

RESPONSE TO REVIEWERS

We thank the reviewers for their valuable comments, and respond point by point below.

ANONYMOUS REFEREE #2

Manuscript entitled 'Analysis of New Particle Formation (NPF) Events at Nearby Rural, Urban Background and Urban Roadside Sites' by Bousiotis et al. reports the occurrence of new particle formation events at three sites of different environments in the United Kingdom: Rural, urban background and near road sites. The authors study parameters of new particle formation such as frequency, growth rates, number concentration of sizes 16 – 20 nm, condensation sink, urban increment, nucleation strength factor and survival probability. The authors also report trajectory cluster analysis as well as the connection of NPF between the three different sites. In general, the manuscript, contains valuable data (three sites of different environments) and treasured statistics (7 years of data). In addition, the manuscript is well written and literature from around the world is acknowledged. However, the authors make big assumptions and conclusions without enough supporting data. The major concerns listed below need to be addressed before the manuscript is considered for publication in ACP.

Major Comments:

1. The authors report the observation of new particle formation events at three sites in the UK based on visual inspection of CPC (> 7 nm) and SMPS (> 16.6 nm). The general character of NPF events is missing. The lowest limit of the instrument is an issue and no big conclusions can be made before ensuring that the observed plume of particles is related to a new particle formation event. Authors should report how these events look like and whether they have a growing mode shape. Also, more characteristics of the growth should be reported such as possible shrinkage (see e.g. Salma et al. (2016)) and the size these particles reach. An example surface plot from each site should be added to the manuscript.

Let's take for example a regional event surface plot from Kerminen et al. (2018): figure 1, if we cannot observe the information below 16 nm, how can the authors prove that the increase in particle concentration is related to NPF, figure 2.

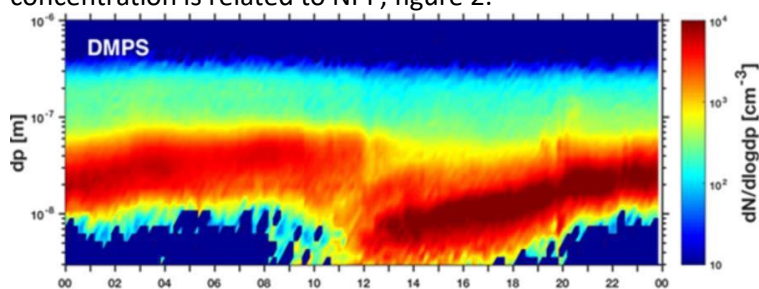


Figure 1 Regional Event example. Figure from Kerminen et al 2018.

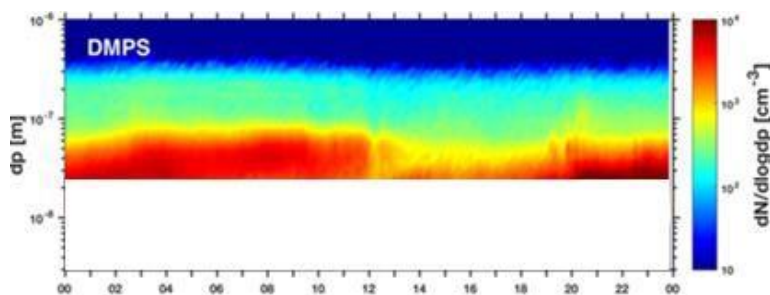


Figure 2 Modified figure 1.

The manuscript refers to many NPF studies from around the world, many of which report NPF starting from 6 nm (Salma et al., 2017), 3 nm (Dal Maso et al., 2005), 1.7 nm (Kirkby et al., 2016) while their measurement starts from 16.6 nm. The authors should present evidence that these observed particles are related to new particle formation events, and not for example a traffic growing mode (Brines et al., 2015).

RESPONSE: The dataset available, as mentioned in the text ranges from 16.6 to 604 nm. To overcome this limitation additional data was used to ensure the correct identification of the NPF events.

To achieve this:

- CPC data was used to provide insight into whether there was an increase on the number of particles of smaller size. An increased number of particles in the size range 7 – 16 nm (provided by the CPC data) right before or at the same time when observed in the SMPS data was a necessary criterion for the occurrence of an event.
- High resolution pollution data was used alongside particle number concentration data in a side by side comparison. A sharp increase in the particle number concentration which was accompanied by a similar increase in the concentrations of pollutants was an indication that these particles were probably associated with pollutant emissions. This was mainly an issue in the roadside (MR) and to a smaller extent with the background sites. Increased particle number concentrations observed at times matching the morning or evening traffic rush hours were also ignored at MR as they always coincided with increased concentrations of pollutants.
- Meteorological data was used. This mainly applies to the urban background site (NK), being in close proximity to London city centre. The possibility of a plume of pollution originating from the London city centre was considered when the site was downwind of it. A power plant to the northeast of the rural background site (HW) was also considered as a possible source of particles, though the distance is larger. Finally, as mentioned in the text, Heathrow airport and its influence were also considered.

In addition to this, the criteria set by Dal Maso et. al. (2005) were fully considered and unless there was a clear new mode of particles at the lower size range of the nucleation mode with a clear growth for at least 3 hours, an NPF event was not assigned. An example of the appearance of the events for each site has been added in the manuscript. Additionally, a discussion of particle shrinkage at later stages, which was observed at MR, is also added to the text.

Due to the limitations of the dataset, events in which the newly formed particles failed to grow to greater than 16 nm could not be seen except in the CPC data. These were rare and due to lack of additional information about their development were ignored. This clarification has been added in the text.

2. Section 2.1: Which years are studied?

RESPONSE: The years studied are 2009 – 2015. This information has been added in the text.

3. Section 2.1: Distance between the three sites should be mentioned.

RESPONSE: The distance between MR and NK is 4.5 km. The distance between HW and London city centre is about 80 km. This information has been added to the text.

4. Section 2.2.1: Authors report a visual inspection of CPC and SMPS data.

- How was this exactly done? Please elaborate.
- Was there any kind of counter-calibration done between these instruments?

RESPONSE: The method used was visual inspection of SMPS data supplemented by the use of CPC data to confirm the increase of the particle number concentration in the smaller size range (7 – 16 nm), as mentioned in (1). The text has been updated to clarify the method used. Both instruments are calibrated by the National Physical Laboratory according to the latest internationally recommended protocols.

5. Section 2.2.2: Calculation of the growth rates:

- Size of growth rates should be mentioned. E.g. growth from 7 to 20 nm? To 50 nm?

- How many points were taken in calculating the GR?
- Line 290: Authors claim that GR in NK (4.4 nm/h) are higher than the regional events GR (3.9 nm/h), what is the error bar on these calculations? Accordingly, these growth rates might be similar.

RESPONSE: As the lower size available was 16 nm, a calculation of the growth rate up to 50 nm was chosen (rather than up to 30 nm, which provided poor results in many cases due to the small range). The number of points taken depended on the development of the event and were considered from the start of the event until a) growth stopped, b) GMD reached 50 nm or c) the day ended. These points were added to the manuscript to clarify the method used.

On the third point made, due to the large variation of the growth rates of the events, the error bars are overlapped for the two groups of events. This has been included as a note in the text.

6. On line 178: the author mention nucleation mode, which is by definition number of particles between 3 and 25 nm, while the authors conduct a large study on a small fraction of this nucleation mode (16 – 25 nm).

RESPONSE: We are not aware of a widely recognised definition of the nucleation mode, with the term taking in different size ranges in the literature. Regardless of that, in the text it is mentioned that *“NPF events are considered when a distinctly new mode of particles which appears in the size distribution at nucleation mode size, prevails for some hours and shows signs of growth”*, which is accurate in relation to the criteria set for NPF event selection in this study.

7. Section 2.2.4: Reference to Kulmala et al. 2017, calculating $P = CS'/GR$. What GR was used here? See point 4.

RESPONSE: The growth rate and condensation sink used are the ones calculated by the methods mentioned in the text. A clarification of this has been added in the text.

8. Section 3.1.1: Reference to Figure S1: cloudiness, and RH.... Is missing. Was cloudiness measured or calculated?

RESPONSE: Cloud amount data, as for all other meteorological data were measurements provided by the Met Office, as mentioned in the text. A plot with average cloud amount for each site has also been added in the supplementary.

9. Section 3.1.2: How can the authors prove that NPF events are happening at the near road site and not transported to the location?

RESPONSE: It cannot be stated with certainty whether the NPF took place at the site or particles were advected. What can be said though with confidence is that regardless of where the particle formation took place (either on the spot or in the close vicinity, as particles of that size range cannot travel to distances greater than some kilometers before either reaching detectable sizes or being diluted – especially in a polluted environment such as the London city centre), the new mode not only persists but it also grows for at least 3 hours. A clarification of this has been added in the text. If the events at the roadside site were due to advection, or a purely regional phenomenon, a much closer correlation of event days and growth rates between MR and NK would be expected than was observed.

10. The authors make big conclusions regarding the SO₂ driving mechanism of NPF which cannot be proved without adequate chemical speciation of the particles formed. These conclusions shall be minimized throughout the manuscript. Authors could try calculating sulfuric acid proxy from SO₂ and CS (Petäjä et al., 2009).

RESPONSE: In the text is stated that SO₂ was found to be lower on event days compared to the average, which logically leads to the conclusion that either the greater concentrations of SO₂ are associated with a more polluted environment with an increased condensation sink (which consequently has a negative effect in the occurrence of an event), or its concentration is adequate and it is not a factor affecting the occurrence of an event (positively or negatively). The calculation of the H₂SO₄ proxy was carried out and provided information that did not help in clarifying this point. It was found that the proxy was higher on event days at the background sites and gave an unclear result for the roadside. This result though provides no additional information as the increased values

of the proxy are the result of the higher solar radiation and the lower condensation sink found during events. Changes were made in the text to “soften” these conclusions.

ANONYMOUS REFEREE #1

The MS mainly deals with the occurrence frequency, particle growth rate, condensation sink, nucleation strength factor, survival parameter and relationships among them at 3 different locations (rural, urban background and urban roadside sites) in the UK over several years. It contains valuable results and conclusions. Some parts of the MS should be elaborated better (some items are given below as examples), and they can definitely be handled and improved. There is, however, a conceptual weakness of the study related to the lower diameter limit of the SMPS system (of 16.6 nm) which can represent the largest source of inconclusive or ambiguous interpretations for the urban sites.

Major comment:

1. New particle formation and growth events are mainly identified, separated from emission sources and classified on the basis of particle number size distributions in the particle diameter range <20 nm (e.g. Kulmala et al., Nat. Protoc., 7, 1651–1667, 2012). The diameter interval available for this in the evaluated work, namely 16.6–20 nm is quite narrow in particular, when you consider the logarithmic scale of the abscissa of size distributions. More importantly, the lower limit is requested to be even smaller (preferably below 10 nm or at 3 nm) for studies in urban atmospheric environments, where huge emission peaks can temporarily dominate the smallest size ranges as well (Nieminen et al., Atmos. Chem. Phys., 18, 14737–14756, 2018). This property (16.6 nm lower limit) of the measuring system and its consequences for the data treatment, results and conclusions at the urban sites should definitely be discussed in detail, explained and resolved before any further opinion could be formed or decision can be made.

RESPONSE: The limitations and consequences due to the available dataset, as well as the additions in the method to ensure the correct selection of NPF events are explained at length in the response to Referee #2 (earlier in this document). As a result of this, clarification of the method and the additional data used have been added to the text.

Some minor comments:

1. Lines 21, 69, etc.: it is advised not to start a sentence with abbreviation.

RESPONSE: Text updated to address the comment.

2. Line 61: consider writing primary particles or emission sources instead of primary emissions.

RESPONSE: Text updated to address the comment.

3. Lines 106–109: it is unusual to attribute particles with a diameter between 1.3 and 3 nm to road traffic emissions, and, therefore, this should be discussed and explained in more detail.

RESPONSE: Text updated to accurately reflect the conclusions of the study mentioned.

4. Lines 149–151 or Table 1: supply more detailed data coverage, e.g. for each year or season of years.

RESPONSE: Table has been added in the S.I. for detailed seasonal data coverage.

5. Lines 198–203: it is requested that the diameter of particles under consideration is specified as the growth rate changes with diameter.

RESPONSE: Text updated to include the size range of the particles considered in the calculation of the growth rate.

6. Lines 262, 263, Table 2: revisit your rounding off strategy.

RESPONSE: Text and tables updated to follow a uniform rounding scheme.

7. Lines 462–463: remove; it is a repetition from lines 235–239.

RESPONSE: Text updated to remove repeated information.

8. Fig. 2: it is unclear from the figure or related text which time interval was considered here. A number of NPF events of 90 at Harwell in summer (JJA, 92 days) should be clarified to avoid any misunderstanding.

RESPONSE: The figure's description has updated to clarify the period plotted.

**Analysis of New Particle Formation (NPF) Events at
Nearby Rural, Urban Background and
Urban Roadside Sites**

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29 ABSTRACT

30 New Particle Formation (NPF) events have different patterns of development depending on the
31 conditions of the area in which they occur. In this study, NPF events occurring at three sites of
32 differing characteristics (rural Harwell (HAR), urban background North Kensington (NK), urban
33 roadside Marylebone Road (MR), London, UK) were studied (seven years of data). The different
34 atmospheric conditions in each study area not only have an effect on the frequency of the events,
35 but also affect their development. The frequency of NPF events is similar at the rural and urban
36 background locations (about 7% of days), with a high proportion of events occurring at both sites
37 on the same day (45%). The frequency of NPF events at the urban roadside site is slightly less (6%
38 of days), and higher particle growth rates (average 5.5 nm h^{-1} at MR compared to 3.4 nm h^{-1} and 4.2
39 nm h^{-1} at HAR and NK respectively) must result from rapid gas to particle conversion of traffic-
40 generated pollutants. A general pattern is found in which the condensation sink increases with the
41 degree of pollution of the site, but this is counteracted by increased particle growth rates at the more
42 polluted location. A key finding of this study is that the role of the urban environment leads to an
43 increment of 20% in $N_{16-20\text{nm}}$ in the urban background compared to that of the rural area in NPF
44 events occurring at both sites. The relationship of the origin of incoming air masses is also
45 considered and an association of regional events with cleaner air masses is found. Due to lower
46 availability of condensable species, NPF events that are associated with cleaner atmospheric
47 conditions have lower growth rates of the newly formed particles. The decisive effect of the
48 condensation sink in the development of NPF events and the survivability of the newly formed

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49 particles is underlined, and influences the overall contribution of NPF events to the number of
50 ultrafine particles in an area. The other key factor identified by this study is the important role that
51 urban pollution plays in new particle formation events.

