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1 Atmospheric ammonia variability and link with PM formation: a
2 case study over the Paris area

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12 Abstract

13 The Paris megacity experiences frequent particulate matter ($PM_{2.5}$, PM with a diameter less than
14 $2.5 \mu m$) pollution episodes in springtime (March-April). At this time of the year, large parts of
15 the particles consist of ammonium sulfate and nitrate which are formed from ammonia (NH_3)
16 released during fertilizer spreading practices and transported from the surrounding areas to
17 Paris. There is still limited knowledge on the emission sources around Paris, their magnitude and
18 seasonality.

19 Using space-borne NH_3 observation records of 10-years (2008-2017) and 5-years (2013-2017)
20 provided by the Infrared Atmospheric Sounding Interferometer (IASI) and the Cross-Track
21 Infrared Sounder (CrIS) instrument, regional pattern of NH_3 variabilities (seasonal and inter-
22 annual) are derived. Observations reveal identical high seasonal variabilities with three major
23 NH_3 hot spots found from March to August. The high inter-annual variability is discussed with
24 respect to atmospheric total precipitation and temperature.

25 A detailed analysis of the seasonal cycle is performed using both IASI and the CrIS instrument
26 data, together with outputs from the CHIMERE atmospheric model. For months of high NH_3
27 concentrations (March to August) the CHIMERE model shows good correspondence with
28 correlation slopes of 0.98 and 0.71 when comparing with IASI and CrIS, respectively. It is found
29 that the model is only able to reproduce half of the observed atmospheric temporal NH_3
30 variability in the domain. In term of spatial variability, the CHIMERE monthly NH_3 concentrations
31 in springtime show a slight underrepresentation over Belgium and the United-Kingdom and
32 overrepresentation in agricultural areas in the French Brittany/Pays de la Loire and Plateau du
33 Jura region, as well as in the north part of Switzerland.

34 Using HYSPLIT cluster analysis of back-trajectories, we show that NH_3 total columns measured in
35 spring over Paris are enhanced when air masses are originated from the Northeast (e. g.,
36 Netherlands and Belgium), highlighting the long-range transport importance on the NH_3 budget
37 over Paris.

38 Finally, we quantify the key meteorological parameters driving the specific conditions important
39 for the $PM_{2.5}$ formation from NH_3 in the Ile-de-France region in springtime. Data-driven results
40 based on surface $PM_{2.5}$ measurements from the Airparif network and IASI NH_3 observations
41 show that a combination of the factors, e. g. a low boundary layer of $\sim 500m$, a relatively low
42 temperature of $5^\circ C$ and a high relative humidity of 70%, contributes to favor $PM_{2.5}$ and NH_3
43 correlation.



44 1. Introduction

45 Ammonia (NH_3) is an atmospheric pollutant and one of the main sources of reactive nitrogen in
46 the atmosphere which is involved in numerous biochemical exchanges impacting all ecosystems
47 [Sutton et al., 2013]. The global budget of reactive N has dramatically increased since the
48 preindustrial era [Holland et al., 2005; Battye et al., 2017] causing major environmental
49 damages such as ecosystems and species extinction, as well as soil and water eutrophication
50 and acidification [Rockström et al., 2009]. NH_3 is a precursor of ammonium salts which can form
51 up to 50% to particulate matter (PM) total mass [Behera et al., 2013]. Large cities such as Paris
52 (which is the most populated area in the European Union with 10.5 million people when its
53 larger metropolitan regions are included) typically experiences strong PM pollution episodes in
54 springtime. These particles are known to be harmful for human health [Pope III et al., 2009]
55 inducing 2000 deaths per year in the Paris megacity [Corso et al., 2016] and to impact the
56 radiative budget of the Earth [Myhre et al., 2013].

57 Because of their impact on the environment, public health, and climate change, NH_3 emissions
58 are regulated in several countries in the world. However, NH_3 emissions of European countries
59 have increased by 2% over the period 2014-2016 [National Emission Ceilings Directive reporting
60 status, 2018], where the Gothenburg Protocol set a reduction of 6% by 2020. In France, where
61 94% of NH_3 emissions come from the agriculture sector [CITEPA, 2018] as a result of extensive
62 fertilizer use to increase crop yields [Erisman et al., 2008], policies have been implemented with
63 the aim to reduce NH_3 emissions by 13% in 2030 related to 2005 [CEIP, 2016]. However NH_3
64 emissions are projected to increase in the future globally with increased population and food
65 demand [van Vuuren et al., 2011] and NH_3 volatilization will be enhanced with climate change
66 [Sutton et al., 2013].

67 Once in the atmosphere, NH_3 is rapidly removed by wet and dry deposition, and by reactions
68 with atmospheric sulfuric and nitric acid, leading to a relatively short lifetime between a few
69 hours and few days [Galloway et al., 2003]. Release of NH_3 in the atmosphere depends on i)
70 agriculture practices: spreading season, fertilizer form (urea, ammonium nitrate), fertilizer
71 application methods, crops, soil conditions such as pH [Hamaoui-Laguel et al., 2014]; and on ii)
72 meteorological conditions (i.e. wind, temperature, and precipitation). Inter-annual variability of
73 PM formation over urban area is poorly understood, since it also depends on many factors such
74 as atmospheric humidity and temperature, which govern the phase equilibrium of secondary
75 aerosols [Fuzzi et al., 2015]. The variety of factors influencing NH_3 volatilization and PM
76 formation illustrates the complexity of predicting their concentrations in the atmosphere
77 [Behera et al., 2013].

78 Atmospheric chemical transport models have difficulty representing both NH_3 and $\text{PM}_{2.5}$
79 distributions due to the challenge of reproducing NH_3 temporal variability [Pinder et al., 2006;



80 Fortems-Cheiney et al., 2016], long-range transport of pollutants [Moran et al., 2014], and
81 secondary aerosol formation in the atmosphere [Petetin et al., 2016]. The GEOS-Chem chemical
82 transport model [Bey et al., 2001] was found to underestimate the observed NH_3 concentrations
83 in most regions of the globe [Zhu et al., 2013; Li et al., 2017]. Heald et al. (2012) compared the
84 IASI observations with the GEOS-Chem model and showed that NH_3 is likely underestimated in
85 California, leading to a local underestimate of ammonium nitrate aerosol. Similarly, the French
86 CHIMERE model [Menut et al., 2013] underestimates the NH_3 budget over Paris [Petetin et al.,
87 2016; Fortems-Cheiney et al., 2016] because of the mis-representation of agricultural emissions
88 in terms of intensity and both spatial and temporal distribution. Often ground and aircraft-
89 based observations are used to provide detailed representation of the atmospheric state that
90 can be used to evaluate and improve the model simulations; however, these can be spatially
91 sparse and/or over short sampling periods, especially globally. Additionally, more recently
92 available (within the last 10-years) sun-synchronous satellite-based infrared sensors have been
93 providing NH_3 observations globally with a spatial resolution of ~ 15 km approximately twice a
94 day. These satellite observations have limited independent vertical information, but do capture
95 the spatiotemporal variabilities needed to help address these issues and improve model
96 simulations, especially in remote locations [Skjøth et al., 2011; Kranenburg et al., 2016].

97 Aside from the Tropospheric Emission Spectrometer (TES, [Beer et al., 2008]), now
98 decommissioned but which was first to demonstrate the capability of thermal infrared
99 instruments to monitoring lower tropospheric NH_3 , 3 missions are able to measure it now : the
100 Atmospheric InfraRed Sounder (AIRS, [Warner et al., 2016]), the Cross-track Infrared Sounder
101 (CrIS, [Shephard and Cady-Pereira, 2015]), and the Infrared Atmospheric Sounding
102 Interferometer (IASI, [Clarisse et al., 2009]). Recent studies have shown the increased capacity
103 of space-borne instruments to derived spatial and seasonal distributions of NH_3 concentrations
104 globally [Clarisse et al., 2009; Shephard et al., 2011; Van Damme et al., 2014a & 2015a],
105 regionally [Beer et al., 2008; Clarisse et al., 2010; Van Damme et al., 2014b] and locally [Van
106 Damme et al., 2018], as well as trends of NH_3 [Warner et al., 2017].

107 Representative measurements of NH_3 concentrations and spatiotemporal variabilities are
108 needed to address the link between NH_3 and $\text{PM}_{2.5}$ formation and improve model simulations.
109 This has been attempted previously in some cities around the world, such as in Shanghai [Ye et
110 al., 2011], Houston [Gong et al., 2013], Santiago City [Toro et al., 2014], and Beijing [Zhao et al.,
111 2016] for instance. However, although the Paris megacity is repeatedly shrouded by particulate
112 pollution episodes, many of studies are limited and performed over relatively short time frame
113 during field campaigns [Petetin et al., 2016; Zhang et al., 2013], or based on numerical
114 simulations [Skylakou et al., 2014]. Our study is a data-driven regional approach and considers a
115 longer time period to study the seasonal/inter-annual variabilities of NH_3 and its impact of $\text{PM}_{2.5}$
116 formation over the Paris megacity. Specifically in this paper we study concentrations and



117 spatiotemporal variability of atmospheric NH₃ from the agricultural sector to gain insights on its
118 effects on megacity air quality using: 1) long-term satellite observations derived from IASI (10
119 years from 2008 to 2017) and CrIS (5 years from 2013 to 2017) at regional scale (400km radius-
120 circle from Paris city center); 2) spatiotemporal patterns of the CHIMERE model evaluated
121 against the IASI and CrIS datasets for 2014 and 2015; and 3) the main meteorological
122 parameters favoring the secondary PM_{2.5} formation from NH₃ in the Paris megacity are
123 analyzed.

