Sources of Submicrometre Particles Near a Major International Airport

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ABSTRACT

The international airport of Heathrow is a major source of nitrogen oxides, but its contribution to the levels of submicrometre particles is unknown, and is the objective of this study. Two sampling campaigns were carried out during warm and cold seasons at a site close to the airfield (1.2 km). Size spectra were largely dominated by ultrafine particles: nucleation particles (<30 nm) were found to be ~10 times higher than those commonly measured in urban background environments of London. Five clusters and 6 factors were identified by applying *k*-means cluster analysis and positive matrix factorization (PMF) respectively to particle number size distributions; their interpretation was based on their modal structures, wind directionality, diurnal patterns, road and airport traffic volumes and on the relationship with weather and other air pollutants. Airport emissions, fresh and aged road traffic, urban accumulation mode and two secondary sources were then identified and apportioned. The fingerprint of Heathrow has a characteristic modal structure peaking at <20 nm and accounts for 30-35% of total particles in both the seasons. Other main contributors are fresh (24-36%) and aged (16-21%) road traffic emissions and urban accumulation from London (around 10%). Secondary sources accounted for less than 6% in number concentrations but for more than 50% in volume concentration. The analysis of a strong regional nucleation event showed that both the cluster categorisation and PMF contributions were affected during the first 6 hours of the event. In 2016, the UK government provisionally approved the construction of a third runway; therefore the direct and indirect impact of Heathrow on local air quality is expected to increase unless mitigation strategies are applied successfully.

Keywords: Airport; black carbon; size distributions; source apportionment; ultrafine particles

1. INTRODUCTION

Emerging markets, developing economies and globalisation have driven a fast and continuing growth of civil aviation in the last decades (Lee et al., 2009); this trend is still growing by ~5.5% y^{-1} (ICAO, 2017). As a consequence, the aircraft and road traffic at airports is also increasing, but the information available on the impact of airport emissions upon air quality at ground level is still inadequate (Webb et al., 2008; Masiol and Harrison, 2014). The quantification of airport impacts on local air quality is complicated by the complexity of multiple mobile and static emission sources, with many airports being located near to major cities, highways or industrial plants. Consequently, the development of successful strategies for emission mitigation and the implementation of measures for air quality improvement to meet regulatory standards require a detailed quantification of the contribution of airport and other emissions to the total air pollution load.

Biological evidence associates the exposure to ultrafine particles (UFPs, <100 nm) with adverse effects upon human health (e.g., Knibbs et al., 2011; Strak et al., 2012; Ostro et al., 2015; Lanzinger et al., 2016). At the current time, there is still limited knowledge of what specific characteristic or association of characteristics may dominate the particle toxicity, and the consequent health outcomes (Atkinson et al., 2010; Strak et al., 2012, Vu et al., 2015a); nevertheless it is well recognised that UFPs can reach the deepest regions of the lung (Salma et al., 2015) and may have orders of magnitude higher surface area to mass ratios compared to larger particles. They offer more surface for the absorption of volatile and semi-volatile species (Kelly and Fussell, 2012; Strak et al., 2012).

Several studies have reported large increases of UFPs near airports (e.g., Westerdahl et al., 2008; Hu et al., 2009; Klapmeyer et al., 2012; Hsu et al., 2012a;b). For example, Hsu et al. (2013) and Stafoggia et al. (2016) detected substantial increases of total particle number concentration (PNC) at the airports of Los Angeles (CA, USA) and Rome Ciampino (Italy), respectively, in the few

minutes after take-offs, especially downwind, while landings made only a modest contribution to ground-level PNC observations. Hsu et al. (2014) observed that departures and arrivals on a major runway of Green International Airport (Warwick, RI, USA) had a significant influence on UFP concentrations in a neighborhood proximate to the end of the runway. In a study carried out at the Los Angeles international airport (CA, USA), Hudda et al. (2014) concluded that emissions from the airport increase PNC by 4- to 5-fold at 8–10 km downwind of the airfield, while Shirmohammadi et al. (2017) reported that the daily contributions of the airport to PNC were approximately 11 times greater than those from three surrounding freeways. Hudda et al. (2016) reported that average PNC were 2- and 1.33-fold higher at sites 4 and 7.3 km from the Boston (MA, USA) airport when winds were from the direction of the airfield compared to other directions.

