1 Measurement report: Atmospheric new particle formation in a peri-

2 urban site in Lille, Northern France

- 3 Suzanne Crumeyrolle¹, Jenni SS Kontkanen^{2,3}, Clémence Rose⁴, Alejandra Velasquez Garcia^{1,5}, Eric
- 4 Bourrianne¹, Maxime Catalfamo¹, Véronique Riffault⁵, Emmanuel Tison⁵, Joel Ferreira de Brito⁵, Nicolas
- 5 Visez⁶, Nicolas Ferlay¹, Frédérique Auriol¹, Isabelle Chiapello¹
- 6 ¹ Univ. Lille, CNRS, UMR 8518 Laboratoire d'Optique Atmosphérique (LOA), 59000 Lille, France
- 7 ² CSC IT Center for Science, Espoo, Finland
- 8 3 Institute for Atmospheric and Earth system Research, University of Helsinki, Helsinki, Finland
- 9 ⁴ Laboratoire de Météorologie Physique, LaMP-UMR 6016, CNRS, Université Clermont Auvergne, 63178, Aubière, France
- 10 ⁵ IMT Nord Europe, Institut Mines-Télécom, Univ. Lille, Centre for Energy and Environment, F-59000 Lille, France
- 11 ⁶Univ. Lille, CNRS, UMR 8516 LASIRE LAboratoire de Spectroscopie pour les Interactions, la Réactivité et
- 12 l'Environnement, F-59000 Lille, France.

14 Correspondence to: Suzanne Crumeyrolle (suzanne.crumeyrolle@univ-lille.fr)

15 Abstract.

- 16 Formation of Ultrafine particles (UFPs) in the urban atmosphere is expected to be less favored than in the
- 17 rural atmosphere due to the high existing particle surface area acting as a sink for newly-formed particles.
- 18 Des the large Condensation Sink (CS) values, previous comparative studies between rural and urban
- 19 site reported higher frequency of New Particle Formation (NPF) events over urban sites in comparison to
- 20 background sites as well as higher particle formation and growth rates attributed to the higher
- 21 concentration of condensable species. The present study aims to better understand the environmental
- 22 factors favoring, or disfavoring, atmospheric NPF over Lille, a large city North of France and to analyze
- their impact on particle number concentration using a 4-year long-term dataset.
- 24 The results highlight a strong seasonal variation of the NPF occurrences with a maximum observe pring
- spring (27 events) and summer (53 events). It was found that high temperature (T > 295K), low RH (RH<

45 %) and high par radiation are ideal to observe NPF events over Lille. Relatively high values of condensation sink (CS ~2.10⁻² s⁻¹) are reported during event days suggesting that high CS does not inhibit the occurrence of NPF over our site. Moreover, the particle Growth Rate (GR_{15.7-30nm}) was positively correlated with the emperature most probably linked to the higher emissions precursors. Finally, the nucleation strength factor (NSF) was calculated to highlight the impact of those NPF events on particle number concentrations. NSF_{15.7-100} eached a maximum of 4 in summer, indicating an enormous contribution of NPF events to particle number concentration at this time of the year.

1 Introduction

New Particle Formation (NPF) leads to the formation of a large number of sub-20nm particles that will contribute significantly to the levels of fine particles observed in ambient air. These particles can have adverse effect on human health as they can penetrate deeply into the pulmonary system (Clifford et al., 2018; Ohlwein et al., 2019). The freshly-formed particles then grow to larger sizes (Dp > 100 nm) at which they may act as cloud condergion nuclei (CCN, (Pierce and Adams, 2009; Ren et al., 2021; Rose et al., 2017; Spracklen et al., 2006). NPF events have been observed around the world (Kerminen et al., 2018; Kontkanen et al., 2017; Kulmala et al., 2004) in various environments from the boundary layer (BL) at urban locations (Kanawade et al., 2022; Roig Rodelas et al., 2019; Tuch et al., 2006; Wehner and Wiedensohler, 2003) as well as remote polar background areas (Dall'Osto et al., 2018) but also within the free troposphere (Rose et al., 2015b, 2015a). NPF events are typically associated to a photochemical origin, thus occurring mostly during daytime (Kulmala et al., 2014), with some scarce events being observed during nighttime (Roig Rodelas et al., 2019; Salimi et al., 2017).

NPF occurrence depends on various factors including precursor emission strength, number concentration of pre-existing aerosol population, meteorological parameters (in particular solar radiation, temperature and relative humidity) oxide on capacity of the atmosphere (Kerminen et al., 2018). Differences were found in both the season and intensity of NPF events according to the site type (urban, traffic, regional background, rural, polar, high altitude (Dall'Osto et al., 2018; Sellegri et al., 2019)). This variability seems

52 to be related to the environmental conditions that are specific to each location, which makes it hard to 53 draw general conclusions on the conditions that trigger NPF events (Berland et al., 2017; Bousiotis et al., 54 2021). However, Nieminen et al. (2018) highlighted a common seasonal occurrence of NPF during spring 55 and summer using datasets from 36 continental sites worldwide. The formation and growth of initial clusters to detectable sizes (Dp > 1-3 nm) compete with their 56 57 simultaneous removal from the ultra-fine particle mode by coagulation with pre-existing particles 58 (Kerminen et al., 2001; Kulmala, 2003). Because of this, the number concentration of particles smaller 59 than 20 nm has been observed to be anti-correlated with the aerosol volume and mass concentration over 60 rural area in Northern Italy (Rodríguez et al., 2005). Indeed, the total aerosol volume is rather small during NPF events (Kerminen et al., 2018; Rodríguez et al., 2008). While the negative effect of increased pre-61 existing particle surface area (often described with the condensation sink, CS) on the occurrence of these 62 events is widely accepted (Kalkavouras et al., 2017), yet cases are found when NPF events occur on days 63 64 with higher CS corred to average conditions (Größ et al., 2018; Kulmala et al., 2017). A recent study (Bousiotis et al., 2021) used large datasets (16 sites) over Europe (6 countries) and 65 66 highlighted that solar radiation intensity, temperature, and atmospheric pressure had a positive relationship with the occurrence of NPF events at the majority of sites (exceptions were found for the 67 southern sites), either promoting particle formation or increasing growth rate. Indeed, solar radiation is 68 considered one of the most in the occurrence of NPF events, as it contributes to the 69 production of NPF precursors. Higher temperatures are considered vorable for the growth of why 70 formed particles (Dada et al., 2017) as can be linked to increased concentrations of organics vapor 71 72 (Wang et al., 2013) that support particle but also reduce the stability of the initial molecular clusters (Deng e, 2020; Kurtén et al., 2007). 73 74 The wind speed, on the other hand, has presented variable effects on the occurrence of NPF events results, appearing to depend on the site location rather than their type (Bousiotis et al., 2021). Additionally, the 75 76

origin of the incoming air masses plays a very important role, since air masses of different origins have different meteorological, physical and chemical characteristics. Therefore, the probability of NPF event occurrence at a given location and time depends not only on local emissions, but also on long range

77

79 transport (Sogacheva et al., 2007, 2005; Tunved et al., 2006) and on synoptic meteorological conditions 80 at the European scale (Berland et al., 2017). 81 Formation of new particles in the urban atmosphere is expected to be less favored than in the rural 82 atmosphere due to the high existing surface area of particles acting as a sink for freshly-femmed particles. Despite the large CS values, previous comparative studies between rural and urban site reported higher 83 frequency of NPF events (Peng et al., 2017) or sites in comparison to background sites as well 84 85 as higher particle formation and growth rates (Nieminen et al., 2018; Salma et al., 2016; Wang et al., 86 2017) attributed to the higher retation of condensable species. This study presents the first 87 observations of new particle formations over Lille, a large city in the north of France. Based on a multi-

annual dataset (2017-2020), the frequency and intensity of the events are analyzed aiming to better

2 Materials and methods

constrain the favorable and unfavorable conditions.

88

89

90

91

92 The ATOLL (ATmospheric Observations in Lille) station is located in Villeneuve d'Ascq, Northern 93 France (50.6114 N, 3.1406 E, 60 m a.s.l.), and only 6 km away from the city center of Lille, which is the 94 core of the metropolis (Métropole Européenne de Lille with more than 1.1 million inhabitants) to which 95 Villeneuve d'Ascq belongs. Observations such as low Single Scattering Albedo (SSA) values (0.75 on 96 average within the PM₁ fraction, Velasquez-Garcia et al., under review) and large particle number 97 concentrations (6140 cm⁻³ on average) suggest that aerosol measurements performed at ATOLL aerosol 98 eonditions are comparable to Global Atmosp C Watch (GAW) sites classified as urban (Laj et al., 2020; 99 Rose et al., 2021). ATOLL is also part of ACTRIS (Aerosols, Clouds, and Trace gases Research 00 InfraStructure Network, http://www.actris.net) program, complementing the high-quality long-term 01 atmospheric data in Northern France. This station is under the influence of many anthropogenic sources, 02 e.g. road traffic, residential sector, agriculture and industries (Chen et al., 2022), as well as maritime 03 emissions, and more episodically under the influence of events of aged volcanic plumes and desert dust 04 (Boichu et al., 2019; Bovchaliuk et al., 2016; Mortier et al., 2013).

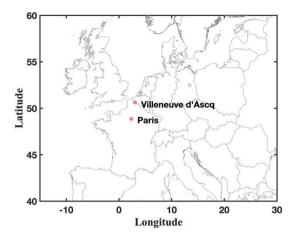




Figure 1: ATOLL location in Villeneuve d'Ascq (Northern France) and picture of the station on the rooftop of the University of Lille P5 building (© LOA).

A large set of *in-situ* and remote sensing instruments are implemented in ATOLL to characterize physical, chemical, optical and radiative properties of particles and clouds. *In-situ* instruments have independent sampling stainless-steel lines located at least 1 meter above the roof top and equipped either with PM₁ cyclone or PM₁₀ inlet. The measurements used for that study were performed between 1st July 2017 and 31st December 2020. The instrument details are described below.

The Scanning Mobility Particle Sizer (SMPS) measures every 5 minutes the particle number size distribution between 15.7-800 nm (divided in more an 100 bins) downstream a Nafion membrane as recommended by ACTRIS standards to keep relative humidity below 40%. The SMPS system consisted of a TSI model 3775-condensation particle counter, a TSI 3081A—type-differential mobility analyzer (DMA) as described by (Villani et al., 2007) and a Nickel aerosol neutralizer (Ni-63 95MBq). The sheath flow rate was controlled with a critical orifice in a closed loop arrangement (Jokinen and Mäkelä, 1997). The scan time was 300 seconds and the particle concentrations were corrected by taking into account charge effects and diffusion calculated using the manufacturer software and algorithms (AIM 10.2.0.11).

21 Accordingly, aerosol number size distribution data from the SMPS measurements were used to classify 22 individual days as NPF event, undefined or non-event days. The classification processe, presented in 23 Dal Maso et al. (2005), is following the decision criteria based on the presence of fine particles (Dp < 25) 24 nm) and their consequent growth to Aitken mode (Dp < 80 nm). Briefly, event days are identified when 25 sub-25nm particle formation and growth are observed. Undefined days are identified when sub-25nm 26 particle formation are observed for more than a ur, but those particles are not growing so the particle 27 diameter remain below 25nm. On non-event days nucleation mode is absent. Finally, undefined days are -28 the days when sub-25nm particles are observed but do not grow subsequently or last less than an hour.

SMPS particle number size distributions were also used for CS (Equation 1, where β_{Mi} is the transitional

correction factor (Fuks and Sutugin, 1970), the Knudsen number is $Kn = 2\lambda_v/d_p$, and α is the

accommodation coefficient and set to unity here) and GR calculations. The CS estimates the loss rate of the condensable vapors (Kulmala et al., 2001) which were assumed to have molecular properties similar to sulfuric acid for CS calculation (Dal Maso et al., 2005). A high CS indicates the properties of large surface area of aerosol particles onto which NPF precursors can condensate. The particle $GR_{15.7-30}$, from

35 15.7 to 30nm, was calculated based on the maximum-concentration method described in (Kulmala et al.,

2012). First, the NPF starting time was identified when the newly formed mode was observable in the first bin of the SMPS (15.7 nm). Then, the time when the concentrations are particles with diameter of 30

nm (N₃₀) peaked was also manually identified. Particle GR_{15.7-30} was then calculated by linear regression

of particle size vs. time span from the NPF start until time when N_{30} reaches a maximum (GR=($D_{p,2}$ –

40 $D_{p,1}$ /(T2-T1)).

