for each instrument. Previous studies showed that the 180 instrument specific corrections are well below 0.1 % for XCO<sub>2</sub> [Frey et al. 2015, Chen et al. 181 2016] and are stable for individual devices. The 1-sigma precision for  $XCO_2$  is in the order of 182 0.01 % - 0.02 % (< 0.08 ppm) e. g. [Gisi et al. 2012, Chen et al. 2016, Hedelius et al. 2016, 183 Klappenbach et al. 2015]. The calibration measurements for this campaign were performed 184 in Karlsruhe w.r.t. the Total Carbon Column Observing Network (TCCON) [Wunch et al. 2011] spectrometer at the Karlsruhe Institute of Technology (KIT), Germany for 7 days 185





before the Paris campaign between April 9<sup>th</sup> and 23<sup>rd</sup>, and after the campaign on May 18<sup>th</sup>
 until 21<sup>st</sup>.

188 Figure 2 (left panel) shows the XCO<sub>2</sub> time series of the calibration campaign, where small 189 offsets between the instruments raw data are visible. As these offsets are constant over 190 time, a calibration factor for each instrument can be easily applied; actually these are the 191 calibration factors previously found for the Berlin campaign [Frey et al. 2015]. These factors 192 are given in Table 2, where all EM27/SUN instruments are scaled to match instrument No. 1. The calibrated XCO<sub>2</sub> values for April 15<sup>th</sup> are shown in Fig. 2 (right panel). None of the five 193 194 instruments that participated in the Berlin campaign show any significant drift; in other words, 195 the calibration factors found one year before were still applicable. This is an impressive 196 demonstration of the instrument stability stated in section 2.2, especially as several 197 instruments (Nos. 1, 3, 5) were used in another campaign in Northern Germany in the 198 meantime. The EM27/SUN XCO<sub>2</sub> measurements can be made traceable to the WMO 199 international scale for in-situ measurements by comparison with measurements of a 200 collocated spectrometer from the TCCON. TCCON instruments are calibrated against in-situ 201 standards by aircraft and aircore measurements [Wunch et al. 2010, Messerschmidt et al. 202 2012] performed using the WMO scale.

During the campaign and for the calibration measurements we recorded double-sided interferograms with 0.5 cm<sup>-1</sup> spectral resolution. Each measurement of 58 s duration consisted of 10 scans using a scanner velocity of 10 kHz. For precise timekeeping, we used GPS sensors for each spectrometer.

207 In-situ surface pressure data used for the analysis of the calibration measurements 208 performed at KIT have been recorded at the co-located meteorological tall tower. During the 209 campaign, a MHD-382SD data-logger recorded local pressure, temperature and relative 210 humidity at each station. The analysis of the trace gases from the measured spectra for the 211 calibration measurements has been performed as described by Frey et al. [2015]. For the 212 campaign measurements we assume a common vertical pressure-temperature profile for all 213 sites, provided by the model, so that the surface pressure at each spectrometer only differs 214 due to different site altitudes. The 3-hourly temperature profile from the European Centre for Medium-Range Weather Forecasts (ECMWF) operational analyses interpolated for site JUS 215 216 located in the centre of the array was used for the spectra analysis at all sites. The individual 217 ground-pressure was derived from site altitudes and pressure measurements performed at 218 each site.

219 Before and after the Paris campaign, side by side comparison measurements were 220 performed with all 5 EM27/SUN spectrometers and the TCCON spectrometer operated in 221 Karlsruhe at KIT. All spectrometers were placed on the top of the IMK office building North of 222 Karlsruhe. The altitude is 133 m above sea level (a.s.l.), coordinates are 49.09° N and 8.43°





E. The processing of the Paris raw observations (measured interferograms) were performed as described by Gisi et al. [2012] and Frey et al. [2015] for the Berlin campaign: spectra were generated applying a DC correction, a Norton-Beer medium apodization function and a spectral resampling of the sampling grid resulting from the FFT on a minimally sampled spectral grid. PROFFWD was used as the radiative transfer model and PROFFIT as the retrieval code.

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### 230 2.3 Atmospheric transport modelling framework

231 We used the chemistry transport model CHIMERE (Menut et al., 2013) to simulate CO2 232 concentrations in the Paris area. More specifically, we used the CHIMERE configuration 233 over which the inversion system of Bréon et al. [2015] and Staufer et al. [2016] was built to 234 derive monthly to 6-hour mean estimates of the CO<sub>2</sub> Paris emissions. Its horizontal grid, and 235 thus its domain and its spatial resolution, are illustrated in Figure 3. It has a  $2 \times 2$  km<sup>2</sup> spatial 236 resolution for the Paris region, and 2 × 10 km<sup>2</sup> and 10 × 10 km<sup>2</sup> spatial resolutions for the 237 surroundings. It has 20 vertical hybrid pressure-sigma (terrain- following) layers that range 238 from the surface to the mid-troposphere, up to 500 hPa. It is driven by operational 239 meteorological analyses of the ECMWF Integrated Forecasting System, available at an approximately 15 × 15 km<sup>2</sup> spatial resolution and 3 h temporal resolution. 240