Despite the strong evidence that airports are major sources of UFPs, their fingerprint within the particle number size distribution (PNSD) may be difficult to identify due to: (i) the nature of semi-volatile compounds emitted by aircraft; (ii) the possible mechanisms of secondary aerosol formation; (iii) the dilution effect; and (iv) the similar modal structures of other emission sources concurrently found in cities, such as road traffic (Masiol and Harrison, 2014). Generally, studies performed within or close to airports have reported increases of particles ranging from 4 to 100 nm in diameter and mostly distributed in the nucleation range (<30 nm). For example, Mazaheri et al. (2009) showed a main nucleation mode and an accumulation mode (40-100 nm) more evident during take-offs; Keuken et al. (2015) reported PNSD dominated by 10-20 nm particles in an area affected by emissions from Schiphol airport (The Netherlands); Hudda and Fruin (2016) found strong increases in particles smaller than 40 nm downwind from the Los Angeles International Airport; Ren et al. (2016) showed that particles peaking at 16 nm dominate the PNSD at various distances from the runway of Tianjin International Airport, China; Masiol et al. (2016) reported that the fingerprint of aircraft emissions sampled under real ambient conditions at the airport of Venice (Italy) has a main mode at approx. 80 nm and a second mode in the nucleation range below 14 nm.

The Greater London area is home to more than 8.5 million inhabitants and is one of the few UK locations not fully achieving the EU and national air quality standards: in 2015 nitrogen dioxide breached the hourly and annual limit values for health, while ozone exceeded the long-term objective (DEFRA, 2016). However, the standards were fully met for both PM_{10} and $PM_{2.5}$.

London Heathrow (LHR) is one of the world's busiest international airports: it is ranked 1st in Europe for total passenger traffic (ACI, 2016). It accommodates more than 1250 flights every day and serves a total of 72.3 million passengers $year^{-1}$. LHR is composed of 5 terminals and 2 runways: northern (3.9 km-long) and southern (3.7 km). Currently, runways operate near their maximum capacity, with a consequent increase in the potential for delays when flights are disrupted. Since 2007, the proposal for expanding LHR with a 3rd runway and a 6th terminal has been intensely debated in the UK. In 2016 the UK government provisionally approved the construction of a third runway (UK Department for Transport, 2017).

LHR is located west of London (Figure SI1). Consequently, air quality in the surroundings of the airport may be affected by the advection of air masses from the city, with the associated high levels of pollutants emitted from traffic, energy demand for domestic heating and local industries. Airport activities may also contribute to air pollution advected to the city when LHR is upwind, with consequent potential impacts upon public health. In addition, as LHR attracts a large number of passengers and workers, the emissions from large volumes of road traffic generated by the airport and the nearby M4 and M25 motorways are difficult to discriminate from non-airport-related road traffic. Due to this complex scenario, the contribution of LHR is difficult to differentiate from the urban background pollution, as already reported by previous modelling and experimental studies (Farias and ApSimon, 2006; Masiol and Harrison, 2015).

Various studies have attempted to quantify the effect of LHR upon air quality, mainly focusing on the nitrogen oxides (NO_x=NO+NO_2), which are well-known tracers for aircraft engine exhausts (e.g., Herndon et al., 2008; Masiol and Harrison, 2014 and references therein), but also arise from other combustion sources. For example, Carslaw et al. (2006) estimated that airport operations in 2001/4 accounted for ~27% of the annual mean NO_x and NO_2 at the airfield boundary and less than 15% (<10 μg m^{-3}) at background locations 2-3 km downwind of the airport. Similar results were found for the 2008/9 period using model evaluation (AEA, 2010) and for the 2005/12 period using experimental data analysis (Masiol and Harrison, 2015). This latter study also reported that PM mass concentrations at eight sites all around LHR were always well below the EU and UK limit.