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1. INTRODUCTION

Ultrafine particles (particles with diameter smaller than 100 nm) typically make the greatest contribution in the total particle count, especially in urban environments (Németh et al., 2018), but a very small contribution to total volume and mass (Harrison et al., 2000). Research studies have indicated that ultrafine particles can cause pulmonary inflammation and may contribute to cardiovascular disease (Oberdörster, 2000) and have increased possibility to penetrate the brain and central nervous system (Politis, ~~Pilinis and Lekkas~~ et al., 2008) compared to fine and coarser particles. Since some studies report that toxicity per unit mass increases as particle size decreases (Penttinen et al., 2001; MacNee et al., 2003; Davidson et al., 2005); it is considered possible that particle number concentrations may be a better predictor of health effects than mass concentrations (Harrison et al., 2000; Atkinson et al., 2010; Kelly et al., 2012; Samoli et al., 2016). Additionally, NPF events have an impact on climate (Makkonen et al., 2012) either by increasing the number of cloud condensation nuclei (Spracklen et al., 2008; Merikanto et al., 2009; Dameto de España et al., 2017; Kalkavouras et al., 2017), or directly affecting the optical properties of the atmosphere (Seinfeld and Pandis, 2012).

The sources of ultrafine particles in urban areas can either be primary ~~particles or emission sources~~ from traffic (Shi et al., 1999; Harrison et al., 2000), airports (Masiol et al., 2017) and other combustion related processes (Keuken et al., 2015; Kecorius et al., 2016), or by new particle formation (NPF) from gaseous precursors. ~~NPF-New particle formation~~ as described by Kulmala et

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74 al. (2014), is the process of production of low-volatility vapours, clustering of these vapours,
75 nucleation, activation of the clusters with a second group of vapours and condensational growth to
76 larger sizes. This process can occur both locally or on a larger scale; in the latter case the events are
77 characterized as regional. Regional events have been found to take place in a scale of hundreds of
78 kilometres (Németh and Salma, 2014; Shen et al., 2018), without being affected by air mass
79 advection (Salma et al., 2016). NPF is one of the main contributors of particles in the atmosphere
80 (Spracklen et al., 2010; Kulmala et al., 2016; Rahman et al., 2017) and this [relative](#) contribution
81 increases moving from a kerbside to a rural area (Ma and Birmili, 2015). While NPF events in rural
82 and remote areas have been widely studied for many years (O'Dowd et al., 2002; Dal Maso et al.,
83 2005; Ehn et al., 2010; Dall'Osto et al., 2017; Kalkavouras et al., 2017), in urban areas intensive
84 studies have started mainly in recent years (Jeong et al., 2010; Minguillón et al., 2015; Peng et al.,
85 2017; Németh et al., 2018). Early studies in Birmingham, UK highlighted the connection of NPF
86 events with solar radiation (Shi et al., 2001) and a low condensation sink (Alam et al., 2003), a
87 measure of pre-existing aerosol loading (Dal Maso et al., 2002). The importance of a low
88 condensation sink was further underlined by later studies, as being one of the most influential
89 variables in the occurrence of NPF in all types of environment (Wehner et al., 2007; Park, Yum and
90 Kim, 2015; Pikridas et al., 2015). An important contributor to many NPF pathways is SO₂ (Woo et
91 al., 2001; Berndt et al., 2006; Laaksonen et al., 2008), which in the presence of solar radiation forms
92 H₂SO₄, often the main component of the initial clusters (Kuang et al., 2008; Kulmala et al., 2013;
93 Bianchi et al., 2016; Kirkby et al., 2016). Dall'Osto et al. (2013) pointed out that the role of SO₂ is

94 less significant in urban areas compared to rural and background areas. SO₂ concentration
95 variability in urban areas was found to have a small impact on the frequency of NPF events (Alam-
96 [Shi and Harrison et al.](#), 2003; Jeong et al., 2010), though it can have an effect on the number of
97 particles formed (Charron, ~~Birmili and Harrison et al.~~, 2007). Furthermore, Dall'Osto et al. (2018)
98 in their research at 24 sites in Europe, pointed out the different role SO₂ seems to play depending on
99 its concentration, and that of other species. Jayaratne et al. (2017) however found that in the heavily
100 polluted environment of Beijing, China, NPF events were more probable in sulphur rich conditions
101 rather than sulphur poor. Apart from its role in the initial formation of the clusters, H₂SO₄ seems to
102 participate in the early stages of growth of the newly formed clusters (Kulmala et al., 2005; Iida et
103 al., 2008; Xiao et al., 2015). In later stages of growth, low or extremely low volatility organic
104 compounds (O'Dowd et al., 2002; Laaksonen et al., 2008; Metzger et al., 2010; Kulmala et al.,
105 2013; Tröstl et al., 2016; Dall'Osto et al., 2018) appear to be more important, while the role of
106 ammonium nitrate in particle growth is also considered (Zhang et al., 2017). While in rural areas the
107 organic compounds are mainly of biogenic origin (Riccobono et al., 2014; Kirkby et al., 2016), in
108 urban areas they mainly originate from combustion ~~procedures~~ [processes](#) (Robinson et al., 2007;
109 Gentner et al., 2012). Many comparative studies have reported higher growth rates in urban areas
110 compared to background sites (Wehner et al., 2007; Jeong et al., 2010; Salma, et al., 2016; Wang et
111 al., 2017), as well as greater particle formation rates (Salma, et al., 2016; Nieminen et al., 2018) and
112 a higher frequency of NPF events (Peng et al., 2017), which was attributed to the higher
113 concentration of condensable species. Salma et al. (2014) however reported fewer NPF events in the

114 city centre of Budapest compared to the urban background, due to the higher condensation sink.
115 Due to the complexity of the conditions and mechanisms within an urban area (Harrison, 2017),
116 NPF events are harder to study and factors to be attributed. ~~A large number of particles of size 1.3 –~~
117 ~~3 nm has been attributed~~ Increased concentrations of particles in the size range 1.3 – 3 nm were
118 measured at a kerbside site when being downwind from the road, following the trends of traffic-
119 related nucleation mode particles, associating them to with traffic emissions at a kerbside site and
120 thus not related to resulting from homogeneous nucleation mechanisms (Rönkkö et al., 2017;
121 Hietikko et al., 2018), and studies in Barcelona, Spain (Dall'Osto et al., 2012; Brines et al., 2014)
122 and Leicester, U.K. (Hama et al., 2017), attributed a larger portion of nucleation mode particles to
123 vehicular emissions compared to photochemically induced nucleation. As the condensation sink is
124 higher within an urban environment, NPF events are less favoured. Their occurrence is attributed to
125 either ineffective scavenging or the higher growth rate of the newly formed particles (Kulmala et
126 al., 2017), when sufficient concentrations of precursors are present in the atmosphere (Fiedler et al.,
127 2005), as particle formation was found to take place on both event and non-event days (Riipinen et
128 al., 2007).

129
130 In this study, NPF events in three areas of different land use in the southern U.K. are analyzed.
131 Studies for NPF events have been conducted in the past for Harwell, Oxfordshire (Charron et al.,
132 2007; 2008) and the effect of NPF upon particle size distributions was also considered for N.
133 Kensington, London (Beddows et al., 2015). A combined study including all three sites has also

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134 been conducted, but in the aspect of ultrafine particle variation (Von Bismarck-Osten et al., 2013).
135 The present study is the first to use a combined long term database for all three sites, focusing on
136 the trends and conditions of NPF events at these sites, as well as the first which identifies NPF
137 events at the highly trafficked Marylebone Road site, as up to this point ultrafine particles were
138 attributed only to traffic (Charron and Harrison, 2003; Dall'Osto et al., 2011). As in this study a
139 rural and an urban background area are studied alongside a kerbside site in the city of London in
140 close proximity, the conditions and development of NPF events in a mid-latitude European region
141 are discussed in relation to the influence of different local environments.

142

143 2. DATA AND METHODS

144 2.1 Site Description and Data Availability

145 This study analysed NPF events in three areas in the southern United Kingdom (Fig. 1). Harwell in
146 Oxfordshire, is located about 80 km west of the greater London area. The site is in the grounds of
147 the Harwell Science Centre in Oxfordshire (51° 34' 15" N, 1° 19' 31" W) and is representative of a
148 rural background area; a detailed description of the site was given by Charron et al. (2013). North
149 Kensington is a suburban area in the western side of London, U.K., 4.5 km west of Marylebone
150 Road. The site is located in the grounds of Sion Manning School (51° 31' 15" N, 0° 12' 48" W) and
151 is representative of the urban background of London. A detailed description of the site was given by
152 Bigi and Harrison (2010). Marylebone Road is located in the centre of London, U.K. The site is
153 located on the kerbside of Marylebone road (51° 31' 21" N; 0° 9' 16" W), a very busy arterial route

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154 within a street canyon. A more detailed description of the area can be found in Charron and
155 Harrison (2003).

156

157 At all three sites, seven years **(2009 – 2015)** of particle number size distributions in the range of
158 16.6 – 604 nm have been measured and recorded as 15-minute averages, using a Scanning Mobility
159 Particle Sizer (SMPS), comprised by an Electrostatic Classifier (EC, TSI model 3080) and a
160 condensation Particle Counter (CPC, TSI Model 3775), operated on behalf of the Department for
161 Environment, Food and Rural Affairs (DEFRA) in the U.K. At all sites the inlet air is dried, and
162 operation is in accord with the EUSAAR/ACTRIS protocol (Wiedensohler et al., 2012). These 15-
163 minute measurements were averaged to an hourly resolution. In Harwell there were 46930 hours of
164 available SMPS data (76.5% coverage), in N. Kensington 51059 (83.3% coverage) and ~~in~~at
165 Marylebone Road 45562 (74.3% coverage). **Detailed data availability is found in ~~Table S1.~~** A free-
166 standing CPC (TSI model 3022A) also operated alongside for most of the years of the survey and
167 was used to give an estimate of particles in the 7-16.6 nm range by difference from the SMPS.

168

169 Additionally, air pollutants and other aerosol chemical composition data were extracted from the
170 DEFRA website (<https://uk-air.defra.gov.uk/>). Meteorological data for Harwell and Heathrow
171 airport (used for N. Kensington and Marylebone road) were available from the Met Office, while
172 solar radiation data from Benson station (for Harwell) and Heathrow airport (for N. Kensington and
173 Marylebone Road), were extracted from the Centre for Environmental Data Analysis (CEDA) site

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174 (<http://www.ceda.ac.uk>). Back trajectory data calculated using the HYSPLIT model (Draxler and
175 Hess, 1998), were extracted by the NOAA Air Resources Laboratory
176 (<https://ready.arl.noaa.gov/READYtransp.php>) and were processed using the Openair package for R
177 (Carslaw and Ropkins, 2012).

178

179 2.2 Methods

180 2.2.1 NPF events selection

181 The identification of the NPF event days was made by visual inspection of SMPS data,
182 supplemented with the use of and CPC data to confirm the formation of a new mode of particles,
183 using the criteria set by Dal Maso et al. (2005). NPF events are considered when a distinctly new
184 mode of particles which appears in the size distribution at nucleation mode size, prevails for some
185 hours and shows signs of growth. Using these criteria, NPF events are classified into two classes, I
186 and II depending on the confidence level. Class I events are further classified to Ia and Ib, with class
187 Ia containing very clear and strong particle formation events, while Ib contains less clear events. In
188 this study the events of class Ia ~~are~~ only are considered as being the most suitable for analysing case
189 studies of NPF events (Figure S1). At this point it should be mentioned that due to the particle size
190 range available, NPF events in which new formed particles failed to grow beyond 16.6 nm (if any)
191 could not be identified. Though such rare formationsoccasions were identified using the CPC data,
192 bursts of new particles in the size range < 16.6 nm that did not appear ~~on~~in the SMPS dataset were
193 ignored as their development was unknown. High time resolution data for gaseous pollutants and

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aerosol constituents was used to identify pollution events affecting particle concentrations and these were removed from the data analysis. -This analysis took account of the fact that nanoparticle emissions from Heathrow Airport affect size distributions at London sites (Harrison et al., 2018), and such primary emission influences were not included as NPF events.

2.2.2 Calculation of the condensation sink and growth rate

For the calculation of the condensation sink the method proposed in Kulmala et al. (2001) was used in which the condensation sink is calculated as

$$CS = 4\pi D \sum \beta_M r N \quad (1)$$

where r is the radius of the particles and N is the number concentration of the particles. D is the diffusion coefficient calculated (for $T = 293$ K and $P = 1013.25$ mbar) according to Polling et al. (2000):

$$D_{vap} = 0.00143 \cdot T^{1.75} \frac{\sqrt{M_{air}^{-1} + M_{vap}^{-1}}}{P \left(D_{x,air}^{\frac{1}{3}} + D_{x,vap}^{\frac{1}{3}} \right)^2} \quad (2)$$

211 where P is air pressure, M is the molar mass and D_x is the diffusion volume for air and H₂SO₄. β_M is
212 the Fuchs correction factor calculated as (Fuchs et al., 1971):

213

$$\beta_M = \frac{1 + K_n}{1 + \left(\frac{4}{3a} + 0.377\right)K_n + \frac{4}{3a}K_n^2}$$

214 (3)

215

216 where K_n is the relation of the particle diameter and the mean free path of the gas λ_m, called the
217 Knudsen number.

