# **Measurement Report: New particle formation characteristics**

# 2 at an urban and a mountain station in Northern China

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# 21 Abstract

Atmospheric new particle formation (NPF) events have attracted increasing attention for their contribution to the global aerosol number budget, and therefore their effects on climate, air quality, and human health. NPF events are regarded as a regional phenomenon, occurring over a large area. Most observations on NPF events in Beijing and its vicinity were conducted in populated areas, whereas observations on NPF events on mountaintops with low anthropogenic emissions are still rare in China. The spatial variation of NPF event 27 intensity has not been investigated in detail by incorporating both urban areas and mountain measurements in Beijing. Here, we provide NPF events characteristics in summers 2018 and 2019 at urban Beijing and a 28 comparison of NPF event characteristics - NPF event frequency, formation rate, and growth rate - by 29 30 comparing an urban Beijing site and a background mountain site separated by ~80 km from June 14 to July 14, 31 2019 as well as giving insights into the connection between both locations. There were no significant difference 32 of formation rates and growth rates observed during the short-term observation in 2019 and longer-term 33 observation in summers 2018 and 2019 at the urban site. During parallel measurements at urban Beijing and 34 mountain background areas, although the median condensation sink during the first two hours of the common NPF events was around  $0.01 \text{ s}^{-1}$  at both sites, there were notable differences in formation rates between the two 35 locations (median of 5.42 cm<sup>-3</sup>s<sup>-1</sup> at the urban site and 1.13 cm<sup>-3</sup>s<sup>-1</sup> at the mountain site during the first two hours 36 37 of common NPF events). In addition, the growth rates in the 7-15 nm range for common NPF events at urban 38 site (median of 7.6 nm.h<sup>-1</sup>) were slightly higher than those at mountain site (median of 6.5 nm.h<sup>-1</sup>). To 39 understand whether the observed events were connected, we compared air mass trajectories as well as 40 meteorological conditions at both stations. Favorable conditions for the occurrence of regional NPF events were 41 largely affected by air mass transport. Overall, our results demonstrate a clear inhomogeneity of regional NPF 42 within a distance of  $\sim 100$  km possibly due to the discretely distributed emission sources.

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44 Keywords: atmospheric aerosols, growth rates, regional new particle formation, sulfuric acid

#### 45 **1 Introduction**

Atmospheric new particle formation (NPF) events resulting from the formation of clusters and stable aerosol particles from gas-phase precursors have been recognized as a major contributor to the global aerosol budget (Kulmala et al., 2004; Zhang et al., 2012). Once the newly formed particles grow to certain sizes, they can act as cloud condensation nuclei (CCN), affecting the regional and global climate (Pierce and Adams., 2009; Yu and Luo., 2009). NPF events were also found to contribute to haze formation and thus can influence air quality, especially in megacities where the precursor concentrations and associated formation rates are rather high (Guo et al., 2014; 2020;Kulmala et al., 2021;Du &Dada et al., 2021).

53 The occurrence of NPF events is a result of the competition between factors promoting and inhibiting cluster 54 formation and their growth. For instance, sufficient sulfuric acid and other low-volatility vapors have been 55 confirmed to be important in particle nucleation and growth in field observations as well as in chamber experiments (Ehn et al., 2014; Wang et al., 2017; Lehtipalo et al., 2018; Yao et al., 2018; Deng et al., 2020b). On 56 57 the other hand, background particles can inhibit new particle formation by acting as condensation sink for vapor 58 precursors and coagulation sink for newly formed particles. Indeed, Cai et al. (2017) found that the Fuchs Surface Area ( $A_{Fuchs}$ ) (which is linearly proportional to condensation sink) determined the occurrence of NPF 59 60 events in urban Beijing. In the atmosphere, ambient conditions, such as air mass trajectories and meteorological 61 conditions, can affect the occurrence of NPF events by modifying the source-sink competition. Wu et al. (2007) 62 summarized favorable conditions for NPF events in Beijing based on a one-year observation as sufficient solar radiation (sunny days), northerly wind, low relative humidity, and less pre-loading large particles. Similarly, in 63 64 other environments, plenty of radiation, intermediate temperatures and low condensation sink favor the 65 occurrence of NPF events (Qi et al., 2015; Dada et al., 2017; Kerminen et al., 2018). Regional NPF events can 66 happen with a spatial extent up to several hundred kilometers and vertical extent from boundary layer to free 67 troposphere under favorable conditions (Hussein et al., 2009; Shen et al., 2011; Dai et al., 2017). Earlier studies 68 have shown that regional NPF events by simultaneous observations at two or more sites had similar features in 69 their occurrence and characteristics. For instance, Komppula et al. (2006) investigated the occurrence of NPF events at two forest stations in northern Finland during 2000-2003. Their results suggested that same air mass 70 71 source regions, favorable weather conditions and clean air at both stations were necessary for NPF events 72 occurring simultaneously at the two stations. Vana et al. (2016) compared observations at three sites over 1000 73 km distance at northern Finland, southern Finland and Estonia in 2013-2014. They found that some events have 74 the same origin. On the other hand, Jun et al. (2014) observed that NPF events occurred less frequently at 75 downtown Toronto than at a nearby background site, and attributed this observation to the high condensation 76 and coagulation sink due to primary particle emission from traffic at urban areas. Moreover, Carnerero et al. 77 (2018) observed horizontal distribution and regional impact of the NPF events with data from three urban, urban 78 background, and suburban stations in the Madrid metropolitan area, Spain in July 2016. Their results indicated 79 that ultra-fine particles were detected quasi-homogenously in an area spanning at least 17 km horizontally and 80 the NPF events extended over the full vertical extension of the mixed layer. Finally, Salma et al. (2016) found that regional NPF events were modified and transformed by urban NPF events during their observation in 81 82 2008-2009 and 2012-2013 in Budapest and at a regional background site 71 km away from it. In comparison to the aforementioned studies in Europe, a similar study was also carried out to understand the regional NPF events in North China Plain. Wang et al. (2013) characterized the NPF events observed at an urban Beijing site and a regional background site about 120 km northeast to the urban site from March to November in 2008. They observed 96 and 87 NPF events at urban Beijing and background site, respectively, among which 52 NPF events were observed simultaneously at both sites. They found that NPF events were slightly weaker in the background site compared to those observed at the urban site. However, the factors that influence the occurrence of NPF events at the two stations simultaneously were left undetermined.

90 In addition to horizontal extension of NPF events, the vertical extension of NPF events also attract attention of 91 researches. It have been confirmed that NPF events can be triggered within the whole low tropospheric column 92 at the same time and even above the planetary boundary layer upper limit (Boulon et al., 2011). Sellegri et al. 93 (2019) reviewed NPF events observed at 6 different altitude stations. They found NPF events was most favored 94 at the altitude close to the interface of the free troposphere (FT) with the planetary boundary layer (PBL) and at 95 the vicinity with clouds. In addition, at high altitude sites, CS may not be the liming factor for NPF occurrence 96 as higher CS associated with more precursors for nucleation and initial growth. Based on observations at two 97 different altitudes (e.g. 340 m and 560 m above sea level) in northern Finland, Komppula et al. (2003) found 98 NPF events had similar formation and growth rates between these two heights, while due to vertical movement 99 of air masses, difference of NPF event start time between these two sites was limited within 30 min. Similar 100 results were also observed at two sites in France that formation and growth rates were similar between two 101 altitudes (e.g., 660 m and 1465 m above sea level) while the contribution of ion-induced nucleation was higher 102 at high altitude (Boulon et al., 2011). Finally, during a recent observation in Spain, growth rates were higher at 103 the mountain site (2500 m a.s.l.) than urban site (680 m a.s.l.), while difference between formation rates varied 104 with altitude.