124 2. Methodology

125 2.1. Region of analysis

126 The domain of analysis covers a circular area of 400 km radius around the Paris city center
127 (Figure 1, larger circle) enabling the study of temporal and spatial variabilities of NH₃ emission
128 sources likely to affect air quality in the Paris megacity. It has been selected for two reasons.
129 First, it includes main regions known for their high NH₃ emissions, which can be transported and
130 affect air quality over the Parisian region (Ile-de-France –IdF-, smaller circle in Figure 1).
131 Emission regions in the Netherlands, North of Germany, Northwest of Belgium, and the Brittany
132 region in France, are highlighted in darker colors in Figure 1 (emissions values are from the
133 European Monitoring and Evaluation Programme -EMEP- 2015). Second, this area corresponds
134 to the transport of 24 hours back-trajectories from Paris generated from the HYSPLIT model for
135 one year, ensuring that NH₃ can indeed be efficiently transported from the emitting sources
136 within the selected domain to the IdF region.

137 2.2. Satellite observations of ammonia

138 For this study we used the available data from IASI and CrIS which are both Fourier transform
139 spectrometers to evaluate the current capacity to observe NH₃ concentrations from space, and
140 study its variability around IdF. Technical information are summarized in Table 1.

141 2.2.1. Infrared Atmospheric Sounding Interferometer (IASI)

142 IASI is a nadir-viewing spectrometer launched on board the Metop-A and Metop-B satellites and
143 operated by EUMETSAT (European Organisation for the Exploitation of Meteorological
144 Satellites), since October 2006 and September 2012, respectively. These satellites are on similar
145 polar orbits with Equator crossing times at 09:30 (21:30) local mean solar time for the
146 descending (ascending) orbit. IASI measures the thermal infrared radiation of the system Earth-
147 atmosphere in the spectral range from 645 to 2760 cm⁻¹ with a spectral resolution 0.5 cm⁻¹
148 apodized. The satellite swath is an area of 2200 km width composed by off-nadir measurements
149 up to 48.3° on both sides of the track. At nadir, the IASI field of view is composed of 4 x 4 pixels
150 of 12 km diameter each [Clerbaux et al., 2009].



151 The NH_3 total columns used here are derived from IASI using an Artificial Neural Network
152 reanalyzed with ERA-interim data (ANNI-NH3-v2.1R [Van Damme et al., 2017]). This dataset is
153 consistent in time and suitable for investigating inter-annual variability, which is one purpose of
154 this study. Note that we have considered here only morning measurements (9:30) since the
155 evening ones (21:30) are associated with larger relative errors [Van Damme et al., 2017]. IASI
156 retrievals provide a robust error estimate for each IASI-NH3 observations, allowing to take into
157 account the variable sensitivity when comparing IASI dataset with independent measurements.
158 Finally, no filter on relative errors of the IASI datasets has been applied following
159 recommendations from Van Damme et al. (2017) and outliers for which concentrations exceed
160 10 standard deviations above the mean in the domain of study have been removed.

161 Over the studied area, Metop-A and Metop-B have an overpass time difference ranging from
162 only a few seconds to 67 minutes depending on the viewing geometry of the satellite scans; the
163 average difference is 26 minutes for the 1325 days of common measurements. Monthly maps
164 for the 10 years of observations between 2008 and 2017 are obtained by averaging Metop-A
165 and whenever Metop-B (the two instruments are considered jointly for their period of common
166 operation from March 2013 to 2017) with more than 10^5 pixels on average over the domain of
167 analysis. The number of available NH_3 columns depends not only on the satellite overpass time
168 but also on the state of the atmosphere being remotely sensed (e.g. thermal contrast and cloud
169 cover). IASI NH_3 has been evaluated using the LOTOS-EUROS model over Europe [Van Damme et
170 al., 2014b] and ground-based and airborne measurements [Van Damme et al., 2015b], showing
171 consistency between the IASI NH_3 and the available datasets. When comparing IASI NH_3
172 (previous IASI-NN version) with ground-based Fourier transform infrared (FTIR) observations, a
173 correlation of 0.8 and a slope of 0.73, with a mean relative difference of $-32.4 \pm (56.3)\%$ have
174 been found [Dammers et al., 2016].

175 2.2.2. Cross-track Infrared Sounder (CrIS)

176 The CrIS instrument [Zavalyov et al., 2013] is a Fourier Transform spectrometer operated by the
177 Joint Polar Satellite System (JPSS) program on Suomi National Polar-orbiting Partnership (NPP)
178 satellite, launched on 28 October 2011. CrIS is in a sun-synchronous orbit with a mean local
179 daytime overpass time of 13:30 (01:30) in the ascending (descending) node. CrIS measures the
180 atmospheric composition over three wavelength bands in the infrared region ($645\text{--}1095\text{ cm}^{-1}$;
181 $1210\text{--}1750\text{ cm}^{-1}$; $2155\text{--}2550\text{ cm}^{-1}$). NH_3 retrievals are performed from the $645\text{--}1095\text{ cm}^{-1}$ band
182 with a spectral resolution of 0.625 cm^{-1} . The CrIS instrument scans a 2200 km swath width (+/
183 50°). At nadir, the CrIS field of view consists of a 3×3 array of circular pixels of 14 km diameter
184 each.

185 The CrIS Fast Physical Retrieval (CRPR) [Shephard and Cady-Pereira., 2015] uses an optimal
186 estimation approach [Rodgers, 2000] that minimizes the difference between the CrIS measured



187 atmospheric spectra and a very fast Optimal Spectral Sampling (OSS) [Moncet et al., 2008]
188 forward model simulated spectrum to retrieve atmospheric profiles of ammonia volume mixing
189 ratios. This physical approach provides direct estimates of the retrieval errors and the vertical
190 sensitivity (averaging kernels) of the satellite observations, which is important as they vary from
191 profile-to-profile depending on the atmospheric state. The retrieved error covariance and
192 averaging kernels are also beneficial for air quality model comparisons and data assimilation
193 into models as any *a priori* information used in the retrieval can be accounted for in a robust
194 manner (i.e. observation operator). CrIS has been shown to retrieve ammonia surface
195 concentrations values down to ~0.2-0.3 ppbv under favorable conditions [Kharol, et al., 2018].
196 CrIS comparisons with ground-based FTIR observations show a correlation of 0.77 with a low
197 CrIS bias of +2% in the total column [Dammers et al., 2017]. Initial evaluation against surface
198 observations from the Ammonia Monitoring Network (AMoN) show that even with the inherent
199 sampling differences between the two surface observations they compare well with a
200 correlation of 0.76 and an overall mean CrIS – AMoN difference of ~+15% [Kharol et al., 2018].

201 For this study, the CrIS quality flag = 4 has been used, ensuring that retrievals provide some
202 information from the measurement (degrees-of-freedom- of-signal > 0.1). In addition, outliers
203 for which concentrations exceed 10 standard deviations above the mean have been removed.

204 2.3. Modelling NH₃ from the CHIMERE model

205 The CHIMERE runs used in this study were obtained in the framework of the Copernicus
206 Atmospheric Monitoring Service (CAMS, <https://atmosphere.copernicus.eu/>), and its annual
207 task devoted to the production of regional reanalysis over Europe. The hindcasts for year 2014
208 and 2015 (raw simulation without data assimilation) were produced over Europe with a
209 horizontal resolution of 0.1° per 0.1° and
210 9 vertical levels stretched from the surface up to 500 hPa (~5000m). The input data to feed
211 CHIMERE [Menuet et al., 2013; Mailler et al., 2017] were the Integrated Forecasting System (IFS)
212 meteorological data from European Centre for Medium-Range Weather Forecasts (ECMWF), the
213 annual emission inventory provided by the Netherlands Organisation for Applied Scientific
214 Research (TNO) [Kuenen et al., 2014] for year 2011 and the fire emissions from the Global Fire
215 Assimilation System (GFAS, [Kaiser et al., 2012]). The model computes hourly concentrations for
216 more than 180 species, among which are the regulated pollutants such as ozone, PM₁₀, and
217 NH₃. Within CHIMERE a comprehensive modelling system allows to compute the evolutions of
218 gaseous species and aerosols taking into account physical and chemical process. More than 30
219 gaseous species are involved in the chemical scheme and an aerosol module assesses the gas-
220 particulate phase equilibrium and compute the aerosol composition (inorganic, organic and
221 natural components). These datasets were evaluated over Europe for several pollutants before
222 being used for air quality studies (<http://policy.atmosphere.copernicus.eu/Reports.html>).



223 The model NH₃ profiles were integrated vertically along the 9 km model layers to provide a
224 column that can be compared to that of the satellite measurements. Concretely this makes the
225 reasonable assumption that all the NH₃ is located within this 0-5km layer (see e.g. Figure 1 in
226 [Whitburn et al., 2016]).

227 2.4. Relative scales and coincidence criteria for dataset comparisons

228 Direct quantitative comparisons of satellite NH₃ products are difficult because of the different
229 overpass times and ground footprint sizes of the 2 space borne instruments, which are not
230 compatible with the high variability of NH₃ in space and time. Therefore, the evaluation of
231 satellite observations is often made with the use of in situ measurements performed at surface
232 and onboard aircrafts [Nowak et al., 2012; Van Damme et al., 2015b], or with ground-based
233 remote-sounding FTIR [Dammers et al., 2016; Dammers et al., 2017].