In this study the CO<sub>2</sub> simulations are based on a forward run over April 25<sup>th</sup> - May 12<sup>th</sup> 2015 241 242 with this model configuration; we do not assimilate atmospheric CO<sub>2</sub> data and so no inversion for surface fluxes was conducted. In the Paris area (the Île-de-France 243 244 administrative region), hourly anthropogenic emissions are given by the IER inventory, see 245 section 2.1. The anthropogenic emissions in the rest of the domain are prescribed from the 246 EDGAR V4.2 database for the year 2010 at 0.1° resolution [Olivier and Janssens-Maenhout 247 et al., 2012]. In the whole simulation domain, the natural fluxes (the Net Ecosystem 248 Exchange: NEE) are prescribed using simulations of C-TESSEL, which is the land-surface component of the ECMWF forecasting system [Boussetta et al., 2013], at a 3 hourly and 15 249 250 × 15 km<sup>2</sup> resolution. Finally, the CO<sub>2</sub> boundary conditions at the lateral and top boundaries of 251 the simulation domain and the simulation CO2 initial conditions on April 25th 2015 are prescribed using the CO<sub>2</sub> forecast issued by the Copernicus Atmosphere Monitoring Service 252 253 (CAMS, http://atmosphere.copernicus.eu/) at a ~15 km global resolution [Agustí-Panareda

254 et al., 2014].

The CHIMERE transport model is used to simulate the  $XCO_2$  data. However, since the model does not cover the atmosphere up to its top, the  $CO_2$  fields from CHIMERE are complemented with that of the CAMS  $CO_2$  forecasts from 500 hPa to the top of the atmosphere to derive total column concentrations. The derivation of modelled  $XCO_2$  at the





259	sites, involves obtaining a kernel-smoothed CO <sub>2</sub> profile of CHIMERE and CAMS and vertical					
260	integration of these smoothed profiles, weighted by the pressure at the horizontal location of					
261	the sites.					
262	The parametrisation used to smooth modelled CO2 profiles approximates the sensitivity of					
263	the EM27/sun $CO_2$ retrieval is a function of pressure and sun elevation. Between 1000 hPa					
264	and 480 hPa, a linear dependency of the instrument averaging kernels on solar zenith angle					
265	$(\Theta)$ is assumed with boundary values following Frey et al. [2015]:					
266						
267	(1a) $k(480 hPa) = 1.125$					
268	(1b) $k(1000 hPa) = 1.0 + 0.45 s^3$					
269						
270	where s=0/90°. k.Approximate averaging kernels are obtained bylinear interpolation to the					
271	pressure levels of CHIMERE and CAMS, respectively. If $p > 1000$ hPa, k is linearly					
272	extrapolated. Above 480 hPa ( $p$ < 480 hPa), the averaging kernels can be approximated by					
273						
274	(2) $k(u,s) = 1.125 - 0.6 u^3 - 0.4 u s^3$					
275						
276	where <i>u</i> is $(480 hPa - p) / 480$ . The kernel-smoothed CO <sub>2</sub> profile, CO <sub>2_model</sub> <sup>s</sup> , is obtained by					
277						
278	(3) $CO_{2\_model}^{s} = \mathbf{K} CO_{2\_model} + (\mathbf{I} - \mathbf{K}) CO_{2}^{a}$					
279						
280						
281	where $CO_{2\_model}$ is the modelled $CO_2$ profile by CHIMERE or CAMS, <i>I</i> the identity matrix and					
282	K is a diagonal matrix containing the averaging kernels k. The a priori CO <sub>2</sub> profile, CO <sub>2</sub> <sup>a</sup> , is					
283	provided by the Whole Atmosphere Community Climate Model (WACCM) model (version					
284	6) and interpolated to the pressure levels of CHIMERE and CAMS. $CO_{2\_model}^{s}$ is the					
285	appropriate CO2 profile to calculate modelled XCO2 at the location of the sites.					
286						
287	For a given site, the simulated XCO <sub>2</sub> data are thus computed from the vertical profile of this					
288	site as:					
289						
290	(3) $XCO_{2\_CHIMERE} = \frac{1}{P_{surf}} \int_{Psurf}^{Ptop\_CHIM} CO_{2\_CHIM}^{s} dp + \int_{Ptop\_CHIM}^{P=0mbar} CO_{2\_CAMS}^{s} dp$					
291						
292	where $p_{surf}$ is the surface pressure, $p_{top_CHIM}$ = 500 hPa the pressure corresponding to the top					
293	boundary of the CHIMERE model, and $\text{CO}_{2\_C\text{HIM}}{}^{s}$ and $\text{CO}_{2\_C\text{AMS}}{}^{s}$ are the smoothed $\text{CO}_{2}$					





294 concentrations of CHIMERE and CAMS respectively. For comparison we also calculated

295 XCO<sub>2</sub> at a lower spatial resolution with the CAMS data alone as:

- 296
- 297

(4)

 $XCO_{2\_CAMS} = \int_{Psurf}^{P=0mbar} CO_{2\_CAMS}^{s} dP$ 

- 298
- 299 3 Results and discussion
- 300 3.1 Observations

# 301 **3.1.1** Meteorological conditions and data coverage/instrument performance.

During the measurement campaign (April 28<sup>th</sup> until May 13<sup>th</sup>, 2015), meteorological 302 303 conditions were a major limitation for the availability of XCO<sub>2</sub> observations. Useful 304 EM27/SUN measurements require direct sunlight and low wind speeds typically yield higher 305 local XCO2. Most of the time during the campaign conditions were partly cloudy and turbid, 306 and so successful measurements at high solar zenith angle (SZA) were rare. Therefore, the data coverage between April 28<sup>th</sup> and May 3<sup>rd</sup> is limited (see Table 3). As is typical for spring 307 periods in Paris, the temperature and the wind direction vary and display less synoptic 308 309 variations than in winter. The dominant wind directions were mostly northeasterly at the 310 beginning of the campaign and mostly southeasterly during the second half of the campaign. 311 We find that the wind speeds during daytime nearly always surpass 3 m s<sup>-1</sup>, which has been identified by Breon et al. [2015] and Staufer et al. [2016] as the cut-off wind speed above 312 313 which the atmospheric transport model CHIMERE performs best in modelling CO<sub>2</sub> 314 concentration gradients in the mixed layer.