29

31

32

33

34

38

41

44

42 $CS = 2\pi D \sum_{i} \beta_{Mi} d_{p,i} N_{i}$

Equation 1

43
$$\beta_{Mi} = \frac{1+K_n}{1+0.337Kn + \frac{4}{3}\alpha^{-1}Kn + \frac{4}{3}\alpha^{-1}Kn^2}$$
 Equation 2

Absorption coefficients (σ_{abs}) were continuously measured with a seven-wavelength aethalometer (AE33,

46 Magee Scientific Inc., Cuesta-Mosquera et al., 2020). According to ACTRIS current guidelines

48 recalculated by 1) multiplying equivalent Black Carbon (eBC) by the mass-specific absorption coefficient 49 (MAC) and then 2) dividing by the suitable harmonization factor to account for the filter multiple 50 scattering effect, i.e. 2.21 (M8020 filter tape) in 2017 and 1.76 (M8060 filter tape) afterwards. The 51 aethalometer samples at 5 L min⁻¹ downstream a PM₁ cyclone (BGI SCC1.197, Mesa Labs). The spectral 52 dependency of σ_{abs} was used to determine the contributions of traffic (fossil fuel - BCff) and wood burning 53 (BCwb) to eBC via a source apportionment model (Sandradewi et al., 2008). 54 Meteorological data including temperature, water vapour mixing ratio, and solar radiation were also measured every minute at the sampling using a weather station (DAVIS Inc weather station, Vantage 55 56 Pro 2). Solar radiation at the surface are measured every minute at the sampling site using a set of Kipp 57 & Zonen pyranometer (CM22, for dime fluxes using a sphere shadower) and Normal Incidence 58 pyrheliometer (CH1 for direct fluxes), the solar radiation being then calculated as the sum of the diffuse 59 and direct fluxes. A sky imager (Cloudcam, CMS) is a camera equipped with a fisheye lens to cover the 60 entire upper half sphere. The cloud cover is estimated from an algorithm, named Findclouds and provided by the manufacturer, comparing the different values of the red, green and blue components of each pixel 61 of the images taken (Shukla et al., 2016). 62

(https://actris-ecac.eu/particle-light-absorption.html), σ_{abs} coefficients at each wavelength have been

Three-day air mass backtrajectories of air masses arriving at ATOLL at half the boundary layer height between July 1, 2017 and December 31, 2020 were computed every hour using the Hybrid Single-Particle Lagrangian Integrated Trajectory (HYSPLIT version 5.1.0) transport and dispersion model from the NOAA Air Resources Laboratory (ARL) (Rolph et al., 2017; Stein et al., 2015) and meteorological input from the Global Data Assimilation System (GDAS) at 1×1° resolution, resulting in 30719

47

63

64

65

66

67

69 3 Results

70

94

3.1 NPF event frequency

71 The seasonal distribution of NPF events at the ATOLL site is displayed in Figure 2. SMPS missing data 72 (in Figure 2) are about 40 % from January to April due to the yearly calibrations at the manufacturer 73 premises and few laboratory campaigns (Oct 2018 – Jun 2019). Over the 4 years of measurements (2017-2020). 96 (11 %) days were classified as NPF events, 355 (40 %) as undefined days and 432 (49 %) as 74 non-event days. The can also note that most of the NPF events identified at the ATOLL site were observed 75 during spring (March-April-May, 27 events corresponding to 15 % of the days when observations were 76 77 available during this season) and summer (June-July-August, 53 events corresponding to 19 %) with a 78 maximum observed in June consistent with a previous study over central Europe (Dall'Osto et al., 2018). 79 During winter, the number of events is extremely limited (only one event observed in February). In the 80 following sections, only observations from spring and summer seasons will be discussed due to the low 81 representativeness of NPF events in fall (n=15) and winter (n=1). Moreover, the undefined event days are 82 seen all year round (frequency around or larger than 20 %) with a clear peak in August (frequency at 62.5 83 %) consistent with observations over boreal forest where undefined days were also observed to be most 84 frequent in early fall (Mazon et al., 2009). 85 Using long-term measurements from 36 sites (polar, rural, high altitude, remote and urban), Nieminen et 86 al., (2018) reported an annual NPF frequency below 15 % for half of the sites (18 sites from all types) 87 and occasionally over 30 % for 10 sites. Moreover, they highlighted a seasonal variation of NPF occurrence with larger (lower) frequency, about 30 % (10 %), during spring (winter). Frequency analysis 88 89 of NPF occurring only over urban or anthropogenically influenced sites show large site-to-site differences for all seasons. Indeed, NPF occurrence frequencies are varying from 20% (Helsinki in Finland, 5a) 90 Paulo in Brazil) to 80 % (Beijing in China, Marikana in South Africa) during spring and from 7 % 91 (Helsinki) to 78 % (Marikana) during winter. Yelly averages of PF occurrence frequencies are between 92 93 11 % (Helsinki) and more than 60 % (Beijing and Marikana).

The ATOLL event frequency (seasonal variation and values) is similar to observations performed in Paris (Dos Santos et al., 2015) while the frequency of undefined and non-event days are quite different, Indeed, in Paris the non-event frequency is larger than 60 % except in July and August whereas over ATOLL the non-event frequency shows a clear seasonal pattern with lower frequency (<40 %) from April to August. Moreover, undefined event frequency in Paris shows a minimum (<5 %) in May and June and remains quite steady during the rest of the year (around 20 %). One can note that the frequency of undefined events (also considered as failed events) is much higher over ATOLL all year long with an average of 40 % while it remains below 40 % over the boreal forest. The quency of undefined events observed at ATOLL is clearly larger than the frequencies observed over more polluted site (Paris) and a pristine site (Boreal forest). This might show that ATOLL is under the influence of air masses or particle and precursor sinks that favor the burst of UFP.

.03

.04

.05

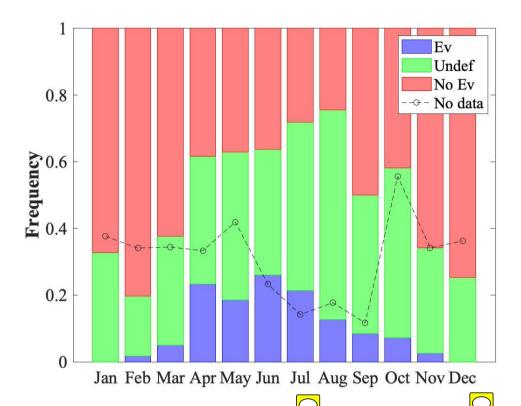


Figure 2: sonal distribution of event days (Diue), undefined days (green), and non-event (rea) days at the ATOLL station, Lille, France, during 2017–2020. Days with missing data are excluded from the total number of days per month and the frequency of missing data are indicated with the black circles.

3.2 Aerosol number size distribution

.08

.09

10

11

.12

13

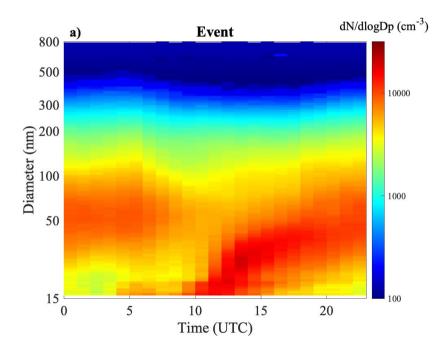
14

.15

Median daily contour plots of the particle number size distributions (PNSD) obtained from the SMPS are shown in Figure 3 separately for NPF event (around 800 PNSD), undefined (around 2300 PNSD) and non-event (around 1700 PNSD) days observed during the warm period (only spring summer). The PNSD were first selected then averaged to one-hour time resolutions median filtering. Atmospheric NPF and subsequent particle growth are seen in Figure 3a as an emergence of new aerosol particles with small diameter followed by the growth of these particles into larger sizes. If this phenomenon is taking



place regionally (few tens of km in radius), a so called 'banana plot' is observed in particle number size distributions as a function of time at a fixed location. The time evolution of the "median NPF day" ure 3a) displays a similar growth pattern for newly formed particles than for individual NPF event days (see supplementary materials). Indeed, one can clearly see a UFP mode appearing from 10:00 to 15:00 (UTC) and growing during the rest of the day. The NPF starting time and the growth rate will be discussed in the following section. By 23:00 UTC, the newly formed particles reach an average diameter of 50 nm, similar to the median diameter of the morning of the pre-existing particles observed during the morning (00:00 – 08:00). The "median undefined day" (Figure 3b) highlights a burst of UFP from 10:00 to 15:00 (UTC) that is not growing and does not last during the whole afternoon. The pavior of the median is again similar to the individual undefined events observed during this period. The "median non-event day" (Figure 3c) shows no sign of particle growth, as expected.



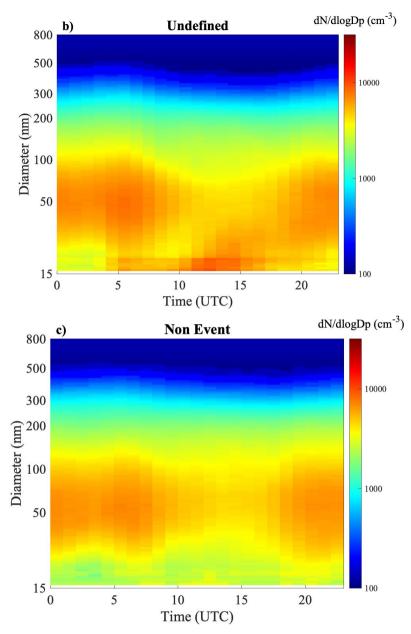


Figure 3: Hourly median particle number size distribution (15.7 nm<Dp<800 nm) observed during NPF event (a), undefined (b) and non-event (c) days in spring and summer from 2017-2020.

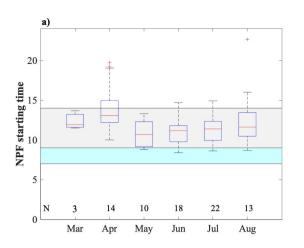
3.3 NPF starting time and growth rate

30

47

48

31 Figure 4 shows the monthly variation of the starting time and growth rate of each event observed at ATOLL. Most NPF events observed in ATOLL were observed to start tween 09:00 and 14:00 UTC .32 (74 %), with fewer events starting in the early morning (07:30 – 09:00UTC, 6 %) and late afternoon .33 (15:00 and 19:30, 20 %). NPF starting time as well as $GR_{15.7-30nm}$ strongly depend on the month during 34 which the event is observed. I ed, the NPF starting time occurs later during spring (also true for fall 35 and winter) and reaches a minimum in May and June (around 08:20). Nocturnal events are rarely 36 observed, with only one occurrence in August. No diurnal NPF event were observed after 16:00 UTC in 37 38 summer. During spring and fall, the average NPF starting time varies between 10:00 and 19:00 UTC. The 39 start time monthly variability is linked to sunrise and sunset times. In the following section, a link between 40 the total solar irradiation and NPF occurrence will be examined. The event ending time was determined as the time when the growth of the freshly formed particles was 41 42 over, i.e. when the diameter reached the diameter of the pre-existing particles. The duration of nucleation events, at ATOLL, was then estimated and varies from an hour up to 28 hours. On average, NPF durator 43 44 is shorter from May to around 8 hours) and increases up to around 13 hours on average during March and November. This seasonal behavior could be due to the presence or availability of condensable 45 46 vapors, air mass origin, and environmental conditions favorable to NPF events (see section 3.2).