This study aims to investigate the impacts of a major airport (LHR) serving a megacity (London) upon the levels of submicrometre particles and to apportion those impacts to aircraft, road traffic and other sources typical of large cities with airports. The main particle size distributions modes are simplified by applying cluster analysis; then, the modal structures of the main potential sources are disaggregated and the submicron particle number concentrations (PNC) are quantified through the positive matrix factorisation (PMF). In addition, the origin of the airport plumes was spatially assessed by matching results with local meteorological data, air mass movements, levels of common air pollutants, $PM_{2.5}$ mass concentration and its chemical speciation as indicators of source location and formation mechanisms.

The atmospheric chemistry and physical properties of UFPs have been extensively investigated in London (e.g., Harrison et al., 2012; Jones et al., 2012; von Bismarck-Osten et al., 2013) with several studies using cluster analysis (Beddows et al., 2009; Brines et al., 2014; 2015) or PMF (Beddows et al., 2015; Vu et al., 2016). However, this study is the first one carried out in South-West London to characterise and quantitatively apportion the impacts of LHR under real ambient conditions. Moreover, only one earlier study (Masiol et al., 2016) has used both cluster analysis and

PMF to directly assess the airport contributions to UFPs. In addition, this study also investigated the effects of a regional nucleation event on the results of the two source apportionment methods.

2. MATERIALS AND METHODS

2.1 Experimental

Two sampling campaigns (each 1 month-long) were carried out during warm (August-September 2014) and cold (December 2014-January 2015) periods at Harlington (Figure SI1). The site was selected as well located to sample the plumes from the airport emissions: it lies 1.2 km N of the northern runway and is located inside a playground, close to a secondary road and near the village of Harlington. This is the location selected for the construction of the 3rd runway. The site is categorised as "urban industrial" by DEFRA and it is therefore more indicative of community exposure rather than direct fresh aircraft emissions. Consequently, it is a good point to quantify the particles generated by the airport after a relatively short ageing and dispersion in the atmosphere, and is more indicative of the fingerprint of aircraft emissions affecting communities than data collected alongside the runway or in the airport apron areas. In addition, previous studies have reported that the site is strongly affected by the plume from the airport (Carslaw et al., 2006; Masiol and Harrison, 2015). Prevailing winds from the 3rd and 4th quadrants are recorded in both summer and winter (Figure SI2): under such circulation regimes, Harlington lies just downwind of LHR. The site is also affected by pollutants arising from the large volumes of road traffic within London, from the local road network as well as those generated by the airport. Tunnel Rd., the main access to LHR from the M4 motorway lies 800 m west, as well as the nearby M4 (640 m north) and M25 (~3.5 km east) motorways, major roads (Bath Rd, part of A4, passes 900 m south; A30 lies 2.8 km SE). The village of Harlington (~400 m west) and advection of air masses from the conurbation of London are other potential external sources.

Ultrafine particle counts and their size distributions from 14.3 to 673.2 nm were measured at 5 min time resolution using a SMPS (scanning mobility particle sizer spectrometer) comprising an electrostatic classifier TSI 3080 with a long differential mobility analyser (TSI 3081) and a CPC (condensation particle counter, TSI 3775) based on condensation of *n*-butyl alcohol (Fisher Scientific, ACS). The SMPS operated at a sheath air to aerosol flow ratio of 10:1 (sheath and sample air flow rates were 3.0 and 0.3 L min^{-1} respectively, voltage 10-9591 V; density 1.2 g/cc; scan time 120 s, retrace 15 s; number of scan 2) while the CPC operated at low flow rate (0.3 L min^{-1}). The use of 5 min resolved spectra has already been used successfully for source apportionment purposes at an airport (Masiol et al., 2016).

Equivalent Black Carbon (eBC) as defined by Petzold et al. (2013) was also measured at 5 min resolution using a 7-wavelength aethalometer (Magee Scientific AE31). The aethalometer operated with an inlet cut-off head to collect PM with aerodynamic diameter of <2.5 μm ($PM_{2.5}$). eBC was derived from the absorbance at 880 nm wavelength (Petzold et al., 2013); raw data were post-processed with the Washington University Air Quality Lab AethDataMasher V7.1 to perform data validation and correct data for non-linear loading effects (Virkkula et al., 2007; Turner et al., 2007).