Despite some periods with unfavourable conditions, more than 10,000 spectra were retrieved among the five deployed instruments. The quality of the spectra for each day was rated according to the overall data availability. The best measurement conditions prevailed for the period between May 7<sup>th</sup> and May 12<sup>th</sup>.

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# 320 3.1.2 Observations of XCO<sub>2</sub> in Paris

321 The observed XCO<sub>2</sub> in the Paris region for all sites (10415 observations) ranges from 397.27 322 to 404.66 ppm with a mean of 401.26 ppm (a median of 401.15 ppm) and a standard 323 deviation of 1.04 ppm for 1-minute averages. We find that all sites exhibit very similar diurnal behaviours with a clear decrease of XCO<sub>2</sub> during daytime and a noticeable day to day 324 325 variability as seen in Figure 4. This is to be expected as they are all subject to very similar 326 atmospheric transport in the boundary layer height and to similar large-scale influences, i.e., 327 surrounding with stronger natural fluxes or air mass exchange with other regions at synoptic 328 time scales. However, observed XCO<sub>2</sub> concentrations at the upwind sites for our network 329 remain clearly higher from sites that are downwind of Paris (see Figure 4). This indicates





that the influence of urban emissions is detectable with this network configuration under favourable meteorological conditions. By comparing the different daily variations in Fig. 5, it is apparent that the day-to-day variations observed at the two southwesterly (typically upwind) sites GIF and RES are approximately 1 ppm, with both sites exhibiting similar diurnal variations throughout the campaign period. This can be expected as their close vicinity would suggest that they are sensitive to emissions from similar areas and to concentrations of air masses arriving from the southwest.

The typical decrease in XCO<sub>2</sub> found over the course of a day is about 2 to 3 ppm. This decrease can only be driven by (natural) sinks of CO<sub>2</sub>, which can be expected to be very strong as our campaign took place after the start of the growing season in Europe for most of southern and central Europe [Roetzer and Chmielewski 2001].

341 The observations at the site located in Paris (JUS) displays similarly low day-to-day 342 variations and a clear decrease in XCO<sub>2</sub> over the course of the day. The latter feature 343 indicates that even in the dense city centre, XCO<sub>2</sub> is primarily representative of a large 344 footprint like in other areas of the globe [Keppel-Aleks, 2011] and is not as critically affected 345 by local emissions than in-situ measurements [Breon et al. 2015, Ammoura et al. 2016]. It is also apparent that the decrease in XCO<sub>2</sub> (the slope) during the afternoon for April 28<sup>th</sup> and 346 29<sup>th</sup> as well as May 7<sup>th</sup> and 10<sup>th</sup> is noticeably smaller than at other days during this 347 348 campaign. As XCO<sub>2</sub> is not sensitive to vertical mixing, this has to be caused by different CO<sub>2</sub> 349 sources and sinks acting upon the total column arriving at JUS.

The two northeasterly (typically downwind) sites PIS and MIT show a markedly larger day-today spread in the background as well as strongly changing slopes for the diurnal XCO<sub>2</sub> decrease. For these sites the exact wind direction is critical as they can be downwind of the city centre that has a much higher emission density or less dense suburbs (see Fig. 1).

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### 355 3.1.3 Gradients in observed XCO<sub>2</sub>

356 In order to focus more on the impact of local emissions and less on that of background influences in our analysis of XCO<sub>2</sub>, we choose to study the spatial gradients ( $\Delta$ ) between 357 different sites. Ideal conditions were sampled during May 7<sup>th</sup>, with predominantly 358 southwesterly winds, and on May 10<sup>th</sup> with southerly winds. We can see in Fig. 6 that all 359 360 sites are, on average, elevated compared to RES, chosen as reference here as it was upwind of Paris during the measurement period. The observations from GIF only show 361 362 minimal differences with RES, and  $\Delta$  increases between PIS, JUS and MIT and RES to 363 reach 1 to 1.5 ppm. During southwesterly winds, MIT is downwind of most of the densest 364 part of the Paris urban area, and JUS is impacted by emissions of neighborhoods to the 365 southwest. The site of PIS is still noticeably influenced by the city centre but, as can be seen 366 in Fig. 1, we likely do not catch the plume of the most intense emissions but rather from the





suburbs. On May 10<sup>th</sup>, with its dominant southerly winds, the situation is markedly different. While GIF is still only slightly elevated, the XCO<sub>2</sub> enhancement at MIT is significantly lower and quite similar to JUS for large parts of the day. The highest  $\Delta$ XCO<sub>2</sub> can now be observed at PIS, again typically ranging from 1 to 1.5 ppm. As seen in Fig. 1, PIS is now directly downwind of the densest emission area, while MIT is only exposed to CO<sub>2</sub> emissions from the eastern outskirts of Paris.