.52

.53

.55

.62

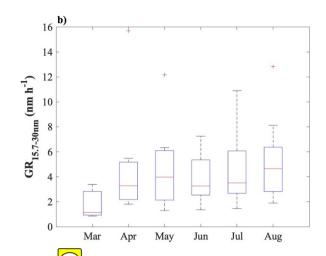


Figure 4: Monthly variation of new particle formation arting time (a) and their Growth Rate (GR_{15.7-30nm}) at the ATOLL station during 2017–2020. The grey area represents the period, from 09:00-14:00, when most of the NPF events occur. The blue area corresponds to the period before the NPF onset (07:00- 09:00). N represents the number of events observed per month.

The Growth Rate from 15.7 to 30nm (GR_{15.7-30nm}) values observed at ATOLL lie within 0.8 to 15.7 nm.h⁻¹ and show a strong monthly variation with the lowest values observed in spring (and fall, not shown here). The largest median values are observed in May and August while the 75th percentile highlights larger values of GR_{15.7-30nm} during summer (Figure 4b). GR_{15.7-30nm} values were in addition plotted as a function of temperature for all years and seasons in Figure 5, which highlights that below 20°C, GR_{15.7-30nm} values are lower than 6 nm.h⁻¹, while, under warn conditions (T >20 °C), GR_{15.7-30nm} reach values up to 16 nm.h⁻¹. These results show a clear temperature dependence of the particle growth. Indeed, higher temperatures have been shown to favor emission of biogenic precursors, including monoterpenes known to favor the occurrence of NPF events (Kulmala 1., 2004). Previous studies (Paasonen et al., 2018; Yli-Juuti et al., 2011) have shown that the growth rate usually has larger values during warm periods and especially during summer. Over urban areas such as Beijing or Shangai, GR_{15-25nm} showed no clear seasonal variation (Yao et al., 2018). However, recent studies also have highlight the link with growth rate seasonal pattern and high abundance of biogenic volatile organic compounds during warmer periods (spring and summer) over boreal forest (Paasonen et al., 2018; Yli-Juuti et al., 2011). Therefore, the

observed seasonal variation of GR_{15.7-30nm} may be related to emissions of organic compounds in the atmosphere varying as a function of temperature variation (Figure 5). This hypothesis is supported by the larger contribution of Organics during NPF event days observed in Figure S3 (Supplementary material).

.72

As previously observed in Figure 3a, the median diameters reached by the end $\frac{1}{3}$ NPF events are similar and averaged around 50 nm. Moreover, the seasonal variation of the NPF event duration could be, then, linked to the $\frac{1}{3}$ GR seasonal variation. As the final diameter is similar in all cases, the lower the $\frac{1}{3}$ GR values will then be associated with the longer NPF duration. The seasonal variation of NPF duration highlighted earlier could then only be a consequence of the $\frac{1}{3}$ GR seasonal variation.

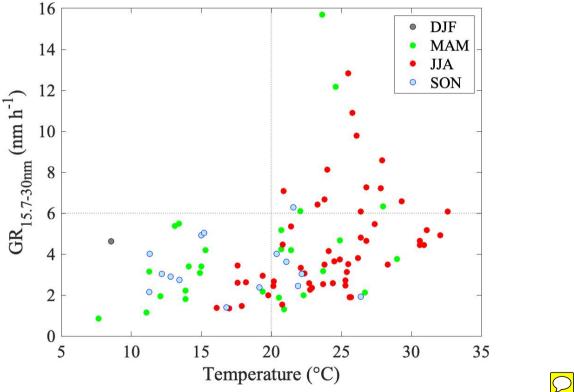


Figure 5: Growth Rate (GR_{15.7-30nm}) values as a function of ambient temperature for different seasons.

3.4 Environmental conditions

.77

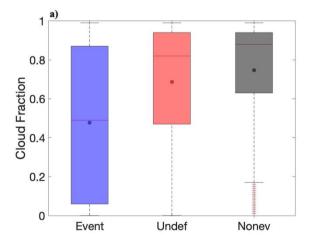
.78

.80

.81

.92

The effect of cloudiness on NPF event occurred is shown in Figure 6a, with a specific focus on measurements collected between 09:00 and 14:00, i.e. the period of time where most NPF tended that. The cloud fraction was calculated from the sky imager dataset following the method by (Shukla et al., 2016) and sorted as a function of event, undefined and non-event days. There is a clear inverse prelation between cloud fraction and NPF occurrences. Average cloud fraction is around 0.47 during event days, 0.68 during unconed days and 0.74 during non-event days. Moreover, the 25th percentiles of the cloud fractions for event, undefined and non-event days, respectively 0.06, 0.47, 0.63, clearly show that the absence of clouds (lower cloud fraction) is mostly associated with NPF events. It is consistent with previous analysis performed over the boreal forest (Dada et al., 2017) and is linked to the fact that radiation seems essential for NPF during the warmer period (spring and summer), as the events occur almost solely during daylight purs (Kulmala et al., 2004). Figure 6b shows the average diel total solar radiation observed during events, non-event and undefined days for spring and summer. As expected, total solar radiation is on average always larger during event days in comparison to non-event days, with a more pronounced difference observed during spring.



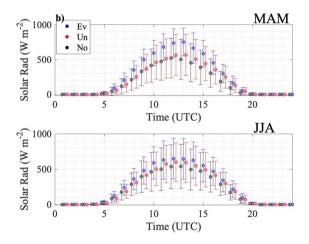


Figure 6: (a) Cloud fraction observed from 09:00 to 14:00 UTC during event, undefined and non-event days. The red line represents the median while the lower and upper edges of the box represent the 25th and 75th percentiles, respectively. The lower and upper edges of the whisker represent 10th and 90th percentiles, respectively. The circles represent the average. (b) Diel variations (UTC) of the mean total solar radiation observed during the event days (blue), undefined days (red) and non-event days (black) during spring (MAM, top) and summer (JJA, bottom) seasons (b). The error bars correspond to one standard deviation.

.93

94

.95

96

97

98

99

00

-01

02

03

04

05

06

07

08

09

10

11

12

13

14

Other environmental parameters known to influence the occurrence of NPF events, such as temperature and humidity were also sorted to highlight diel and seasonal various (Figure 7). Our results (Figure 7a) indicate that NPF is favored by low values of ambient relative humidity, especially during spring, consistently with previous studies (Duplissy et al., 2016; Hamed et al., 2011; Merikanto et al., 2016). A few reasons can explain this tendency: (1) high RH values (RH > 90 %) observed at the surface are usually associated to the presence of low altitude clouds reducing incoming total radiation and then preventing NPF formation, (2) at moderately high RH (RH >40 %), hydrophilic aerosols could grow which will enlarge the sink for precursors and (3) high RH values may limit the formation of some Volatile Organic Compounds (VOC) through ozonolysis reactions, inhibiting the formation of condensable vapors necessary for condensation (Fick et al., 2003; Tillmann et al., 2010). Figure 7b shows the diel median temperature conditions (T) during NPF events, nonevents and undefined days. NPF events occurred within temperatures ranging between 3°C and 33.5°C. During bom seasons, averaged temperatures during event days are always larger than during non-event days, with, again larger differences during spring. One should note that days with high temperatures in spring and summer are usually also days with high solar radiation, consistently with conclusions from Figure 6. The temperature difference between undefined days and event days is clearly marked during spring and fade away during summer. As previously discussed, higher temperatures fargemission of biogenic precursors, including monoterpenes known to favor the occurrence of NPF event (Kulmala et al., 2004). Isoprene emission is also larger at higher temperature but according to Heinritzi et al., (2020) it

. Moreover, high temperature can also lead to evaporation of molecular clusters which may inhibit NPF events (Dada et al., 2017; Deng et al., 2020).

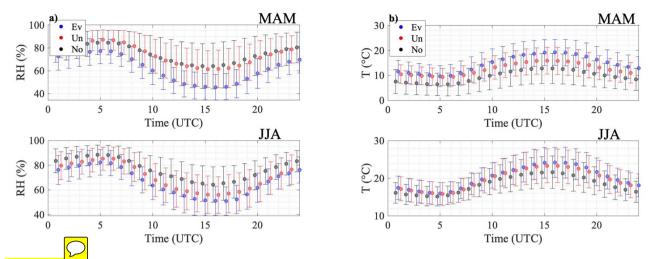


Figure 7: Diel variation (UTC) of mean Relative Humidity (RH, a) and mean temperature (b) observed during the event days (blue), undefined days (red) and non-event days (black) during spring (MAM) and summer (JJA) seasons. The error bars correspond to one standard deviation.

3.5 Condensation sink

The CS characterizes the loss rate of atmospheric variety to aerosol particles. The diel variations of CS calculated for spring and summer and for NPF event, undefined and non-event days are shown in Figure 8a. Hourly averaged CS values are high (larger than 2×10^{-2} s⁻¹) during vent days occurring during spring and summer (Figure 8a). During NPF event days and over different urban sites (Beijing, Nanjing or Hong Kong), CS values ranging from 0.6 up to $10.7.10^{-2}$ s⁻¹ were reported (Xiao et al., 2015). For pristine sites, such as Hyytiälä, the CS values are between $0.05 - 0.35\times10^{-2}$ s⁻¹. As events occur anyway, low values of CS, often considered as the major limiting factor in the NPF occurrence do not inhibit the occurrence of NPF events in ATOLL consistently to previous observatory (Größ et al., 2018) or over Chinese megacities (Xiao et al., 2015). One can assume that the presence of large concentrations of precursors could explain the formation of particles over polluted sites such as ATOLL. Unfortunately, precursors were not measured over the 4-year period of interest here therefore this assumption would require further investigation beyond the scope of this study. Recent studies (Marten et al., 2022; Wang et al., 2020), performed in the CLOUD chamber demonstrate that the

presence of nitric acid (HNO₃) and ammonia (NH₃), typical within urban environment, contribute to freshly particles survival by increasing dramatically their growth rate.

In the afternoon S during event days increases due to the growth of freshly emitted particles, especially during summer. Contribution of newly formed particles (Dp < 50 nm) to the CS is about 36 % and 27 %, during summer and spring respectively, while the contribution of pre-existing particles (Dp > 150 nm) to the CS is below 20 % for both seasons. Moreover, during non-event days, the size resolved median CS is shifted to larger particle diameters with a maximum observed around 100 nm for all seasons.

To period the impact of the background CS on NPF occurrence, all CS values obsered from 07:00-09:00, period before NPF starting time (green area on Figure 4a), were averaged during event, non-event and undefined days. It was found that the total CS_{07-09h} was larger (around 16 %) during non-event days in comparison to undefined and event days. Moreover, this difference is mostly due to particles larger than 70 nm according to size resolved CS_{07-09h} (Figure 8b). The difference between non-event and event days is lower than what is usually observed over pristine sites (Lyubovtseva et al., 2005) but significant enough to trigger the NPF event occurrence.



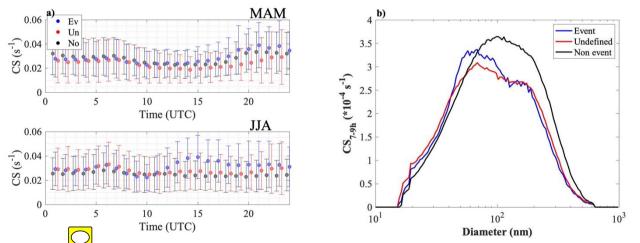


Figure 8: (a) Diel variation of Concensation Sink (CS) during spring (MAM) and summer (JJA) seasons. (b) Median size resolved CS for MAM and JJA during event days (blue), undefined (red) and non-event days (black).

45

- Moreover, Du et al (2022) highlights the emical composition of the particles that contribute to the CS 46 47 during period when the NPF occur the most (10:00-15:00) over Beijing. As the CS observed over 48 ATOLL is largely influenced by the freshly formed particles, the chemical composition of the particle as a function of CS will be presented dum; two specific periods: before (07:00 – 09:00) and ming (09:00 49 - 14:00) the period when the NPF occur the most for NPF event and non-event days (Figure 1). During 50 51 non-event days, both periods (07:00-09:00 and 09:00 - 14:00) highlight similar mass fraction of all compounds with on average 41% Organis, 16% of nitrate, 21% of sulfate, 11% of ammonium, less than 52 1% of chloride and around 10% of Black carbon. As the aerosol sources during non-event days are 53 54 supposed to be the same throughout the day, this result was actually expected.
- The same comparison (before -07:00-09:00- and during -09:00-14:00- the NPF periods), for event days, 55 56 highlight a larger contribution of Organics to the CS during the NPF period (54% in average) in 57 comparison to the period right before the start of the NPF events (46% in average). Indeed, for large 58 values of CS (> 0.045) the contribution of Organics is larger than 50%, reaching a maximum of 69% for 59 CS of 0.085 s⁻¹. This result suggests that organic vapors may be involved particle growth.
- 60 The comparison of chemical mass fraction before the NPF period (Figure 2a) during NPF event (top) and non-event days (bottom) highlight a decrease of sulfate mass fraction down to 15% before and during the 61 NPF period of event days in comparison to non-event days. Therefore, the most effective aerosol chemical -62 63 composition for removal of gaseous compounds needed for NPF would be with less organics and higher 64 SO₄.