218

219 The growth rate of the particles on nucleation event days was also calculated as proposed by
220 Kulmala et al. 2012, using the formula

221

$$GR = \frac{D_{P_2} - D_{P_1}}{t_2 - t_1}$$

222 (4)

223

224 for the size range 16.6 – 50 nm. The number of points taken depended on the development of the
225 event and were considered from the start of the event until a) growth stopped, b) GMD reached 50
226 nm or c) the day ended.
227 ~~for the period of each event day when growth was observed.~~

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230 **2.2.3 Calculation of the urban increment (U.I.)**

231 The urban increment is defined as the ratio of the number concentration of particles below 20 nm
232 for event days to the average (for the period April – October, when the majority of the events take
233 place) for North Kensington to that at Harwell. This provides with a measure of the new particles
234 formed in each area in comparison to the average conditions, and is calculated by

$$236 \text{ U.I.} = \frac{NK_{\text{Nuc Max}} - NK_{\text{Bg}}}{HW_{\text{Nuc Max}} - HW_{\text{Bg}}} \quad (5)$$

237
238 where $NK_{\text{Nuc Max}}$ is the maximum concentration of particles below 20 nm found in the diurnal cycle
239 on event days (found at 13:00) and NK_{Bg} is the average mean concentration at the same time (same
240 for Harwell in the denominator).

242 **2.2.4 Calculation of nucleation strength factor (NSF) and the P parameter**

243 The Nucleation Strength Factor (NSF) was proposed as a measure of the effect nucleation events
244 have in the composition of ultrafine particles in an area. Two factors were proposed. First is the
245 NSF_{NUC} . This is calculated as

$$246 \text{ NSF}_{\text{NUC}} = \frac{\left(\frac{N_{(\text{smallest size available}-100)}}{N_{(100-\text{largest size available})}} \right)_{\text{nucleation days}}}{\left(\frac{N_{(\text{smallest size available}-100)}}{N_{(100-\text{largest size available})}} \right)_{\text{non-nucleation days}}} \quad (6)$$

248

249 and provides of a measure of the concentration increment on nucleation days exclusively caused by
 250 new particle formation (NPF). The second factor is NSF_{GEN} calculated as

251

$$\text{NSF}_{\text{GEN}} = \frac{\left(\frac{N_{\text{smallest size available}} - 100}{N_{100 - \text{largest size available}}} \right)_{\text{all days}}}{\left(\frac{N_{\text{smallest size available}} - 100}{N_{100 - \text{largest size available}}} \right)_{\text{non-nucleation days}}} \quad (7)$$

253 and gives a measure of the overall contribution of NPF on a longer span (Salma et al. 2017).

254 The dimensionless survival parameter P, as proposed in Kulmala et al. (2017), was calculated as

255

$$P = \frac{CS'}{GR'}$$

256

257 where CS' = CS/(10⁻⁴ s⁻¹) and GR' = GR/(1 nm hour⁻¹). CS and GR values used as were calculated

258 with the methods mentioned at 2.2.2. An increased P parameter is an indication that a smaller

259 percentage of newly formed particles will survive to greater sizes. Hence this is the inverse of

260 particle survivability, and values of P<50 are typically required for NPF in clean or moderately

261 polluted environments, although higher values of P are observed in highly polluted atmospheres

262 (Kulmala et al, 2017).

263

264

265

266 3. RESULTS AND DISCUSSION

267 3.1 NPF Events ~~at~~in the Background Areas

268 3.1.1 Conditions and trends of NPF events

269 The number of NPF event days for each site per year, those that took place simultaneously ~~on~~at
270 both urban and rural background sites, as well as those events that took place ~~in~~at all three sites
271 simultaneously appear in Table 1. Given that overall data recovery was in the range of 74-83%,
272 results from individual years are unreliable, but the seven-year runs should average out most of the
273 effects of incomplete data recovery. The number of events is similar for Harwell and N.
274 Kensington, with a frequency of about 7% of all days with data. There is a clear seasonal variation
275 favouring summer and spring (Figure 2) for both areas of the study. A similar pattern of variation
276 was found for N. Kensington by Beddows et al. (2015). In general, higher solar radiation, lower
277 relative humidity, low cloud cover and higher pressure conditions, lower concentrations of
278 pollutants as well as lower condensation sink are found when NPF events took place in all areas
279 (Figure S2~~4~~), as was also reported by Charron et al. (2007) for Harwell. While SO₂ is one of the
280 main factors for NPF events to occur, concentrations are lower when events take place. This is
281 indicative that SO₂ concentrations in these areas are sufficient for events to take place, and higher
282 concentrations are likely to be associated with higher pollution and a higher condensation sink. The
283 proxy for [H₂SO₄] was calculated for the background sites using the method outlined in (Petäjä et
284 al., (2009) and was found to be higher on event days for both background sites (results not
285 included). This indicates the possible positive effect of increased concentrations of H₂SO₄ in the

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286 occurrence of NPF events as well as, since SO₂ concentrations were found lower, the increased role
287 of either the solar radiation (via the formation of OH radical) or the reduced condensation sink to its
288 formation. This is also similar to the case for of gaseous ammonia (results not included) for
289 Harwell where data was available, as there was no distinct variation found between event and non-
290 event days, but as the concentration of ammonia in the U.K. is in the range of few ppb (Sutton et al.,
291 1995), it is sufficient according to ternary nucleation theory (Korhonen et al., 1999) for NPF events
292 not to be limited by ammonia. The average growth rate for Harwell was found to be 3.37-4 nm h⁻¹,
293 within the range given by Charron et al. (2007) and higher at N. Kensington at 4.2-2 nm h⁻¹, a trend
294 found for all seasons (Figure 3). The increased growth rate in the urban area can be related to the
295 greater presence of organic matter and other condensable species. In both areas NPF events had
296 higher growth rates in summer than in spring, as was also found in previous studies (Kulmala et al.,
297 2004; Nieminen et al., 2018). This may be associated with the higher presence-concentration of
298 organic compounds emitted by trees during summer (Riipinen et al., 2007), or faster oxidation rates
299 due to higher concentrations of hydroxyl radical and ozone (Harrison et al., 2006).

300

301 About 45% of the events took place simultaneously in both background areas. These events are
302 characterized as regional, as NPF takes-took place in-on a larger scale, regardless of the local
303 conditions on-of the given area. In this case, meteorological conditions were even clearer, indicative
304 of the greater dependence of regional events on synoptic conditions rather than local. While most
305 chemical constituents were also lower in concentration during regional events, different patterns

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306 were found for organic compounds and sulphate for each background area. In Harwell sulphate was
307 higher during regional events, while in N. Kensington organic compounds were higher during
308 regional events. This may be indicative of the variable role that specific chemical species have in
309 condensational nanoparticle growth (Yue et al., 2010). In all cases though, the concentrations of
310 these species were lower compared to the average conditions. Despite these differences, the growth
311 rate of particles was found to be higher for local events in N. Kensington (4.4 nm h^{-1}) compared to
312 regional events (3.9 nm h^{-1}), ~~though within the margin of error~~ though within the margin of uncertainty. In Harwell, no difference
313 was found in the growth rate between regional and local events.

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314

315 3.1.2 Variability of the origin of the air masses on NPF events

316 As both background ~~areas-sites~~ are relatively close to each other (about 80 km) and had similar
317 number of event days, a combined clustering of back trajectories for the event days (only) in these
318 two areas was attempted. This would provide an insight into the origin of air masses for local and
319 regional events, as well as the conditions for these air masses. The data for local N. Kensington
320 events and both local and regional events in Harwell were clustered together and the results along
321 with the characteristics of the air mass clusters are found in Figure S32.

322

323 Cluster C3, which is placed between C2 and C4 among those originating from the Atlantic Ocean,
324 has the highest percentage for both area specific and regional events. Specifically, for regional
325 events the percentage is over 35%, much higher compared to all other, showing a clear “preference”

326 of regional events for cleaner and faster moving air masses from mid-latitudes of the Atlantic
327 Ocean. This “preference” explains the lower production and growth rate of the new particles found
328 for regional events, compared to local ones, as air masses from this area have lower organic carbon
329 and SO₂ concentrations. Cluster C5, originating straight from the north but representing air masses
330 that have crossed the Irish Sea and have not extensively gone over land presents a similar case.
331 These cold and clean air masses are associated with a low growth rate and survivability of the
332 newly formed particles. Local events for both sites apart from those in Cluster C3 are highly
333 associated with Clusters C1 and C2. C1, which contains slow and polluted air masses, presents the
334 highest growth rate and as a result high particle survivability, as given by the P parameter (see
335 later). On the other hand, C2 which consists of warm and moist air masses from lower latitudes is
336 the least common for regional events and presents high growth rate and survival probability of the
337 particles. Apart from the weak relation found with particulate organic carbon concentrations and
338 growth rate (Figure S32), there appears to be an inverse relation between the temperature and
339 survivability of the particles. Warmer air masses seem to be related to higher particle survival
340 probability, which may be attributable to greater growth rates as temperature increases (Yli-Juuti et
341 al., 2011).

342

343 **3.1.3 Urban increment and particle development**

344 The urban environment, depending on the conditions, may have a positive or negative effect in the
345 number of the particles formed and their consequent survival and growth. Both Harwell and N.

346 Kensington are in background areas, rural and urban respectively. As a result, while the
347 concentrations of pollutants are higher in N. Kensington than Harwell, their effect is smaller
348 compared to that of Marylebone Road. A comparison of the particles smaller than 20 nm, gives
349 insight into the formation and survival of the newly formed particles in the initial stages.
350 Calculating the urban increment (equation 5) using the two background sites showed around 20%
351 more particles of size 16 - 20 nm in N. Kensington than Harwell for event days, an increment that is
352 even stronger when solely local events are considered (Figure 4). As the sizes of the particles in the
353 calculation are relatively large and due to the higher condensation sink found in N. Kensington, this
354 increment is expected to be larger for smaller size particles. A possible explanation for this result
355 may be the greater concentration of organic compounds which is observed in N. Kensington, as
356 discussed earlier, which leads to more rapid formation of secondary condensable species that
357 enhances the nucleation process in the more polluted area.

358

359 Considering the local events, most of the pollutant [concentration](#) data available appear to be higher
360 which is reflected in the condensation sink as well. The role of the polluted background appears to
361 be decisive in the further growth of the newly formed particles, especially for Harwell. This, at both
362 sites causes the number of particles of greater size to be smaller for the later hours in the days of
363 local events (Figure S43). Another possible reason for this difference in the larger size ranges can
364 be the higher concentration of organic content on the days of regional events at N. Kensington (as
365 discussed earlier). On the other hand, for Harwell all hydrocarbons with available data are lower

366 throughout the day (apart from ethane) during regional events. Unlike N. Kensington, at Harwell
367 particles smaller than 20 nm as well as the growth rate of the newly formed particles are almost the
368 same for regional and local events.

369

370 The calculation of the increment in Marylebone Road provided negative results; particles smaller
371 than 20 nm were less abundant on event days compared to the average, throughout the day. This is
372 due to the fact that Marylebone road is heavily affected by traffic pollution and on average,
373 conditions do not promote NPF events due to the high condensation sink, unless clear conditions
374 prevail, which are also associated with a low particle load.

375

376 **3.2 NPF Events at Marylebone Road**

377 For many years, NPF events were thought not to take place in heavily polluted urban areas, as the
378 effect of the increased condensation sink was considered ~~detrimental~~crucial in suppressing the
379 formation and growth of new particles. Recent long term analyses have shown this is not the case
380 and nowadays an increasing number of studies ~~studies~~ confirm the occurrence of NPF events in
381 urban areas. In this study, for the same period of seven years as for the two background areas, NPF
382 events were found to occur for 6.1% of days at Marylebone Road, lower than in the background
383 areas. Though, due to the particle size range available there cannot be a definitive answer to
384 whether the formation of the particles takes place in the specific ~~area~~ locality of the sampling site,
385 due to the observed increase in particle concentrations in the range 7 – 16 nm (provided by the CPC

386 data) and the increased growth rates found in urban areas in general, it can be assumed that the
387 formation takes place either in the area of the measuring site or in its close vicinity, while the
388 growth of the particles persists in the area for several hours, despite the high condensation sink.

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389 Seasonal variation is similar to that at the background sites, but day of the week variation is stronger
390 at Marylebone Road further favouring weekends (Figure S54), as on these days traffic intensity is
391 lower.

392

393 In general, similar conditions found ~~in the background areas~~ to affect NPF events at the background
394 sites are also found at Marylebone Road, despite a much larger condensation sink. (Figure S24). As
395 a result, less particles of size smaller than 20 nm were found on NPF event days than the average
396 for the site, as the sum of background particles plus those formed on these days were less than that
397 on an average day. The growth rate of the newly formed particles (5.5 nm h⁻¹), is higher than that of
398 the background sites (~~5.5 nm h⁻¹~~), which is in agreement with the findings in the study of the
399 background areas on the possible role of the condensable species, the concentrations of which are
400 even greater at the urban kerbside. About 15% of NPF event days at Marylebone Road presented
401 particle shrinkage after the initial growth; the study of these cases though is outside of the context
402 of the present work. At Marylebone Road, the number of NPF days which were common with the

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403 background sites was fewer, as local conditions (high condensation sink) are detrimental to the
404 occurrence of NPF events and thus the days of regional events including Marylebone Road were
405 separately studied for this site. The regional event days that were common for all three sites were 37

406 (31% of events at Marylebone Road) (Table 1). As with the other two areas, the growth rate is
407 higher during local events, but the conditions are mixed, with lower concentrations of sulphate and
408 organic compounds but higher SO₂, NO_x and elemental carbon. The relationship with higher wind
409 speed (mainly western) (Figure S65), solar radiation (which results in greater H₂SO₄ formation) and
410 lower relative humidity, indicate the stronger relation of the regional events with synoptic
411 conditions than the local events in the heavily polluted environment of Marylebone Road.