In addition to largely populated urban areas, there is a large mountain area within the Beijing-Tianjin-Hebei (BTH) region, where to our best knowledge, the characteristics of NPF events are understudied. In this study, we conducted simultaneous measurements of NPF event characteristics at an urban site in Beijing and a background mountain site about 80 km west to urban Beijing from June 14 to July 14 2019.

Based on our observations, we aim to (i) compare the characteristics of the NPF events between the two sites,
including the frequency, formation rate, and growth rate; (ii) figure out the connections and differences between

NPF events at these two sites; (iii) identify the favorable conditions for regional NPF events. Due to the profound participation of NPF events in the global aerosol number loading and air quality degradation, identifying the conditions those promote or inhibit the occurrence of regional scale NPF events could help to minimize its adverse effects.

### 115 2 Experiment and methodology

#### 116 2.1 Measurement sites' description

Urban site: The Beijing University of Chemical Technology - BUCT (39.94° N, 116.31° E) station is located on the fifth floor of a university building inside the west campus of BUCT. The station is surrounded by several main roads with heavy traffic and residential areas and thus, can be considered a typical urban station. The altitude of the west campus of BUCT is around 20 m above sea level and the urban site is around 12 m above ground level. More details of this station can be found in Zhou et al. (2020). Observations at the urban site are continuous since January 17, 2018 and were only interrupted for necessary instrument maintenance. The location is referred to as '**UB**' from here after and is shown on the map in Figure 1.

Mountain site: The Beijing Forest Ecosystem Research Station (39.96° N, 115.43° E) is located in the west of Beijing, referred to as '**MT**' from here after, which is part of the Chinese Ecological Research Network (CERN). It is located in the mountain areas west of Beijing, about 80 km from the urban site; see also in Figure 1. The altitude of the station is 1170 m above sea level and it is surrounded by forests. The closest anthropogenic activities are associated with small villages located in the valley nearby the MT station. Observations at MT station are from June 14 to July 14, 2019. For comparison reasons, we only used the data collected simultaneously at both stations.

Longquan station: The Longquan national monitoring station sits in Longquan town, Mengtougou District, Beijing. It is 20 km west to UB site and 60 km east to MT site and considered as a suburban station. The location is referred to as 'LQ' from here after and is shown on the map in Figure. 1.

#### 134 2.2 Instrumentation

As shown in Figure 2, the data qualities of particle number size distribution at both sites during the short-term parallel observations was good in general. Particle number size distribution data in the size range of 6-840 nm 137 were collected using a differential mobility particle sizer (DMPS) at the UB station. The instrument consists of one Hauke-type DMA (differential mobility analyzer, home-built by university of Helsinki) in different flow 138 139 rates and one CPC (condensation particle counter, TSI Model 3772). Details of this instrument can be found in 140 Salma et al. (2011) and Kangasluoma et al. (2020). At MT station, a scanning mobility particle sizer (SMPS, 141 consists of a TSI Differential Mobility Analyzer model 3081) and a fast mobility particle sizer (FMPS, TSI 142 Model 3091) were used to measure particle number size distribution from June 14 to June 28 and from June 29 143 to July 14, respectively. The size ranges of the SMPS and FMPS are 7-1218 nm and 6.04-856 nm, respectively. 144 The total number concentration from 4-3000 nm, measured by Condensation Particle Counter (CPC; TSI Model 145 3775), was used to calibrate the particle number size distributions from FMPS according to the method 146 suggested by Zimmerman et al. (2015). More details about the instrument are found in the previous studies 147 (Wang et al., 2019; Gao et al., 2020). The particle number size distribution measured by FMPS correlated well 148 with SMPS during the comparison in laboratory after being calibrated (Lee et al., 2013).

To ensure high quality of particle number size distribution data at UB site, a particle number size distribution system (PSD) also sampled in parallel with DMPS from June 1 to August 31, 2019 (summer 2019). It included a nano-scanning mobility particle sizer (nano-SMPS, 3–55 nm, mobility diameter), a long SMPS (25–650 nm, mobility diameter) and an aerodynamic particle sizer (APS, 0.55–10  $\mu$ m, aerodynamic diameter). Details of this instrument can be seen at Liu et al. (2016) and Deng et al. (2020b).

154 The PSD was used as a reference. As shown in Figure 3, median particle number size distribution obtained from 155 PSD and DMPS matched well in data trend. Varying with particle diameter, particle number size distribution 156 data measured by DMPS can be higher or lower than PSD within a factor of 2.

157 We cannot compare particle number size distribution data obtained from DMPS, SMPS and FMPS as we did not 158 sample with these three instruments in parallel at the same site. However, it is reasonable to assume that particle 159 number size distribution obtained from FMPS were comparable with those from DMPS as on one hand the 160 measurement techniques of particle number size distribution in the size range of these two instruments have 161 been well developed and be applied in quite a lot observations (Wang et al., 2017; Kangasluoma et al., 2020), on 162 the other hand, the particle number size distribution from FMPS was carefully calibrated and the FMPS was 163 properly operated during the observation as discussed above. Similar conclusions apply for the SMPS as well where we can rely on using the measurement from this instrument to discuss at least NPF event frequency at MT 164

site during June 14 to June 28, 2019, during which parameters of only one NPF event are calculated.

Sulfur dioxide (SO<sub>2</sub>) concentration data were collected by Thermo Environmental Instrument model 43i-TLE with a time resolution of 5-min at the UB station. There were no direct measurement of SO<sub>2</sub> concentrations at the MT station, but the SO<sub>2</sub> measurement at the closest national monitoring station (Longquan station, around 60 km from MT station and 20 km from UB station, see Figure 1) was used to indicate the strong decline of SO<sub>2</sub> concentration from urban Beijing towards the west areas. Time series of SO<sub>2</sub> concentration at UB station and Longquan station during the whole observation is shown in Figure 4. Due to the lower emission, the SO<sub>2</sub> concentration at the MT station is expected to be even lower than that in Longquan station.

The sulfuric acid concentration was measured at UB station by a chemical ionization-atmospheric interface-time of flight mass spectrometers (CI-APi-ToF, Aerodyne Research Inc.) equipped with a nitrate chemical ionization at UB station (Lu et al., 2019). There were no sulfuric acid data available at MT station and since no SO<sub>2</sub> concentrations were available, a sulfuric acid proxy concentration could not be derived.

The meteorological conditions such as relative humidity (RH, %), temperature (°C) and solar radiation (UVA and UVB, W/m<sup>2</sup>) were measured using a Vaisala Weather station data acquisition system (AWS310, PWD22, CL51), Metcon at UB station and using Vaisala MAWS301 automatic weather station at MT station. The measurements at the MT station were carried out at the height of 1.5 m. The wind speed (m/s) and wind direction (°) data were also measured by the weather station at UB site, while at MT site, we obtained with reanalyzed data from ERA5 model (Olauson, 2018).

# 183 2.3 Air mass back trajectories

Air mass back trajectories were calculated using a Lagrangian particle dispersion model FLEXPART (FLEXible PARTicle dispersion model) version 9.02 (Stohl et al., 2005). As the meteorological input, we used ECMWF (European Centre for Medium-Range Weather Forecast) operational forecast data with 0.15° horizontal and 1-hour temporal resolution. Particle retroplume simulations were performed hourly for both sites during the whole study period. For each retroplume simulation, we used 50 000 model particles distributed evenly between 0–100 m above the measurement site. The released model particles were traced backwards in time for 72 h, unless they exceeded the model grid (20–60°N, 95–135°E, resolution: 0.05°).