These results are different from pu et al (2022) showing a large increase of nitrate and a decrease of organics with the CS values for event and non-event days.

68 69

65

66

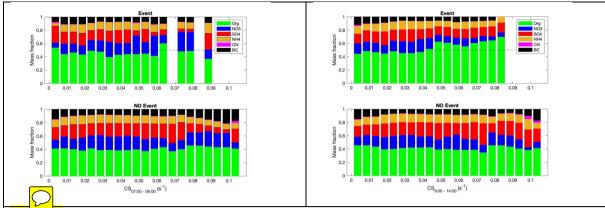


Figure 1: Mass fraction of the major compound measured before (07:00 – 09:00) and during (09:00 – 14:00) NPF period at the ATOLL station during event and non-event days. The black dashed line corresponds to 50% mass fraction.

Additionally, the correlation coefficients between meteorological parameters and pollutants (gas and particles) are reported in Table 1 for the entire period of measurements (all seasons). Hourly average over a time window between 09:00 - 14:00 (NPF event starting time period) of few variables (total CS, T, RH and BCwb) were used to calculate those correlation coefficients (corresponding to 7025 and 35433 data points for NPF event and Non-event days, respectively).

The correlation of Black Carbon from wood burning (BC_{wb}) during non-event days with the condensation sink is high (R = 0.67). This collation between these parameters is clearly absent during event days (R = 0.19). One can also note that NOx concentrations have a positive correlation (0.30) with CS during NPF non-event days while the same correlation is negative (-0.17) during NPF event days. The NOx sources over urban area are mostly anthropogenic (house heating, traffic and industries) sources, which is consistent its relatively high correlation coefficients with BC_{wb} (0.47 and 0.65). As highlighted in (Barreira et al., 2020), BC_{wb} and NOx are evolving through the year showing a minimum in summer and

a maximum in winter when sources are stronger due to colder temperatures and residential heating emissions. As non-event days are mostly (62 %) observed during cold months (fall and winter) and NPF events are largely (82 %) observed during warmer months (spring and summer), the correlation between BC_{wb}, NOx and CS during non-event is not surprising. However, during spring, air masses observed during NPF events are clearly "cleaner" (in terms of NOx and BC_{wb}) than non-event cases. Indeed, NO_x and BC_{wb} concentrations are lowered by 18 % and 36 % respectively during spring NPF event days in comparison to non-event days. During summer, NO_x and BC_{wb} concentrations reach an annual minimum and there both pollutant concentrations are similar between NPF event and non-event days (lowered by 0.04 % and 0.01 % during NPF event days).

Table 1: Correlation coefficients between different meteorological projects F, RH), Nitrogen oxide (NOx), Black carbon concentrations (BCwb from wood burning and total condensation sink during event and non-event for the 4 years period (2017-2020) and in a time window (09:00 – 14:00). High positive or negative correlations are marked in bold.

		CS	Т	RH	NOx	BCwb
Event days	CS	1				
	Т	0.55	1			
	RH	-0.39	-0.40	1		
	NOx	-0.17	-0.24	0.48	1	
	BC _{wb}	0.19	-0.04	0.11	0.47	1
Non- event days	CS	1				
	Т	0.06	1			
	RH	-0.03	-0.50	1		
	NOx	0.30	-0.44	0.44	1	
	BC _{wb}	0.67	-0.37	0.28	0.65	1

Moreover, during event days the temperature is positively correlated (0.55) with the CS, while, during non-event days, this correlation is clearly not observed during non-event days (0.06). Over boreal forest, CS and temperature are correlated during event day (Liao et al., 2014). Indeed, this coupling comes from the enhanced growth of particles due larger monoterpene emissions at higher temperature, which naturally leads to higher concentration of larger particles and thus higher CS. As the particle growth during event days is clearly related to temperature increase (Figure 5) most probably due to higher concentration of condensable gasses, it is not surprising to observe this temperature and CS coupling.

3.6 Air mass trajectories

simultaneously (35 % of cases).

01

02

03

04

05

06

07

.08

09

-10

-11

.12

13

.14

.15

16

.17

.18

19

20

21

22

23

24

25

26

One can note that environmental conditions (CS, Temperature and RH) observed during undefined events are mostly between event and non-event days. A deeper analysis on undefined days reveals that on these days, particle growth stopped due to (i) a decrease of the total irradiance due to a cloud passage over the site (20 % of cases), (ii) a shift of the wind direction (17 % of cases), (iii) or both parameters changing

The shift of the wind orientation leading to a stop of the particle growth indicates that NPF events are associated with certain wind directions or air mass origins. To investigate this, HYSPLIT back trajectories were first sorted as a partition of event, non-event and undefined days. Only the back-trajectories arriving between 09:00-14:00 (period of NPF high occurrences) were selected for further analysis. During the NPF events, the predominant air masses were tracked back along the Eastern North Sea region. Comparing these results to back trajectories during non-event days highlight more continental influence. Indeed, most of the back trajectories during non-event days pass over large cities (Dunkirk, Paris, London, Rotterdam) before reaching Lille metropolis. Those air masses might then have been slightly enriched in primary precursor vapors. This result is consistent with primary precursor vapors. This result is consistent with primary precursor vapors air masses are associated with NPF event cases observed during spring.

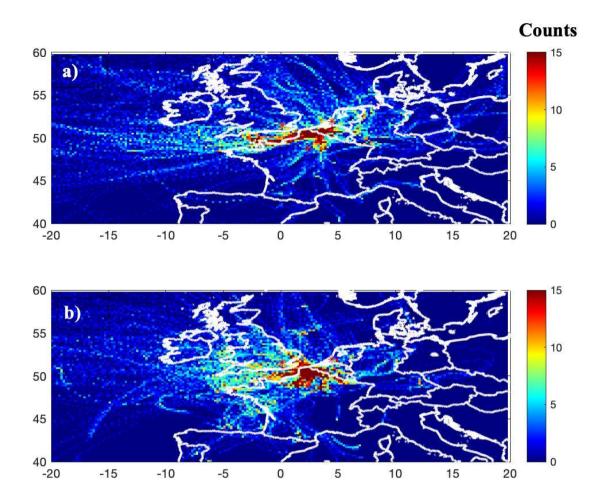


Figure 9: 3 days hourly back trajectories arriving in ATOLL between 09:00-14:00 UTC during (a) New Particle Formation (NPF) events and (b) non-events days. The back trajectories were calculated for each hour at ATOLL at half the boundary layer height. The blor contour represents the back trajectories crossing counts in each grid cell (resolution 0.2° 0.2°).

3.7 Nucleation strength factor

The nucleation strength factor (NSF_{15.7-100}) is calculated as the ratio of fine to accumulation particle concentrations observed during nucleation dazer the same ratio observed during non-event day (Salma et al., 2017). Fine and accumulation mode particle number concentrations ($N_{15.7-100}$ and $N_{100-800}$) were

retrieved from the SMPS data. The limited atmospheric residence time of fine particles (typically lower than 10 h) means that a large portion of the N_{15.7} concentration can also be related to local emissions and/or formation processes, including NPF events. On the contrary, due to a longer residence within the atmosphere (up to 10 days), $N_{100-800}$ is more related to large spatial and temporal scales. Therefore, the numerator represents the increase of $N_{<100}$ relative to $N_{100-800}$ caused by all sources while the denominator represents the same property due to all sources except NPF. The NSF method is based on the hypothesis that aerosol sources are similar from day to day of from season to season, excepting the sporadic occurrence of NPF. Considering the large number of event (96) and non-event (432) days used to calculate NSF_{15.7-1}pne can assume that the sporadic/occasional (i.e. not observed on daily basis) sources of UF particles other than NPF events (e.g. volcanic plumes) have little impact on the NSF_{15,7-100} in comparison to the sources always active (such as traffic, industries etc...). NSF is generally used to better assess the contribution of NPF to fine particle number concentrations (represented by N<100) relative to the regional background particle number concentrations. If the NSF \approx 1, then the relative contribution of NPF to particle number concentration with respect to other sources is negligible, like in Granada (Spain) urban site (Casquero-Vera et al., 2021). Moreover, Salma et al. (2017) also defined two thresholds for NSF₆₋₁₀₀ to description as a single source: a considerable contribution ($1 < NSF_{6-100} < 2$) or larger than of any other source sectors together (NSF_{6-100} >2). One should keep in mind that these thresholds were defined accordingly to the lower cut off diameter originally set at 6nm. As the lower cut off diameter used in this study is a bit larger (15.7 nm instead of 6nm) than the one used by Salma et al. (2017), the calculated NSF_{15,7-100} would necessarily be underestimated in comparison to NSF₆₋₁₀₀ from Salma et al. (2017). The hourly median of fine to accumulation particle concentration ratio was computed for NPF event and non-event days. Figure 10 shows the NSF_{15,7-100} diel variation observed at the ATOLL platform over 4 years of measurements. During spring, the NSF_{15.7-100} factor remound quite constant (about 1.5) during night and morning and peaks at 16:00 UTC to reach a maximum at 2.5. This indicates that NPF has a significant effect on particle number concentration only a few (2-3) hours after the averaged NPF starting time. During summer, the

32

-33

34

35

36

37

38

39

40

41

42

43

44

45

46

47

48

49

-50

51

-52

53

-54

-55

-56

57

-58

tendency of the NSF_{15,7-100} is quite similar with a unique peak at 13:00 UTC (again 2-3 hours after the

59 averaged NPF starting time). At that time the median NSF_{15.7-100} values reach 4 while from 21:00 to 06:00 60 UTC the NSF_{15.7-100} remains low (averaged at 1.08). Therefore, during summer, the NPF contribution to particle number concentration is extremely high from 10:00 to 18:00 and then negligeable for the rest of 61 62 the day in comparison to other sources. Such NSF₁₀₋₁₀₀ variations were observed in other European cities (Budapest, Vienna and Prague) with 63 64 maximum reaching 2.7, 2.3 and 3.4 respectively with a lower cut-off diameter set at 10nm (Németh et al., 65 2018). Moreover, Salma et al. (2017) reported NSF₆₋₁₀₀ peaks at midday varying from 2.2 and 2.7 for 66 Budapest city center and from 2 to 7.2 for near city background for each season with NSF₆₋₁₀₀ maximum 67 reached during winter. The nucleation frequency during winter in Budapest is low (<10 %), similarly to 68 our observations, however, the impact of these limited number of events on particle number 69 concentrations is high. For the record, the NSF_{15,7-100} factor peaked at 3.5 and 2.3 during winter and fall .70 respectively. As previously shown (Sebastian et al., 2021), NPF events can also play a major role on Earth's radiative 71 budget when the newly formed particles grow to climate-relevant sizes (50-100nm). In order to .72 .73 understand the NPF influence on these particles the NSF₅₀₋₁₀₀ was also calculated (see supplementary 74 figures). The results show a large increase up to 1.6 of the NSF_{50-100} in the early afternoon for both seasons. .75 This suggest a potential impact on the CCN concentration that needs to be further studied with proper 76 instrumentation.