412

413 **3.3 Connection of NPF Events with Incoming Air Masses**

414 **3.3.1 Air mass back trajectory clustering and connection with NPF events**

415 The origin of the air masses plays a very important role in the occurrence of NPF events, as shown
416 in Section 3.1.2. Air masses of different origins have different characteristics. Back trajectories
417 provide excellent insight into the source of the air masses. Air mass back trajectories were
418 calculated both for all days and for NPF event days for each site separately, with the aim of
419 complementing the analysis in Section 3.1.2 which addressed only the event days. The additional
420 analysis gives a view of the frequency of NPF events within different air mass types. The initial air
421 mass back trajectory clustering ended up with an optimal solution of 9 clusters of different air
422 masses. As many of these clusters had similar characteristics and origin, solutions with fewer
423 clusters were attempted. As the number of clusters was decreasing clusters became a mixture of
424 different origins, thus making the distinction of different sources harder. As a result, the method

425 chosen was to merge clusters of similar origin and characteristics, which kept the detail of the large
426 number of clusters and made the separation of the different origins more distinct.

427

428 The resulting four merged clusters (Figure 5), using the characterisation proposed by McIntosh et
429 al. (1969), are:

- 430 • An **Arctic** cluster, which originates mainly from the northerly sector. It occurs about 10% of
431 the time and consists of cold air masses, which either passed over northern parts of the U.K. or
432 through the Irish Sea.
- 433 • A **Tropical** cluster, which originates from the central Atlantic. It occurs 25% of the time and
434 contains warmer air masses. A small percentage of this cluster contains masses that have
435 passed over countries south of the U.K. Even though these days were more polluted, the
436 clustering method was unable to clearly distinguish these days as it does not take into account
437 particle numbers or composition, even when the 9-cluster solution was applied.
- 438 • A **Polar** cluster, which originates from the north Atlantic. It is the most common type of air
439 mass arriving in the areas of study and occurs about 40% of the time bringing fast moving,
440 “clean” air masses with increased marine components (Cl, Na, Mg) from the west. This cluster
441 also contains airmasses that have passed through Ireland, though an effect on particle size and
442 chemical composition is not distinct.
- 443 • A **Continental** cluster, which originates from the east. It occurs about 25% of the time and
444 consists mainly of slow moving air masses, originating from the London area (for the

background areas) and/or continental Europe. It has higher concentrations of most pollutants as well as the highest condensation sink.

The occurrence of each air mass class for average and event days for Harwell and London (both sites) can also be found in Figure 5, while their main characteristics for each site can be found in Table S2†. Though in this case the air mass grouping for each site was done in a different analysis, the resulting groups are almost identical in their characteristics and frequency, as the sites are close to each other.

The Polar cluster is the one prevailing on both average and event days. This consists of clean fast-moving air masses originating mainly from mid and high latitudes of the Atlantic, and this cluster presents favourable conditions for NPF events. The association of NPF events with air masses from the mid-Atlantic at N. Kensington was also found by Beddows et al. (2015). Cool Arctic air masses on average are not clean as they may have passed over the northern U.K. The event days associated with this air mass type have the lowest concentrations of the pollutants within available data for all areas. The increased percentage of events with this air mass at all sites indicates that lower temperatures, in a clear atmosphere with sufficient solar radiation are favourable for NPF events as found in previous studies (Napari et al., 2002; Jeong et al., 2010; Kirkby et al., 2011). A similar trend of increased probability with polar and arctic maritime air masses was also found for Hyytiälä, Finland by Nilsson et al. (2001). Tropical air masses have a lower probability for NPF events,

465 which is associated with the fact that a number of these days are associated with air masses which
466 have passed from continental areas south of the U.K. (France, Spain etc.). Specifically for
467 Marylebone Road the NPF probability is a lot lower (11% versus 17% for N. Kensington and 20%
468 for Harwell). This is due to the fact that these air masses are more related to southerly winds which,
469 in Marylebone Road are associated with a street canyon vortex which causes higher pollutant
470 concentrations at this site. Finally, the Continental cluster presents the lowest probability for NPF
471 events. The air masses in this group originate from continental Europe and for the background areas
472 in most cases have passed over the London region as well. This results in both a higher
473 condensation sink and concentration of pollutants, which limits the number of days with favourable
474 conditions for NPF events. Growth rate for all sites though appears to be higher for air masses
475 originating from more polluted areas (Figure 6), which appear to enhance the growth process due to
476 containing a higher concentration of condensable species (after oxidation).

477

478 **3.4 Nucleation Strength Factor (NSF)**

479 The NSF (equations 6 and 7) is used to describe the effect nucleation events have on the number of
480 particles at a site. The values of NSF for each site and for seasons spring and summer are shown in
481 Table 2. The decrease of the contribution of NPF events to particle number, moving from the rural
482 area to the kerbside was also found in previous studies (Salma et al., 2014; 2017). This is explained
483 by the increased contribution to the particle number concentrations of other sources, mainly
484 combustion in the urban environment, compared to rural areas. Apart from this trend, in the

485 background areas the increase of N_{16-100} was greater in spring than summer. This effect seems
486 stronger in the urban background area compared to the rural, as in that area the variability of N_{16-100}
487 is greater for event days compared to that of the rural area. On the other hand, the contribution of
488 NPF events in the longer span, as is illustrated by the NSF_{GEN} appears to favour summer for all
489 areas, showing the increased formation and survivability of particles in this season.

490

491 For Marylebone Road the result for the increase of the N_{16-100} is greater in summer than in spring, in
492 contrast to what was found for the background sites. This is due to the fact that in summer the
493 traffic intensity is decreased, giving the contribution from NPF events a stronger effect compared to
494 the other sources. The very small increase found on NPF events in Marylebone Road, with a factor
495 of just 1.26, a lot lower than that found in the urban area of Seoul, South Korea (Park et al., 2015),
496 is indicative of the reduced effect of NPF events in an area which is heavily affected by traffic, as
497 also pointed out by Von Bismarck-Osten et al. (2013) in their study on particle composition in
498 Marylebone Road.

499

500 3.5 The Survival Parameter P

501 ~~The survival parameter P is a measure of the probability for newly formed particles to survive to~~
502 ~~detectable sizes.~~ The average values of the P parameter for each of the areas of this study are 10.5
503 for Harwell, 15.8 for N. Kensington and 28.9 for Marylebone Road. The values found put
504 Marylebone Road to the upper end of heavily polluted areas in Europe, North Kensington to the

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505 same level as many other urban areas in Europe, while Harwell had somehow higher values
506 compared to other rural background areas in Europe, as calculated by Kulmala et al. (2017). The
507 seasonal, air mass origin and local versus regional variations can be found in Figure 7 (winter is
508 excluded due to very low number of events). While the increasing trend of the P parameter as we
509 move from rural background to kerbside was expected, it can be seen that there is a clear seasonal
510 pattern in all three areas, with summer having the lowest P parameter (greatest survivability)
511 compared to the other two seasons. This is associated with the higher growth rate found in summer
512 for all areas of this study, as the differences in the condensation sink on event days are negligible
513 between seasons. The case is similar for regional and local events. The result per air mass origin is
514 related to the different conditions and parameters of each incoming air mass in each area. For
515 example, the higher P parameter for Tropical air masses at Marylebone Road, is associated with the
516 higher condensation sink found for this kind of air masses, due to the street canyon effect which is
517 specific for Marylebone Road for southerly wind directions with which these air masses are mainly
518 related, while the higher values for the rather clean Arctic air masses for the other two areas are
519 associated with the lower growth rates found for this kind of air mass in these areas. The more
520 polluted Continental air masses seem to have a different effect for rural and urban areas. Their
521 higher condensation sinks and concentrations of pollutants have a negative effect on P-values for
522 the rural site and a positive effect at the urban sites. The exact opposite is found for the cleaner air
523 masses of the Polar cluster, which appear to result in reduced P-values of the newly formed

524 particles at the urban sites. This is related to the lower condensation sink associated with this air
525 mass type.

526

527 **4. CONCLUSIONS**

528 Seven years of data from three distinct areas (regional background, urban background, kerbside) in
529 the southern U.K. were analysed and the conditions associated with NPF events were studied. NPF
530 events were found to occur on about 7% of days at background sites and less at the kerbside site.

531 The conditions on event days for all three areas were similar, with clear atmospheric conditions and
532 a lower condensation sink. While the condensation sink appears to be the most important factor

533 limiting NPF events at the kerbside site, SO₂ was found to have smaller concentrations on event
534 days for all areas, which indicates that either on average it is in sufficient concentrations for NPF

535 events to occur, or that other variables that participate in the production mechanism of H₂SO₄ are

536 more important. The growth rate of the newly formed particles increases from the rural site to the

537 kerbside and is greater in summer compared to other seasons for all three sites. Almost half of the

538 NPF events at the rural and urban background sites were found to happen simultaneously. In these

539 cases, the atmospheric conditions were cleaner, which resulted in slower growth rates. While most

540 of the chemical species available were at lower concentrations in regional events, a difference in the

541 behaviour with respect to sulphate and organic compounds was found between the two background

542 site types.

543

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(2nd)

544 The prevailing origin of air masses in the southern U.K. is from mid and high latitudes of the
545 Atlantic Ocean. These fast-moving air masses present an increased probability for NPF to occur.
546 The case is similar for the cooler and cleaner arctic air masses, while air masses from the tropics
547 and continental Europe, having greater pollutant content, have decreased NPF probability, but a
548 higher growth rate of particles when NPF events occurred. Regional events appear to be more
549 associated with cleaner air masses, presenting a smaller growth rate and condensation sink
550 compared to local events. The difference in growth rate is probably related to the greater content of
551 condensable species; a positive relation of particle survival probability with temperature was also
552 found.

553
554 Comparing the background areas in this study, particles of 16-20 nm were found to be about 20%
555 greater in concentration (above long-term average) on NPF event days at the urban [background](#)
556 [background](#) site compared with the rural site. This is associated with a higher abundance of
557 condensable species in the urban environment, which enhances the nucleation and growth process.
558 This effect though is limited as particle size increases and NPF events have a greater effect on the
559 overall $N_{<100\text{ nm}}$ in the rural areas, compared to urban, as calculated by the NSF. The effect becomes
560 even smaller at the kerbside as the number of background particles emitted by traffic is a lot greater.
561
562 The occurrence of NPF events at the highly polluted Marylebone Road site is at first sight
563 surprising given the elevated condensation sink. This must be counteracted by an abundance of

condensable material, which is surprising given the generally modest rate of atmospheric oxidation processes in comparison to residence times in a street canyon (Harrison, 2017). However, Giorio et al. (2015), using Aerosol Time-of-Flight Mass Spectrometry, reported rapid chemical processes within the Marylebone Road street canyon leading to production of secondary particulate matter from road traffic emissions. They postulated that this resulted from very local gas to particle conversion from vehicle-emitted pollutants. Condensation of such reaction products upon pre-existing particles could explain the enhanced particle growth rates observed at Marylebone Road (Figure 3).

Finally, particle survival probability was found to decrease moving from rural to urban areas. While formation and initial growth of new particles is increased in urban areas, their survivability reduces as their size increases. The probability of particles to survive to greater sizes was found to be increased in summer for all areas, which is also explained by the higher growth rate. The probability is also different depending upon the origin of the air masses and is related to conditions specific for each area.

In the present work, the effects of atmospheric conditions upon the NPF process are studied. NPF is a complex process, highly affected by meteorological conditions (local and synoptic), the chemical composition as well as the pre-existing conditions in an area. For this reason, the study of NPF events in one area cannot provide safe assumptions for other areas, as the mixture of conditions

584 found in different places is unique and alters the occurrence and development of NPF events. Thus,
585 more studies on the conditions and the trends in NPF events should be conducted to better
586 understand the effect of the numerous variables that affect those processes.

587

588 **DATA AVAILABILITY**

589 Data supporting this publication are openly available from the UBIRA eData repository at
590 <https://doi.org/10.25500/edata.bham.00000307>.

591

592 **AUTHOR CONTRIBUTIONS**

593 This study was conceived by MD and RMH who also contributed to the final manuscript. The data
594 analysis was carried out by DB with guidance from DCSB, and DB also prepared the first draft of
595 the manuscript. FDP provided advice on the analysis.

596

597 **COMPETING INTERESTS**

598 The authors have no conflict of interests.

599

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604

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1044 **TABLE LEGENDS:**

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1046 **Table 1:** Number of NPF events per site.

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1048 **Table 2:** Annual and seasonal NSF for all areas of study.

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1052 **FIGURE LEGENDS:**

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1054 **Figure 1:** Map of the measuring stations.

1055 **Figure 2:** Number of NPF events per season [for all seven years of the present study](#) (Winter –
1056 DJF; Spring – MAM; Summer – JJA; Autumn – SON) at Harwell (rural), N.
1057 Kensington (urban background) and Marylebone Road (urban roadside).

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1059 **Figure 3:** Growth rate per season at the three sites.

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1061 **Figure 4:** Diurnal variation of $N_{16-20nm}$ at each site: annual average and NPF event days.

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1063 **Figure 5:** Map and frequency of incoming air mass origin – average and for NPF events per site.

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1065 **Figure 6:** Growth rate per incoming air mass at each of the sites.

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1067 **Figure 7:** Survival parameter P (a) per season, (b) for regional and local events (for Marylebone
1068 Road) is regional for all 3 sites and (c) by incoming air mass origin.

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1072 **Table 1:** Number of NPF events per site.
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| | Harwell | N. Kensington | Marylebone Road | Regional (Background sites)* | Regional (All 3 sites)** |
|---------|---------|------------------|--------------------|------------------------------------|--------------------------------|
| 2009 | 9 | 0 | 4 | 0 | 0 |
| 2010 | 29 | 22 | 22 | 11 | 9 |
| 2011 | 15 | 10 | 23 | 4 | 1 |
| 2012 | 8 | 28 | 12 | 3 | 0 |
| 2013 | 25 | 23 | 27 | 13 | 11 |
| 2014 | 29 | 34 | 13 | 18 | 6 |
| 2015 | 25 | 22 | 18 | 11 | 10 |
| Overall | 140 | 139 | 119 | 60 | 37 |

1074 * Refers to events occurring simultaneously at Harwell and N. Kensington
1075 ** Refers to events which occur simultaneously at all three sites

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1079 **Table 2:** Annual and seasonal NSF for all areas of study.
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| | Harwell | N. Kensington | Marylebone Road |
|--------------------------------|---------|------------------|--------------------|
| NSF _{NUC} (Spring) | 2.04 | 2.03 | 1.20 |
| NSF _{NUC} (Summer) | 2.01 | 1.72 | 1.26 |
| NSF _{NUC} (Year) | 2.25 | 1.86 | 1.26 |
| NSF _{GEN} (Spring) | 1.10 | 1.07 | 1.02 |
| NSF _{GEN} (Summer) | 1.18 | 1.11 | 1.01 |
| NSF _{GEN} (Year) | 1.10 | 1.06 | 1.02 |

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Figure 1: Map of the measuring stations.