Instruments were installed into a plastic/metal case designed for sampling purposes: (i) air inlets were ~1.8 m above the ground and were composed of conductive materials to avoid particle losses and sampling artefacts; (ii) the case was cooled by fans in summer and was warmed by an electrical tubular heater in winter for maintaining an indoor air temperature within an acceptable range for running the equipment (temperature inside the case was recorded and periodically checked); (iii) instruments were isolated from vibration using rubber pads and foam foils. Devices were fully serviced, calibrated by authorised companies and underwent internal cross-calibrations with other similar instruments under lab conditions. Moreover, frequent periodic checks, maintenance of instruments and cleaning of inlets was performed throughout the sampling campaign.

Routine air pollutants (NO, NO_2, NO_x, O_3, PM_{10}, $PM_{2.5}$) were measured at Harlington with 1 h time resolution by the UK Automatic Urban and Rural Network under the auspices of the UK Department for Environment, Food and Rural Affairs (DEFRA; http://uk-air.defra.gov.uk/). Gaseous species were analysed using automatic instruments according to European standards and National protocols: EN 14211:2012 for nitrogen oxides and EN 14625:2012 for ozone. PM_{10} and $PM_{2.5}$ were analysed using tapered element oscillating microbalance and filter dynamics measurement system (TEOM-FDMS) to provide measurements accounting for volatile (VPM_{10}, $VPM_{2.5}$) and non-volatile ($NVPM_{10}$, $NVPM_{2.5}$) fractions. Quality assurance and quality control procedures followed the standards applied for the Automatic Urban and Rural Network (AURN) and the London Air Quality Network (LAQN). Instruments were routinely calibrated, and every six months were fully serviced and underwent intercalibration audits.

Some additional variables are also computed from the air pollutants to help the interpretation of results. The NO_2/NO_x ratio is indicative of the partitioning of nitrogen oxides, while the levels of oxidants ($OX=O_3+NO_2$, expressed in ppbv) can be used to roughly assess the oxidative potential in the atmosphere (Kley et al., 1999; Clapp and Jenkin, 2001). These two new variables are useful in investigating the atmospheric chemistry behind the $NO\text{-}NO_2\text{-}O_3$ system. Delta-C (the difference between absorbance at 378 and 880 nm, also called UVPM) was also computed. This variable was largely used as a proxy to estimate the fraction of carbonaceous material emitted by biomass burning (e.g., Sandradewi et al., 2008; Wang et al., 2011). However, Delta-C results should be used with caution: Harrison et al. (2013) showed that there are probably other UV absorbing contributors than wood-smoke to the aethalometer signal. Consequently, Delta-C is used here only for qualitative purposes.

Weather data were measured hourly by the Met Office at LHR; met data include wind direction and speed, atmospheric pressure, air temperature, relative humidity (RH), visibility, rain and solar irradiance.

During the two campaigns, 24-h $PM_{2.5}$ samples were also collected on quartz filters using a high volume air sampler (TE-6070, Tisch Environmental, Inc.) and analysed for the daily concentrations of major $PM_{2.5}$ components: organic carbon (OC) and elemental carbon (EC) by thermo-optical analysis (EUSAAR_2 protocol) and major inorganic ions (Na^+, K^+, ammonium, nitrate, sulphate, oxalate) by ion chromatography. Analytical methods are reported in detail in Yin et al. (2010). The results of the chemical speciation of $PM_{2.5}$ are presented in a companion paper (in preparation) and are used in this study only to assist the interpretation of PMF results.

2.2 Data Handling and Chemometric Approaches

Data were analysed using R version 3.3.1 (R Core Team, 2015) and a series of supplementary packages, including 'Openair' (Carslaw and Ropkins, 2012). Preliminary data handling and clean-up were carried out to check the robustness of the dataset, detect anomalous records and to delete extreme outliers. SMPS data with unreliable behaviour or instrument errors were completely deleted. An in-depth analysis of the dataset revealed few records with anomalously high PNC, which were likely related to probable instrumental issues, extreme weather conditions (e.g., high wind gusts, heavy rain striking the inlet), or infrequent local emissions, e.g., maintenance, painting and recreational activities (including fires) on the playground where the site is located, road maintenance close the site and probable short-term parking of high-emission vehicles near the site. Since this study aims to investigate the overall contributions of LHR, all data are used for descriptive statistics, but data greater than the 99.5th percentile were further removed for explorative, cluster and PMF analyses. This data exclusion successfully removed the extremely high events occurring during the sampling campaigns and significantly improved the stability and

physical meaning of PMF solutions. Missing data for other variables were linearly interpolated between the nearest values of the time series.