234 The purpose here of comparing CrIS and IASI is to assess qualitatively the spatiotemporal
235 patterns of the NH₃ sources derived from the two datasets and use these regional observations
236 to evaluate the CHIMERE model in the domain of analysis at the local time for their respective
237 overpasses: 9:30 and 13:30. CHIMERE outputs, in terms of NH₃ concentrations, have already
238 been compared to the IASI observations at regional scale (Europe, [Fortems-Cheiney et al.,
239 2016]), and to surface measurements at local scale (Paris, [Petetin et al., 2016]), but have never
240 been evaluated against the CrIS observations.

241 One aspect that needs to be considered when comparing concentration amounts inferred from
242 infrared satellite observations is the importance of the algorithm and the a priori information
243 used in the retrieval, especially for NH₃ which has limited vertical information. Some differences
244 between the IASI and CrIS observations might arise due to instrument measurement differences
245 (e.g. sensitivity), difference sampling period (e.g. overpass times of morning/evening vs middle
246 of day/night), and retrieval algorithm differences, but they have both been validated and shown
247 to capture well the spatiotemporal variations in lower tropospheric ammonia. Since the purpose
248 of our study is not to quantitatively compare IASI and CrIS NH₃ data, but rather to use these
249 independent datasets to assess NH₃ sources patterns over the domain and qualitatively
250 evaluate the CHIMERE model in term of NH₃ concentrations and variabilities, a standardization
251 procedure was applied to their retrieved absolute NH₃ columns. We computed “standardized
252 columns” for each independent dataset (IASI, CrIS, and CHIMERE, separately) for 2014 and 2015
253 over the domain of study in such a way that the corresponding values have a standard deviation
254 of 1 and a mean of 0, as in [Wilks, 2011].

255 In addition, to compare CHIMERE outputs with satellite data/columns, spatial and temporal
256 coincidence criteria have been applied. To compare satellite observations, all CrIS pixels located
257 within a 25-km radius circle from the center of the IASI ground pixels have been considered



258 within the same day of measurements. A spatial criterion of 25 km has been chosen because it
259 optimizes the number of pairs involved in the statistics and improves the correlations. As for the
260 comparisons between the model and the observations: all CHIMERE outputs located within the
261 same $0.15^\circ \times 0.15^\circ$ grid box than the satellite and within 1 hour from its measurement have been
262 selected.

263 3. Results

264 3.1. NH₃ regional observations derived from IASI (10-years) and CrIS (5- 265 years)

266 3.1.1. Seasonal variabilities

267 First the seasonal variability was investigated over the IdF area. On a monthly basis, the 10-year
268 and 5-year averaged regional NH₃ total column distributions derived from IASI and CrIS were
269 found to exhibit a high seasonality over the domain (Figures 2 and 3). Note that the distributions
270 in Figures 2 and 3 have been obtained by averaging satellite NH₃ observations in $0.25^\circ \times 0.25^\circ$
271 grid boxes. Both satellite datasets exhibits the same variability over the domain even if the time
272 period is different (10-years versus 5-years) and the sampling hour differs (~ 9.30 versus ~ 13.30).
273 One note that CrIS and IASI NH₃ columns present small differences in term of NH₃ total columns
274 in low concentration regimes in the domain of study.

275 In these figures (2 and 3) high NH₃ concentrations (up to 2.10^{16} molecules/cm²) can be observed
276 from March to August at different locations of the domain:

- 277 • The French Champagne-Ardennes region in March and April (Figures 2 and 3, box A),
- 278 • The northern part of the domain corresponding to the Netherlands and the North of Belgium
279 from April to August (Figures 2 and 3, box B), and
- 280 • The Brittany/Pays de la Loire regions (West of France) mainly in April and August but still
281 persistent from March to August (Figures 2 and 3, box C).

282 The observed seasonality is related to agricultural practices (fertilizer application period varying
283 as function of the crop types and farming species) and changes in temperatures, with higher
284 temperatures favoring volatilization. This explains the high concentration in July and August.

285 In the Champagne-Ardennes region, areas of hotspots do not correspond to vineyards but to
286 field vegetables and root crops (from the Institut National de la Recherche Agronomique INRA
287 https://odr.inra.fr/intranet/carto/cartowiki/index.php/OTEX_et_Orientation_Agricole_des_terri
288 toires, and AGRESTE, Service Central d'Enquêtes et d'Études Statistiques, 2015
289 <http://agreste.agriculture.gouv.fr/IMG/pdf/R4215A15.pdf>). This is a leader region for mineral
290 fertilization used for sugar industry in France [Ramanantenasoa et al., 2018]. Hamaoui-Laguel et
291 al. (2014) and Fortems-Cheiney et al. (2016) have previously noted that NH₃ emissions in this



292 region, mainly due to fertilizer over barley, sugar beet, and potato starch in early March, were
293 higher than what have been reported in the EMEP inventory.

294 NH₃ concentrations are high from April to August in the northern part of the domain that is
295 known for its animal farming (Eurostat 2014, [http://ec.europa.eu/eurostat/statistics-
296 explained/index.php?title=File:Livestock_density_by_NUTS_2_regions,_EU-28,_2013.png](http://ec.europa.eu/eurostat/statistics-explained/index.php?title=File:Livestock_density_by_NUTS_2_regions,_EU-28,_2013.png), [Van
297 Damme et al., 2014a]).

298 In the Pays de la Loire, NH₃ concentrations are high in April and August and remain relatively
299 high from March to September. Hotspots are found in areas of livestock farming, mainly poultry
300 and granivorous, which explains the high and relatively constant NH₃ concentrations over
301 warmer periods in this region.

302 3.1.2. Inter-annual variabilities

303 As can be seen in Figures 2 and 3, NH₃ concentrations are enhanced between March and August
304 in the domain. In this section, inter-annual variabilities are discussed regarding meteorological
305 conditions and agricultural practices during this time period.

306 Inter-annual variability of NH₃ is higher in springtime than in summer, e.g. in June the variance is
307 8 times lower than for the other months. To illustrate the inter-annual variability in springtime,
308 maps of monthly mean NH₃ total columns derived in March-April period from IASI (2008-2017
309 time period) and from CrIS (2013-2017 time period) are shown in Figure 4. Both satellite
310 distributions exhibit the same inter-annual variability from 2013 to 2017 with higher NH₃
311 concentrations in 2015 over the northern part of the domain than the other years. NH₃
312 concentrations derived from IASI in 2011 are 150% higher in spring (March and April) compared
313 to 2016 (Figure 4). This inter-annual variability is partly driven by meteorological conditions and
314 specific agricultural constrains (crop type and phenological stage for instance).

315 To investigate the impact of meteorological conditions on atmospheric NH₃ variability, we
316 computed the monthly mean anomalies of total precipitation versus skin temperature derived
317 from ECMWF ERA-interim [Dee et al., 2011], color coded by NH₃ total columns anomalies
318 derived from IASI, as shown in Figure 5. Monthly mean anomalies have been calculated relative
319 to the 10-years averages (in %). In this figure, monthly NH₃ total columns are at least 10% higher
320 (positive anomalies, red dots) when skin temperatures are higher and total precipitation are
321 lower than the 10-year average. In contrast, negative monthly NH₃ total columns anomalies
322 (blue dots, Figure 5) are associated with higher total precipitation and lower skin temperatures
323 than the 10-years average. To further detail this analysis, Figure 1 of the supplement
324 information shows bar plots of monthly mean NH₃ total columns derived from IASI, total
325 precipitation and skin temperature derived from ECMWF from March to August, plotted in
326 different colors for the different years of measurements from 2008 to 2017. NH₃ total columns



327 are larger by more than 300% in March-April 2012 compared to 2013 (Figure S1a). Total
328 precipitation is higher (0.4 mm compared to 1 mm, Figure S1b) and skin temperature is lower
329 (281 compared to 288 K, Figure S1c) in March 2013 than in March 2012 on average over the
330 domain. Overall, total precipitation is anti-correlated with NH_3 concentrations in the
331 atmosphere ($R = -0.52$ from March to August for all years, not shown here) because of a) the
332 wet deposition importance in the atmospheric NH_3 removal and b) the absence of fertilization
333 during rainy periods. Skin temperature is relatively correlated with NH_3 concentrations ($R = 0.33$
334 from March to August for all years) since higher temperature increases volatilization of NH_3
335 from the surface to the atmosphere.

336 In addition, NH_3 concentration is maximum in March 2011 whereas it peaks later in April for
337 2012 (Figure S1a). Springtime is a spreading fertilizer period depending on many agricultural and
338 meteorological constrains. When temperature are mild, such as in 2012 (Figure S1b), fertilizer
339 spreading occurs sooner because the phenological growth stage is more advanced. Fertilizing
340 process period also varies in function of the sowing date which depends on agricultural
341 practices and crop types: corn is fertilized in early spring whereas rapeseed is in late spring.

342 Overall, all these meteorological (precipitation and temperature) and agricultural (fertilizer and
343 manure applications) parameters account for the high NH_3 inter-annual variabilities revealed by
344 both IASI and CrIS in the domain of study.

345 3.2. Comparisons of NH_3 columns derived from IASI, CrIS, and CHIMERE

346 for 2014 and 2015

347 To discuss the representation of agricultural emissions in the models in terms of intensity and
348 both spatial and temporal distributions, regional satellite observations derived from IASI and
349 CrIS have been compared to the CHIMERE model in the region of analysis.