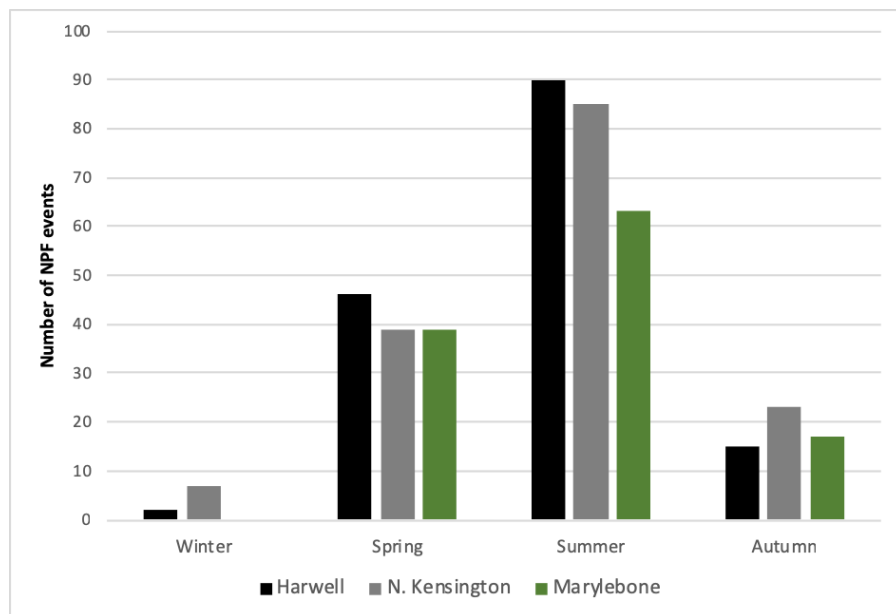


Figure 2: Number of NPF events per season for all seven years of the present study (Winter – DJF; Spring – MAM; Summer – JJA; Autumn – SON) at Harwell (rural), N.Kensington (urban background) and Marylebone Road (urban roadside).

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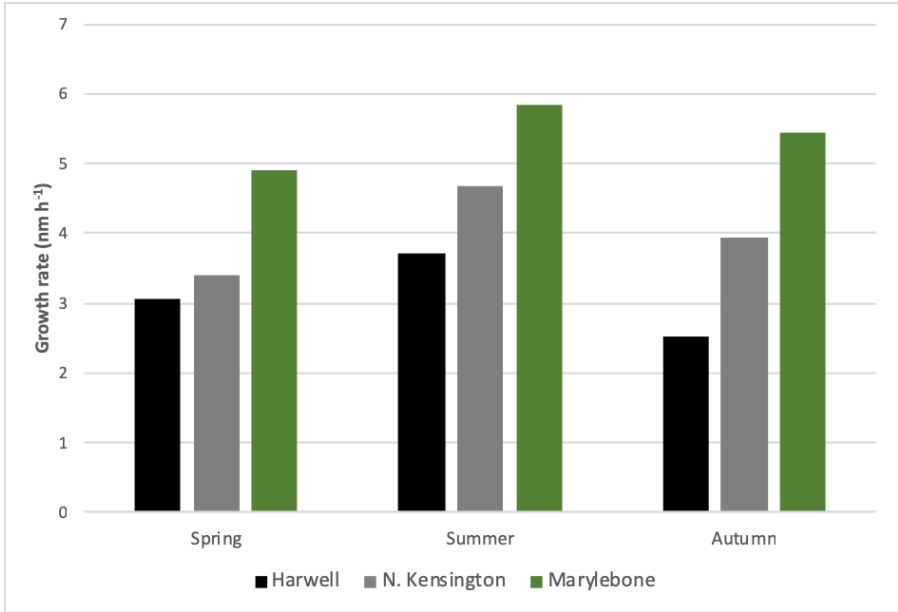


Figure 3: Growth rate per season at the three sites.

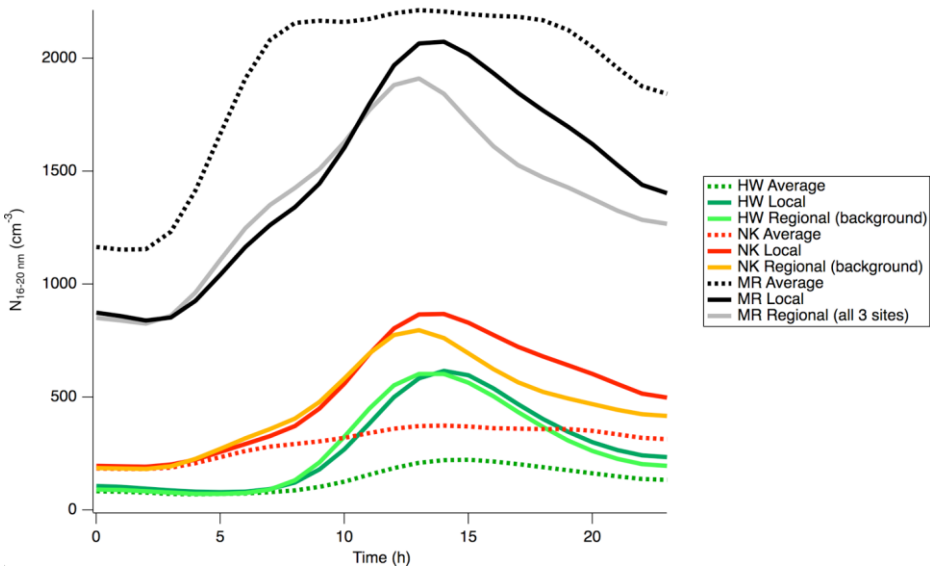
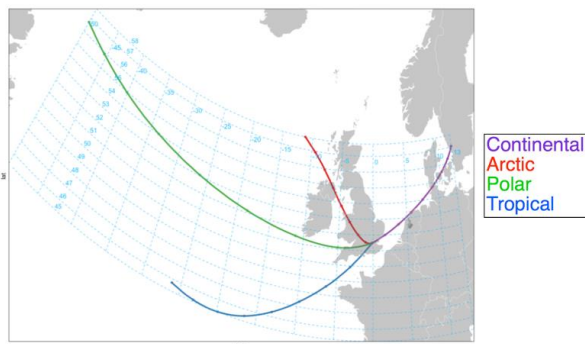
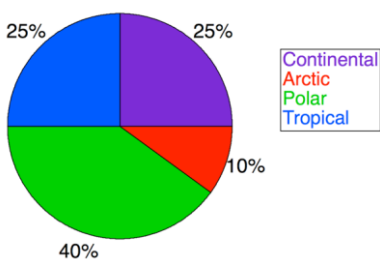


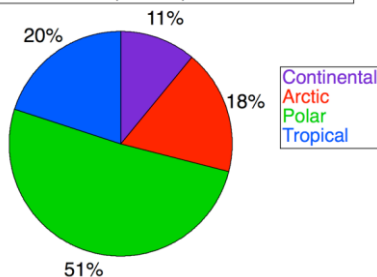
Figure 4: Diurnal variation of $N_{16-20\text{nm}}$ at each site: annual average and NPF event days.



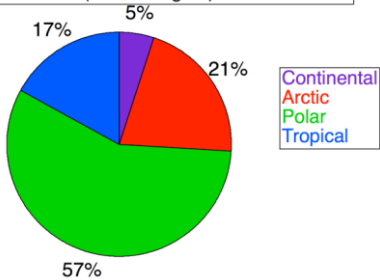
Frequency per air mass trajectory



Frequency of event days per air mass trajectory (Harwell)



Frequency of event days per air mass trajectory (N. Kensington)



Frequency of event days per air mass trajectory (Marylebone Road)

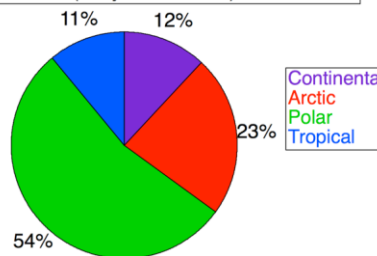
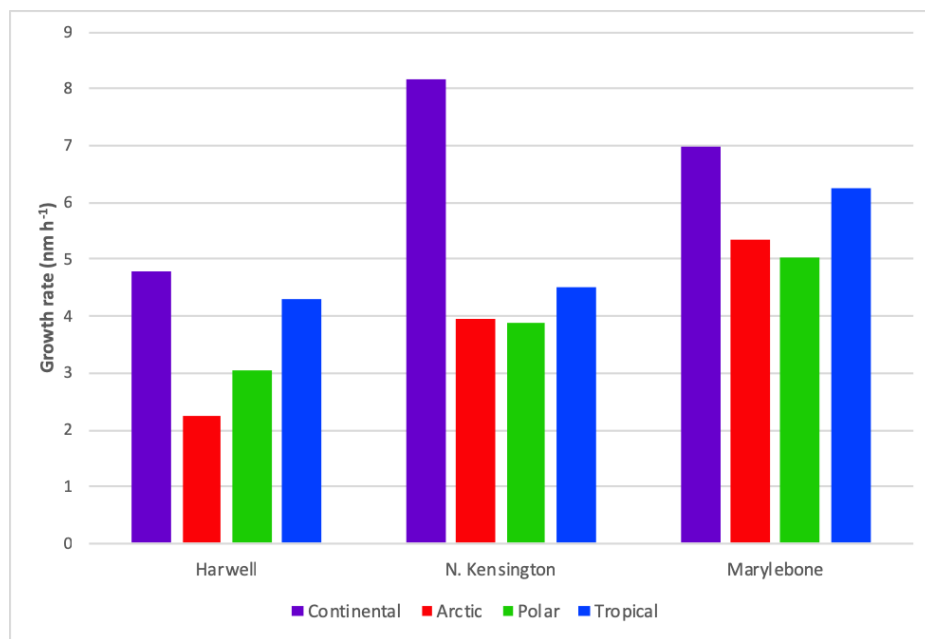


Figure 5: Map and frequency of incoming air mass origin – average and for NPF events per site.

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1139 **Figure 6:** Growth rate per incoming air mass origin at each of the sites.

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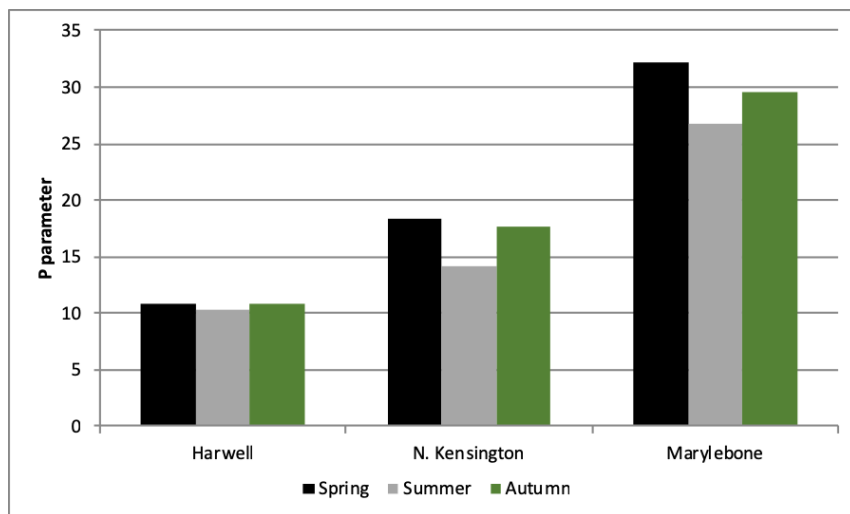
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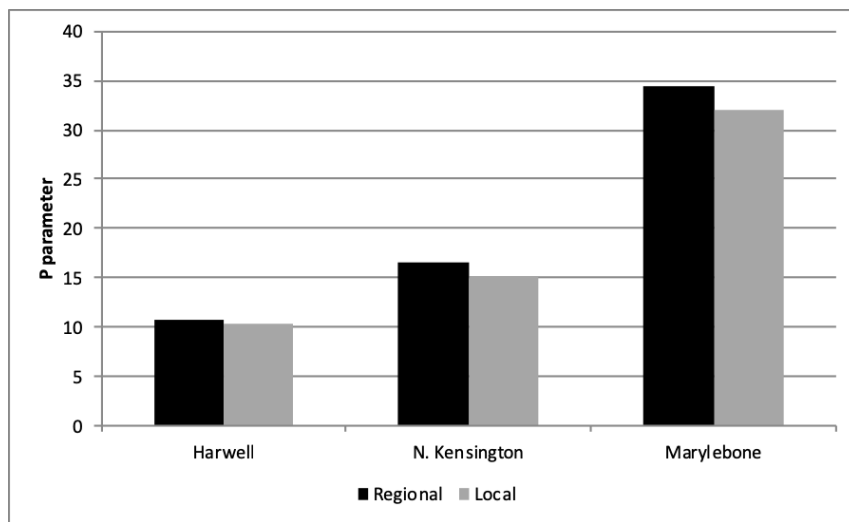
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(a)

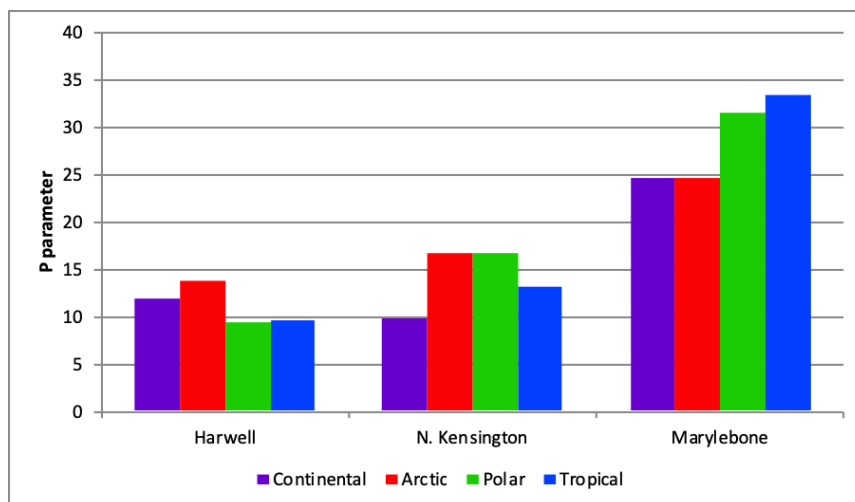


(b)



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(c)



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1152 **Figure 7:** Survival parameter P (a) per season, (b) for regional and local events (for Marylebone
1153 Road regional is for all 3 sites) and (c) by incoming air mass origin.