The particle number size distributions (PNSDs) were firstly grouped by applying a *k*-means cluster analysis. The full method is exhaustively discussed in Beddows et al. (2009; 2014) and aims to assemble single spectra into *k* clusters. The clustering groups observations with spectra similar to their cluster centroids (means), i.e. observations that are likely generated by the same set of formation processes or emission sources. The optimum number of clusters (*k*) was determined by an optimisation algorithm based on the spectral shapes (Beddows et al., 2009). The choice to apply the *k*-mean clustering method was based on several reasons: (i) Salimi et al. (2014) reported that *k*-means is the best performing clustering among others methods tested on PNSD data; (ii) *k*-means is a well-established method which has been widely applied over a number of different sites (e.g., Dall'Osto et al., 2012; Wegner et al., 2012; Beddows et al., 2014; Brines et al., 2014; 2015); and (iii) the method was previously applied successfully to airport data (Masiol et al., 2016).
PMF analysis was performed by applying the USEPA PMF5 model. Details of the PMF model are reported elsewhere (Paatero and Tapper, 1994; Paatero, 1997; USEPA, 2014), while the best practice and standards are extensively reviewed in several papers (e.g., Reff et al., 2007; Belis et al., 2014; Brown et al., 2015; Hopke, 2016). SMPS data at 5 min resolution were used as the PMF input matrix. Uncertainties associated with SMPS data were estimated according to the empirical method proposed by Ogulei et al. (2007). Uncertainty for the total variable (total particle number concentration, PNC) was set at 300% of the PNC concentration and also marked as "weak" to avoid it driving the profiles.

The best PMF solutions were identified: (i) by investigating solutions between 3 and 10 factors; (ii) by considering the minimization of the objective function *Q* with respect to the expected (theoretical) value and its stability over multiple (n=100) runs, (iii) by obtaining low values for the

sum of the squares of the differences in scaled residuals for each base run pair by species; (iv) by minimizing the number of absolute scaled residuals over ±3 and by keeping them symmetrically distributed; (v) by keeping the result uncertainties calculated by bootstrap (BS, n=200) and displacement (DISP) methods within an acceptable range (Paatero et al., 2014); (vi) by obtaining modelled total variable (PNC) successfully predicted ($R^2 > 0.9$ and slopes ≈1); and (vii) by avoiding the presence of edges in the G-space plots (Paatero et al., 2002) and, then, the presence of hidden/unresolved sources.

A series of additional tools were used to analyse the raw data, link source apportionment results to other variables, such as local atmospheric circulation and regional/transboundary transport of air masses. Briefly, polar plots aim to map pollutant average concentrations by wind speed and direction as continuous surfaces (Carslaw et al., 2006), while polar annuli plot by wind direction and hours of the day. The potential locations of distant sources were assessed using back-trajectory analysis and a concentration weighted trajectory (CWT) model (Stohl, 1998). Back-trajectories were computed with the HYSPLIT4 model (Stein et al., 2015; Rolph, 2016) using NCEP/NCAR reanalysis gridded meteorological data. Set-up: -96 h with a starting height of 500 m a.g.l. CWT is a method of weighting trajectories with associated concentrations to detect the most probable source areas of long-range transports of pollutants; it has been used and reviewed in a number of prior studies (e.g., Stohl, 1996; Lupu and Maenhaut, 2002; Squizzato and Masiol, 2015).