373 It is also important to note that the impact of the local biosphere that is assumed to cause 374 the strong decrease in  $XCO_2$  during the day is not seen on both days for these spatial 375 gradients. For a more comprehensive interpretation of these observations the use of a 376 transport model (as described in section 2.3) is necessary.

377

### 378 3.2 Modelling

### 379 3.2.1 Model performance

380 Before interpreting the modelled XCO<sub>2</sub> we need to evaluate the performance of the chosen 381 atmospheric transport model framework as described in section 2.3. Comparing it to meteorological observations (wind speed and wind direction) at GIF in Fig. 7, we find that 382 383 CHIMERE predicts these variables well throughout the duration of the campaign. Changes in 384 wind speed direction and speed are reproduced with a slight overestimation at low wind-385 speeds (>1m/s). Besides the meteorological forcing, the model performance can also be 386 expected to depend on the chosen model resolution. Therefore, we compared XCO<sub>2</sub> at JUS 387 calculated based on the coarser resolution atmospheric transport and flux framework CAMS 388 (15 km), and the higher resolution emission modelling input for the framework based on 389 CHIMERE (2 km) for the inner domain and on CAMS boundary conditions (see Fig. 8). We 390 find that the coarser model displays similar inter-daily variations, but that the high-resolution 391 model modifies the modelling results on shorter time-scales. We find that the afternoon 392 XCO<sub>2</sub> decreases are often more pronounced in CHIMERE. Only the high-resolution will be 393 considered and referred to in the following.

394

The impact of using different flux maps (fossil fuel  $CO_2$  and biosphere models) on the modelled  $XCO_2$  can unfortunately not be explicitly investigated here as only one highresolution (1 km) emission product was available for this study (see section 2.3).

398

### 399 3.2.2 Modelled XCO<sub>2</sub> and its components

The modelled XCO<sub>2</sub> for the five sites (Fig. 9) co-evolves over the period of the campaign with occurences of significant differences. This was already seen with the measurements, but the model allows looking at the full time series. The model reveals clear daily cycles of XCO<sub>2</sub>, with an accumulation during night-time and a decrease during daytime. Despite a good





general agreement of modelled XCO<sub>2</sub> at all sites for, e.g., the timing of daily minima and
their synoptic changes, differences in XCO<sub>2</sub> are observed between the sites for many days.
Typically the northeasterly sites (PIS, MIT) show an enhancement in XCO<sub>2</sub> compared to the
southwesterly sites (GIF, RES).

408 To understand the synoptic and diurnal variations of XCO<sub>2</sub>, we analyzed the contribution of 409 different sources (and sinks) of CO<sub>2</sub>, namely the net ecosystem exchange (NEE), the fossil 410 fuel CO<sub>2</sub> emissions (FFCO2), and the boundary conditions (BC), i.e., the variations of CO<sub>2</sub> 411 not caused by fluxes within our domain (the example of JUS is given in Fig. 10; see the 412 supplement for the other sites). The day-to-day variability of modelled XCO<sub>2</sub> is dominated by 413 changing boundary conditions and coincides with synoptic weather changes. The strong 414 daily decrease in XCO<sub>2</sub> can be directly linked to NEE, which leads to a decrease of  $\sim$ 1 ppm 415 (but up to 4 ppm) during the day. The XCO<sub>2</sub> from fossil fuel emissions causes significant 416 enhancements compared to the background, but is often compensated by NEE. During short 417 periods, fossil fuel emissions can however lead to enhancements of up to 4 ppm.

418

### 419 **3.2.3 Modelled** Δ**XCO**<sub>2</sub> gradients and its components

To be able to assess the impact of local sources and reduce the influence of NEE and BC on 420 421 the modelled signals, we analyse the XCO<sub>2</sub> gradient (i.e. station-to-station difference) with 422 RES being taken as reference. In Fig. 11 we compare  $\Delta$  along a south-north direction. For 423 the modelled  $\Delta$  we can see that MIT shows a positive value during the campaign period 424 whenever the predominant wind direction was southwesterly (grey shaded areas). We also 425 find that  $\Delta$  between JUS and RES was both negative and positive during the campaign, and 426 predominantly negative between MIT and JUS. When split into FFCO2, BC and NEE 427 components, we can clearly see that the total  $\Delta$  is dominated by FF causing XCO<sub>2</sub> offsets of 428 up to 4 ppm, but more typically 1 ppm gradients are observed. Gradients can also change 429 rapidly (within a few hours) if the wind direction changes, for example on May 1<sup>st</sup> and May 430 12<sup>th</sup>. This highlights the fact that, during such conditions, we cannot assume a simple 431 upwind-downwind interpretation of our sites. As expected, the contributions from BC and 432 NEE are generally greatly reduced when analysing ∆XCO<sub>2</sub>. The most important impact of NEE on the XCO<sub>2</sub> gradients of -1ppm and +1ppm can be seen on May 8<sup>th</sup> and May 11<sup>th</sup>, 433 respectively. This means that, despite greatly reducing the impact of NEE on average, the 434 435 contribution of NEE cannot be fully ignored. BC is an overall negligible contribution to  $\Delta XCO_2$ , even though it reaches -0.4 ppm on May 11<sup>th</sup>. 436

437

#### 438 3.3 Model data and observations comparison

439 3.3.1 XCO<sub>2</sub>





440 A comparison of modelled and observed XCO<sub>2</sub> is of course limited to the relatively short 441 periods when observations are available. Over these periods. we observe that the main 442 features of the modelled and observed XCO<sub>2</sub> are comparable. The daytime variations are 443 well reproduced by the model and the general relative concentrations between sites are preserved, e.g., the highest values for XCO<sub>2</sub> at MIT are on May 9<sup>th</sup> and highest XCO<sub>2</sub> for PIS 444 are later on May 10<sup>th</sup> and May 11<sup>th</sup> as seen in Figure 12. We can also see a general issue in 445 446 reproducing the background XCO<sub>2</sub> for each day in the model as observed XCO<sub>2</sub> is 447 significantly lower by typically between 1 to 2 ppm. We also see that the timing of the daily minima is not fully covered in the observed data as it typically happens after sunset and 448 449 cessation of biosphere uptake.