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1 **SUPPLEMENTARY INFORMATION**

2

3 **Analysis of New Particle Formation (NPF) Events at Nearby**
4 **Rural, Urban Background and Urban Roadside Sites**

5

6 **Dimitrios Bousiotis, Manuel Dall'Osto, David C.S. Beddows and**
7 **Roy M. Harrison**

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Table S1: Data availability per season (all numbers are percentages of available data)

| | Harwell | | | | N. Kensington | | | | Marylebone Road | | | |
|------|---------|--------|--------|--------|---------------|--------|--------|--------|-----------------|--------|--------|--------|
| | Winter | Spring | Summer | Autumn | Winter | Spring | Summer | Autumn | Winter | Spring | Summer | Autumn |
| 2009 | 15 | 97 | 10 | 80 | 57 | 97 | 100 | 100 | 100 | 65 | 86 | 68 |
| 2010 | 37 | 53 | 100 | 95 | 58 | 87 | 93 | 100 | 46 | 100 | 87 | 86 |
| 2011 | 72 | 75 | 99 | 73 | 89 | 87 | 73 | 89 | 79 | 99 | 100 | 67 |
| 2012 | 82 | 86 | 100 | 95 | 56 | 88 | 99 | 86 | 0 | 0 | 87 | 66 |
| 2013 | 91 | 70 | 99 | 100 | 84 | 92 | 98 | 98 | 57 | 92 | 84 | 100 |
| 2014 | 97 | 62 | 99 | 99 | 84 | 78 | 97 | 98 | 89 | 79 | 76 | 99 |
| 2015 | 77 | 100 | 61 | 70 | 80 | 99 | 65 | 100 | 74 | 100 | 98 | 100 |

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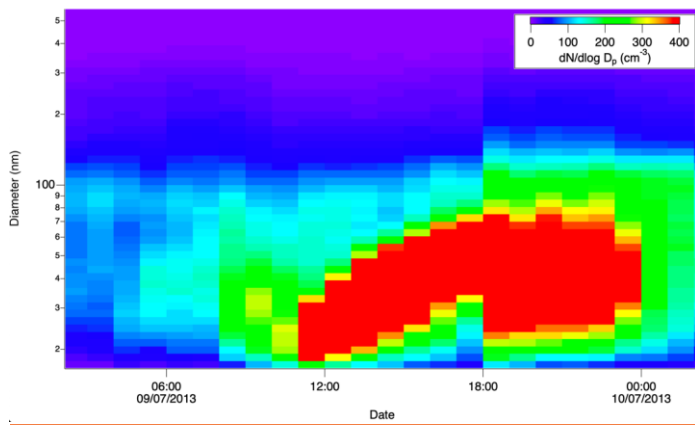
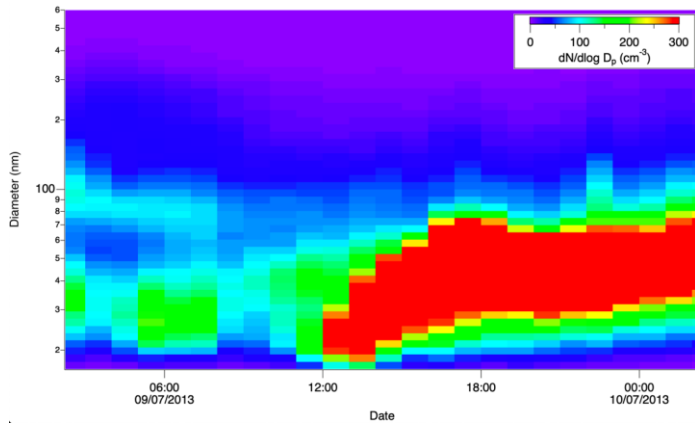
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Table S21: Conditions per air mass origin for NPF event days (April – October average in parenthesis) for all areas of study.

| Harwell | | | | |
|--|------------------|---------------------|---------------------|---------------------|
| | Continental | Arctic | Polar | Tropical |
| Condensation sink (s^{-1}) | 5.05E-03 (5E-03) | 2.71E-03 (3.32E-03) | 2.57E-03 (2.87E-03) | 3.19E-03 (2.87E-03) |
| Wind speed ($m s^{-1}$) | 3.52 (3.63) | 3.87 (3.47) | 3.64 (3.69) | 3.74 (4.17) |
| Temperature ($^{\circ}C$) | 15.5 (13.6) | 12.2 (11.5) | 13.6 (13.1) | 16.3 (15) |
| SO ₂ ($\mu g m^{-3}$) | 1.87 (1.81) | 1.11 (1.82) | 1.11 (1.27) | 1.22 (1.36) |
| NO _x ($\mu g m^{-3}$) | 9.58 (13.9) | 5.49 (8.01) | 4.66 (7.2) | 5.81 (7.69) |
| SO ₄ ²⁻ ($\mu g m^{-3}$) | 2.70 (3.3) | 1.37 (2.05) | 1.44 (1.64) | 1.37 (1.57) |
| Particulate OC ($\mu g m^{-3}$) | 2.85 (2.88) | 1.35 (1.59) | 1.52 (1.63) | 1.98 (1.76) |

| North Kensington | | | | |
|--|---------------------|---------------------|---------------------|---------------------|
| | Continental | Arctic | Polar | Tropical |
| Condensation sink (s^{-1}) | 7.20E-03 (9.35E-03) | 5.20E-03 (6.37E-03) | 5.40E-03 (6.38E-03) | 4.89E-03 (6.32E-03) |
| Wind speed ($m s^{-1}$) | 3.89 (3.44) | 3.92 (3.65) | 4.46 (4.2) | 4.74 (4.44) |
| Temperature ($^{\circ}C$) | 18.4 (15) | 12.7 (13.1) | 15.5 (14.6) | 17 (16.4) |
| SO ₂ ($\mu g m^{-3}$) | 1.68 (2.23) | 1.33 (1.89) | 1.73 (1.75) | 1.74 (1.72) |
| NO _x ($\mu g m^{-3}$) | 33.5 (55) | 28.5 (39.2) | 30.3 (39.4) | 24 (34.9) |
| SO ₄ ²⁻ ($\mu g m^{-3}$) | 1.93 (2.23) | 0.95 (1.36) | 0.98 (1.13) | 1.30 (1.47) |
| Particulate OC ($\mu g m^{-3}$) | 3.84 (4.90) | 2.24 (2.95) | 2.81 (2.96) | 2.43 (3.03) |

| Marylebone | | | | |
|--|---------------------|---------------------|--------------------|---------------------|
| | Continental | Arctic | Polar | Tropical |
| Condensation sink (s^{-1}) | 1.65E-02 (1.96E-02) | 1.16E-02 (1.57E-02) | 1.4E-02 (2.14E-02) | 1.82E-02 (2.39E-02) |
| Wind speed ($m s^{-1}$) | 3.92 (3.41) | 3.50 (3.64) | 3.84 (4.13) | 4.77 (4.4) |
| Temperature ($^{\circ}C$) | 17.4 (15.2) | 13.4 (13.4) | 15.3 (14.8) | 16.9 (16.3) |
| SO ₂ ($\mu g m^{-3}$) | 4.99 (6.39) | 4.31 (5.63) | 5.38 (7.43) | 6.95 (8.17) |
| NO _x ($\mu g m^{-3}$) | 172 (250) | 139 (214) | 191 (303) | 269 (336) |
| SO ₄ ²⁻ ($\mu g m^{-3}$) | 3.24 (3.35) | 1.47 (1.6) | 1.52 (1.61) | 1.24 (1.8) |
| Particulate OC ($\mu g m^{-3}$) | 6.03 (6.91) | 3.81 (4.73) | 4.67 (5.97) | 5.31 (6.6) |



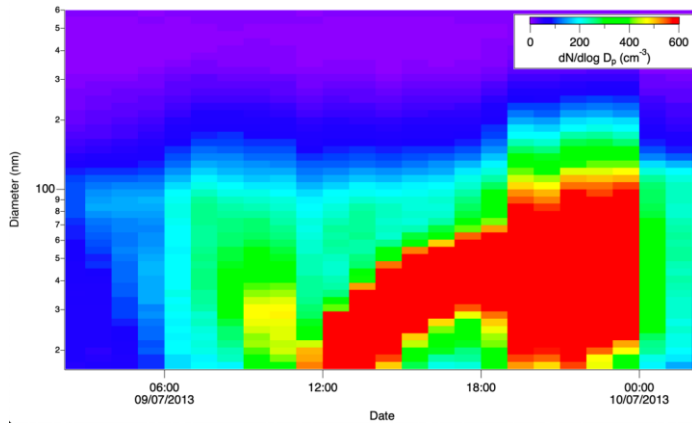
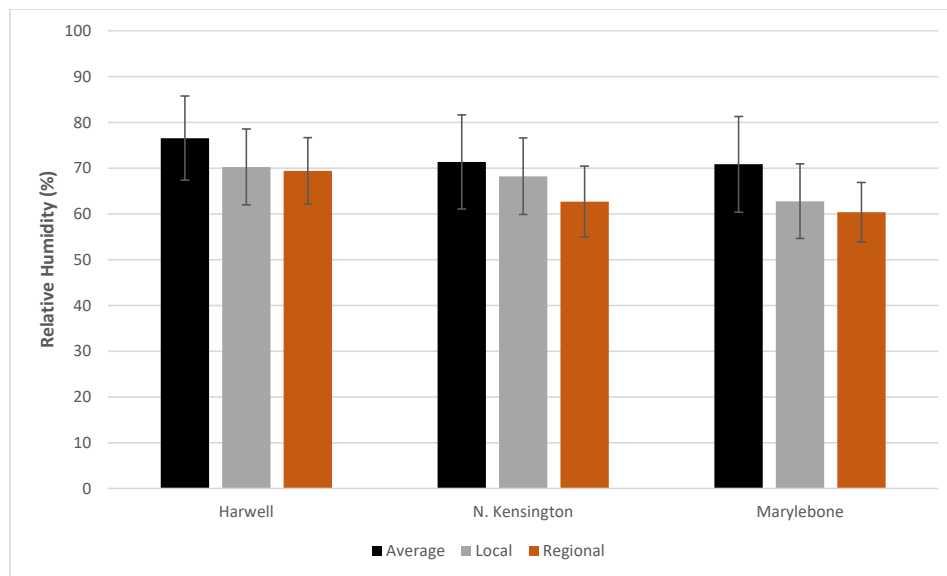
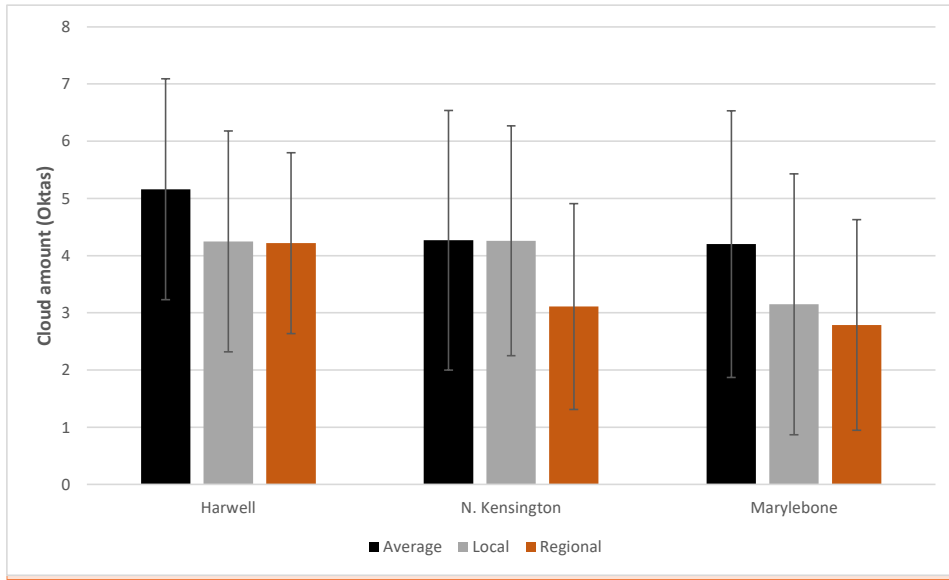


Figure S1: Example of a regional NPF event for all the sites of the present study (from top to bottom is HW, NK, MR)

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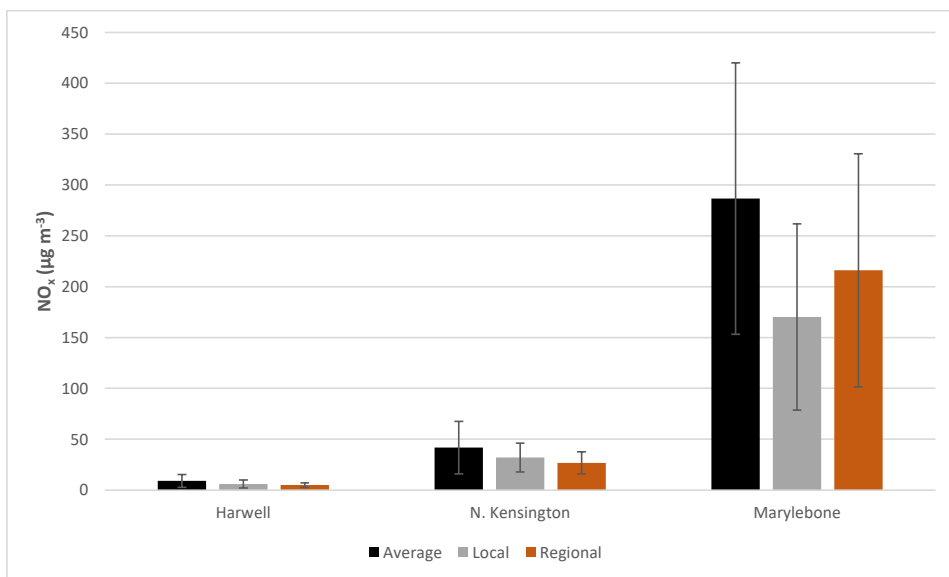