191 Based on the arrival direction of the 72-h backward trajectories, the prevailing air mass transport conditions at

192 each site were classified into 5 groups: North group, West group, East group, South group and Local group. Air 193 masses arriving from north, north-west and north-east including Mongolia, Inner-Mongolia and north-east China 194 were classified into the North group. Air masses from Shanxi province, Inner-Mongolia and further west were 195 classified into the West group. Air masses from the ocean east of Beijing were classified into the East group and 196 air masses from southern areas were classified into the South group. Stagnant air masses that had only travelled 197 short distances and/or were circulating around the measurement site were classified into the Local group. 198 Examples of air mass trajectories belonging to these five groups are shown in Figure 5. In general, air masses 199 from the north and west supply clean air from the mountainous areas to both stations, whereas air masses from 200 the east and south travel over highly populated areas, thus accumulating air pollutants. However, the impact of 201 local air masses on the pollution levels at the two sites can be different; at UB station, local air masses are 202 polluted by the urban emissions, while at MT station stagnant air could cause a clean situation due to low local 203 emissions. More details on the relationship between air mass transport conditions and the extent of pollution is 204 discussed in later sections.

# 205 2.4 Estimating the spatial extent of NPF

206 The observation of regional new particle formation events, where the growth of newly formed particles can be 207 followed for several hours, is a result of NPF taking place over a large spatial area. This is because as time 208progresses, the particles observed at a measurement site should have originated from further and further away 209 due to non-zero wind conditions. Following the progression of the observed NPF event and using air mass back 210 trajectories, we can estimate where the particles observed at different stages of the NPF event were initially 211 formed by calculating the air mass locations at the onset time of the NPF event (assuming that NPF occurs 212 simultaneously over the larger area). Typically, the mode related to the NPF event disappears from the 213 observations after some time. This is an indication of the currently observed air mass arriving from an area 214 where NPF was no longer taking place due to unfavorable local conditions. If the shift in the air mass origin 215 towards unfavorable conditions occurs gradually over time, the mode related to the NPF event can enter a stage 216 of growth stagnation (or even decrease in size) before disappearing completely (Kivekäs et al., 2016). This is 217 because the increasing transport time between NPF onset and observation of the particles at the measurement 218 site provides less and less additional 'material' for aerosol growth towards the more unfavorable conditions. 219 Calculating the locations where NPF is assumed to have taken place for longer data sets including several regional NPF events can give an estimation of the typical spatial extent of NPF around the measurement location. It should be noted that even in relatively clear cases, the subjective determination of NPF event onset and end times can easily lead to uncertainties of few tens of kilometers in the estimations. In locations with strong primary pollution sources, such as urban Beijing, objective determination of NPF event start and stop times becomes even more difficult. More details and discussion related to the method and its uncertainties can be found in Kristensson et al. (2014).

### 226 2.5 NPF event classification

227 Particle number size distribution data from both stations were used for classifying individual days into new 228 particle formation (NPF) event days and non-event days. This classification followed procedures presented by Dal Maso et al. (2005) and later adapted for urban locations (Chu et al., 2021) in which a day is classified as a 229 NPF event day if (a) a new mode in the size range smaller than 25 nm appeared and (b) the new mode kept 230 231 growing over several hours. On the other hand, non-event days are the days which do not fit any of the abovementioned criteria and undefined days are the days which fit either one of the abovementioned criteria. At 232 233 UB site, we also observed some cases in which nucleation mode particle number concentration burst without 234 mode diameter increase. It could be related to non-regional NPF events (Dai et al., 2017). We did not observe 235 such cases at the MT site. Actually, the abundant anthropogenic emissions in the megacity could provide enough 236 precursors for non-regional NPF events. However, traffic emissions can also provide abundant primary 237 nucleation mode particles, making it difficult to distinguish whether the new mode was from NPF event or 238 traffic. So we classified such events as "undefined" also.

# 239 2.6 Characteristics of NPF events

#### 240 2.6.1 Condensation sink

The condensation sink (CS) was calculated from particle size distribution data using the method described byKulmala et al. (2012):

243 
$$CS = 2\pi D \sum_{dp'} \beta_{m,dp'} dp' N_{dp'}$$

244 (1)

where *D* is the diffusion coefficient of the condensing vapor, sulfuric acid in our case, and  $\beta_{m,dp'}$  represents the transition-regime correction,  $N_{dp'}$  is the particle number concentration with diameter dp'. As shown in Figure 6, particles in size range of 20-800 nm dominated the total CS at UB station and particles in the size range of 50-800 nm dominated the total CS at MT station. Although the size ranges of DMPS, FMPS and SMPS slightly differ, all of them cover the main size range which constituted the CS and thus the calculation of CS should not be significantly influenced by differences in the instrument size ranges.

As shown in Figure 6(c&e) on NPF event days, particles smaller than 100 nm built a CS of  $3.7 \times 10^{-3}$  s<sup>-1</sup>, contributing 37% to the total CS. While at MT site, particles smaller than 100 nm only built a CS of  $1.2 \times 10^{-3}$  s<sup>-1</sup>, contributing less than 12% to the total CS (Figure 6 (d&f)). Although 100-840 nm particle number concentration at UB site was much less than that at MT site, 1-100 nm (especially 25-100 nm) particles compensated total CS by higher number concentration on NPF event days (CS at each site will be discussed in section 3.1.2 ).

The data on rainy days were discarded from analysis at both sites, hence the precipitation was considered to have minor effects on our CS calculation. We calculated CS at both sites assuming RH as 0%. It should be noted that the CS may have been underestimated by a factor of 1.12-1.33 at MT site when we include RH in the CS

calculation, e.g. 30%-70% during 9:00-15:00, as shown in Figure 10.

260 2.6.2 Particle growth rates

261 Growth rates were calculated for the size range of 7-15 nm (GR<sub>7-15 nm</sub>) using the 50% appearance time method

introduced by Lehtipalo et al. (2014) and Dada et al. (2020a) according to

$$GR = \frac{dp_2 - dp_1}{t_2 - t_1}$$

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264 (2)

where  $t_2$  and  $t_1$  are the appearance times of particles with sizes of  $dp_2$  and  $dp_1$ , respectively. The appearance time is defined as the time at which the concentration of particles at size  $d_p$  reaches 50% of its maximum.

# 267 2.6.3 Particle formation rates

The formation rates of particles of diameters 7 nm ( $J_7$ ) were calculated from particle number size distribution data using the method presented by Kulmala et al. (2012) and modified for urban environments by Cai and Jiang

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$$J_{k} = \frac{dN_{[d_{k},d_{u})}}{dt} + \sum_{d_{g}=d_{k}}^{d_{u}=1} \sum_{d_{i}=d_{\min}}^{+\infty} \beta_{(i,g)} N_{[d_{i},d_{i+1})} - \frac{1}{2} \sum_{d_{g}=d_{\min}}^{d_{u}=1} \sum_{d_{g}=d_{\min}}^{d_{i}^{3}+d_{g+1}^{3} \leq d_{u}^{3}} \beta_{(i,g)} N_{[d_{i},d_{i+1})} N_{[d_{g},d_{g+1})} + \frac{dN}{dd_{i}} \bigg|_{d_{i}=d_{u}} \bullet GR_{u}$$

$$(3)$$

Here,  $J_k$  is the formation rate at size  $d_k$ , cm<sup>3</sup>·s<sup>-1</sup>, (7 nm in this study);  $d_u$  is the upper size limit of the targeted aerosol population (10 nm in this study);  $d_{min}$  is the smallest particle size detected by particle size spectrometers (to make the results comparable, the  $d_{min}$  was set to 7 nm);  $N_{[dk,du]}$  is the number concentration of particles from size  $d_k$  to  $d_u$  (particles with diameters of du are not accounted for);  $d_i$  represents the lower limit of the  $i^{th}$  size bin;  $\beta_{(i,g)}$  is the coagulation coefficient for the collision of two particles with the size of  $d_i$  and  $d_g$ ; and  $GR_u$  refers to the growth rate at size  $d_u$ , nm·h<sup>-1</sup> Deng et al. (2020b).