450

### 451 3.3.2 ΔXCO<sub>2</sub>

452 Due to the prevailing southeasterly wind conditions, we can compare XCO<sub>2</sub> at the typical 453 downwind sites (PIS, MIT) relative to the mostly upwind sites (RES, GIF) and expect 454 elevated XCO<sub>2</sub> downwind. We find that the  $\Delta$ XCO<sub>2</sub> of PIS relative to RES falls along the 1:1 line with a slope of 1.07 $\pm$ 0.09 with a Pearson's R of 0.8. Negative  $\Delta$ XCO<sub>2</sub> values, seen in 455 456 Fig. 13, are associated with meteorological conditions when winds come from northerly or 457 easterly directions, i.e., the roles of normal upwind and downwind sites are reversed. The 458 gradient of XCO<sub>2</sub> MIT relative to RES has a significantly lower range for modelled XCO<sub>2</sub> while the observed range of XCO2 is similar to PIS. The slope of observed to modelled 459 460  $\Delta XCO_2$  is 1.72±0.06 with a Pearson's R of 0.96. This points to a significant underestimation of the impact of urban sources on the MIT-RES gradient, which is also visible in the more 461 462 negative  $\Delta XCO_2$  during northerly wind conditions. This could indicate that the spatial 463 distribution of our emissions prior should be improved, i.e., emissions in the eastern outskirts/suburbs are likely underestimated in the IER emissions model. The low modelled 464 465  $\Delta XCO_2$  could also be due to overestimated horizontal dispersion in the model, which seems 466 less likely. When comparing the mean modelled daily cycle of the days when observations 467 exist with the mean diurnal cycle for all days within the field campaign period, we find that 468 the days with observations do not significantly differ from those without observations (see 469 Fig. 14). An investigation of typical diurnal variations of modelled  $\Delta XCO_2$  can only be 470 performed to a limited degree with the observational data available. Within the large 471 uncertainties, the modelled and observed  $\Delta XCO_2$  agree throughout the day. When analysing 472 the modelled  $\Delta XCO_2$  components we also find that the observed daytime increases of 473  $\Delta XCO_2$  are driven by  $CO_2$  added by urban FF  $CO_2$  burning and that the impact of FF is significantly higher at PIS (up to 1 ppm) then at MIT site (0.5 ppm) in the model., Our 474 475 observations indicate that both sites have strong diurnal variations. As the biogenic





- 476 contribution is expected to be similar, this is another indication that the impact of FF477 emissions on the MIT site is larger than predicted by our modelling framework.
- 478 Different ∆XCO<sub>2</sub> diurnal variations can be found for other upwind-downwind site pairs, but
- 479 they are all systematically driven by the locally-added CO<sub>2</sub> from FFCO<sub>2</sub>.
- 480

## 481 5 Conclusion and Outlook

482 For the two-weeks field campaign we demonstrated the ability of a network of five 483 EM27/SUN spectrometers, placed in the outskirts of Paris, to successfully track the XCO2 484 changes due to the urban plume of the city. However, we also found that XCO<sub>2</sub> cannot be 485 easily linked to local emissions as, even in such a densely populated area, XCO2 is still 486 significantly influenced by natural CO<sub>2</sub> uptake during the growing season. Using a gradient approach, i.e., analysing the difference between XCO<sub>2</sub> measured at upwind and downwind 487 488 stations, greatly reduced the impact of remote CO<sub>2</sub> sinks. Overall, the XCO<sub>2</sub> variability modelled using our ECMWF-CHIMERE system with IER (1 x 1 km<sup>2</sup>) emissions data was 489 490 found to be comparable with the observed variability and diurnal evolution of XCO<sub>2</sub>, despite 491 a significantly enhanced background for modelled XCO<sub>2</sub>. Our modelling framework, run at a 492 2 x 2 km<sup>2</sup> resolution over Paris also predicts that NEE and BC only significantly impacts 493  $\Delta$ XCO<sub>2</sub> during a few situations, specifically when meteorological conditions change making 494 the concept of 'upwind' and 'downwind' not applicable. When comparing modelled and measured  $\Delta XCO_2$  we find strong correlations (Pearson's R) of 0.8 and 0.96 for PIS-RES and 495 496 MIT-RES, respectively. This can be considered as an excellent degree of correlation as even 497 a model simulation which used optimised fluxes, based on surface observation, showed 498 correlations of 0.91 for its posterior results [Breon et al. 2015]. The offset between model 499 and observations also diminished for  $\Delta XCO_2$  and the slope found between observed and 500 modeled PIS-RES gradient is statistically in accordance with a 1:1 relationship (1.07±0.09). 501 However, the slope of the MIT-RES XCO<sub>2</sub> gradient of 1.72±0.06 suggests that the emission 502 model could potentially be improved, unless this underestimation of  $\Delta XCO_2$  by the model is 503 caused by overestimated dispersion in the model. It seems rather likely that the dispersion 504 would cause a general spreading of emission plumes and not systematically underestimate 505 emissions impacts from less densely populated, parts of Île-de-France. The data also 506 confirm previous results by models that XCO<sub>2</sub> gradients caused by a megacity do not 507 exceed 2 ppm, which supports the previous requirement for satellite observations of less than 1 ppm precision on individual soundings, and biases lower than 0.5 ppm (Ciais et al. 508 509 2015). The gradients are mainly caused by the transport of FFCO<sub>2</sub> emissions but, 510 interestingly, during specific episodes, a significant contribution comes from biogenic fluxes, 511 suggesting that these fluxes cannot be neglected even when using gradients.