350 3.2.1. Annual cycle

351 Standardized monthly mean concentrations derived from IASI, CrIS, and CHIMERE for 2014 and
352 2015 are shown in Figure 6. These years were selected as NH_3 total columns were found to vary
353 a lot, reaching 10% higher in March and 50% lower in May than the 10-years average

354 As can be seen from the plot, the 3 datasets exhibit similar patterns in terms of seasonality: all
355 are enhanced in March-April and in summer, and show a decrease in May. However two major
356 differences can be noted. First, CrIS standardized NH_3 columns are higher in winter (November,
357 December, and January) compared to the other dataset which can be also be seen in Figure 3.
358 This could be attributed to a higher number of outliers, given the larger standard deviation
359 (shaded areas, Figure 6) and no attempt to account for potential non-detects when
360 concentrations fall below the instrument detection limits. For these months, NH_3 levels are low



361 and undetectable by satellite observations (Figures 2 and 3) so these high values could be
362 interpreted as observational noise. The detection limit depends on the instrument
363 characteristics and atmospheric state, with IASI minimum detection limit of $\sim 2\text{-}3$ ppbv ($\sim 4\text{-}6 \cdot 10^{15}$
364 $\text{molecules}\cdot\text{cm}^{-2}$) [Clarisse et al., 2010] and CrIS $\sim 0.5\text{-}1.0$ ppbv ($\sim 1\text{-}2 \cdot 10^{15}$ $\text{molecules}\cdot\text{cm}^{-2}$)
365 [Shephard and Cady-Pereira, 2015; Kharol et. al., 2018]. Second, the CHIMERE standardized NH_3
366 columns are enhanced in September 2014, which is not supported by the observations. It has
367 been recently shown that CHIMERE overestimated NH_3 emissions in autumn over Europe
368 [Couvidat et al., 2018]. Generally, the amplitude of the modelled seasonal cycle exceeds the
369 measured ones, which could be explained by higher concentrations measured in winter due to
370 the observational noise and lower emissions.

371 Over the whole period, the coefficient of determination (r^2) between the standardized monthly
372 mean NH_3 columns derived from IASI (CrIS), and the CHIMERE model is 0.58 (0.18) for the
373 annual cycles of 2014 and 2015 (not shown here). If we only consider months of high NH_3 in the
374 domain from March to August, the correlation between the observational datasets and the
375 model is good with linear regression slope values between IASI (CrIS) and CHIMERE of 0.98
376 (0.71), as shown in Figure 7. The seasonal cycle is thus well reproduced by the model, which is
377 encouraging given the fact that annual total emissions are simply disaggregated with a monthly
378 profile in the model. However, the values of the r^2 lower than 0.5 indicate that the CHIMERE
379 model only reproduces at most half of the observed monthly temporal NH_3 variabilities in the
380 domain. Similar variabilities are found between the observations and the model outputs since
381 the coefficients of correlation of the standard deviations are 0.4 and 0.6 between CHIMERE and
382 IASI and CrIS, respectively.

383 3.2.2. Spatial variability of NH_3 in springtime

384 The IASI and CrIS regional maps have been compared to the CHIMERE model for the March-April
385 period in 2014 and 2015 to evaluate the model's capacity to reproduce the spatial distribution
386 of the episodic emissions from fertilizer spreading practices in springtime, as well as their inter-
387 annual variability. Satellite NH_3 measurements in springtime have been gridded at $0.15^\circ \times 0.15^\circ$
388 spatial resolution, and the associated CHIMERE maps have been computed following the
389 coincident criteria described in section 2.4 at the same spatial resolution (Figures 8 and 9).

390 First one can notice that the spatial distribution of NH_3 observed in springtime by both satellite
391 instruments are in good agreement, even though their overpass time is different (~ 4 hours
392 apart). This was already seen in the inter-annual variability agreement seen in Figure 4. In spring
393 2014, IASI and CrIS both reveal three main regions of enhanced NH_3 concentrations (North,
394 Champagne-Ardennes, and Brittany/Pays de la Loire region) already identified by the 10-years
395 and 5-years of IASI and CrIS observation maps (Boxes A, B, and C of Figures 2 and 3). In 2015,
396 concentrations of NH_3 in the northern part of the domain are higher than in 2014, as indicated



397 by both IASI and CrIS observations (Figure 9, upper panels). Overall, satellite observations are
398 able to capture similar spatial distributions of high NH_3 concentrations in springtime, and their
399 evolution in time.

400 In spring 2014, the CHIMERE model reproduces the high concentrations in the three regions of
401 the domain identified in Figures 2 and 3. Additional NH_3 hot spots in the southeastern part of
402 the domain including the Po Valley, Switzerland, and the wine region between Besancon and
403 Lyon (blue box in Figure 8) are indicated by the CHIMERE model. NH_3 emissions in this latter
404 region are comparable to average agricultural plains over France. Only dispersion conditions
405 related to wind speed and boundary layer height can explain high NH_3 concentrations over this
406 area.

407 In spring 2015, satellite observations and the CHIMERE model outputs exhibit very similar
408 patterns in term of high NH_3 distributions, with however higher NH_3 concentrations indicated by
409 the model in the southern part of the domain (blue box in Figure 9).

410 Finally, the (model - observations) differences between the standardized NH_3 column derived
411 from the satellite instruments in springtime 2014-2015 and the corresponding NH_3 columns
412 derived from the CHIMERE model are shown in Figure 2 of the supplement information. One
413 can see that very similar patterns are presented when comparing the model to independent
414 satellite observations from IASI and CrIS: the modelled NH_3 concentrations are systematically
415 lower for both years over Belgium and United Kingdom, and higher in the southern part of the
416 domain (green square, Figure S2) including the Pays de la Loire region (box C in Figures 2 and 3),
417 and in the southeastern part of the domain (over the North part of Switzerland and the Plateau
418 du Jura region - between Besancon and Lyon cities – blue box in Figure 8). Reasons of enhanced
419 NH_3 columns derived from the model in this latter region are not clear yet. An explanation could
420 be that the temporal distribution of the emissions is misrepresented in the model since the
421 modelled concentrations are enhanced in April whereas the two satellite observations are
422 enhanced earlier in March for both years. It is worth noting that there are no EMEP stations
423 measuring surface NH_3 concentrations in these regions. As for the Brittany/Pays de la Loire
424 region, it has already been shown that the LOTOS-EUROS atmospheric model [Schaap et al.,
425 2008] using similar chemistry schemes and NH_3 emissions shows higher columns each year in
426 this area [Van Damme et al., 2014b].

427 3.3. Conditions for PM formation in the Paris megacity

428 To investigate the impact of intensive agriculture practices on the Paris megacity air quality, we
429 need to better understand the role of NH_3 in the formation of $\text{PM}_{2.5}$ that depends, among
430 others, on specific meteorological conditions such as atmospheric temperature and humidity
431 that alter the gas-particle partitioning. The link between high NH_3 concentrations inducing $\text{PM}_{2.5}$



432 formation in the Paris megacity is known [Petetin et al., 2016; Zhang et al., 2013] but
433 quantification of such phenomena is difficult due the lack of long-term NH_3 monitoring in the
434 IdF region. $\text{PM}_{2.5}$ is however measured hourly at several locations in Paris by the Airparif
435 network (<https://www.airparif.asso.fr/>, Figure 1). Thanks to the 10 years of IASI observations,
436 an observational evidence of $\text{PM}_{2.5}$ formation in the IdF region (100 km around Paris - black box
437 in Figure 1) is represented in Figure S3. Simultaneous enhancements in March of $\text{PM}_{2.5}$
438 measured at the surface and NH_3 columns derived from the IASI observations over the IdF
439 region are clearly visible. However, high concentrations of NH_3 observed in summer are not
440 associated with high $\text{PM}_{2.5}$ concentrations. This reflects the complexity of the $\text{PM}_{2.5}$ formation
441 depending on various factors, such as NH_3 emissions, atmospheric chemistry (acidic content of
442 the atmosphere), transport, and specific meteorological conditions involved in the gas to solid
443 phase conversion between NH_3 and ammonium salts.

444 To evaluate the impact of long-range transport on NH_3 levels observed over the Parisian region
445 (IdF) in spring, back-trajectory analysis was performed. In total 231 24-hours back-trajectories
446 ending in Paris (period from February 15th to May 15th from 2013 to 2016) were classified into 8
447 clusters using HYSPLIT (<https://ready.arl.noaa.gov/HYSPLIT.php>). Figure 10 shows the mean
448 trajectories for each cluster associated with the average NH_3 total columns measured by IASI
449 over the IdF region. In this figure, higher NH_3 columns are found under the influence of air
450 masses transported from the northern part of the domain (over Belgium and the Netherlands,
451 clusters 4 and 5) and from the Brittany region (cluster 8), which are the major sources regions of
452 NH_3 in spring in the domain as previously identified (Figures 2 and 3). Clusters 2 and 3 (Figure
453 10) are associated with intermediate NH_3 levels since air masses moved slowly transporting
454 NH_3 -rich air from rural regions near IdF (such as the Champagne-Ardennes region - Box A in
455 Figures 2 and 3) to Paris. Finally, low NH_3 concentrations are measured when air masses
456 originated from ocean regions passing through continental areas with minor NH_3 sources in
457 spring (clusters 1, 6 and 7, Figure 10). This reflects the importance of long-range transport in the
458 NH_3 budget observed over the Paris megacity in spring.