.77

.78

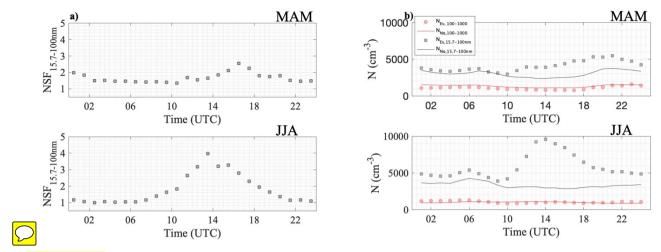


Figure 10: (a) Diel variation of the Nucleation Strength Factor (NSF_{15.7-100}) during MAM and JJA calculated from number concentration during the 2017-2020 period. (b) Diel variation of particle number concentrations (N) for each season within the diameter ranges from 15.7 to 100 nm (N_{15.7-100}, black) and from 100 1000 nm (N₁₀₀₋₁₀₀₀, red) at the ATOLL site during the 2017-2020 period. The dots correspond to event days while the line correspond to non-event days.

This study was based on 4-years (2017- 2020) measurements performed at the ATOLL site, in the close

4 Conclusions

.79

vicinity of the city of Lille, Northern France. This paper is dedicated to studying New Particle Formation (NPF) occurrence over a peri-urban site. The results highlight a strong sponal variation of the NPF event frequency, with a maximum occurrence observed during spring (15 %) and summer (19 %). Lee undefined cases, which correspond to bursts of UFP that do not grow, are much more frequent (40 % on average) than NPF events all year long. The highest frequency (68 %) is observed in August and the lowest one (17 %) in February. The interruption of the particle growth during undefined events can be mostly attributed to changes of environmental conditions (irradiance and wind direction).

Seasonal variation of NPF parameters was also clearly observed and associated with environmental parameters. High temperature (T > 295K), low RH (RH< 45 %) and high solar radiation favor the occurrence of NPF events at ATOLL. The presence of clouds, linked to a decrease of solar radiation, is limiting the NPF event occurrences. Moreover, NPF events start earlier in the morning during from May

to September most probably related to variations in sunrise time. The Growth Rate calculated between 15.7 and 30 nm (GR_{15.7-30nm}) ranges from 1.8 nm.h-1 in March up to 10.9 nm.h-1 in July. The GR_{15.7-30nm} was also found to be positively correlated with temperature. This correlation might be related to larger emissions of biogenic pursors at higher temperatures, including monoterpenes known to favor the occurrence of NPF event (Kulmala et al., 2).

.98

.99

Relatively high values of Condensation Sink (averaged CS > 2.10^{-2} s⁻¹) are reported during NPF events as well as during non-event days. These results suggest that high CS values are not limiting the NPF event occurrence, consistent with recent studies focusing on NPF events over urban sites (Deng et al., 2020; Hussein et al., 2020; Pushpawela et al., 2018). Looking more closely before the NPF onset (from 07:00 – 09:00 UTC), CS_{07-09h} values are larger by 16 % during non-event days. Interestingly, CS tends to increase during event days (especially in sumper) and size resolved CS clearly shows a peak shift from 150 nm during non-event days to 50 nm during event days highlighting the strong contribution of newly formed particles on CS.

Air masses trajectories (HYSPLIT) arriving over ATOLL during event days highlight a specific path along the Eastern North Sea region with only a small fraction passing over any continental area and therefore not crossing many anthropogenic sources, while, most of the back trajectories during non-event days pass over large cities (Dunkirk, Paris, London, Rotterdam) before reaching Lille. The precursor vapor concentration and probably their nature might differ from both "clean" and "polluted" air masses and therefore promote or inhibit NPF event occurrences, a point which requires further investigation.

The impact of NPF events on particle number concentrations has been estimated through the nucleation strength factor (NSF; Salma et al., 2017). The NSF_{15.7-100nm} diel variation was calculated for spring and summer occurring 2 to 3 hours after the average NPF starting time and reaching 1.5 and 4 during spring and summer respectively. The extremely large NSF_{15.7-100nm} value observed during summer highlights the very high NPF contribution to the fine parties (Dp < 100 nm) number concentration in comparison to other regional sources. Recently, (Ren et al., 2021) highlighted the strong impact of newly formed particles from NPF on Cloud Condensation Nuclei (CCN) especially at sites close to anthropogenic sources, such as ATOLL. In future studies, the impact of local vertical dynamics such as the effect of

boundary layer dynamics as in Lampilahti et al. (2020 and 2021) as well as the CCN enhancement factor will be analysed.

2223

24

20

21

Acknowledgements

25 This research was supported by the French national research agency (ANR) under the MABCaM (ANR-16-CE04-0009) 26 contract. Part of the instrumental system has been financially supported by the CaPPA project (Chemical and Physical 27 Properties of the Atmosphere), which is funded by the French National Research Agency (ANR) through the PIA (Programme 28 d'Investissement d'Avenir) under contract "ANR-11-LABX-0005-01", and by the Regional Council "Hauts-de-France". «This 29 work has benefited from the support of the research infrastructure ACTRIS-FR, registered on the Roadmap of the French 30 Ministry of Research. The authors also thank the Région Hauts-de-France, and the Ministère de l'Enseignement Supérieur et 31 de la Recherche (CPER Climibio), and the European Fund for Regional Economic Development for their financial support. 32 The authors gratefully acknowledge the NOAA Air Resources Laboratory (ARL) for the provision of the HYSPLIT transport 33 and dispersion model and/or READY website (https://www.ready.noaa.gov) used in this publication. We thank Francois

34

3536

Data availability

Thieuleux for ECMWF data sharing during this work.

- 38 ATOLL measurements are available through the EBAS database (https://ebas.nilu.no) and SMPS data
- 39 before 2020 through https://doi.org/10.5281/zenodo.6794562. GDAS files for back-
- 40 trajectory calculation are available at https://www.arl.noaa.gov/hysplit/hysplit/. NOx data are
- 41 available from the ATMO open data website: https://data-atmo-hdf.opendata.arcgis.com.

42 43

44

References:

46

- 47 Barreira, L.M.F., Helin, A., Aurela, M., Teinilä, K., Friman, M., Kangas, L., Niemi, J.V., Portin, H.,
- 48 Kousa, A., Pirjola, L., Rönkkö, T., Saarikoski, S., Timonen, H., 2020. In-depth characterization of
- 49 submicron particulate matter inter-annual variations at a street canyon site in Northern Europe (preprint).
- 50 Aerosols/Field Measurements/Troposphere/Chemistry (chemical composition and reactions).
- 51 https://doi.org/10.5194/acp-2020-908
- Berland, K., Rose, C., Pey, J., Culot, A., Freney, E., Kalivitis, N., Kouvarakis, G., Cerro, J.C., Mallet, M.,
- 53 Sartelet, K., Beckmann, M., Bourriane, T., Roberts, G., Marchand, N., Mihalopoulos, N., Sellegri, K.,
- 54 2017. Spatial extent of new particle formation events over the Mediterranean Basin from multiple ground-
- 55 based and airborne measurements. Atmospheric Chem. Phys. 17, 9567-9583.
- 56 https://doi.org/10.5194/acp-17-9567-2017
- 57 Boichu, M., Favez, O., Riffault, V., Petit, J.-E., Zhang, Y., Brogniez, C., Sciare, J., Chiapello, I., Clarisse,
- 58 L., Zhang, S., Pujol-Söhne, N., Tison, E., Delbarre, H., Goloub, P., 2019. Large-scale particulate air
- 59 pollution and chemical fingerprint of volcanic sulfate aerosols from the 2014–2015 Holuhraun flood lava
- 60 eruption of Bárðarbunga volcano (Iceland). Atmospheric Chem. Phys. 19, 14253-14287.
- 61 https://doi.org/10.5194/acp-19-14253-2019
- Bousiotis, D., Brean, J., Pope, F.D., Dall'Osto, M., Querol, X., Alastuey, A., Perez, N., Petäjä, T.,
- 63 Massling, A., Nøjgaard, J.K., Nordstrøm, C., Kouvarakis, G., Vratolis, S., Eleftheriadis, K., Niemi, J.V.,
- Portin, H., Wiedensohler, A., Weinhold, K., Merkel, M., Tuch, T., Harrison, R.M., 2021. The effect of
- 65 meteorological conditions and atmospheric composition in the occurrence and development of new
- 66 particle formation (NPF) events in Europe. Atmospheric Chem. Phys. 21, 3345-3370.
- 67 https://doi.org/10.5194/acp-21-3345-2021
- 68 Bovchaliuk, V., Goloub, P., Podvin, T., Veselovskii, I., Tanre, D., Chaikovsky, A., Dubovik, O., Mortier,
- 69 A., Lopatin, A., Korenskiy, M., Victori, S., 2016. Comparison of aerosol properties retrieved using
- 70 GARRLiC, LIRIC, and Raman algorithms applied to multi-wavelength lidar and sun/sky-photometer
- 71 data. Atmos Meas Tech 9, 3391–3405. https://doi.org/10.5194/amt-9-3391-2016
- 72 Casquero-Vera, J.A., Lyamani, H., Titos, G., Minguillón, M.C., Dada, L., Alastuey, A., Querol, X.,
- 73 Petäjä, T., Olmo, F.J., Alados-Arboledas, L., 2021. Quantifying traffic, biomass burning and secondary
- source contributions to atmospheric particle number concentrations at urban and suburban sites. Sci. Total

- 75 Environ. 768, 145282. https://doi.org/10.1016/j.scitotenv.2021.145282
- 76 Chen, G., Canonaco, F., Tobler, A., Aas, W., Alastuey, A., Allan, J., Atabakhsh, S., Aurela, M.,
- 77 Baltensperger, U., Bougiatioti, A., De Brito, J.F., Ceburnis, D., Chazeau, B., Chebaicheb, H.,
- Daellenbach, K.R., Ehn, M., El Haddad, I., Eleftheriadis, K., Favez, O., Flentje, H., Font, A., Fossum, K.,
- 79 Freney, E., Gini, M., Green, D.C., Heikkinen, L., Herrmann, H., Kalogridis, A.-C., Keernik, H., Lhotka,
- 80 R., Lin, C., Lunder, C., Maasikmets, M., Manousakas, M.I., Marchand, N., Marin, C., Marmureanu, L.,
- Mihalopoulos, N., Močnik, G., Necki, J., O'Dowd, C., Ovadnevaite, J., Peter, T., Petit, J.-E., Pikridas,
- M., Matthew Platt, S., Pokorná, P., Poulain, L., Priestman, M., Riffault, V., Rinaldi, M., Różański, K.,
- 83 Schwarz, J., Sciare, J., Simon, L., Skiba, A., Slowik, J.G., Sosedova, Y., Stavroulas, I., Styszko, K.,
- 84 Teinemaa, E., Timonen, H., Tremper, A., Vasilescu, J., Via, M., Vodička, P., Wiedensohler, A., Zografou,
- 85 O., Cruz Minguillón, M., Prévôt, A.S.H., 2022. European Aerosol Phenomenology 8: Harmonised
- 86 Source Apportionment of Organic Aerosol using 22 Year-long ACSM/AMS Datasets. Environ. Int.
- 87 107325. https://doi.org/10.1016/j.envint.2022.107325
- 88 Clifford, S., Mazaheri, M., Salimi, F., Ezz, W.N., Yeganeh, B., Low-Choy, S., Walker, K., Mengersen,
- 89 K., Marks, G.B., Morawska, L., 2018. Effects of exposure to ambient ultrafine particles on respiratory
- 90 health and systemic inflammation in children. Environ. Int. 114, 167-180.
- 91 https://doi.org/10.1016/j.envint.2018.02.019
- 92 Cuesta-Mosquera, A., Močnik, G., Drinovec, L., Müller, T., Pfeifer, S., Minguillón, M., Björn, B.,
- 93 Buckley, P., Dudoitis, V., Fernández-García, J., Fernández Amado, M., Brito, J., Flentje, H., Heffernan,
- 94 E., Kalivitis, N., Kalogridis, C., Keernik, H., Marmureanu, L., Luoma, K., Wiedensohler, A., 2020.
- 95 Intercomparison and characterization of 23 Aethalometers under laboratory and ambient air conditions:
- Procedures and unit-to-unit variabilities. https://doi.org/10.5194/amt-2020-344
- 97 Dada, L., Paasonen, P., Nieminen, T., Buenrostro Mazon, S., Kontkanen, J., Peräkylä, O., Lehtipalo, K.,
- 98 Hussein, T., Petäjä, T., Kerminen, V.-M., Bäck, J., Kulmala, M., 2017. Long-term analysis of clear-sky
- 99 new particle formation events and nonevents in Hyytiälä. Atmospheric Chem. Phys. 17, 6227-6241.
- 00 https://doi.org/10.5194/acp-17-6227-2017
- Dal Maso, M., Kulmala, M., Riipinen, I., Wagner, R., Hussein, T., Aalto, P.P., Lehtinen, K.E.J., 2005.
- Formation and growth of fresh atmospheric aerosols: Eight years of aerosol size distribution data from