512 Unfortunately, the duration of the campaign was relatively short, so that an in-depth analysis 513 of mean daily cycles or the impact of ambient conditions (traffic conditions, temperature, 514 solar insolation, etc.) on the observed gradient and underlying fluxes could be investigated 515 here. Hence, future studies in Paris and elsewhere should aim to perform longer-term observations during different seasons, which will allow better understanding changes in 516 517 biogenic and anthropogenic CO<sub>2</sub> fluxes. A remotely-controllable shelter for the EM27/SUN instrument is currently under development [Heinle and Chen, 2017]. This will considerably 518 519 facilitate the establishment of permanent spectrometer arrays around cities and other 520 sources of interest. Nevertheless, our study already indicates that such observations of 521 urban XCO<sub>2</sub> and  $\Delta$ XCO<sub>2</sub> contain original information to understand local sources and sinks 522 and that the modelling framework used here is suitable to support their interpretation. An 523 improved model will also be able to adjust or better model the background conditions and 524 potentially use this type of observations to estimate local CO<sub>2</sub> fluxes using a Bayesian 525 inversion scheme similar to the existing system based on in-situ observations for Paris 526 [Staufer et al. 2016].

527 We expect that the previous successful collaboration in the framework of the Paris campaign 528 will mark the permanent implementation of COCCON as a common framework for a French-529 Canadian-German collaboration on the EM27/SUN instrument. The acquisition of additional 530 spectrometers is planned by several partners.

531

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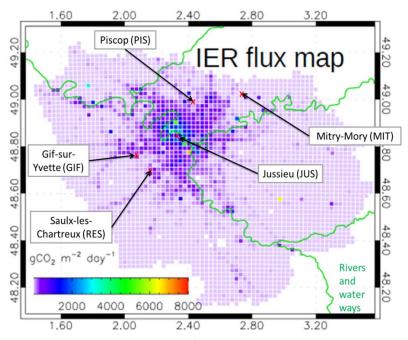




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Figure 1.  $CO_2$  emissions in the Île-de-France region according to the IER emission

681 inventory. Measurement sites are indicated by red crosses





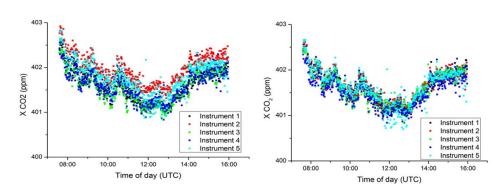
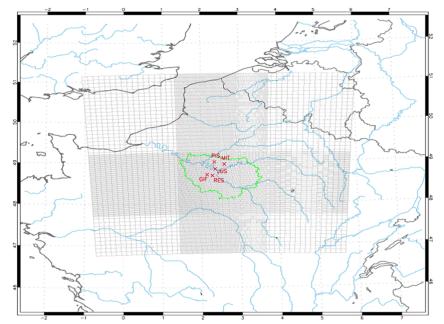




Figure 2. Sample parallel measurements of the five EM27/SUN instruments in Karlsruhe for raw data (left panel) and the data with the applied correction (right panel) taken on April 15<sup>th</sup>, 2015.





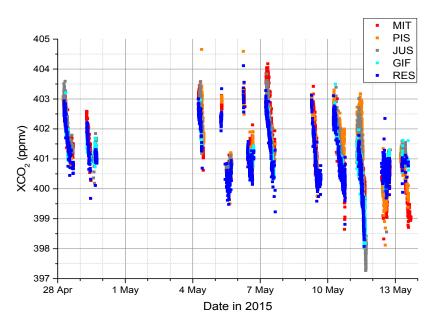


686

Figure 3. Modelling domain and numerical grid configuration of CHIMERE with a zoom
 on the Île-de-France region at 2 x 2 km<sup>2</sup>.







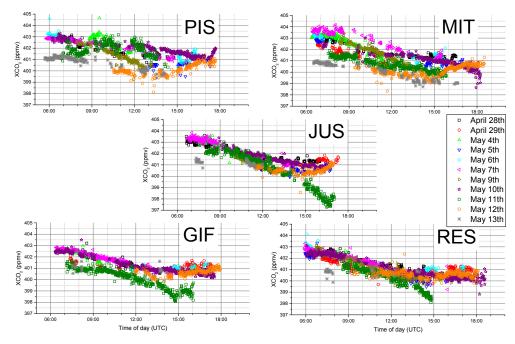
689

690 Figure 4. Time series of observed XCO<sub>2</sub> in the Parisian region for all five sites (all valid

691 data of~1 minute averages).