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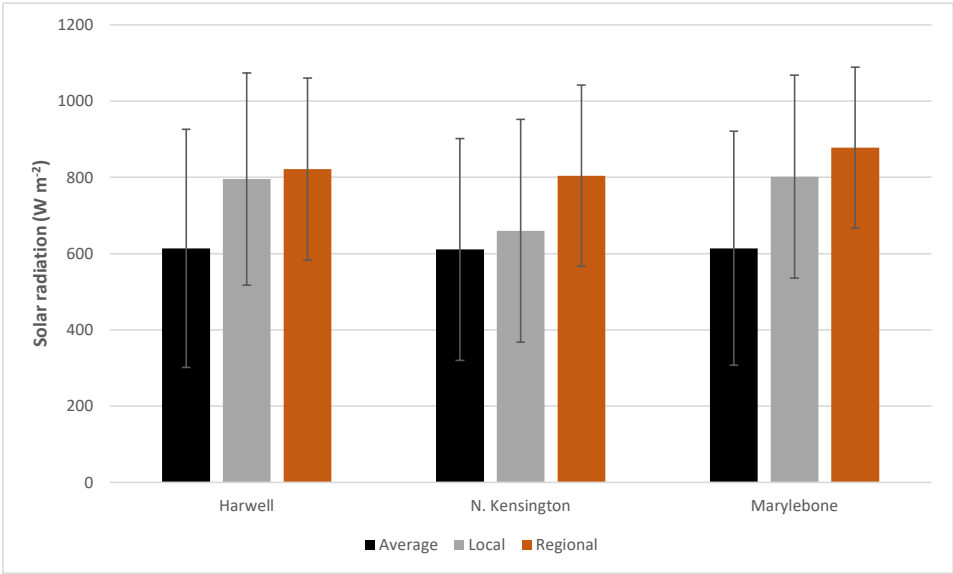


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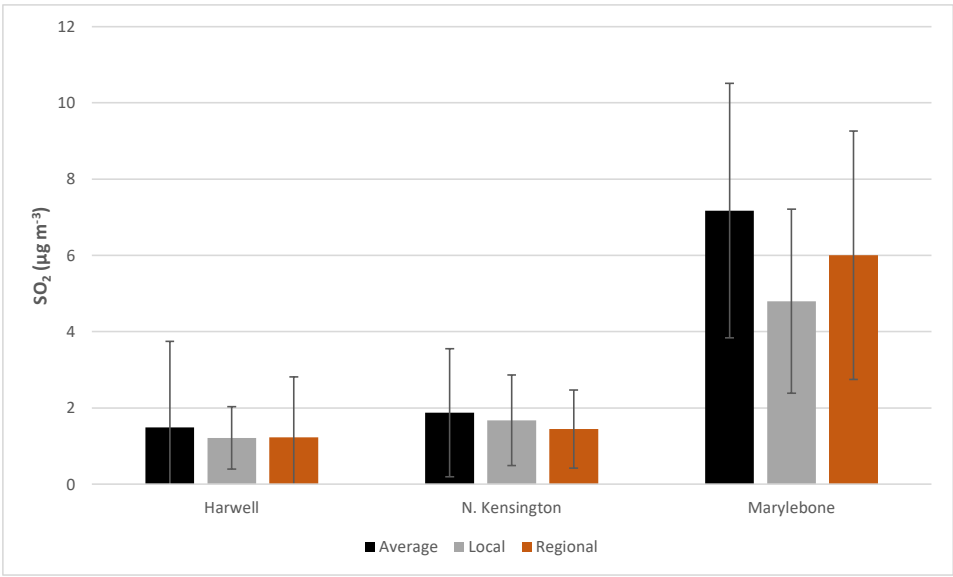
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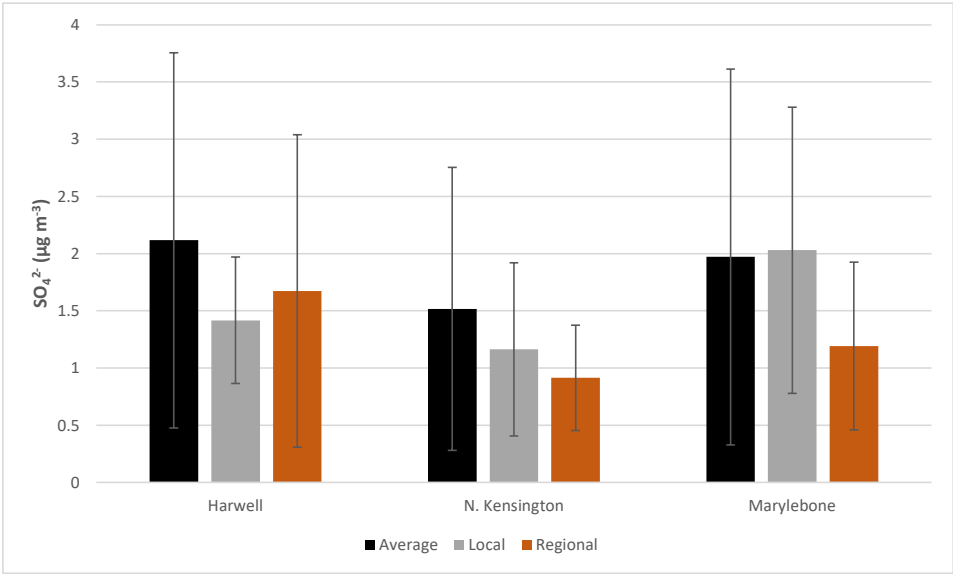


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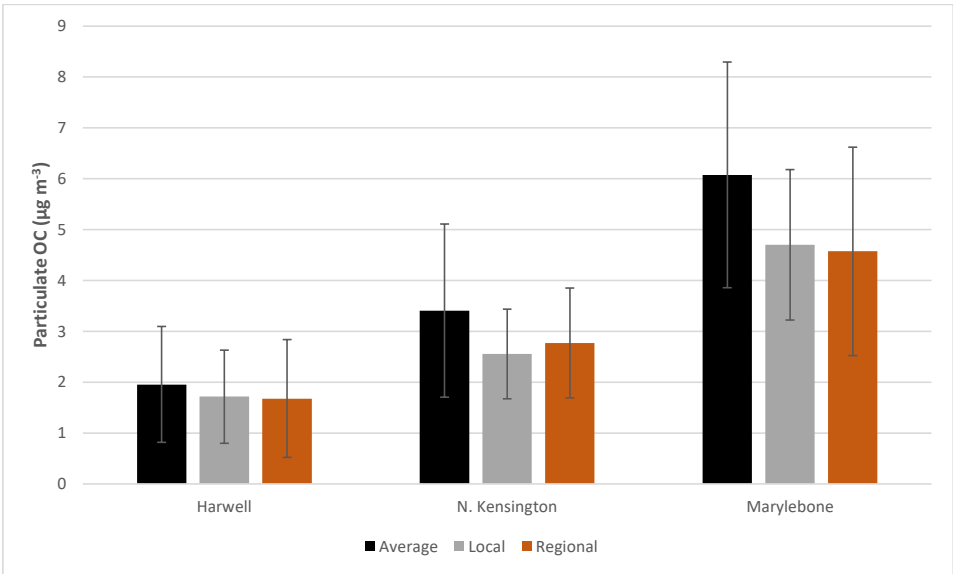


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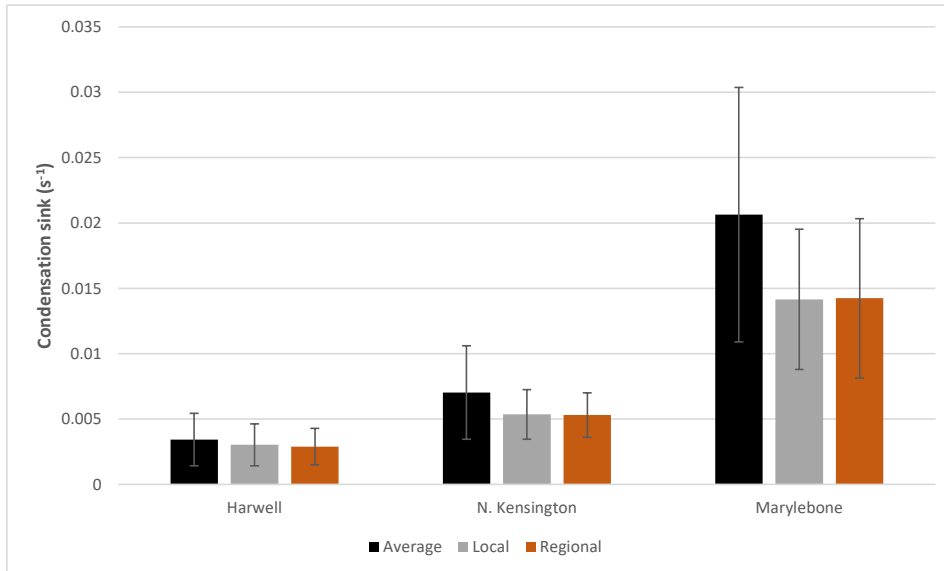
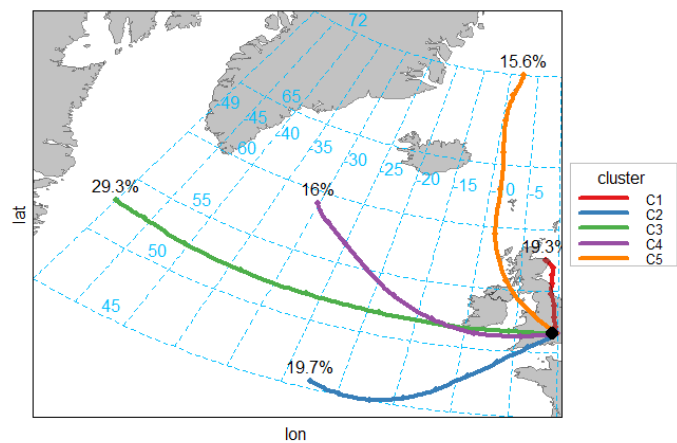
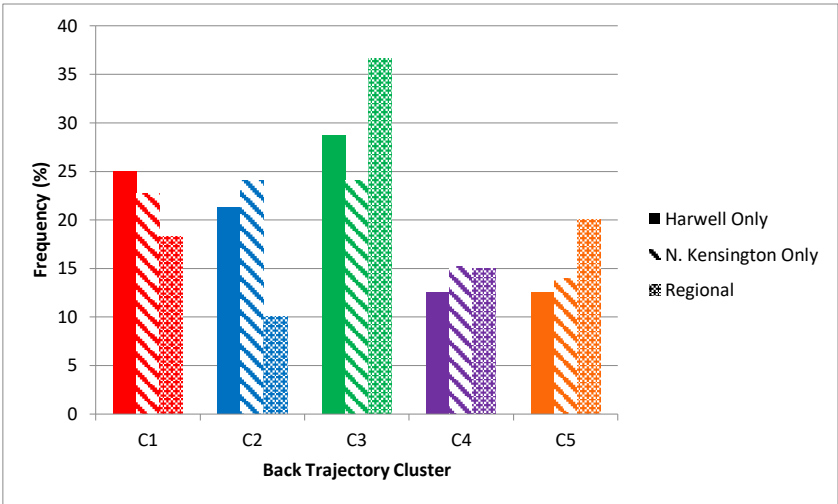


Figure S24: Average and NPF event conditions for Harwell and N. Kensington and Marylebone Road. On Marylebone Road, Regional events' conditions refer to the Regional event days for all 3 sites.