459 To quantitatively assess the influence of meteorological parameters on the formation of $\text{PM}_{2.5}$
460 from NH_3 in the IdF region, timeseries of NH_3 total columns, $\text{PM}_{2.5}$ surface concentrations, and
461 four meteorological parameters (temperature at 2 m, boundary layer height, total precipitation
462 and relative humidity) derived from ECMWF - ERA-Interim [Dee et al., 2011] were analyzed. To
463 compute daily and monthly means, IASI NH_3 total columns have been averaged over IdF (black
464 box in Figure 1), $\text{PM}_{2.5}$ concentrations measured between 9 AM and 11 AM have been averaged
465 over the 14 stations (dark points in Figure 1), and ECMWF data have been averaged over a 300
466 km region around Paris (the blue box in Figure 1). Figure 11 shows all these parameters for
467 spring 2014.



468 We have flagged pollution episodes in both time series ($\text{PM}_{2.5}$ and NH_3) by selecting data above
469 1-sigma standard deviation over the mean of the datasets from 2013 to 2016. This time period
470 was selected to have enough IASI observations in the IdF region. Then two cases have been
471 defined to study the temporal correlation between NH_3 and $\text{PM}_{2.5}$: case A in which both NH_3 and
472 $\text{PM}_{2.5}$ pollution episodes appear simultaneously, i.e. within the same day or 2 days apart
473 (shaded in red in Figure 11); case B in which pollution episodes appear at least 3 days apart
474 (shaded in blue in Figure 11). In Figure 11, a strong relationship between peaks of NH_3 , $\text{PM}_{2.5}$
475 and meteorological parameters can be seen. For example, between March 3rd and March 19th
476 2014 (case A), the boundary layer height is exceptionally low (456 m; compared to 760 m on
477 average); the temperature is relatively low (280 K; 282 K on average); and there is no
478 precipitation (0.01 mm/h; 0.11 mm/h on average). One note that peaks of maximum NH_3
479 observed in IdF on March 11th and 12th are associated with air masses coming from the northern
480 part of the domain (clusters 4 and 5 in Figure 10). In contrast, for the case B in which
481 appearance of peaks of NH_3 and $\text{PM}_{2.5}$ is not simultaneous, meteorological conditions are
482 different: the boundary layer is thicker (908 m on April 23rd 2014), or temperature is higher (285
483 K on April 11th 2014).

484 To further investigate the influence of meteorological parameters on the pollution episodes in
485 the IdF region, detailed analysis have been made over the whole dataset. Figure 12 shows the
486 statistical distribution of meteorological parameters corresponding to case A, case B, and all
487 observations. One can see that the boundary layer height is significantly lower in case A ($550 \pm$
488 205 m) than in case B (751 ± 276 m), and that precipitations are absent in case A (0,019 mm/h)
489 compared to case B (0,085 mm/h). The temperature at 2 meters also differs between the two
490 cases (case A: 278 ± 3 K; case B: 282 ± 4 K), but the humidity is almost the same ($70\% \pm 17\%$
491 versus $75\% \pm 18\%$). Thus the combination of the following three meteorological parameters favors
492 simultaneous appearances of NH_3 and of $\text{PM}_{2.5}$ in Paris (i.e. case A): low surface temperatures
493 (5°C), with thin boundary layers ($\sim 500\text{m}$), and rare precipitations. In addition, the Wilcoxon-
494 Mann-Whitney test ([Wilks, 2011], not shown here) indicates that each single parameter has no
495 significant influence on the NH_3 - $\text{PM}_{2.5}$ correlation. Therefore only a combination of these
496 different parameters has an impact on secondary aerosol formation from NH_3 .

497 An explanation of these findings might be that anticyclonic conditions (low planetary boundary
498 layer), preventing pollutant dispersions in the lower atmosphere [Salmond and McKendry,
499 2005], along with moderate wind fields allow NH_3 plumes to be transported from rural to urban
500 regions [Petit et al., 2015]. In addition, thanks to relatively low atmospheric temperatures and a
501 moderate relative humidity, conversion of gas phase NH_3 to ammonium salts is then
502 accentuated via optimal phase equilibrium [Watson et al., 1994; Nenes et al., 1998]. Finally,
503 with the absence of rain, ammonium salts are stabilized in the aerosols.



504 Our observations are in agreement with previous studies [Bessagnet et al., 2016; Wang et al.,
505 2015], which have shown that the formation of ammonium salt needs a specific humidity of 60 -
506 70%, because it corresponds to the deliquescence point of NH_4NO_3 in ambient air. This is in
507 agreement with our results since the mean of relative humidity in case A is 70%. Our results also
508 support the idea that a relatively low atmospheric temperature favor $\text{PM}_{2.5}$ formation since the
509 phase equilibrium leads to NH_4NO_3 decomposition above 30 °C.

510 4. Conclusions

511 This study focuses on seasonal and inter-annual variabilities of NH_3 concentrations in a 400 km
512 radius-circle area around Paris to assess the evolution of major NH_3 agricultural sources and its
513 key role in the formation of the secondary aerosols that affect air quality over the Paris
514 megacity.

515 Thanks to 10-years and 5-years of regional NH_3 observations derived from IASI and CrIS, three
516 main regions of high NH_3 occurring between March and August were identified. Observed inter-
517 annual variabilities of NH_3 concentrations have been discussed with respect to total
518 precipitations and atmospheric temperature, showing that total precipitations are anti-
519 correlated with high NH_3 concentrations, and that mild temperature in late winter causes
520 precocious fertilizer spreading due to advanced phenological growth stage.

521 To evaluate our knowledge on agricultural emissions in terms of intensity and both spatial and
522 temporal distributions, coincident CHIMERE model outputs have been compared to satellite
523 observations of IASI and CrIS for 2014 and 2015. The annual cycle is well reproduced by the
524 model (correlation slopes of 0.98 and 0.71 between the model and IASI and CrIS, respectively)
525 but the model is only able to reproduce half of the observed atmospheric NH_3 variability.
526 Focusing on spring periods (March-April 2014 and 2015) of episodic NH_3 emissions, the two
527 independent satellite observations derived from IASI and CrIS show very similar spatial
528 distributions of high NH_3 concentrations, as well as their evolution in time. The comparison
529 between CHIMERE NH_3 columns and coincident satellite observations highlights the same
530 difference spatial patterns with a systematic underestimation of NH_3 concentrations from the
531 model over Belgium and an overestimation in the southern part of the domain (French
532 Brittany/Pays de la Loire and Plateau du Jura regions, as well as North of Switzerland).

533 Focusing on the Ile-de-France (IdF, 100 km around Paris) region, we found that air masses
534 originated from rich- NH_3 areas, mainly the northern part of the domain over Belgium and the
535 Netherlands, increase the observed NH_3 total columns measured by IASI over the urban area of
536 Paris.

537 To assess the link between NH_3 and $\text{PM}_{2.5}$ over the Parisian (IdF) region, the main
538 meteorological parameters driving the optimal conditions involved in the $\text{PM}_{2.5}$ formation have



539 been identified. The results show that relatively low temperature, thin boundary layer, coupled
540 with almost no precipitation, favor the PM_{2.5} formation with the presence of atmospheric NH₃ in
541 the IdF region. Based on a more observational approach over large time scale, this work is in
542 agreement with previous studies.

543 This study highlights the need for a better representative NH₃ monitoring to improve numerical
544 simulation of spatial and temporal NH₃ variabilities, especially at fine scales. In order to
545 compare IASI and CrIS data in absolute values, it would be recommended to derive both
546 datasets using the same retrieval algorithm. Thus, by combining these datasets bi-daily NH₃
547 total columns in absolute values at regional scale would be provided. This would help inferring
548 variability of top-down NH₃ emissions. Complementarily, long term quantification of NH₃ diurnal
549 cycle inside Paris would improve comparisons with local PM_{2.5} needed to understand secondary
550 aerosols formations. For this purpose, an ongoing activity consists in the deployment of a mini-
551 DOAS instrument [Volten et al., 2012] used for long-term and continuous monitoring of
552 atmospheric NH₃ concentrations in the center of Paris from the QUALAIR platform
553 (<https://www.ipsl.fr/en/Our-research/Atmospheric-chemistry-and-air-quality/Tropospheric-chemistry/QUALAIR>). Finally, the geostationary-orbit sounder IRS-MTG ([Stuhlmann et al.,
554 2005], to be launched after 2022) will provide NH₃ columns at very high sampling rate (every 0.5
555 hour over Europe) with an unprecedented spatial resolution (pixel size of 4 km).

557

558 Author contribution:

559 CV wrote the paper with contributions of all coauthors. CV and CC designed the study. MV, LC,
560 and SW performed IASI retrievals and ED, MWS, and KEC performed the CrIS retrievals. FM ran
561 the CHIMERE simulations. CV and TW analyzed the data with guidance from CC and PFC. All
562 authors discussed the results and contributed to the final paper.