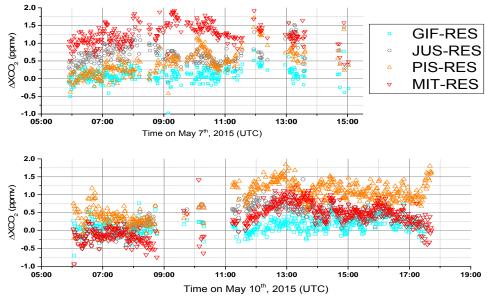


692 693

Figure 5. Time series of observed XCO<sub>2</sub> in the Parisian region sorted by station





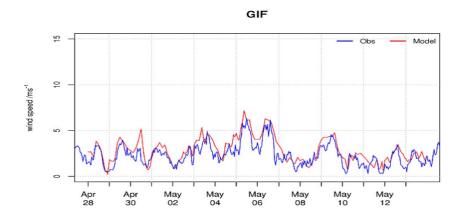


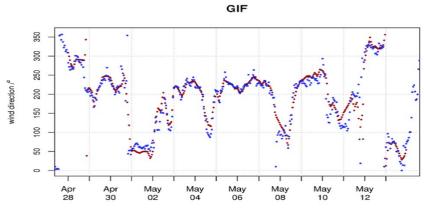
Time on May 10<sup>th</sup>, 2015 (UTC)695Figure 6. Observed spatial gradients of XCO2 for May 7th (southwesty winds) and May

696 10th (southerly winds)







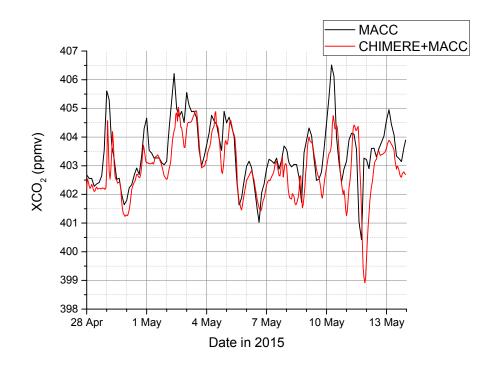


697

Figure 7. Comparison of modelled and observed wind speeds and directions at the Gif-Sur-Yvette measurement site





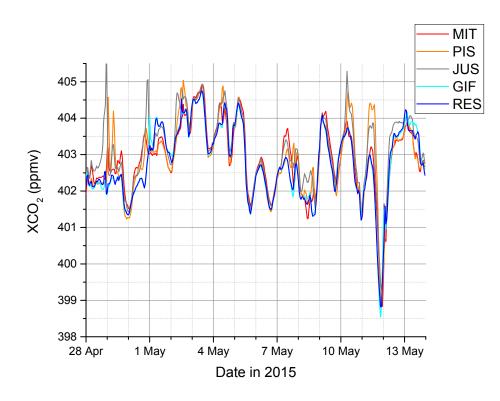


700 701 Figure 8. Comparison of modelled XCO<sub>2</sub> from ECWMF-CAMS (15 x 15 km<sup>2</sup>) with

CHIMERE simulation (inner domain, 2 x 2 km<sup>2</sup>) for JUS. 702





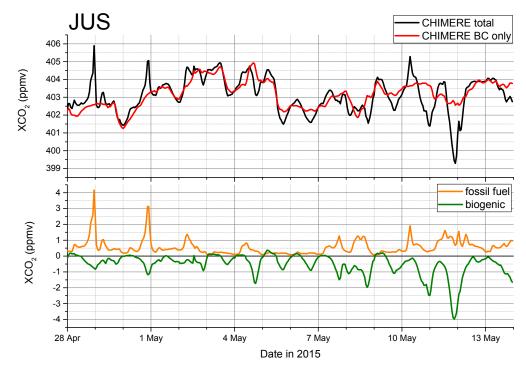


703 704

Figure 9. Modelled XCO<sub>2</sub> for all stations







705 706 Figure 10: Time series of XCO<sub>2</sub> and related fluxes for JUS. The topp panel provides a 707 comparison of modelled total XCO<sub>2</sub> and XCO<sub>2</sub> variations due to changes in boundary 708 conditions. The lower panel shows the contribution of the different flux components,

709 namely fossil fuel CO<sub>2</sub> emissions and biogenic fluxes.





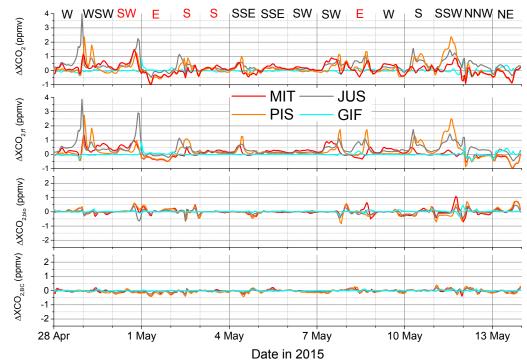
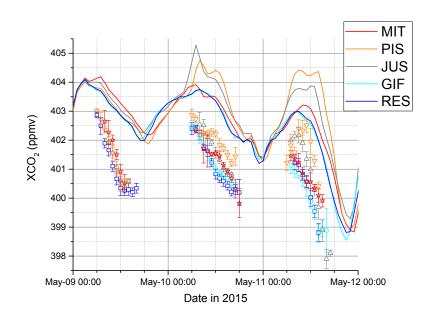


Figure 11. Modelled XCO<sub>2</sub> gradients for each station relative to RES. The dominant
wind conditions for each day given at the top of the figure and days without
observations due to precipitation are in red.