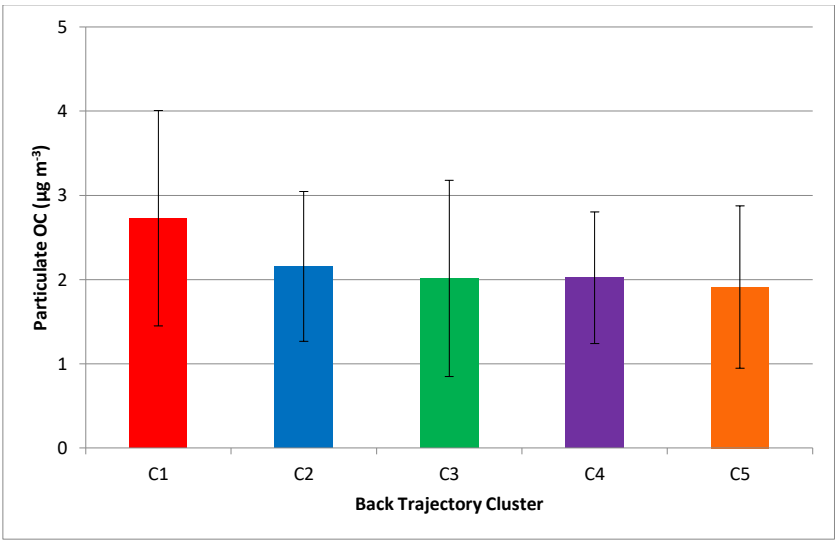
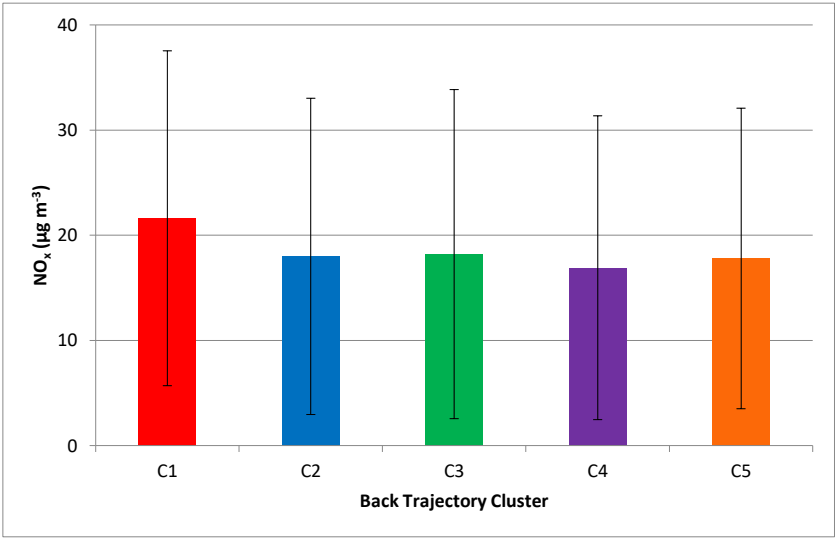
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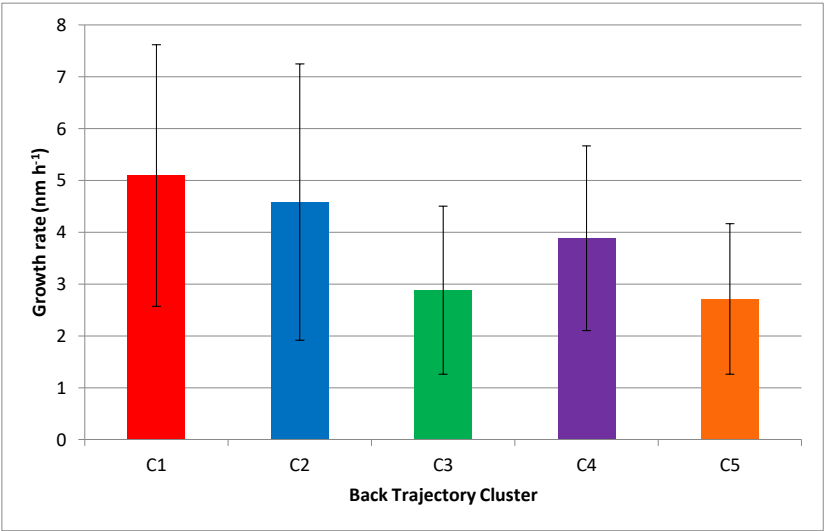
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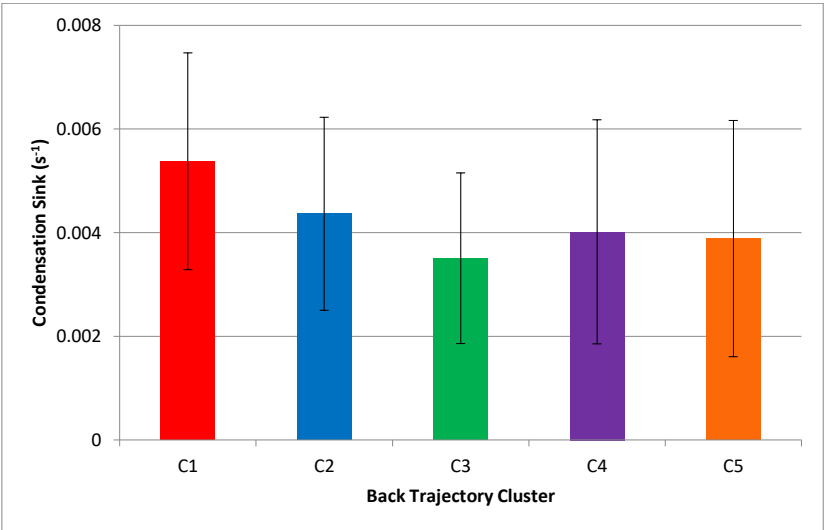
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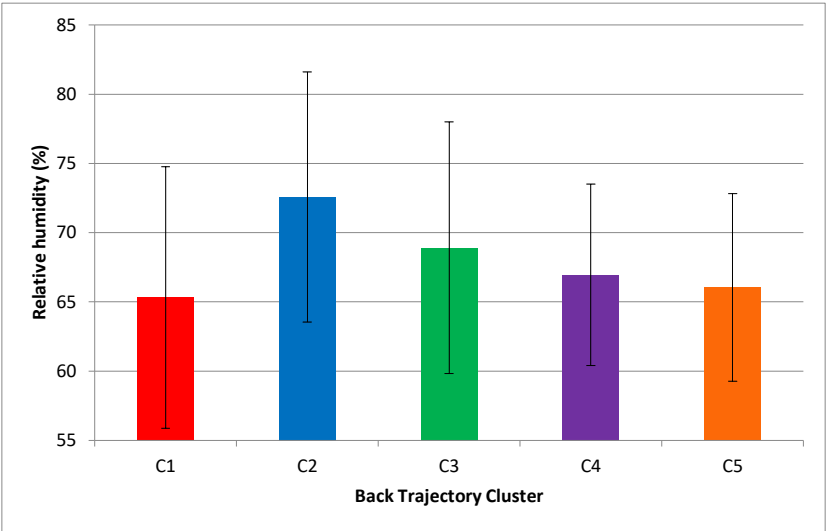


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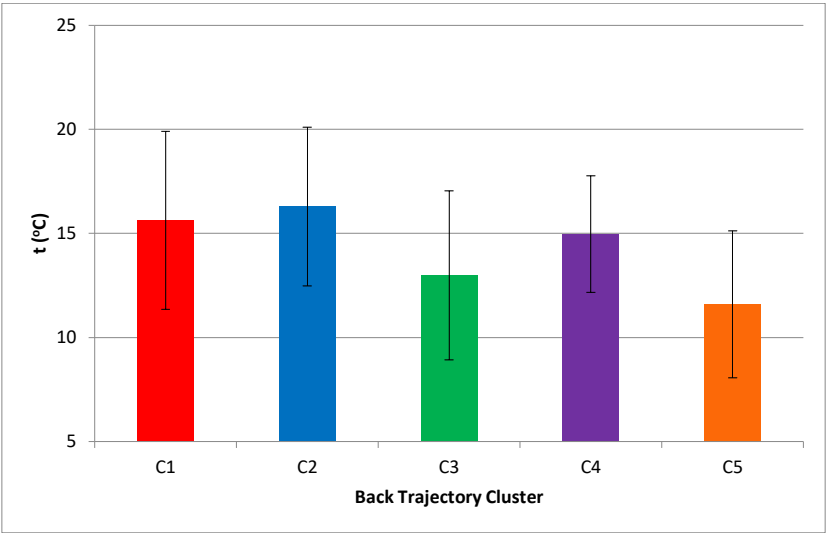


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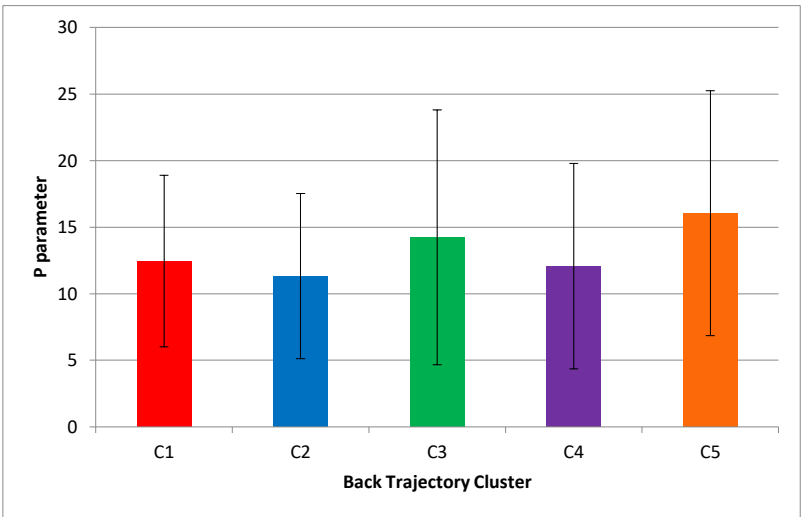
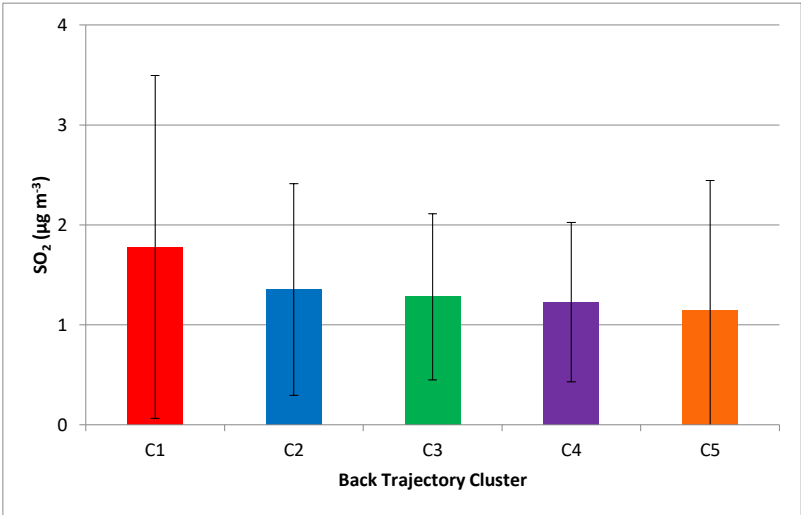


Figure S32: Air mass origin frequency and conditions for NPF events in Harwell and N. Kensington.

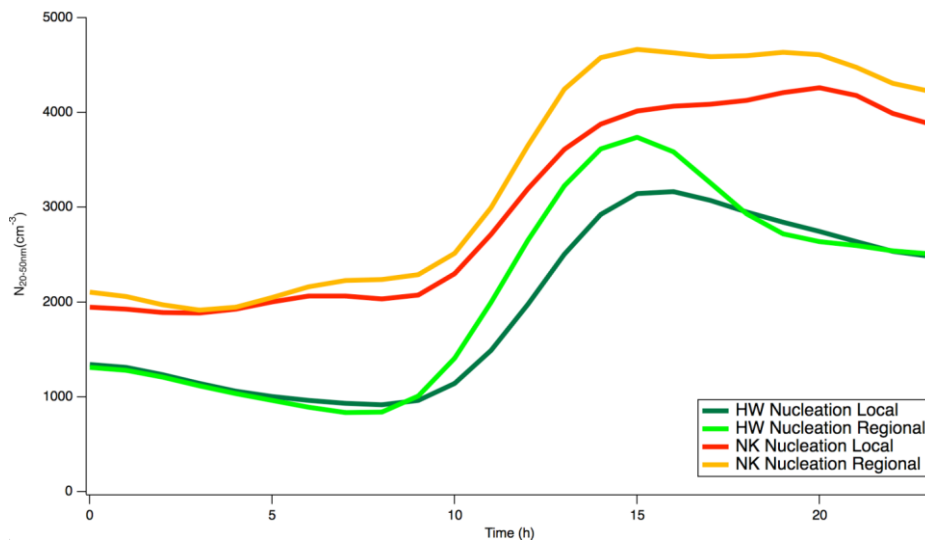


Figure S3: Diurnal variation of $N_{20-50nm}$ for Harwell and N. Kensington during local and regional NPF events.

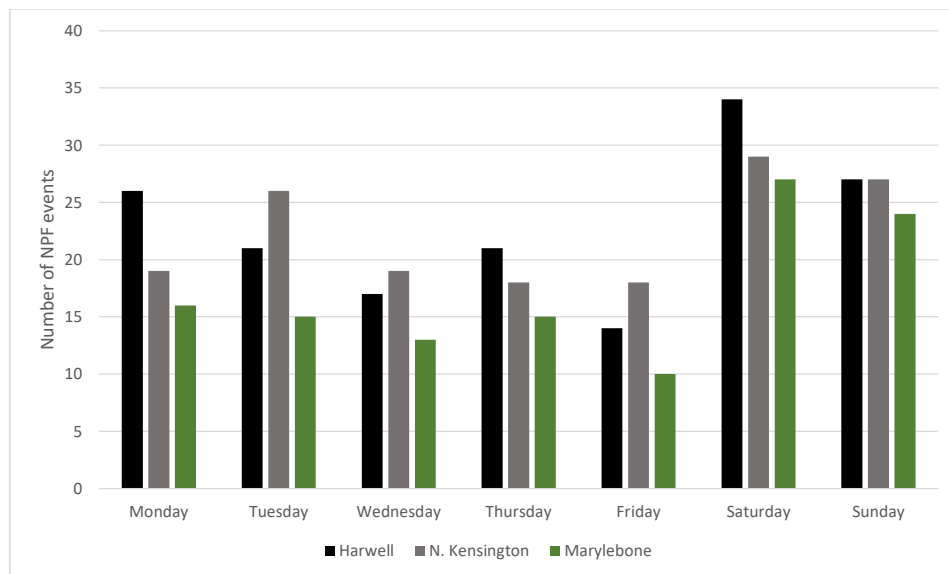
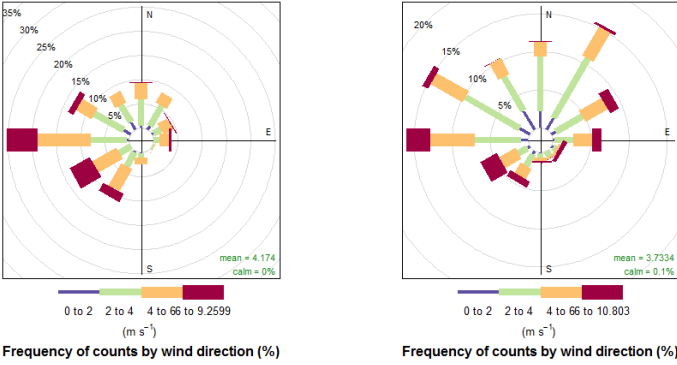


Figure S4: Weekly variation of NPF events in Marylebone Road.

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70 **Figure S65:** Wind profile for Local (left) and Regional (right) NPF events in Marylebone Road.

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