Determination of nucleation start and stop times was affected by traffic emissions at UB station. Hence, we chose a time window of the first 2 hours of NPF event for formation rates calculation at both sites. During the time window, we always observed 7-10 nm particle number concentration burst significantly from the background level at both sites.

### 283 **3 Results and discussion**

# 284 3.1 Origin of NPF events at both sites

During our observation in summer 2018 and 2019 (from June to August of each year) at UB station, there were 155 days with valid data, 53 days of which were classified as NPF event days, corresponding to an NPF event frequency of 34%. This NPF event frequency was consistent with an earlier observation in summer in urban Beijing from 2004 to 2008 while smaller than other seasons especially winter during that observation and another one-year observation in UB station (Wu et al., 2007; Wang et al., 2013;Deng et al., 2020b).

For comparison of NPF characteristics between UB and MT stations, a parallel short-term observation was conducted at MT station from June 15 to July 14, 2019. In Figure 2, we show the particle number size distribution and CS during our short-term observations at both stations. There were a total of 12 and 13 NPF events observed at the UB station and the MT station, corresponding to an NPF event frequency of 48% (12 of 25) and 52% (13 of 25), respectively. Data were considered as valid when visual inspection of the particle number size distribution data and the instrument status did not indicate problems in the measurements. Only days with valid data at both stations were taken into consideration in our analysis. In addition, 9 NPF events were observed at both stations on the same day (referred to as common NPF events). Detailed information on the classified NPF event and non-event days, including the formation rates, growth rates, as well as their associated air mass origins during the short-term observation are provided in Table 1.

300 In order to understand the conditions favoring NPF events at both stations, we analyzed various ambient 301 parameters including air mass trajectories, meteorological variables, condensation sink as well as sulfuric acid 302 concentration.

#### 303 3.1.1 Favorable air mass origin for NPF events at individual locations

304 In Figure 7, we show frequencies of air masses arriving at UB station from different directions during our 305 observation in summer 2018 and 2019. The most frequent air masses arriving at UB station belonged to the 306 South group. During our observation in the two summers, out of 155 days were 52 days belonging to the South 307 group and 39, 32, 9 and 23 days in air masses belong to North, East, West and Local groups, respectively. NPF event frequency with respect to air masses is also shown in Figure 7. It is noticeable that air mass origin 308 309 influenced the occurrence of NPF events at UB site as the majority of NPF events occurred when the air masses 310 were coming from the north. During our observation in summer 2018 and 2019, 34 (out of 55) NPF events 311 occurred in air masses from the North group and 9, 2, 2 and 6 NPF events in the South, East, West and Local 312 groups, respectively (Figure 7a). One prominent feature of these air masses is their difference in CS. The CS of 313 the air masses classified as the North group (with median values of 0.01 s<sup>-1</sup> at UB station) is substantially lower 314 than that in other air mass classes (CS = 0.03, 0.025, 0.017, 0.03 s<sup>-1</sup>, for south, east, west and local, respectively), which might explain the high NPF event frequency associated with this air mass class. During the observation 315 316 from June 14 to July 14 in summer 2019, the most frequent air masses arriving at both sites belonged to the North group as shown in Table 1. Out of 25 days, there were 8 and 9 days belonging to the North group, at UB 317 318 and MT sites, respectively. The highest frequency of NPF events also occurred when the air masses were coming from the north. The high NPF events frequency during our observation form June 14 to July 14 could 319 320 also be attributed to the frequent air masses arriving at both sites from north to Beijing.

321 As shown in Table 1, NPF events occurring simultaneously at both sites only happened when air masses arrived

at both sites from the same directions, suggesting that most of the observed NPF events took place over the whole studied area, extending for several hundreds of kilometers (Dai et al., 2017;Du et al., 2021). The occurrences of common NPF events also closely connected with air mass origins that 7 (out of 9) common NPF events occurred under air masses in the North group, with the other two NPF events in the South group.

### 326 *3.1.2 The role of condensation sink in NPF event occurrence*

Figure 8a shows the difference in CS between NPF event and non-event days during our observation in summer 327 328 2018 and 2019 (two whole summers) at UB site and short-term parallel observations at both sites. The 'NPF1' 329 and 'non-event1' referred to NPF and non-event days during the two whole summers, respectively, while 330 'NPF2' and 'non-event2' referred to NPF and non-event days during the short-term parallel observation period 331 from June 14 to July 14, 2019 at both sites, respectively. The longer-term periods are used for confirming the 332 representativeness of the short-term overlapping period for the whole summer. As shown in Figure 8a, the median CS on NPF1 or NPF2 days is equivalent for UB station ( $CS_{NPF1} = 0.010s^{-1}$ ;  $CS_{NPF2} = 0.009s^{-1}$ ) and less 333 than a factor of 1.2 different between non-event1 and non-event2 in UB station ( $CS_{nonevent1} = 0.023s^{-1}$ ; 334 335  $CS_{nonevent2} = 0.020s^{-1}$ ), which confirms the representativeness of our short-term measurement period to the overall 336 urban Beijing summer.

Our results in Figure 8a show that the median CS was ~  $0.01 \text{ s}^{-1}$  during the first 2 hours of the NPF events, at both stations. On common NPF event days, the median CS was  $0.009 \text{ s}^{-1}$  at UB station and ~ $0.01\text{ s}^{-1}$  at MT station, respectively. In comparison, on non-event days, during roughly the same time period (9:00–11:00 LT), the CS was substantially higher, with median values of  $0.02 \text{ s}^{-1}$  and  $0.014 \text{ s}^{-1}$ , at UB and MT stations, respectively. Figure 8b presents the median CS during the first 2 hours of NPF events on common NPF event days measured at both stations, and shows the high correlation between the two.

Figure 8c shows the NPF event frequency as a function of CS during our observation at UB site in summer 2018 and 2019 and how the NPF event frequency decreased with increasing CS. When CS was smaller than  $0.01 \text{ s}^{-1}$ , all days were classified as NPF event days, and when CS was larger than  $0.035 \text{ s}^{-1}$ , no day was classified as NPF event day. This shows the major role of background particles in controlling the occurrence or inhibition of NPF events as shown in several previous studies in China and internationally (Deng et al., 2020a; Cai et al., 2017; Kulmala et al., 2017). While we cannot present a similar figure from the MT station, the same conclusion applies where CS does play a role in inhibiting NPF observation owing to the difference in the CS values observed between NPF and nonevents at MT as shown in figure 8a. Yet, since the overall preexisting particle
concentration at the MT is rather on the low end, the role of CS might not be as vital at the MT station as for the
UB station.

Different from NPF events under low CS ( $<0.01 \text{ s}^{-1}$ ), these NPF events under high CS were characterized by a relatively high H<sub>2</sub>SO<sub>4</sub> concentration ( $>10^7 \text{ cm}^{-3}$ ) or low formation rates (Figure 9a), discussed in further details in the coming sections. In comparison, at MT station, when CS was smaller than ~ 0.013 s<sup>-1</sup>, most (10 out of 14) days were classified as NPF event days as shown in Figure 9d. When CS was larger than ~0.013 s<sup>-1</sup>, we only observed one local NPF event and another two non-local NPF events (Table 1). The local NPF event under high CS at the MT station was characterized as high UV (>30 W/m<sup>2</sup>) and low formation rate ( $J_7$  were too small to be reliably calculated) as well.