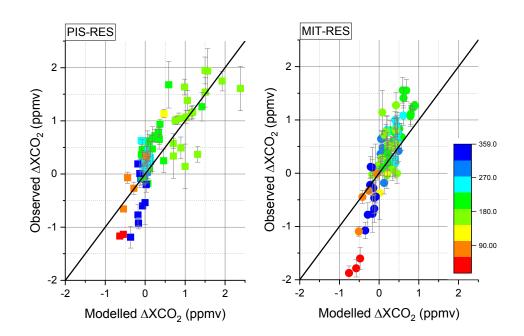
714

715 Figure 12. Comparison of modelled (solid lines) and observed hourly averaged XCO<sub>2</sub>

716 (symbols) with standard deviations as error bars.





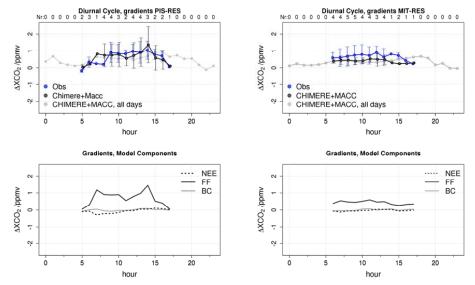


717

Figure 13. Comparison of modelled and observed hourly averaged  $\Delta XCO_2$  for gradients between PIS and RES (left) and MIT and RES (right), with standard deviations as vertical bars and the points color coded by wind direction from 0 to 359 degrees.







722

723 Figure 14. Comparison of modelled and observed mean daily cycle of hourly averaged

724 ΔXCO<sub>2</sub> of PIS with RES (top left) and of MIT with RES (top right) during the campaign

725 and the modelled contribution of different CO<sub>2</sub> sources/sinks (bottom panels).





Location	ID	Lat (deg)	Lon (deg)	Position
Piscop	PIS	49.019	2.347	20 km NNW of JUS
Mitry-Mory	МІТ	48.984	2.626	25 km NW of JUS
Jussieu	JUS	48.846	2.356	Paris city centre
Saulx-les- Chartreux	RES	48.688	2.284	20 km SSW of JUS
Gif-Sur-Yvette	GIF	48.708	2.148	20 km SW of JUS

726 Table 1. Location of FTIR measurement instruments during the field campaign





Instrument	XCO <sub>2</sub> factor Berlin	XCO <sub>2</sub> factor before	XCO <sub>2</sub> factor after
		Paris	Paris
1	1.0000 (0.0003)	1.0000 (0.0003)	1.0000 (0.0003)
2	0.9992 (0.0003)	0.9991 (0.0003)	0.9992 (0.0003)
3	1.0002 (0.0003)	1.0001 (0.0004)	1.0000 (0.0005)
4	0.9999 (0.0003)	1.0000 (0.0004)	1.0000 (0.0004)
5	0.9996 (0.0003)	0.9995 (0.0003)	0.9995 (0.0003)

727 Table 2. Normalisation factors for the five EM27/SUN instruments derived during

728 measurements before and after the Paris field campaign. Values in parentheses are

729 standard deviations. Measurements of instrument 1 were arbitrarily chosen as

reference from which the others were scaled. The calibration factors from a previous
 field campaign in Berlin [Hase et al. 2015] are also shown. Calibration factors between

the two field campaigns agree well within 0.02 % (~0.08 ppm) for all instruments.





Date	No. of observations	Quality	Wind speed (m s <sup>-1</sup> )	Wind direction
	MIT GIF PIS RES JUS	•		
28 Apr 2015 (Tu)	179 102 178 199 234	++	4	W
29 Apr 2015 (We)	110 124 0 161 53	+	5	SW-W
04 Mai 2015 (Mo)	194 85 96 163 83	+	6	S-SE
05 Mai 2015 (Tu)	77 27 85 185 92	+	8	S-SW
06 Mai 2015 (We)	81 88 87 139 0	+	8	SW
07 Mai 2015 (Th)	169 313 252 286 238	+++	3	SW
09 Mai 2015 (Sa)	179 0 181 289 149	++	6	W
10 Mai 2015 (Su)	325 478 362 542 282	++++	3	S
11 Mai 2015 (Mo)	410 431 251 298 413	++++	3	SSW
12 Mai 2015 (Tu)	324 222 230 326 203	+++	4	NNW
13 Mai 2015 (We)	159 18 182 28 56	+	4	NE

733 Table 3. Summary of all measurement days with the number of observations at each

734 of the sites, Mitry Mory (MIT) , Gif Sur Yvette (GIF), Piscop (PIS), Saulx-les-Chartreux

735 (RES), Jussieu (JUS), the overall quality ranking of each day according to the number

736 of available observations and temporal coverage (with classification from poor to

737 great: +, ++, +++, ++++), and the ground-level wind speed and direction.





	Total obs	Mean (ppm)	STD (ppm)	Quartile1 (ppm)	Median (ppm)	Quartile3 (ppm)
RES	2616	401.11	0.93	400.44	400.88	401.96
GIF	1888	401.05	0.92	400.94	400.94	401.58
JUS	1803	401.33	1.17	401.31	401.31	402.04
PIS	1904	401.62	0.95	401.03	401.59	402.44
MIT	2207	401.26	1.15	401.11	401.11	401.95

738 Table 4. Statistics of observed XCO<sub>2</sub> 1-minute averages for all sites