- O3 SMEAR II, Hyytiälä, Finland. Boreal Environ. Res. 10, 323–336.
- Dall'Osto, M., Beddows, D.C.S., Asmi, A., Poulain, L., Hao, L., Freney, E., Allan, J.D., Canagaratna,
- M., Crippa, M., Bianchi, F., de Leeuw, G., Eriksson, A., Swietlicki, E., Hansson, H.C., Henzing, J.S.,
- Granier, C., Zemankova, K., Laj, P., Onasch, T., Prevot, A., Putaud, J.P., Sellegri, K., Vidal, M., Virtanen,
- A., Simo, R., Worsnop, D., O'Dowd, C., Kulmala, M., Harrison, R.M., 2018. Novel insights on new
- 08 particle formation derived from a pan-european observing system. Sci. Rep. 8, 1482.
- 09 https://doi.org/10.1038/s41598-017-17343-9
- Deng, C., Fu, Y., Dada, L., Yan, C., Cai, R., Yang, D., Zhou, Y., Yin, R., Lu, Y., Li, X., Qiao, X., Fan,
- 11 X., Nie, W., Kontkanen, J., Kangasluoma, J., Chu, B., Ding, A., Kerminen, V.-M., Paasonen, P.,
- Worsnop, D.R., Bianchi, F., Liu, Y., Zheng, J., Wang, L., Kulmala, M., Jiang, J., 2020. Seasonal
- 13 Characteristics of New Particle Formation and Growth in Urban Beijing. Environ. Sci. Technol. 54, 8547–
- 8557. https://doi.org/10.1021/acs.est.0c00808
- Dos Santos, V.N., Herrmann, E., Manninen, H.E., Hussein, T., Hakala, J., Nieminen, T., Aalto, P.P.,
- Merkel, M., Wiedensohler, A., Kulmala, M., Petäjä, T., Hämeri, K., 2015. Variability of air ion
- concentrations in urban Paris. Atmospheric Chem. Phys. 15, 13717–13737. https://doi.org/10.5194/acp-
- 18 15-13717-2015
- Duplissy, J., Merikanto, J., Franchin, A., Tsagkogeorgas, G., Kangasluoma, J., Wimmer, D., Vuollekoski,
- H., Schobesberger, S., Lehtipalo, K., Flagan, R.C., Brus, D., Donahue, N.M., Vehkamäki, H., Almeida,
- J., Amorim, A., Barmet, P., Bianchi, F., Breitenlechner, M., Dunne, E.M., Guida, R., Henschel, H.,
- Junninen, H., Kirkby, J., Kürten, A., Kupc, A., Määttänen, A., Makhmutov, V., Mathot, S., Nieminen, T.,
- Onnela, A., Praplan, A.P., Riccobono, F., Rondo, L., Steiner, G., Tome, A., Walther, H., Baltensperger,
- U., Carslaw, K.S., Dommen, J., Hansel, A., Petäjä, T., Sipilä, M., Stratmann, F., Vrtala, A., Wagner, P.E.,
- Worsnop, D.R., Curtius, J., Kulmala, M., 2016. Effect of ions on sulfuric acid-water binary particle
- formation: 2. Experimental data and comparison with QC-normalized classical nucleation theory:
- BINARY PARTICLE FORMATION EXPERIMENTS. J. Geophys. Res. Atmospheres 121, 1752–1775.
- https://doi.org/10.1002/2015JD023539
- Fick, J., Pommer, L., Nilsson, C., Andersson, B., 2003. Effect of OH radicals, relative humidity, and time
- on the composition of the products formed in the ozonolysis of α -pinene. Atmos. Environ. 37, 4087–

- 4096. https://doi.org/10.1016/S1352-2310(03)00522-3
- Fuks, N.A., Sutugin, A.G., 1970. Highly Dispersed Aerosols. Ann Arbor Science Publishers.
- Größ, J., Hamed, A., Sonntag, A., Spindler, G., Manninen, H.E., Nieminen, T., Kulmala, M., Hõrrak, U.,
- Plass-Dülmer, C., Wiedensohler, A., Birmili, W., 2018. Atmospheric new particle formation at the
- research station Melpitz, Germany: connection with gaseous precursors and meteorological parameters.
- 36 Atmospheric Chem. Phys. 18, 1835–1861. https://doi.org/10.5194/acp-18-1835-2018
- Hamed, A., Korhonen, H., Sihto, S.-L., Joutsensaari, J., Järvinen, H., Petäjä, T., Arnold, F., Nieminen,
- T., Kulmala, M., Smith, J.N., Lehtinen, K.E.J., Laaksonen, A., 2011. The role of relative humidity in
- 39 continental new particle formation. J. Geophys. Res. 116, D03202.
- 40 https://doi.org/10.1029/2010JD014186
- Heinritzi, M., Dada, L., Simon, M., Stolzenburg, D., Wagner, A.C., Fischer, L., Ahonen, L.R.,
- 42 Amanatidis, S., Baalbaki, R., Baccarini, A., Bauer, P.S., Baumgartner, B., Bianchi, F., Brilke, S., Chen,
- D., Chiu, R., Dias, A., Dommen, J., Duplissy, J., Finkenzeller, H., Frege, C., Fuchs, C., Garmash, O.,
- Gordon, H., Granzin, M., El Haddad, I., He, X., Helm, J., Hofbauer, V., Hoyle, C.R., Kangasluoma, J.,
- Keber, T., Kim, C., Kürten, A., Lamkaddam, H., Laurila, T.M., Lampilahti, J., Lee, C.P., Lehtipalo, K.,
- Leiminger, M., Mai, H., Makhmutov, V., Manninen, H.E., Marten, R., Mathot, S., Mauldin, R.L.,
- Mentler, B., Molteni, U., Müller, T., Nie, W., Nieminen, T., Onnela, A., Partoll, E., Passananti, M., Petäjä,
- T., Pfeifer, J., Pospisilova, V., Quéléver, L.L.J., Rissanen, M.P., Rose, C., Schobesberger, S., Scholz, W.,
- Scholze, K., Sipilä, M., Steiner, G., Stozhkov, Y., Tauber, C., Tham, Y.J., Vazquez-Pufleau, M., Virtanen,
- A., Vogel, A.L., Volkamer, R., Wagner, R., Wang, M., Weitz, L., Wimmer, D., Xiao, M., Yan, C., Ye,
- P., Zha, Q., Zhou, X., Amorim, A., Baltensperger, U., Hansel, A., Kulmala, M., Tomé, A., Winkler, P.M.,
- Worsnop, D.R., Donahue, N.M., Kirkby, J., Curtius, J., 2020. Molecular understanding of the suppression
- of new-particle formation by isoprene. Atmospheric Chem. Phys. 20, 11809–11821.
- https://doi.org/10.5194/acp-20-11809-2020
- Jokinen, V., Mäkelä, J.M., 1997. Closed-loop arrangement with critical orifice for DMA sheath/excess
- flow system. J. Aerosol Sci. 28, 643–648. https://doi.org/10.1016/S0021-8502(96)00457-0
- Kalkavouras, P., Bossioli, E., Bezantakos, S., Bougiatioti, A., Kalivitis, N., Stavroulas, I., Kouvarakis,
- G., Protonotariou, A.P., Dandou, A., Biskos, G., Mihalopoulos, N., Nenes, A., Tombrou, M., 2017. New

- particle formation in the southern Aegean Sea during the Etesians: importance for CCN production and
- cloud droplet number. Atmospheric Chem. Phys. 17, 175–192. https://doi.org/10.5194/acp-17-175-2017
- Kanawade, V.P., Sebastian, M., Hooda, R.K., Hyvärinen, A.-P., 2022. Atmospheric new particle
- 62 formation in India: Current understanding and knowledge gaps. Atmos. Environ. 270, 118894.
- 63 https://doi.org/10.1016/j.atmosenv.2021.118894
- Kerminen, V.-M., Chen, X., Vakkari, V., Petäjä, T., Kulmala, M., Bianchi, F., 2018. Atmospheric new
- particle formation and growth: review of field observations. Environ. Res. Lett. 13, 103003.
- 66 https://doi.org/10.1088/1748-9326/aadf3c
- Kerminen, V.-M., Pirjola, L., Kulmala, M., 2001. How significantly does coagulational scavenging limit
- atmospheric particle production? J. Geophys. Res. Atmospheres 106, 24119–24125.
- 69 https://doi.org/10.1029/2001JD000322
- Kontkanen, J., Lehtipalo, K., Ahonen, L., Kangasluoma, J., Manninen, H.E., Hakala, J., Rose, C., Sellegri,
- K., Xiao, S., Wang, L., Qi, X., Nie, W., Ding, A., Yu, H., Lee, S., Kerminen, V.-M., Petäjä, T., Kulmala,
- M., 2017. Measurements of sub-3 nm particles using a particle size magnifier in different environments:
- from clean mountain top to polluted megacities. Atmospheric Chem. Phys. 17, 2163–2187.
- https://doi.org/10.5194/acp-17-2163-2017
- Kulmala, M., 2003. Atmospheric science. How particles nucleate and grow. Science 302, 1000–1001.
- https://doi.org/10.1126/science.1090848
- Kulmala, M., Dal Maso, M., Mäkelä, J.M., Pirjola, L., Väkevä, M., Aalto, P., Miikkulainen, P., Hämeri,
- K., O'Dowd, C.D., 2001. On the formation, growth and composition of nucleation mode particles. Tellus
- Ser. B Chem. Phys. Meteorol. 53, 479–490. https://doi.org/10.1034/j.1600-0889.2001.530411.x
- Kulmala, M., Kerminen, V.-M., Petäjä, T., Ding, A.J., Wang, L., 2017. Atmospheric gas-to-particle
- 81 conversion: why NPF events are observed in megacities? Faraday Discuss. 200, 271–288.
- 82 https://doi.org/10.1039/C6FD00257A
- Kulmala, M., Petäjä, T., Ehn, M., Thornton, J., Sipilä, M., Worsnop, D.R., Kerminen, V.-M., 2014.
- 84 Chemistry of Atmospheric Nucleation: On the Recent Advances on Precursor Characterization and
- Atmospheric Cluster Composition in Connection with Atmospheric New Particle Formation. Annu. Rev.
- Phys. Chem. 65, 21–37. https://doi.org/10.1146/annurev-physchem-040412-110014