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578

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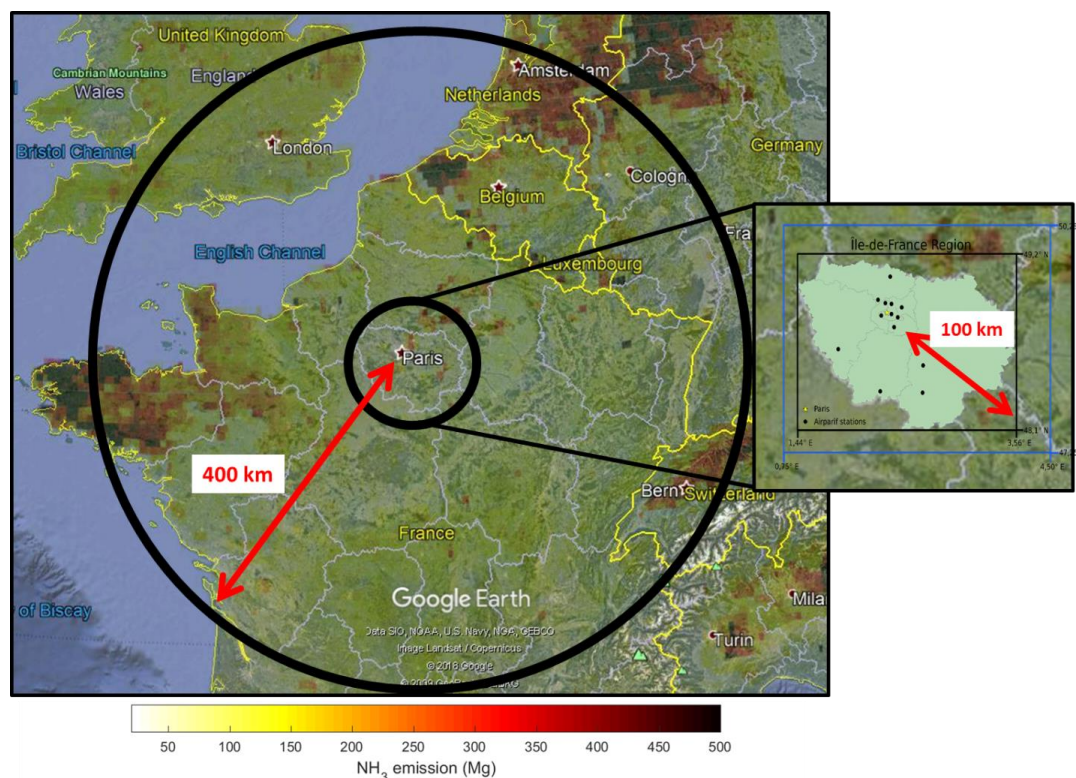


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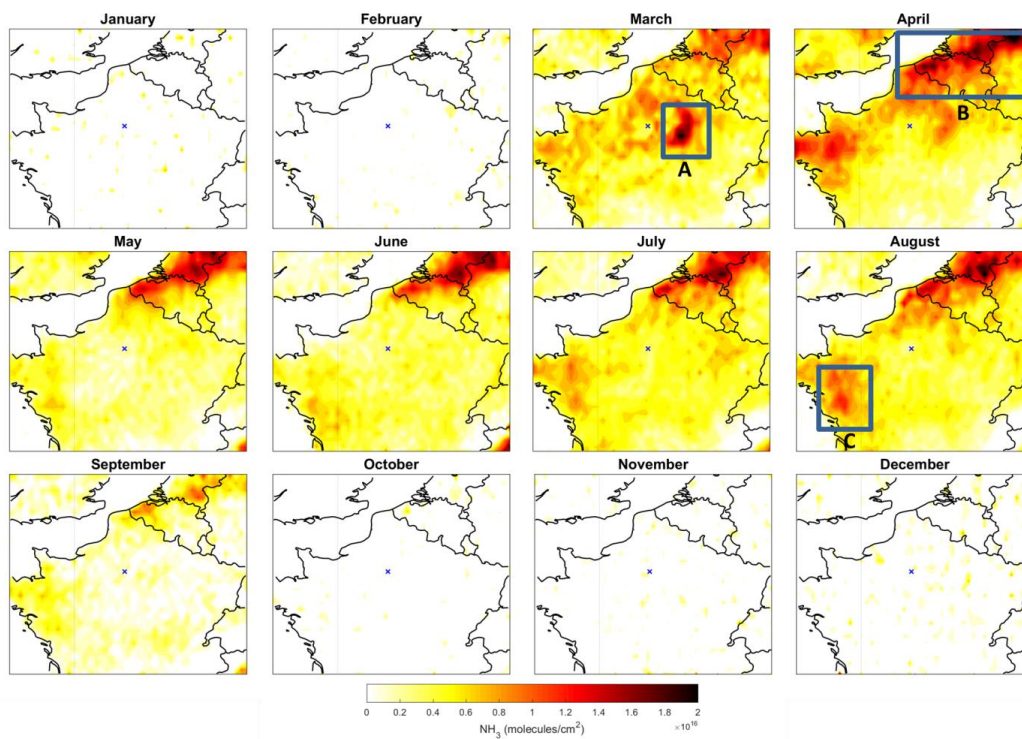
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FIGURES



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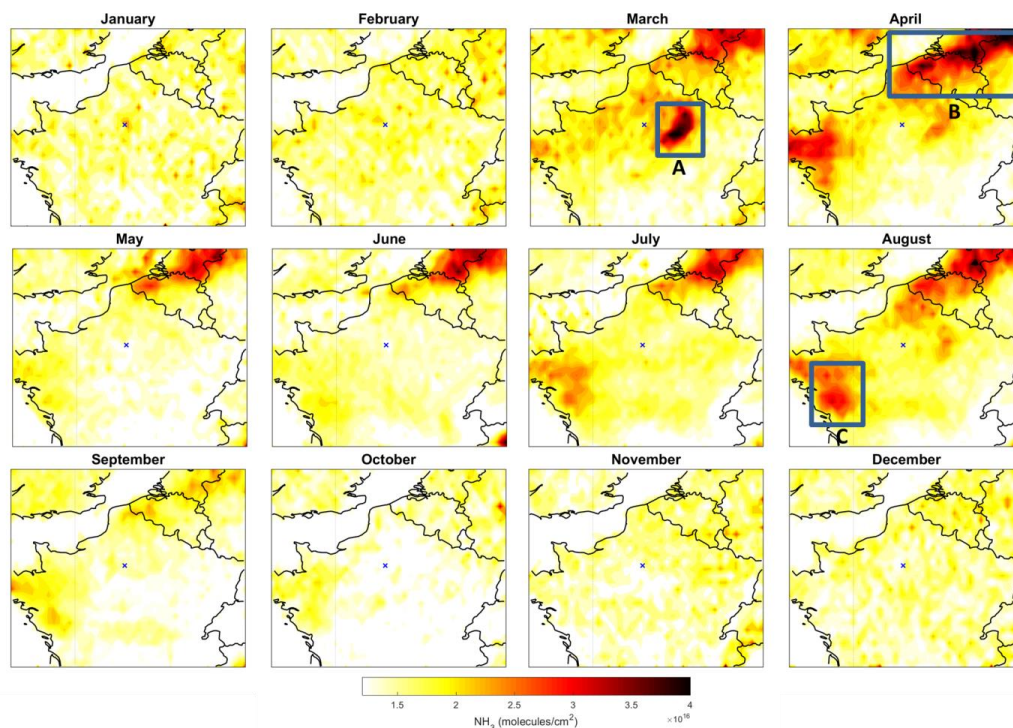
845 Figure 1: Region of analysis: 400 km radius-circle around the Paris megacity and 100 km
846 Paris. The latter is representative of the Ile-de-France (IdF) region where the Airparif PM
847 observational network is located. Black points are the locations of the Airparif stations
848 measuring hourly $\text{PM}_{2.5}$ concentration at the surface. The black (blue) box delimitates the IdF
849 region in which the IASI NH_3 (ECMWF) data have been considered. The overlay represents NH_3
850 emissions (in Mg per year and per cell of $0.1^\circ \times 0.1^\circ$) derived from the EMEP inventory for 2015.



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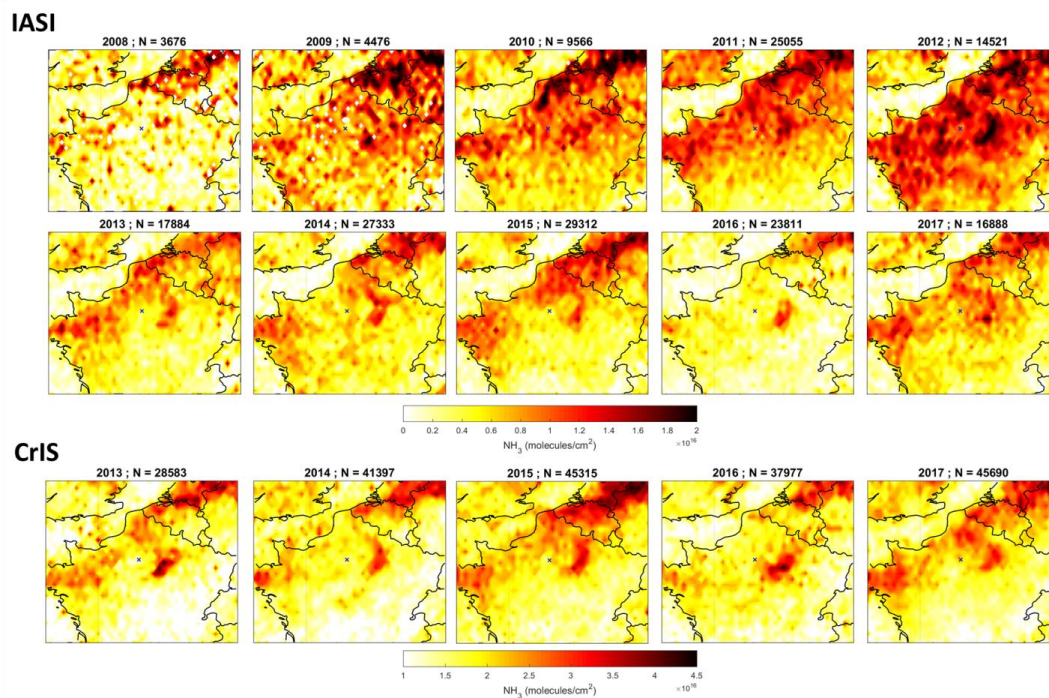
852 Figure 2: Monthly means of NH_3 total columns ($\text{molecules}/\text{cm}^2$) derived from 10 years (2008-
853 2017) of IASI NH_3 -retrieved columns. The blue cross indicates Paris location.