360 *3.1.3 Role of meteorological variables in NPF event occurrence* 

While the air mass source regions, and their connection to the CS, seem to explain the general picture of NPF event occurrences at the two sites well, we still have some cases unexplained. For example, as shown in Table 1, there were several non-event days observed at MT station with air masses belonging to North and West groups, which were connected to low CS. This indicates that a further investigation into other NPF-related variables is still required.

In Figure 10, we show diurnal variation of meteorological variables during our observation in summer 2018 and 2019 at UB site and observations from June 14 to July 14 in 2019 at UB and MT sites. It is noticeable that the short-term observation compared well with the long-term observation and therefore is representative of summer at UB site as shown in Figure 10.

370 First, the intensity of solar radiation is considered to be one of the most important parameters deciding NPF 371 event occurrence as it translates into photochemistry strength (Chu et al., 2019). The median UV (UVA+UVB) 372 intensity at the UB station on NPF event and non-event days was 38.3 and 32.9 W/m<sup>2</sup>, respectively. The UV 373 intensity was on average ~15% higher on NPF event days than on non-event days at UB station. Although UV 374 intensity was important for NPF event occurrence, we still observed NPF events at UB station under low UV intensity, e.g. cases on June 30 and July 6. These two events all started immediately after sunrise (6:30 LT on 375 376 June 30 and 7:00 LT on July 6, see Table 1) and median UV intensity during the first two hours of NPF events 377 was only 13.2 and 14.1 W/m<sup>2</sup>, respectively. However, sulfuric acid concentration was higher than  $10^7$  cm<sup>-3</sup> at the same time, the possible reason is high SO<sub>2</sub> concentration and low CS (~0.003 s<sup>-1</sup>) outcompeted the low UV intensity (Dada et al., 2020b) as well as the possibility of having other H<sub>2</sub>SO<sub>4</sub> sources (Yao et al., 2020).

At MT station, the median UV intensity on NPF event and non-event days was 28.4 and 14.2 W/m<sup>2</sup>, respectively. The lower UV at MT station, in general might be related to the higher RH (Figure 10c&d) and thus more cloudiness and fog at the MT station (Hamed et al., 2010;Dada et al., 2018). The UV intensity was on average ~100% higher on NPF event days than on non-event days at UB station. All local NPF events happened when UV intensity was higher than 15 W/m<sup>2</sup> as shown in Figure 9d.

On the other hand, as shown in Figure 10c&d, the median relative humidity (RH) was lower on NPF event days than non-event days at both stations. This is consistent with earlier results that high RH suppressed NPF events by increasing CS and coagulation sink (CoagS), as it can enhance the particle hygroscopic growth (Hamed et al., 2010; Hamed et al., 2011). In addition, high RH was also found to be associated with more clouds resulting in less solar radiation (Dada et al., 2018).

The median temperatures at UB on event and non-event days were 31 °C and 29 °C, respectively, and at MT station 23 °C and 19 °C, respectively. The median temperature was lower at the MT station than at the UB station, due to the higher altitude of the station and likely also the weaker solar radiation (Figure 10e&f). At both stations, the median temperature was very similar on NPF event and non-event days, suggesting that temperature was not a crucial factor for NPF event occurrence during the measurement in summer.

# 395 3.1.4 Role of sulfuric acid concentrations in NPF event occurrence

396 Sulfuric acid has been found to be the main precursor vapor participating in NPF in China and in many locations 397 around the world due to its low volatility (Yao et al., 2018; Chu et al., 2019). In Figure 9a, we show the 398 concentration of sulfuric acid as a function of CS during summer 2018 and 2019 at UB site. As shown in Figure 399 9b, the median sulfuric acid (H<sub>2</sub>SO<sub>4</sub>) concentrations at UB station were  $8.1 \times 10^6$  cm<sup>-3</sup> and  $4.5 \times 10^6$  cm<sup>-3</sup> on NPF 400 event days and non-event days, respectively, during observation from June 14 to July 14 in 2019 and  $7.9 \times 10^6$ 401 cm<sup>-3</sup> and  $3.4 \times 10^6$  cm<sup>-3</sup> on NPF event days and non-event days, respectively, during the observation in summer 402 2018 and 2019. This suggests that  $H_2SO_4$  was important for NPF events at the UB station (Deng et al., 2020b; 403 Dada et al., 2020b). On the other hand, as shown in Figure 9a, the  $H_2SO_4$  concentration during 9:00-11:00 (local 404 time) on non-event days could be comparable with that on NPF event days, especially when CS was high. The 405 H<sub>2</sub>SO<sub>4</sub> concentration during 9:00- 11:00 (local time) on non-event days could be comparable with that on NPF 406 event days, especially when CS was high. Altogether, our observation shows that the occurrence of NPF events 407 was controlled by both  $H_2SO_4$  and CS at the UB station (Cai et al., 2020).

408 In addition, although we did not perform the measurement of  $H_2SO_4$  at the MT station, concentration of  $H_2SO_4$ 409 is expected to be much lower than that at the UB station. First, the SO<sub>2</sub> concentration at measurement at 410 Longquan Town was always below the detection limit ( $\sim 0.5$  ppb) during our observation period. In comparison, 411 median SO<sub>2</sub> concentration at UB station was 0.87 ppb for all days and 0.65 ppb for NPF event days during our short-term parallel observation period. The spatial decreasing trend of SO<sub>2</sub> concentration from urban Beijing to 412 413 the west implies a low SO<sub>2</sub> concentration at the MT station, especially when the nearby anthropogenic sources 414 are sparse (Liu, 2008; Ying, 2010; Wang, 2011; Yang-Chun et al., 2013). Second, the oxidation of  $SO_2$  by 415 photochemistry reactions could also be limited by the low solar radiation at the MT station as we discussed in 416 3.1.3. Third, CS, as the main sink of H<sub>2</sub>SO<sub>4</sub>, was comparable at the MT station to that in the UB station on NPF 417 event days (as shown in Figure 8a). Altogether, the lower production rate and the equivalent loss rate of H<sub>2</sub>SO<sub>4</sub> 418 at the MT station likely result in the lower H<sub>2</sub>SO<sub>4</sub> concentration, in comparison to UB station.

Due to the lack of  $H_2SO_4$  measurements, the NPF mechanism at the MT station cannot be inferred. Nevertheless, we show that the occurrence of NPF is a response to photochemistry (and very likely to  $H_2SO_4$ ) and CS in Figure 9d. It is clear that high UV intensity and low CS favored the occurrence of NPF. However, there existed exceptions. For example, two NPF events were observed even when the UV intensity was low and the CS was high, besides, it was an undefined day on June 28 despite of the high UV intensity and low CS. These exceptional cases will be discussed in detail in Section 3.6.1 and 3.6.2, respectively.

#### 425 3.2 NPF event start time at both stations

There was no significant difference in NPF event start times between the long-term and short-term parallel observations at UB site. In this section, we only compare NPF event start times of coincident events at UB and MT sites during the short-term parallel observations.

During our observation, there was no advection of air masses between the two sites on common NPF event days, indicating that the NPF events occurred at each site independently. As shown in Table 1, all common NPF events started after sunrise and prior to noon except the two non-local NPF events at MT station. However, NPF event start time was different between the two sites. Earlier researches in Nanjing, China and Nordic stations showed the similar results that NPF events can be observed simultaneously at two or more sites, but the start
time can be different, local meteorology, source strength and background aerosols could drive temporal behavior
of NPF events at each sites (Hussein et al., 2009;Dai et al., 2017).