- Kulmala, M., Petäjä, T., Nieminen, T., Sipilä, M., Manninen, H.E., Lehtipalo, K., Maso, M.D., Aalto,
- P.P., Junninen, H., Paasonen, P., Riipinen, I., Lehtinen, K.E.J., Laaksonen, A., Kerminen, V.M., 2012.
- Measurement of the nucleation of atmospheric aerosol particles. Nat. Protoc. 7, 1651–1667.
- 90 https://doi.org/10.1038/nprot.2012.091
- 81 Kulmala, M., Vehkamäki, H., Petäjä, T., Dal Maso, M., Lauri, A., Kerminen, V.-M., Birmili, W.,
- McMurry, P.H., 2004. Formation and growth rates of ultrafine atmospheric particles: a review of
- observations. J. Aerosol Sci. 35, 143–176. https://doi.org/10.1016/j.jaerosci.2003.10.003
- Kurtén, T., Torpo, L., Ding, C.-G., Vehkamäki, H., Sundberg, M.R., Laasonen, K., Kulmala, M., 2007.
- A density functional study on water-sulfuric acid-ammonia clusters and implications for atmospheric
- cluster formation. J. Geophys. Res. Atmospheres 112. https://doi.org/10.1029/2006JD007391
- Laj, P., Bigi, A., Rose, C., Andrews, E., Lund Myhre, C., Collaud Coen, M., Lin, Y., Wiedensohler, A.,
- Schulz, M., Ogren, J.A., Fiebig, M., Gliß, J., Mortier, A., Pandolfi, M., Petäja, T., Kim, S.-W., Aas, W.,
- Putaud, J.-P., Mayol-Bracero, O., Keywood, M., Labrador, L., Aalto, P., Ahlberg, E., Alados Arboledas,
- 00 L., Alastuey, A., Andrade, M., Artíñano, B., Ausmeel, S., Arsov, T., Asmi, E., Backman, J.,
- Baltensperger, U., Bastian, S., Bath, O., Beukes, J.P., Brem, B.T., Bukowiecki, N., Conil, S., Couret, C.,
- Day, D., Dayantolis, W., Degorska, A., Eleftheriadis, K., Fetfatzis, P., Favez, O., Flentje, H., Gini, M.I.,
- Gregorič, A., Gysel-Beer, M., Hallar, A.G., Hand, J., Hoffer, A., Hueglin, C., Hooda, R.K., Hyvärinen,
- O4 A., Kalapov, I., Kalivitis, N., Kasper-Giebl, A., Kim, J.E., Kouvarakis, G., Kranjc, I., Krejci, R., Kulmala,
- M., Labuschagne, C., Lee, H.-J., Lihavainen, H., Lin, N.-H., Löschau, G., Luoma, K., Marinoni, A.,
- Martins Dos Santos, S., Meinhardt, F., Merkel, M., Metzger, J.-M., Mihalopoulos, N., Nguyen, N.A.,
- Ondracek, J., Pérez, N., Perrone, M.R., Petit, J.-E., Picard, D., Pichon, J.-M., Pont, V., Prats, N., Prenni,
- A., Reisen, F., Romano, S., Sellegri, K., Sharma, S., Schauer, G., Sheridan, P., Sherman, J.P., Schütze,
- M., Schwerin, A., Sohmer, R., Sorribas, M., Steinbacher, M., Sun, J., Titos, G., Toczko, B., Tuch, T.,
- Tulet, P., Tunved, P., Vakkari, V., Velarde, F., Velasquez, P., Villani, P., Vratolis, S., Wang, S.-H.,
- Weinhold, K., Weller, R., Yela, M., Yus-Diez, J., Zdimal, V., Zieger, P., Zikova, N., 2020. A global
- analysis of climate-relevant aerosol properties retrieved from the network of Global Atmosphere Watch
- 13 (GAW) near-surface observatories. Atmospheric Meas. Tech. 13, 4353-4392.
- 14 https://doi.org/10.5194/amt-13-4353-2020

- Liao, L., Kerminen, V.-M., Boy, M., Kulmala, M., Dal Maso, M., 2014. Temperature influence on the
- natural aerosol budget over boreal forests. Atmospheric Chem. Phys. 14, 8295–8308.
- 17 https://doi.org/10.5194/acp-14-8295-2014
- Lyubovtseva, Y.S., Sogacheva, L., Maso, M.D., Bonn, B., Keronen, P., Kulmala, M., 2005. Seasonal
- variations of trace gases, meteorological parameters, and formation of aerosols in boreal forests 10, 18.
- Marten, R., Xiao, M., Rörup, B., Wang, M., Kong, W., He, X.-C., Stolzenburg, D., Pfeifer, J., Marie, G.,
- Wang, D.S., Scholz, W., Baccarini, A., Lee, C.P., Amorim, A., Baalbaki, R., Bell, D.M., Bertozzi, B.,
- Caudillo, L., Chu, B., Dada, L., Duplissy, J., Finkenzeller, H., Carracedo, L.G., Granzin, M., Hansel, A.,
- Heinritzi, M., Hofbauer, V., Kemppainen, D., Kürten, A., Lampimäki, M., Lehtipalo, K., Makhmutov,
- V., Manninen, H.E., Mentler, B., Petäjä, T., Philippov, M., Shen, J., Simon, M., Stozhkov, Y., Tomé, A.,
- Wagner, A.C., Wang, Y., Weber, S.K., Wu, Y., Zauner-Wieczorek, M., Curtius, J., Kulmala, M., Möhler,
- O., Volkamer, R., Winkler, P.M., Worsnop, D.R., Dommen, J., Flagan, R.C., Kirkby, J., Donahue, N.M.,
- 27 Lamkaddam, H., Baltensperger, U., Haddad, I.E., 2022. Survival of newly formed particles in haze
- conditions. Environ. Sci. Atmospheres 2, 491–499. https://doi.org/10.1039/D2EA00007E
- Mazon, S.B., Riipinen, I., Schultz, D.M., Valtanen, M., Maso, M.D., Sogacheva, L., Junninen, H.,
- Nieminen, T., 2009. Classifying previously undefined days from eleven years of aerosol-particle-size
- distribution data from the SMEAR II station, Hyytia"la", Finland. Atmos Chem Phys 10.
- Merikanto, J., Duplissy, J., Määttänen, A., Henschel, H., Donahue, N.M., Brus, D., Schobesberger, S.,
- Kulmala, M., Vehkamäki, H., 2016. Effect of ions on sulfuric acid-water binary particle formation: 1.
- Theory for kinetic- and nucleation-type particle formation and atmospheric implications: BINARY
- PARTICLE FORMATION THEORY. J. Geophys. Res. Atmospheres 121, 1736–1751.
- 36 https://doi.org/10.1002/2015JD023538
- Middlebrook, A.M., Bahreini, R., Jimenez, J.L., Canagaratna, M.R., 2012. Evaluation of Composition-
- 38 Dependent Collection Efficiencies for the Aerodyne Aerosol Mass Spectrometer using Field Data.
- Aerosol Sci. Technol. 46, 258–271. https://doi.org/10.1080/02786826.2011.620041
- Mortier, A., Goloub, P., Podvin, T., Deroo, C., Chaikovsky, A., Ajtai, N., Blarel, L., Tanre, D., Derimian,
- 41 Y., 2013. Detection and characterization of volcanic ash plumes over Lille during the Eyjafjallajökull
- eruption. Atmos Chem Phys 13, 3705–3720. https://doi.org/10.5194/acp-13-3705-2013

- Németh, Z., Rosati, B., Zíková, N., Salma, I., Bozó, L., Dameto de España, C., Schwarz, J., Ždímal, V.,
- Wonaschütz, A., 2018. Comparison of atmospheric new particle formation events in three Central
- European cities. Atmos. Environ. 178, 191–197. https://doi.org/10.1016/j.atmosenv.2018.01.035
- Nieminen, T., Kerminen, V.-M., Petäjä, T., Aalto, P.P., Arshinov, M., Asmi, E., Baltensperger, U.,
- Beddows, D.C.S., Beukes, J.P., Collins, D., Ding, A., Harrison, R.M., Henzing, B., Hooda, R., Hu, M.,
- Hõrrak, U., Kivekäs, N., Komsaare, K., Krejci, R., Kristensson, A., Laakso, L., Laaksonen, A., Leaitch,
- W.R., Lihavainen, H., Mihalopoulos, N., Németh, Z., Nie, W., O'Dowd, C., Salma, I., Sellegri, K.,
- Svenningsson, B., Swietlicki, E., Tunved, P., Ulevicius, V., Vakkari, V., Vana, M., Wiedensohler, A.,
- Wu, Z., Virtanen, A., Kulmala, M., 2018. Global analysis of continental boundary layer new particle
- formation based on long-term measurements. Atmospheric Chem. Phys. 18, 14737–14756.
- 53 https://doi.org/10.5194/acp-18-14737-2018
- Ohlwein, S., Kappeler, R., Kutlar Joss, M., Künzli, N., Hoffmann, B., 2019. Health effects of ultrafine
- particles: a systematic literature review update of epidemiological evidence. Int. J. Public Health 64, 547–
- 56 559. https://doi.org/10.1007/s00038-019-01202-7
- Paasonen, P., Peltola, M., Kontkanen, J., Junninen, H., Kerminen, V.-M., Kulmala, M., 2018.
- Comprehensive analysis of particle growth rates from nucleation mode to cloud condensation nuclei in
- 59 boreal forest. Atmospheric Chem. Phys. 18, 12085–12103. https://doi.org/10.5194/acp-18-12085-2018
- 60 Peng, Y., Dong, Y., Li, X., Liu, X., Dai, J., Chen, C., Dong, Z., Du, C., Wang, Z., 2017. Different
- Characteristics of New Particle Formation Events at Two Suburban Sites in Northern China. Atmosphere
- 62 8, 258. https://doi.org/10.3390/atmos8120258
- Pierce, J.R., Adams, P.J., 2009. Uncertainty in global CCN concentrations from uncertain aerosol
- 64 nucleation and primary emission rates. Atmospheric Chem. Phys. 9, 1339–1356.
- 65 https://doi.org/10.5194/acp-9-1339-2009
- Ren, J., Chen, L., Fan, T., Liu, J., Jiang, S., Zhang, F., 2021. The NPF Effect on CCN Number
- Concentrations: A Review and Re-Evaluation of Observations From 35 Sites Worldwide. Geophys. Res.
- 68 Lett. 48, e2021GL095190. https://doi.org/10.1029/2021GL095190
- Rivellini, L.-H., Chiapello, I., Tison, E., Fourmentin, M., Féron, A., Diallo, A., N'Diaye, T., Goloub, P.,
- 70 Canonaco, F., Prévôt, A.S.H., Riffault, V., 2017. Chemical characterization and source apportionment of

- submicron aerosols measured in Senegal during the 2015 SHADOW campaign. Atmos Chem Phys 17,
- 72 10291–10314. https://doi.org/10.5194/acp-17-10291-2017
- Rodríguez, S., Cuevas, E., González, Y., Ramos, R., Romero, P.M., Pérez, N., Querol, X., Alastuey, A.,
- 74 2008. Influence of sea breeze circulation and road traffic emissions on the relationship between particle
- number, black carbon, PM1, PM2.5 and PM2.5–10 concentrations in a coastal city. Atmos. Environ. 42,
- 76 6523–6534. https://doi.org/10.1016/j.atmosenv.2008.04.022
- Rodríguez, S., Van Dingenen, R., Putaud, J.-P., Martins-Dos Santos, S., Roselli, D., 2005. Nucleation
- and growth of new particles in the rural atmosphere of Northern Italy—relationship to air quality
- 79 monitoring. Atmos. Environ. 39, 6734–6746. https://doi.org/10.1016/j.atmosenv.2005.07.036
- Roig Rodelas, R., Chakraborty, A., Perdrix, E., Tison, E., Riffault, V., 2019. Real-time assessment of
- wintertime organic aerosol characteristics and sources at a suburban site in northern France. Atmos.
- 82 Environ. 203, 48–61. https://doi.org/10.1016/j.atmosenv.2019.01.035
- 83 Rolph, G., Stein, A., Stunder, B., 2017. Real-time Environmental Applications and Display sYstem:
- 84 READY. Environ. Model. Softw. 95, 210–228. https://doi.org/10.1016/j.envsoft.2017.06.025
- Rose, C., Collaud Coen, M., Andrews, E., Lin, Y., Bossert, I., Lund Myhre, C., Tuch, T., Wiedensohler,
- 86 A., Fiebig, M., Aalto, P., Alastuey, A., Alonso-Blanco, E., Andrade, M., Artíñano, B., Arsov, T.,
- Baltensperger, U., Bastian, S., Bath, O., Beukes, J.P., Brem, B.T., Bukowiecki, N., Casquero-Vera, J.A.,
- 88 Conil, S., Eleftheriadis, K., Favez, O., Flentje, H., Gini, M.I., Gómez-Moreno, F.J., Gysel-Beer, M.,
- Hallar, A.G., Kalapov, I., Kalivitis, N., Kasper-Giebl, A., Keywood, M., Kim, J.E., Kim, S.-W.,
- 90 Kristensson, A., Kulmala, M., Lihavainen, H., Lin, N.-H., Lyamani, H., Marinoni, A., Martins Dos
- 91 Santos, S., Mayol-Bracero, O.L., Meinhardt, F., Merkel, M., Metzger, J.-M., Mihalopoulos, N., Ondracek,
- J., Pandolfi, M., Pérez, N., Petäjä, T., Petit, J.-E., Picard, D., Pichon, J.-M., Pont, V., Putaud, J.-P., Reisen,
- 93 F., Sellegri, K., Sharma, S., Schauer, G., Sheridan, P., Sherman, J.P., Schwerin, A., Sohmer, R., Sorribas,
- 94 M., Sun, J., Tulet, P., Vakkari, V., van Zyl, P.G., Velarde, F., Villani, P., Vratolis, S., Wagner, Z., Wang,
- 95 S.-H., Weinhold, K., Weller, R., Yela, M., Zdimal, V., Laj, P., 2021. Seasonality of the particle number
- oncentration and size distribution: a global analysis retrieved from the network of Global Atmosphere
- 97 Watch (GAW) near-surface observatories. Atmospheric Chem. Phys. 21, 17185–17223.
- 98 https://doi.org/10.5194/acp-21-17185-2021