3. RESULTS AND DISCUSSION

3.1 Overview of Data

The wind roses during the two sampling periods are provided in Figure SI2. Descriptive statistics of all collected variables are reported as boxplots in Figure SI3. PNSDs were initially split into 3 ranges: nucleation (14-30 nm), Aitken nuclei (30-100 nm) and accumulation (>100 nm). On average the total PNC during the warm season was 1.9 x10^4 particles cm^{-3}, of which 1.1 x 10^4, 6.4 x 10^3 and 1.5 x 10^3 particles cm^{-3} were classified as nucleation, Aitken and accumulation ranges, respectively (Figure SI3). During the cold season, the total average PNC was 2.2 x 10^4 particles cm^{-3}, composed of 1.4 x 10^4, 6.3 x 10^3 and 1.4 x 10^3 particles cm^{-3} as nucleation, Aitken and accumulation ranges, respectively (Figure SI3). Concentrations lie between those of London, Marylebone Road (kerbside) and London, North Kensington (background), and nucleation particles were ~10 times higher than the annual average measured in North Kensington as reported by Vu et al. (2016), while Aitken particles were 1.9 times higher. It is therefore evident that the main difference lies in the concentration of the finest size ranges: in both seasons, spectra were dominated by UFP (D_p<100 nm) particles (~92% of total PNC), which only accounted for ~12% of total particle volume concentration (PVC, computed by approximation to spherical particles). On the other hand, accumulation mode particles accounted for ~8% of PNC and ~88% of PVC volume. The high levels of total PNC are not surprising: several studies carried out into or close to airports (e.g., Hsu et al., 2013;2014; Hidda et al., 2014; 2016; Stafoggia et al., 2016; Shirmohammadi et al., 2017) reported significant increases in the concentrations of UFPs.

During the two sampling campaigns, air pollutants measured in Harlington (Figure SI3) were similar to the average concentrations measured over an 8 year period (2005-2012) in the vicinity of LHR (Masiol and Harrison, 2015). Consequently, despite the two short campaigns carried out in this study, results may be considered representative of the average levels of air pollution recorded at Harlington. The average concentrations of eBC were 2.4 and 2.1 μg m^{-3} during the warm and cold season, respectively. The average concentration of Delta-C was 0.1 μg m^{-3} during the warm season and 0.36 μg m^{-3} in winter.

Analysis of the data showed a non-normal distribution for most of the variables: the nonparametric Kruskal-Wallis one-way analysis of variance was therefore used to test the difference of concentrations over the two periods (Kruskal and Wallis, 1952): almost all variables are different at the 0.05 significance level, except NO, NO_x and O_3. This result indicates a seasonal effect upon air quality in the LHR area and suggests investigating the sources over the two periods separately.

The average PNSDs are shown in Figure 1 as well as their median distributions and interquartile ranges. Spectra are categorised by time of day (7am-7pm and 7pm- 7am local time). In addition, the particle volume size distributions (PVSDs) are also provided. Results for the warm season show that the average daytime PNSD is dominated by a main peak in the nucleation range (extending below 14 nm) and a second mode in the Aitken range (between 30 and 50 nm). The nocturnal spectrum is characterised by a drop of the nucleation mode to concentration values similar to the Aitken peak (mode around 35 nm). During the cold season, the average diurnal and nocturnal PNSDs present a main peak at 15-25 nm and a second mode at 70-100 nm. In summary, both seasons show reductions of the finest modes during nighttime, while the second mode is almost constant throughout the day. As a consequence, the modal structure of PNVDs is also almost constant throughout the day.

The diurnal cycles of the 3 particle ranges, eBC, solar irradiation and airport movements are shown in Figure 2. A comprehensive overview of the patterns for all the variables is provided in Figure SI4. Generally, diurnal cycles derive from the interplay of emissions, dispersion and atmospheric chemical processes. Consequently, they need to be investigated along with patterns for airport and motorway traffic (Figure 2 and Figure SI5, respectively), and as polar annuli (Figures SI6 and SI7) and polar plots (Figures SI8 and SI9), which give preliminary insights into the origin and spatial location of most probable emission sources. During nighttime, airport traffic is restricted to limit

noise and community disturbance: flights are generally constant from 6 am to 8 pm and are kept at minimum overnight, with no departures normally scheduled between 11 pm and 6 am (Figure 2). Road traffic is more difficult to define. Data for M4 and M25 motorways are provided by the UK Department for Transport: data for the M4 motorway show typical morning (7-8 am) and evening (5-6 pm) peaks due to rush hours, but this pattern is not well-resolved for the M25 (Figure SI5). In addition, despite it being likely that traffic on minor and local roads also follows patterns dominated by rush hours, traffic generated by the airport is more difficult to characterise, with Tunnel Rd. and other busy roads serving LHR being frequently congested.