# 436 **3.3** Particle formation and growth rates at both stations

437 The formation rates  $(J_7)$  at the two stations during the measurements are presented in Figure 11a.  $J_7$  observed during the short-term parallel observation (NPF2) at UB site were in the range of 3.0-10.0 cm<sup>-3</sup> s<sup>-1</sup> with a median 438 of 5.4 cm<sup>-3</sup> s<sup>-1</sup>, comparable with those observed in summer 2018 and 2019 (NPF1 = 2-14.0 cm<sup>-3</sup> s<sup>-1</sup> with a 439 median of 4.9 cm<sup>-3</sup> s<sup>-1</sup>) and significantly higher than the values in the MT station (0.75-3.0 cm<sup>-3</sup> s<sup>-1</sup> with a median 440 441 of  $0.82 \text{ cm}^{-3}\text{s}^{-1}$ ) for common NPF events (Figure 11b). These values are comparable to earlier observations in 442 urban Beijing and another regional background station in North China Plain (NCP) (Wang et al., 2013). Earlier 443 observations in NCP and Yangtze River Plain also show higher formation rates at urban sites than corresponding 444 background sites by roughly a factor of 2 due to lower anthropogenic emissions at background sites (Wang et al., 445 2013; Dai et al., 2017; Shen et al., 2018). The much lower  $J_7$  observed at MT station is very likely associated with 446 the low H<sub>2</sub>SO<sub>4</sub> concentration at this station, as we discussed above. However, other reasons, such as the low 447 concentration of  $H_2SO_4$  stabilizers, e.g., amines, cannot be ruled out either. Also, the  $J_7$  at UB station could be 448 affected by particle emissions due to the proximity of the location to the highway, while compared with NPF 449 events, the effect of traffic emissions is shown to be minor (Kontkanen et al., 2020; Zhou at al., 2020). In 450 addition, Boulon et al. (2011) observed that new particles could be formed at low altitude and transported to the 451 higher altitude sites, however, to confirm whether the phenomenon can happen at MT site, we still need 452 observation on vertical wind conditions or vertical evolution of potential temperature.

The growth rates in size range of 7-15 nm (GR<sub>7-15nm</sub>) at the UB station (4.8-12.9 nm/h with a median of 7.8 nm/h) during NPF2 was also comparable with the whole summers (NPF1) (4.8-12.9 nm/h with a median of 8.5 nm/h). While the difference in  $J_7$  was 7 times higher in UB than in MT, the observed GR were only a slightly higher at UB than at the MT station (5.7-10.5 nm/h with a median of 6.5 nm/h) for common NPF events (Figure 11c&d), implying that precursors needed for particle formation were much more abundant in the polluted urban environment (Wang et al., 2013), while those needed for growth are rather comparable. The GR at UB station was comparable to other long-term observation at UB station (1.1-8.0 nm/h) in 2018, and other urban areas in 460 China (Herrmann et al., 2014;Chu et al., 2019;Deng et al., 2020b). Consistent with earlier observations showing 461 that H<sub>2</sub>SO<sub>4</sub> could only contribute to a small fraction of the particle growth in this size range (Paasonen et al., 462 2018;Qi et al., 2018;Guo et al., 2020), the growth rates at both stations cannot be explained by the H<sub>2</sub>SO<sub>4</sub> 463 concentration. This implies that other condensable species, very likely low-volatility organic vapors, play an 464 important role in particle growth at both stations. At the UB station, anthropogenic VOCs are dominant 465 precursors of these low-volatility organic vapors (Guo et al., 2020; Deng et al., 2020a), while VOCs at MT 466 station, with rare anthropogenic sources, are likely dominated by biogenic emissions.

# 467 3.4 Ending diameters of newly-formed grown particles

468 Earlier observations have shown that diameters of newly-formed particles should be larger than 70 nm to contribute to cloud condensation nuclei significantly (Man et al., 2015; Ma et al., 2021) and will be considered 469 470 as haze particles when their size reaches larger than 100 nm (Kulmala et al., 2021). In Figure 12, we show 471 ending diameters (End Dp) of newly formed grown particles during our observations at both sites. End Dp during the observation from June 14 to July 14 at UB site (21-105 nm, with a median of 49 nm, Figure 12a) had 472 473 similar characteristics as those during the long-term observation in summer (21-126 nm, with a median of 56 nm, 474 Figure 12a) where most of End Dp were in the range of 25-70 nm. As shown in Fig.12b, 61% of End Dp were in 475 the range of 25-70 nm, and only 9% of End Dp were larger than 100 nm during our observation in summer 2018 476 and 2019 at UB site. We found that the ending diameters slightly higher at UB site than MT site, but the 477 difference is not significant (49 nm vs 45 nm) as shown in Figure 12c.

478 Earlier research has pointed out that in order to observe particle growth until 100 nm at a measurement station 479 under typical conditions, simultaneous NPF should happen over a very large area (e.g. with wind speed 5 m/s 480 and growth rate of 3 nm/h from the station to roughly 600 km upwind from the station) (Paasonen et al., 2018). During our observation in summer 2018 and 2019, most of the newly formed modes kept growing for about 20 481 482 hours after an NPF event started, and the maximum horizontal extension of the observed NPF events in the 483 growth stage is restricted to within about 200 km ( $\sim 2^{\circ}$  in latitude) north of UB site (Figure 13). As shown in 484 Figure 13, the population density is also higher within the area extending  $\sim 200$  km north than beyond this limit. 485 Therefore, it seems that NPF events were limited to the regions with some contribution from anthropogenic 486 emissions during air mass transport from north to Beijing. Roughly similar extent of the NPF area is also seen in 487 other directions. However, towards the south it is more likely that increasing condensation sink from 488 accumulating pollution becomes the limiting factor for NPF occurrence rather than decreasing strength in 489 emission sources. NPF events at MT station had similar characteristics as those at UB station with the NPF 490 event region extending a few hundred kilometers towards the north. The NPF events in this direction were 491 disrupted after a relatively similar distance (or they enter the growth stagnation phase). The limited NPF event 492 area could possibly explain why most End Dp we observed were smaller than 70 nm.

#### 493 3.5 Effect of topography

494 In Figure 14, we show average particle number size distributions and particle number concentrations on NPF 495 event and non-event days during our short-term parallel observation at both sites. On NPF event days, 496 nucleation (6-25 nm) and Aitken (25-100 nm) mode particle number concentrations were much smaller at MT 497 station than those at UB station due to smaller formation rates and less anthropogenic emissions. Interestingly, 498 accumulation (100-840 nm) mode particle number concentrations were higher at MT station (701-2900 cm<sup>-3</sup>, 499 with a median of 1500 cm<sup>-3</sup>) than those at UB station (350-1416 cm<sup>-3</sup> with a median of 700 cm<sup>-3</sup>) (Figure 14b). 500 Due to the close proximity of the two measurement sites, the air mass arrival directions and source regions were 501 (mostly) similar at both sites throughout the measurement period, hence the regional and transported air masses 502 cannot explain the higher accumulation mode particle number concentration at MT site. As there were few 503 primary emissions at MT site, the accumulation mode particles could be attributed to secondary particles 504 (Kulmala et al., 2021), indicating particles at MT station were more aged than those at UB station (Figure 14a). 505 The possible reasons is that mountains block pollution diffusion, which in the end resulted in comparable CS at 506 MT station as UB station.