- Rose, C., Sellegri, K., Asmi, E., Hervo, M., Freney, E., Colomb, A., Junninen, H., Duplissy, J., Sipilä,
- M., Kontkanen, J., Lehtipalo, K., Kulmala, M., 2015a. Major contribution of neutral clusters to new
- 01 particle formation at the interface between the boundary layer and the free troposphere. Atmospheric
- 02 Chem. Phys. 15, 3413–3428. https://doi.org/10.5194/acp-15-3413-2015
- Rose, C., Sellegri, K., Freney, E., Dupuy, R., Colomb, A., Pichon, J.-M., Ribeiro, M., Bourianne, T.,
- 04 Burnet, F., Schwarzenboeck, A., 2015b. Airborne measurements of new particle formation in the free
- 05 troposphere above the Mediterranean Sea during the HYMEX campaign. Atmospheric Chem. Phys. 15,
- 06 10203–10218. https://doi.org/10.5194/acp-15-10203-2015
- 07 Rose, C., Sellegri, K., Moreno, I., Velarde, F., Ramonet, M., Weinhold, K., Krejci, R., Andrade, M.,
- Wiedensohler, A., Ginot, P., Laj, P., 2017. CCN production by new particle formation in the free
- troposphere. Atmospheric Chem. Phys. 17, 1529–1541. https://doi.org/10.5194/acp-17-1529-2017
- 10 Salimi, F., Rahman, M.M., Clifford, S., Ristovski, Z., Morawska, L., 2017. Nocturnal new particle
- 11 formation events in urban environments. Atmospheric Chem. Phys. 17, 521-530.
- 12 https://doi.org/10.5194/acp-17-521-2017
- Salma, I., Németh, Z., Kerminen, V.-M., Aalto, P., Nieminen, T., Weidinger, T., Molnár, Á., Imre, K.,
- Kulmala, M., 2016. Regional effect on urban atmospheric nucleation. Atmospheric Chem. Phys. 16,
- 8715–8728. https://doi.org/10.5194/acp-16-8715-2016
- Sandradewi, J., Prévôt, A.S.H., Szidat, S., Perron, N., Alfarra, M.R., Lanz, V.A., Weingartner, E.,
- Baltensperger, U., 2008. Using aerosol light absorption measurements for the quantitative determination
- of wood burning and traffic emission contributions to particulate matter. Environ. Sci. Technol. 42, 3316–
- 19 3323.
- 20 Sebastian, M., Kanawade, V.P., Soni, V.K., Asmi, E., Westervelt, Daniel.M., Vakkari, V., Hyvärinen,
- A.-P., Pierce, J.R., Hooda, R.K., 2021. New Particle Formation and Growth to Climate-Relevant Aerosols
- 22 at a Background Remote Site in the Western Himalaya. J. Geophys. Res. Atmospheres 126,
- e2020JD033267. https://doi.org/10.1029/2020JD033267
- Sellegri, K., Rose, C., Marinoni, A., Lupi, A., Wiedensohler, A., Andrade, M., Bonasoni, P., Laj, P., 2019.
- New Particle Formation: A Review of Ground-Based Observations at Mountain Research Stations.
- 26 Atmosphere 10, 493. https://doi.org/10.3390/atmos10090493

- 27 Shukla, K.K., Niranjan Kumar, K., Phanikumar, D.V., Newsom, R.K., Kotamarthi, V.R., Ouarda,
- T.B.M.J., Ratnam, M.V., 2016. Identification of the cloud base height over the centralHimalayan region:
- 29 Intercomparison of Ceilometer and DopplerLidar (preprint). Clouds/Remote Sensing/Validation and
- Intercomparisons. https://doi.org/10.5194/amt-2016-162
- 31 Sogacheva, L., Hamed, A., Facchini, M.C., Kulmala, M., Laaksonen, A., 2007. Relation of air mass
- history to nucleation events in Po Valley, Italy, using back trajectories analysis. Atmos Chem Phys 15.
- 33 Sogacheva, L., Maso, M.D., Kerminen, V.M., Kulmala, M., 2005. Probability of nucleation events and
- aerosol particle concentration in different air mass types arriving at Hyytiala southern Finland, based on
- back trajectories analysis. Boreal Environ. Res. 10, 479–491.
- 36 Spracklen, D.V., Carslaw, K.S., Kulmala, M., Kerminen, V.-M., Mann, G.W., Sihto, S.-L., 2006. The
- 37 contribution of boundary layer nucleation events to total particle concentrations on regional and global
- scales. Atmospheric Chem. Phys. 6, 5631–5648. https://doi.org/10.5194/acp-6-5631-2006
- 39 Stein, A.F., Draxler, R.R., Rolph, G.D., Stunder, B.J.B., Cohen, M.D., Ngan, F., 2015. NOAA's
- 40 HYSPLIT Atmospheric Transport and Dispersion Modeling System. Bull. Am. Meteorol. Soc. 96, 2059–
- 41 2077. https://doi.org/10.1175/BAMS-D-14-00110.1
- Tillmann, R., Hallquist, M., Jonsson, Å.M., Kiendler-Scharr, A., Saathoff, H., Iinuma, Y., Mentel, Th.F.,
- 43 2010. Influence of relative humidity and temperature on the production of pinonaldehyde and OH radicals
- from the ozonolysis of α -pinene. Atmospheric Chem. Phys. 10, 7057–7072. https://doi.org/10.5194/acp-
- 45 10-7057-2010
- Tuch, T.M., Herbarth, O., Franck, U., Peters, A., Wehner, B., Wiedensohler, A., Heintzenberg, J., 2006.
- Weak correlation of ultrafine aerosol particle concentrations <800 nm between two sites within one city.
- 48 J. Expo. Sci. Environ. Epidemiol. 16, 486–490. https://doi.org/10.1038/sj.jes.7500469
- Tunved, P., Hansson, H.-C., Kerminen, V.-M., Ström, J., Dal Maso, M., Lihavainen, H., Viisanen, Y.,
- Aalto, P.P., Komppula, M., Kulmala, M., 2006. High natural aerosol loading over boreal forests. Science
- 312, 261–263. https://doi.org/10.1126/science.1123052
- 52 Villani, P., Picard, D., Marchand, N., Laj, P., 2007. Design and validation of a 6-volatility tandem
- 53 differential mobility analyzer (VTDMA). Aerosol Sci. Technol. 41, 898–906.
- https://doi.org/10.1080/02786820701534593

- Wang, M., Kong, W., Marten, R., He, X.-C., Chen, D., Pfeifer, J., Heitto, A., Kontkanen, J., Dada, L.,
- Kürten, A., Yli-Juuti, T., Manninen, H.E., Amanatidis, S., Amorim, A., Baalbaki, R., Baccarini, A., Bell,
- 57 D.M., Bertozzi, B., Bräkling, S., Brilke, S., Murillo, L.C., Chiu, R., Chu, B., De Menezes, L.-P., Duplissy,
- J., Finkenzeller, H., Carracedo, L.G., Granzin, M., Guida, R., Hansel, A., Hofbauer, V., Krechmer, J.,
- 59 Lehtipalo, K., Lamkaddam, H., Lampimäki, M., Lee, C.P., Makhmutov, V., Marie, G., Mathot, S.,
- Mauldin, R.L., Mentler, B., Müller, T., Onnela, A., Partoll, E., Petäjä, T., Philippov, M., Pospisilova, V.,
- Ranjithkumar, A., Rissanen, M., Rörup, B., Scholz, W., Shen, J., Simon, M., Sipilä, M., Steiner, G.,
- 62 Stolzenburg, D., Tham, Y.J., Tomé, A., Wagner, A.C., Wang, D.S., Wang, Y., Weber, S.K., Winkler,
- P.M., Wlasits, P.J., Wu, Y., Xiao, M., Ye, Q., Zauner-Wieczorek, M., Zhou, X., Volkamer, R., Riipinen,
- 64 I., Dommen, J., Curtius, J., Baltensperger, U., Kulmala, M., Worsnop, D.R., Kirkby, J., Seinfeld, J.H.,
- 65 El-Haddad, I., Flagan, R.C., Donahue, N.M., 2020. Rapid growth of new atmospheric particles by nitric
- acid and ammonia condensation. Nature 581, 184–189. https://doi.org/10.1038/s41586-020-2270-4
- 67 Wang, Z., Wu, Z., Yue, D., Shang, D., Guo, S., Sun, J., Ding, A., Wang, L., Jiang, J., Guo, H., Gao, J.,
- Cheung, H.C., Morawska, L., Keywood, M., Hu, M., 2017. New particle formation in China: Current
- 69 knowledge and further directions. Sci. Total Environ. 577, 258–266.
- 70 https://doi.org/10.1016/j.scitotenv.2016.10.177
- Wang, Z.B., Hu, M., Wu, Z.J., Yue, D.L., Zheng, J., Zhang, R.Y., Pei, X.Y., Paasonen, P., Dal Maso, M.,
- Boy, M., Wiedensohler, A., 2013. Investigation of the connections between atmospheric new particle
- 73 formation and organics at an urban site of Beijing (preprint). Aerosols/Field
- 74 Measurements/Troposphere/Chemistry (chemical composition and reactions).
- 75 https://doi.org/10.5194/acpd-13-3419-2013
- Wehner, B., Wiedensohler, A., 2003. Long term measurements of submicrometer urban aerosols:
- statistical analysis for correlations with meteorological conditions and trace gases. Atmospheric Chem.
- Phys. 3, 867–879. https://doi.org/10.5194/acp-3-867-2003
- Xiao, S., Wang, M.Y., Yao, L., Kulmala, M., Zhou, B., Yang, X., Chen, J.M., Wang, D.F., Fu, Q.Y.,
- 80 Worsnop, D.R., Wang, L., 2015. Strong atmospheric new particle formation in winter in urban Shanghai,
- 81 China. Atmospheric Chem. Phys. 15, 1769–1781. https://doi.org/10.5194/acp-15-1769-2015
- 82 Yao, L., Garmash, O., Bianchi, F., Zheng, J., Yan, C., Kontkanen, J., Junninen, H., Mazon, S.B., Ehn,

- M., Paasonen, P., Sipilä, M., Wang, M., Wang, X., Xiao, S., Chen, H., Lu, Y., Zhang, B., Wang, D., Fu,
- Q., Geng, F., Li, L., Wang, H., Qiao, L., Yang, X., Chen, J., Kerminen, V.-M., Petäjä, T., Worsnop, D.R.,
- 85 Kulmala, M., Wang, L., 2018. Atmospheric new particle formation from sulfuric acid and amines in a
- 86 Chinese megacity. Science 361, 278–281. https://doi.org/10.1126/science.aao4839
- 87 Yli-Juuti, T., Nieminen, T., Hirsikko, A., Aalto, P.P., Asmi, E., Hõrrak, U., Manninen, H.E., Patokoski,
- J., Dal Maso, M., Petäjä, T., Rinne, J., Kulmala, M., Riipinen, I., 2011. Growth rates of nucleation mode
- 89 particles in Hyytiälä during 2003–2009: variation with particle size, season, data analysis method and
- 90 ambient conditions. Atmospheric Chem. Phys. 11, 12865–12886. https://doi.org/10.5194/acp-11-12865-
- 91 2011

92