Nucleation particles are likely associated with aircraft movements. The daily pattern shows high and almost constant concentrations between 7 am and 11 pm (Figure 2): hourly averages ranged from 10 x 10^3 to 15 x 10^3 particles cm^{-3} during the warm season and from 10 x 10^3 to 21 x 10^3 particles cm^{-3} during the cold season. On the contrary, the concentrations of nucleation particles significantly (Kruskal-Wallis at $p<0.05$) drop overnight (hourly averages ranging from 5 x 10^3 to $6 \cdot 10^3$ particles cm^{-3} and from 1 x 10^3 to $5 \cdot 10^3$ particles cm^{-3} during the warm and cold season, respectively); the maximum average concentrations are recorded for winds blowing from the SW quadrant (polar plots and polar annuli in Figures SI6-9), i.e. the airfield and, in particular, the location of the main LHR terminals (Figure SI1). As a consequence of the dominance of nucleation particles over size spectra, also total PNC follows the pattern (Figures 2) and wind directionality (Figures SI8-9) of nucleation particles. On the contrary, accumulation particles appear to be more associated with road traffic. These particles increase for winds blowing from northern sectors (Figures SI6-9), i.e. toward the M4. Accumulation particles also present the morning (6-8 am) and evening (6-11 pm) rush hour peaks during the warm season, but only the evening peak (from 6 pm to the night) was found in the cold season (Figure 2). Generally, the evening peaks start around 6 pm, which is consistent with the peak of traffic (Figure SI5) but they extend late in the evening and night probably because the drop of the mixing layer top and the consequent concentration of

pollutants close to the ground level. Aitken nuclei exhibit a mixed behaviour between nucleation and accumulation particles (Figure 2): two different patterns can be found, which are more consistent with road traffic in summer and with aircraft traffic in winter.

Despite some studies indicating that airports are strong sources of black carbon (Dodson et al., 2009), other studies report no strong relationships with the flight activity (Masiol et al., 2016; Hsu et al.,2016). Similarly to NO_2 (Figure SI4) and accumulation particles (Figure 2), aethalometer data also shows typical patterns of road traffic-influenced sites for all wavelengths, with two daily peaks corresponding to the hours with higher traffic (Figure 2). However, Delta-C does not present any evident pattern (Figure SI4). eBC shows increased concentrations when winds blow from northern sectors (plus SE in winter, Figure SI7 and SI9); which excludes airport activities as being a dominant source in the study area.

Particulate matter mass concentration (PM_{10} and $PM_{2.5}$) has very weak diurnal patterns (Figure SI4). Its wind directionality shows evident increases for northerly winds (Figure SI8-9). It is therefore evident that PM mass concentrations are dominated by non-airport sources, i.e. regional secondary pollutants, traffic from the nearby M4 or background pollution from London. $PM_{2.5}$ concentrations normally do not exceed the Limit Values in the Greater London area (DEFRA, 2016).

3.2 *k*-means Cluster Analysis

The clustering algorithm extracted 5 clusters for both periods. The number of clusters was selected according to the optimisation algorithm, i.e. local maxima in the Dunn indices and silhouette (Beddows et al., 2009). The extraction of 5 clusters represents a good compromise for the interpretation of spectral observations. Hussein et al. (2014) reported that is not prudent to describe the spectra with few clusters (2-4), which are not sufficient to explain variations and detailed

differences in the PNSD observed in the urban atmosphere. On the other hand, they also reported that extracting too many (>10) clusters may make the aerosol source attribution more challenging.

The cluster centroids (mean spectra of each cluster), the 10th, 25th, 75th and 90th percentile, the hourly counts patterns and resulting wind roses are shown in Figures 3 and 4 for the warm and cold season campaigns, respectively. Despite extracted clusters exhibiting significantly different modal structures for PNC, no differences can be observed for the particle volume size spectra, which all show a unimodal peak at approx. 200-300 nm. Clusters accounted for 14%-25% of total observations for both the seasons: Table SI1 summarises the percentage of the total observations for each cluster.