Figure 15 shows an example of the wind distribution before and during NPF event on June 30, 2019 at 850 hPa (close to the altitude of MT station) and 10 m above ground level. As shown in Figure 15, the reanalyzed wind directions at 850 hPa were similar as those at 10 m above the ground level at MT station. Actually, the wind conditions on other NPF event days at MT station during our observation had similar characteristics that the wind directions were similar between 850 hPa and 10 m above ground level indicating air masses well mixed during NPF events. Earlier observations also found NPF event happened uniformly within the mixing layer at their observation stations and particle number size distribution remains roughly constant within the mixing layer 514 (Shen et al., 2018;Lampilahti et al., 2021).

#### 515 4 Summary and conclusions

516 We conducted observations of NPF events at an urban site (UB) and a background mountain site (MT) in 517 Beijing and fully analyzed the favorable conditions for NPF event occurrences at each of the sites. In order to 518 identify the similarities and differences between NPF events at both stations in terms of frequency, intensity, we 519 compared certain NPF events' characteristics including formation rate, growth rate as well as NPF event start 520 time and ending diameters of newly-formed growing particles at both stations. We found that NPF events are 521 most of the time a regional phenomenon occurring over the studied areas and connected closely with air masses 522 source regions during our observation. The air masses from north favored common NPF events more than any 523 other mass trajectories due to their associated clean air masses and thus low CS. Additionally, air masses from 524 the north group always resulted in an NPF event at UB station, while other factors still suppressed their 525 occurrence at the MT station. For example, we found that sufficiently high solar radiation, e.g. UV (UVA+UVB) 526 intensity larger than 15 W/m<sup>2</sup> is required for an NPF event to occur at MT and NPF events observed under solar radiation conditions smaller than 15 W/m<sup>2</sup> were rather transported NPF events from areas upwind to MT station. 527 Moreover, we found that the CS limit for NPF event occurrence at UB station was  $\sim 0.032$  s<sup>-1</sup>, which is 528 529 consistent with earlier observations in urban Beijing. In comparison, at MT station the CS limit could be ~0.013 s<sup>-1</sup>, above which local-NPF events could possibly be suppressed associated with the lower SO<sub>2</sub> 530 531 concentration.

532 Although NPF events could happen simultaneously at both stations, the NPF event strength (formation rates) 533 was significantly higher at UB than MT station, possibly due to spatial inhomogeneity in the sources of aerosol 534 precursor compounds as well as solar radiation. In addition, the growth rates in size range of 7-15 nm were also 535 slightly higher at UB than MT station. Regional NPF events were observed to occur with the horizontal extent 536 within around 200 kilometers when air masses arriving at Beijing from the north, as a result, only a few NPF 537 events were observed to end with mode diameters larger than 70 nm. The upwind extension of regional NPF events was limited to the areas with some anthropogenic emissions. There should not be any discrete boundary 538 539 between the regions that NPF event is or is not occurring, but with decreasing anthropogenic emissions, the 540 strength (formation rates and growth rates) should decrease. Overall, our results highlight the importance of 541 anthropogenic emissions for NPF events occurrence and subsequently growth in north China plain during 542 summer. However, there are still some uncertainties due to the limited data set. For more robust knowledge on 543 NPF events in north China plain, and to figure out the effect of urban emissions on regional NPF events, we still 544 need long-term observation including particle number size distribution down to sub-3 nm, gas and particle phase 545 chemistry upwind and downwind urban Beijing. Such observations can shed light into the regionality of NPF 546 events and the dynamical development of the aerosol population influenced by radical chemistry in the plume of 547 a megacity.

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549 Author contributions: YZ, CY, YG, XY performed the measurements. YZ, SH, CY, YG, LD, XY analyzed the 550 data. YZ, CY, SH, LD wrote the manuscript. All authors reviewed the paper and contributed to the scientific 551 discussions.

552 **Data availability:** The data displayed in this manuscript will be available online at <u>zenodo.com</u> once the 553 manuscript is in its final publication format.

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# 823 Tables and Figures

824	Table 1: NPF event and non- event days during our observation at both stations.
021	Tuble 1, 111 1 Crent and non crent days during our observation at both stations

Date	Туре	Air masses (9:00-15:00)		GR7-15nm (nm/h)		<b>J</b> <sub>7</sub> (cm <sup>-3</sup> s <sup>-1</sup> )		Event Start (LT)		Ending diameter (nm)		CS (s <sup>-1</sup> )	
		UB	МТ	UB	МТ	UB	MT	UB	МТ	UB	MT	UB	МТ
2019/06/14	a	North	North	8.61	-	4.97	-	9:00	8:00	71	-	0.017	0.008
2019/06/15*	a	Local	Local	12.63	-	5.56	-	11:00	15:00	82	60	0.013	0.029
2019/06/17	d	East	Local									0.031	0.011
2019/06/18	с	Local	West		10.5		0.17		12:00		45	0.039	0.008
2019/06/19	d	South	Local									0.037	0.047
2019/06/21	d	East	Local									0.035	0.018
2019/06/23	e	East	East									0.033	0.013
2019/06/24	f	Local	Local		8.21		-		12:00		50	0.027	0.014
2019/06/25*	a	Local	Local	-	-	-	-	12:00	15:00	-	53	0.032	0.027
2019/06/28	g	West	West	-		-		11:00				0.022	0.006
2019/06/29	a	North	North	12.93	7.14	6.93	2.28	9:00	8:00	21	19	0.008	0.011
2019/06/30	a	North	North	4.82	6.57	9.86	1.37	6:30	9:30	31	25	0.003	0.008
2019/07/01	a	North	North	7.31	5.82	3.84	0.82	9:00	8:30	105	102	0.006	0.009
2019/07/02	d	Local	West									0.013	0.014
2019/07/03	а	North	North	7.89	6.52	3.25	0.75	9:00	8:00	72	46	0.015	0.006
2019/07/04	b	Local	Local	-		-		10:00		53		0.012	0.012
2019/07/06	а	North	North	7.39	6.51	9.21	1.75	7:00	9:30	25	19	0.004	0.011
2019/07/07	b	North	North	7.61		3.61		9:00		32		0.008	0.005
2019/07/08	d	East	East									0.019	0.012
2019/07/09	d	East	East									0.021	0.015
2019/07/10	h	East	East									0.017	0.013
2019/07/11	d	East	East									0.039	0.014

2019/07/12	f	East	East		5.57		0.37		9:30		24	0.018	0.014
2019/07/13	с	Local	North		6.32		0.70		10:00		30	0.037	0.012
2019/07/14	a	North	North	12.04	9.86	3.91	0.89	9:30	9:30	63	47	0.023	0.017

\*a' means NPF event observed at both stations, 'b' means NPF event day at UB station while non-event day at MT station, 'c' means NPF event day at MT station while non-event day at UB station, 'd' means non-event day at both stations on the same day, 'e' means undefined day at both stations, 'f' means undefined day at UB station while NPF event day at MT station, g means undefined day at MT station while NPF event day at UB station, h means undefined day at UB station while non-event day at MT station, \* means NPF event observed at MT station was transported from somewhere else. – means the values cannot be reliably calculated. Only days when particle number size distribution were valid are included in this table.



835 Figure 1: Map showing locations of urban station (UB), Longquan station (LQ), and mountain station (MT). Image is

836 produced using © Google Maps.



839 Figure 2: Time series of particle number size distribution and CS (the blue line) at UB and MT stations during our

841 at MT station, respectively.

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<sup>840</sup> observations. Time resolutions for particle number size distribution data and CS were 8 min at UB station and 4 min



844 Figure 3: Median particle number size distribution in 5-900 nm measured by DMPS and PSD during our observation





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Figure 4: Time series of SO<sub>2</sub> concentration (ppb) at UB station and Longquan station (LQ) during our observation (left axis) as well as H<sub>2</sub>SO<sub>4</sub> concentration measured at UB station (right axis). Data under detection limit are set as zero at both stations. Data on NPF event days were marked in red at UB station and black at MT station. Time resolution for SO<sub>2</sub> data was 5 min at UB station and 1h at LQ station, respectively.