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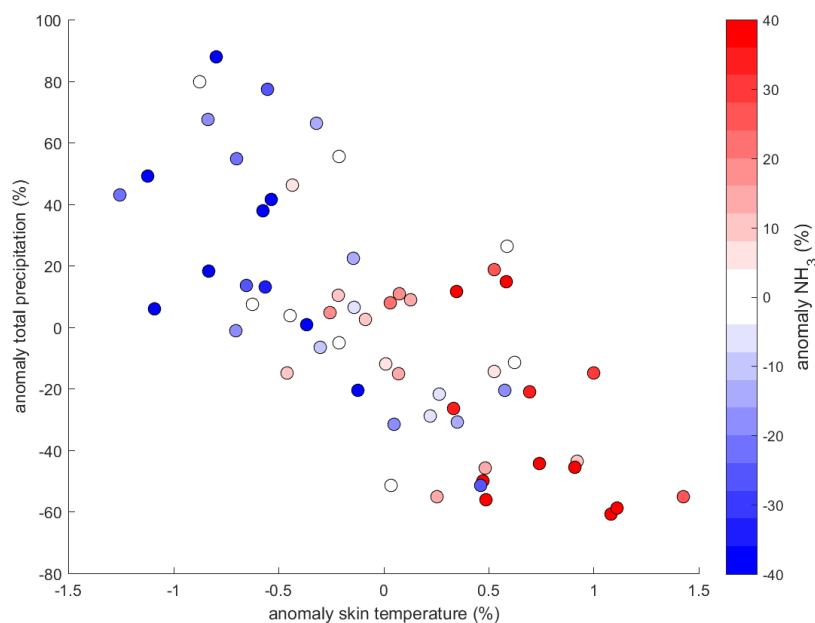
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856 Figure 3: Monthly means of NH_3 total columns (molecules/ cm^2) derived from 5 years (2013-
857 2017) of CrIS NH_3 -retrieved columns. The blue cross indicates Paris location.



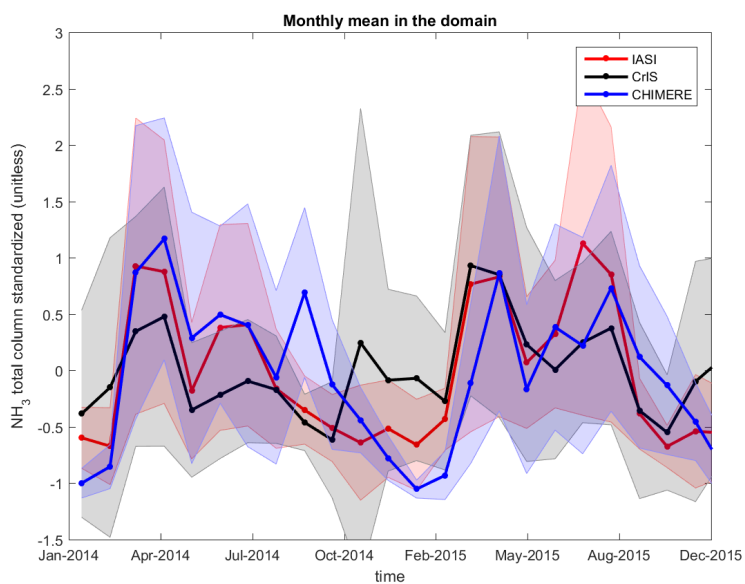
858

859 Figure 4: Maps of monthly mean NH₃ total columns (molecules/cm²) in March-April period
860 derived from IASI from 2008 to 2017 and CrIS from 2013 to 2017.



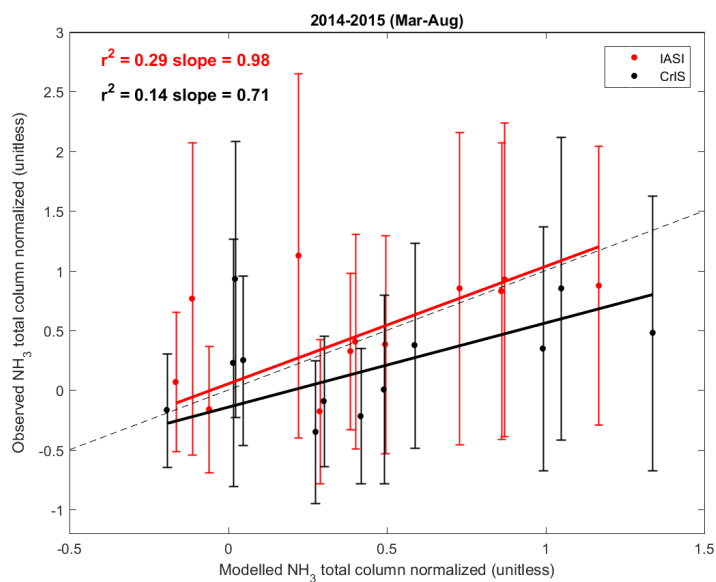
861

862 Figure 5: Scatter plot of monthly mean anomaly (relative to the 10-years – 2008 to 2017 -
863 monthly average) of total precipitation versus skin temperature derived from ECMWF from
864 March to August in the domain, and color coded by the NH₃ total columns anomaly derived
865 from IASI.



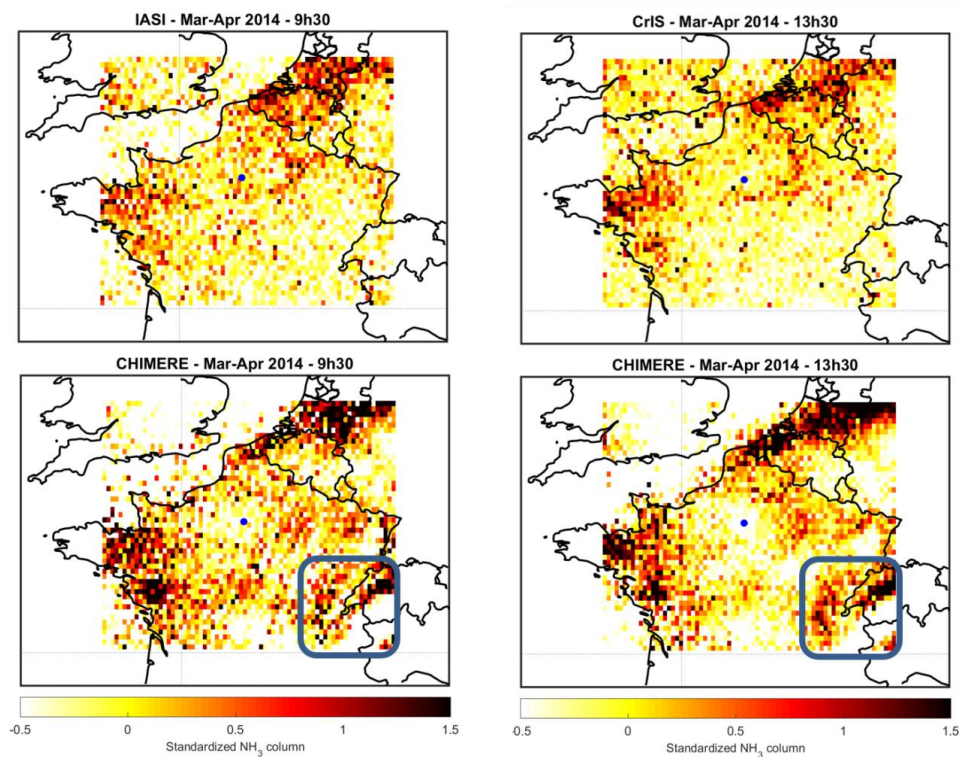
866

867 Figure 6: Standardized monthly mean concentrations derived from IASI (red), CrIS (black) and
868 CHIMERE (blue) for 2014 and 2015. Shaded areas correspond to the one-sigma standard
869 deviation around the means.