853 Figure 5: Examples of air masses arrived at both stations from (a) North group, (b) West group, (c) East group, (d)

854 South group and (d) Local group during 9:00-15:00 (local time, LT). Both stations are under the same marker.

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Figure 6: Median CS size distribution (a&b), accumulated CS contributed by particles from 6 nm and the ratio between accumulated CS and total CS (c&d); Contribution of size-segregated particles to total CS (e&f) at each site on NPF and non-event days during 9:00-15:00 (local time, LT). Figures on the left and right panels represented data observed at UB and MT site, respectively. The time resolutions for CS and particle number concentration data were 8 min at UB station and 4 min at MT station, respectively.

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Figure 7: Occurrence of NPF events and non-events under air masses arriving from different directions (a) as well as medians and percentiles of condensation sink (CS, s<sup>-1</sup>) during the 9:00-11:00 (local time) under different air masses (b) during our observation in summer 2018 and 2019 at UB station. The red line represents the median of the data and the lower and upper edges of the box represent 25<sup>th</sup> and 75<sup>th</sup> percentiles of the data, respectively. The length of the whiskers represents 1.5× interquartile range which includes 99.3% of the data. Data outside the whiskers are considered outliers and are marked with red crosses. The time resolution of CS was 8 min.



Figure 8: (a) Median and percentiles of condensation sink (CS, s<sup>-1</sup>) during our observations at both stations. The 873 874 'NPF1' and 'non-event1' referred to NPF and non-event days during summer 2018 and 2019, while 'NPF2' and 875 'non-event2' referred to NPF and non-event days during the short-term parallel observation from June 14 to July 14, 876 2019 at both sites. The red line represents the median of the data and the lower and upper edges of the box represent 25th and 75th percentiles of the data, respectively. The length of the whiskers represents 1.5× interquartile range which 877 878 includes 99.3% of the data. The time resolution of CS was 8 min. (b) Median CS during the first 2 hours of NPF 879 events on common NPF event days measured at both stations (MT vs. UB) . (c) Numbers of NPF event, non-event and 880 undefined days as well as NPF event frequency as a function of CS during our observation in summer 2018 and 2019 881 at UB station.



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Figure 9: (a) Median condensation sinks (CS, s<sup>-1</sup>) and H<sub>2</sub>SO<sub>4</sub> concentration (SA, cm<sup>-3</sup>) and (b) (b) medians and 884 885 percentiles of H<sub>2</sub>SO<sub>4</sub> concentration observed at UB station during the first 2 hours of NPF events and 9:00-11:00 on non-event days. (c) solar radiation (UVA+UVB, W/m<sup>2</sup>) during the first 2 hours of every NPF event and 9:00-11:00 on 886 887 every non-event day at UB station. The 'NPF1' and 'non-event1' referred to NPF event and non-event days in 888 summer 2018 and 2019 and the 'NPF2' and 'non-event2' referred to NPF event and non-event days during the 889 observation from June 14 to July 14, 2019. (d) Median condensation sinks (CS, s<sup>-1</sup>) and solar radiation (UVA+UVB, 890 W/m<sup>2</sup>) during the first 2 hours of every NPF event and 9:00-11:00 on every non-event day at MT station. Transported 891 NPF event cases and one non-event day with air masses belonging to west group (Jun 28) were all pointed out in the 892 figure. Size of data points on NPF event days means formation rate  $(J_7, cm^{-3}s^{-1})$  when it can be calculated reliably. The 893 time resolution of CS was 8 min at UB station and 4 min at MT station, respectively. The time resolution was 30 min 894 for SA data at UB station and 1h for solar radiation data at both stations.

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Figure 10: (a, b) Diurnal pattern of solar radiation (UV, W/m<sup>2</sup>), (c, d) Temperature (T, °C), and (e, f) Relative humidity (RH, %), at UB (left panel) and MT (right panel) stations on both NPF event and non-event days. Time resolutions for all data points here were 1h. The 'NPF1' and 'non-event1' referred to NPF event and non-event days in summer 2018 and 2019 and the 'NPF2' and 'non-event2' referred to NPF event and non-event days during the observation from June 14 to July 14, 2019.



Figure 11: Median and percentages of formation rates of 7 nm (J<sub>7</sub>, cm<sup>-3</sup>s<sup>-1</sup>) (a) and growth rates from 7 to 15 nm (GR<sub>6-15 nm</sub>, nm/h) (c) measured at both stations during our observation as well as comparison between J<sub>7</sub> (b) and GR<sub>6-15 nm</sub> (d) of common NPF events. The red line represents the median of the data and the lower and upper edges of the box represent 25<sup>th</sup> and 75<sup>th</sup> percentiles of the data, respectively. The length of the whiskers represents 1.5× interquartile range which includes 99.3% of the data. The 'NPF1' and 'non-event1' referred to NPF event and non-event days in summer 2018 and 2019 and the 'NPF2' and 'non-event2' referred to NPF event and non-event days during the observation from June 14 to July 14, 2019.



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Figure 12: (a) Median and percentiles of end diameters (End Dp, nm) of NPF events measured at both sites. The red 914 915 line represents the median of the data and the lower and upper edges of the box represent 25th and 75th percentiles of 916 the data, respectively. The length of the whiskers represents 1.5× interquartile range which includes 99.3% of the 917 data. The 'NPF1' and 'non-event1' referred to NPF event and non-event days in summer 2018 and 2019 and the 918 'NPF2' and 'non-event2' referred to NPF event and non-event days during the observation from June 14 to July 14, 919 2019. (b) Frequencies of end diameters in the size range of smaller than 25 nm, 25-70 nm, 70-100 nm and above 100 920 nm during our observation at UB station in summer 2018 and 2019. (c) Comparison between end diameters of 921 common NPF events at both stations.



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925 Figure 13: Spatial extent of the area where new particle formation events are estimated to have taken place based on 926 air mass back trajectories and the observed NPF events at both sites. Each line represents a single NPF event and 927 extends to the point beyond which continuation of the mode formed in an NPF event was no longer observed at the 928 measurement site. In other words, if an air mass is located outside the area roughly outlined by the colored lines 929 during the typical onset time of NPF and then transported to our measurement sites, NPF is unlikely to have occurred 930 in said air mass. The lines change color from pink to light blue if the observed NPF event enters a stage of growth 931 stagnation, which can indicate a less favorable environment for the formation and growth of new particles. The lines 932 are overlaid on top of a population density map (Gridded Population of the World; GPWv4.10; CC BY 4.0), which is 933 used to illustrate the level of anthropogenic activities and emissions.



938Figure 14: Median particle number size distribution as well as CS (blue lines) on NPF event and non-event days at939UB (a&b) and MT (c&d) stations and median and percentiles of nucleation (e), Aitken (f) and accumulation (g)940modes particle number concentration on NPF event and non-event days during our observation from June 14 to July94114, 2019 at both stations. The red line represents the median of the data and the lower and upper edges of the box942represent 25<sup>th</sup> and 75<sup>th</sup> percentiles of the data, respectively. The length of the whiskers represents 1.5× interquartile943range which includes 99.3% of the data. Data outside the whiskers are considered outliers and are marked with red944crosses.



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947 Figure 15: Wind distribution at 8:00, 9:00 and 10:00 on June 30, 2019 at 10 m above the ground level (lower panel)

948 and 850 hPa (close to the altitude of MT station, upper panel). The blue and black points on the figures represent MT

949 and UB stations, respectively.