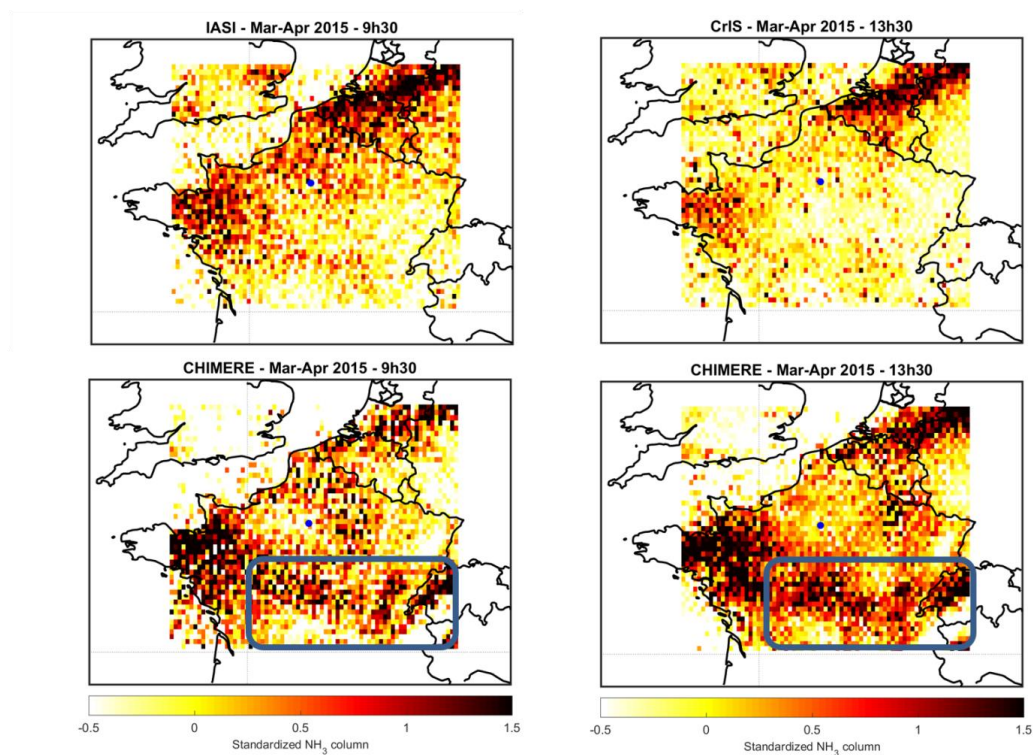
870

871 Figure 7: Correlation plots between monthly means NH_3 standardized concentrations derived
 872 from satellite observations (IASI in red and CrIS in black) and the CHIMERE outputs for the
 873 March to August months of 2014 and 2015. The 1:1 line is represented in the dashed line.



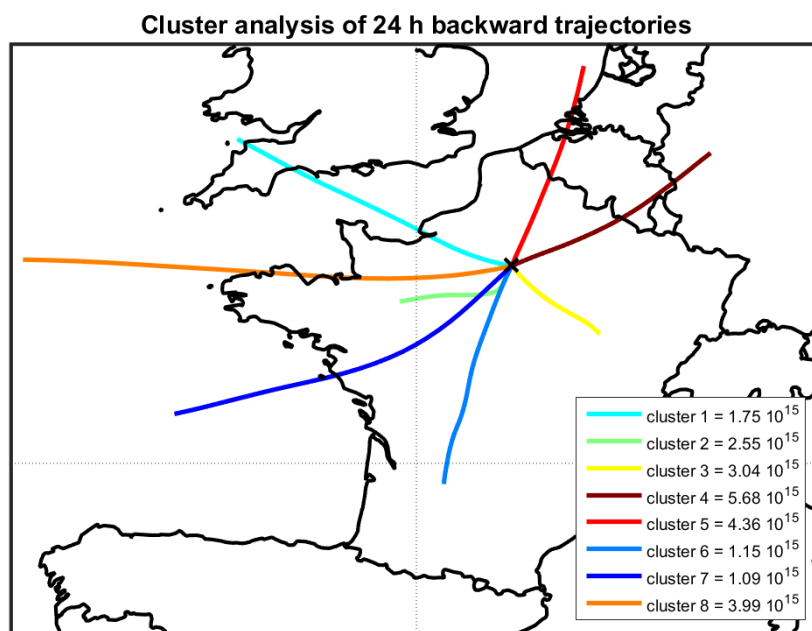
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875 Figure 8: Standardized NH_3 column derived from the satellite instruments (IASI = top left panel,
876 and CrIS = top right panel) and the corresponding NH_3 column derived from the CHIMERE model
877 (coincident with IASI – bottom left panel, and coincident with CrIS – bottom left panel) for
878 March-April 2014. Blue dots indicate Paris location.



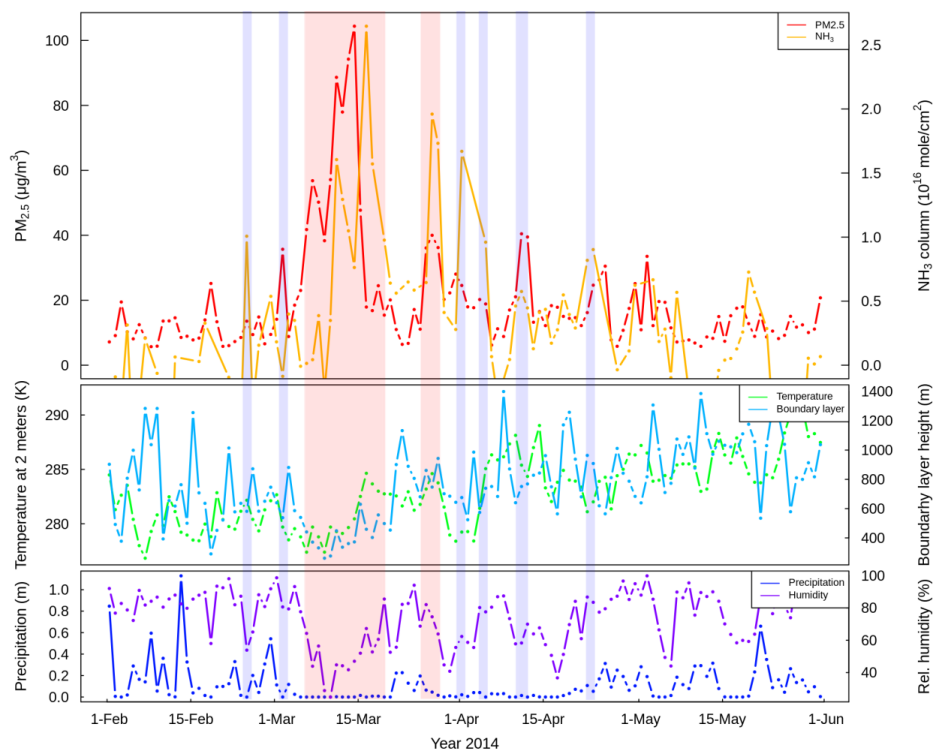
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880 Figure 9: Same than Figure 7 but for March-April 2015.

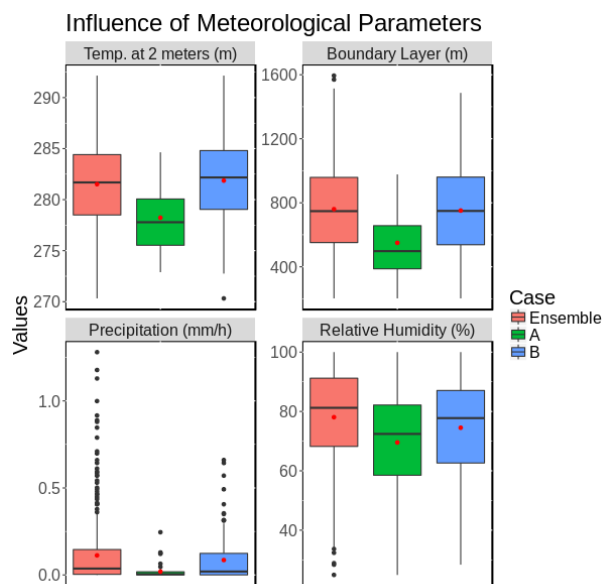


881

882 Figure 10: Cluster analysis of 24-h backward trajectories arriving in spring in Paris (from
883 February 15th to May 15th for the 2013-2016 period) using HYSPLIT-4 model obtained from the
884 NOAA Air Resources Laboratory. Mean trajectories of the 8 clusters are shown in different
885 colors, associated with the NH₃ concentrations measured by IASI in the IdF region (in
886 molecules/cm⁻²).



887
888 Figure 11: Average concentrations of NH_3 total columns derived from IASI (in molecules/ cm^2 ;
889 orange, upper panel) and $\text{PM}_{2.5}$ derived from the Airparif network selected within 2 hours from
890 the IASI overpass (in $\mu\text{g}/\text{m}^3$; red, upper panel) for 2014 as example. Periods of simultaneous
891 (independent) enhancements of NH_3 and PM concentrations are represented with red (blue)
892 areas, i.e. case A (case B). Temperature at 2 meters (in Kelvin; green, middle panel), boundary
893 layer height (in meter; blue, middle panel), precipitation (in meter; dark blue, lower panel), and
894 relative humidity (in percent; purple, lower panel) derived from the ECMWF ERA-interim.



895

896 Figure 12: Statistical distributions of meteorological parameters corresponding to case A, case B,
897 and all observations derived from 2013 to 2016. The medians and the quartiles are presented by
898 center lines and borders of the boxes, respectively. The mean values are indicated by red points,
899 and the extreme values (i.e. those beyond $Q1 - 1.5 \text{ IQR}$ and $Q3 + 1.5 \text{ IQR}$) by black points.



900

TABLE

Satellite	Overpass time (LT)	Time coverage	Nadir spatial resolution (km)	Spectral range (cm ⁻¹)	Spectral resolution (cm ⁻¹)	Spectral Noise* (K) @270K @ 970 cm ⁻¹	References
IASI Metop-A/B	9.30 (AM/PM)	2006-present	12	645–2760	0.5 (apodized)	~0.2	Clerbaux et al., 2009
CrIS Suomi-NPP	1.30 (AM/PM)	2011-present	14	645–1095; 1210–1750; 2155–2550	0.625; (unapodized)	~0.05	Zavyalov et al., 2013

*Spectral noise comparison values in main ammonia spectral region (~970 cm⁻¹) obtained from Zavyalov et al., 2013.

901 Table 1: Instrumental specifications for the IASI and CrIS satellite instruments.