Three clusters (*cluster 1* during the warm season and *clusters 1 and 5* in winter) are likely shaped by the airport emissions. The modal structures present sharp peaks for nucleation particles which extend below the SMPS detection limit (14 nm) and drop at 30-40 nm; no secondary modes are present in the Aitken or accumulation ranges. These clusters show a large increase in frequency during the afternoon and evening hours (cluster 1 for the warm season and cluster 5 for the cold season) or extended over the daytime (cluster 1 for the cold season), similarly to the airport aircraft movement profiles (Figure 2). Aircraft are known to emit particles in the nucleation range (e.g. Mazaheri et al., 2009;2013; Masiol and Harrison, 2014; and references therein; Lobo et al., 2015) and the wind roses are also compatible with an origin from the airfield and the main LHR terminals (Figures 3 and 4). However, daytime regional photochemical nucleation events in London occur around noon-2 pm and are mostly recorded from June to September (Vu et al., 2016). Consequently, the modal structure of cluster 1 for the warm season could be additionally shaped by regional photochemical nucleation. The reasons driving the split of the spectra likely shaped by LHR into two clusters during the cold season are unclear. A further comparison of the cluster and PMF results will help in interpreting this outcome.

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442 The modal structures of the *clusters 4* for both seasons peak for nucleation particles and extend
443 below 14 nm, but also show probable modes between 50 and 200 nm (Figures 3 and 4). They
444 represent the typical spectra associated with aged anthropogenic emissions, mostly due to road
445 traffic. It is recognised that road traffic contributes to a large range (30-200 nm) of PNSD in the
446 urban atmosphere (e.g., Yue et al., 2008; Costabile et al., 2009; Harrison et al., 2011), which is
447 compatible with these spectra. The directional analysis for the warm season shows increased levels
448 when air masses move from the sectors more affected by traffic, i.e. London (NE), M4 (N) and M25
449 (W) motorways and Tunnel Rd (W), while the hour count profile presents a huge maximum during
450 daytime. In winter, this modal structure mostly occurred for westerly winds: the atmospheric
451 circulation during the cold season mostly experienced winds blowing from the SW quadrant, with
452 NE sectors poorly represented (Figure SI1). As a consequence, the limited number of observations
453 for air pollution advected from the Greater London area may have affected the detection of the
454 urban background from London. This lack of data is also reflected by diurnal profile, which shows a
455 marked peak in the late afternoon, concurrent to the peak of traffic on M4 and M25 (Figure SI5).
456

457 Three clusters (*clusters 2* and *3* during the warm season and *cluster 2* in winter) exhibited similar
458 hourly profiles with most of the counts occurring overnight (Figures 3 and 4). This pattern is largely
459 attributable to the dynamics of the mixing layer, since the diurnal cycles are the mirror image of the
460 ambient air temperature (Figure SI4). Because of this, these clusters could be potentially affected by
461 the reduced height of the mixing layer occurring overnight. These clusters exhibit bimodal
462 structures with the coarser modes with respect to the remaining clusters: cluster 2 for the warm
463 season shows a main peak in number concentrations at 30-40 nm and a second peak in the finest
464 range (<16 nm), clusters 3 for the warm season peaks at 14 and 60-70 nm, and cluster 2 for the cold
465 season extends over a wide size range with two modes around 20-30 nm and 100-150 nm.
466 Consequently, these clusters are likely representative of spectra mostly shaped by the drop of the

mixing layer height and the formation of secondary aerosols. In this context, the potential role of nighttime nitrate formation through condensation of NH_4NO_3 and the heterogeneous reactions of N_2O_5 and NO_3 on pre-existing particles cannot be ignored (Seinfeld and Pandis, 2006; Bertram and Thornton, 2009; Brown and Stutz, 2012). The wind roses reveal that both clusters 2 occur under similar westerly wind regimes. Regional aerosols appear to be the most probable source. On the contrary, cluster 3 for the warm season occurs with winds from London (NE) and likely represents particle size spectra mainly shaped by primary and secondary aerosols advected from the most urbanised areas, i.e. it is most likely associated with the urban background of London.

Cluster 5 for the warm season and *cluster 3* for the cold season may be associated with road traffic. They reveal modal structures with a dominant peak around 20-35 nm (cluster 5 also shows a possible second peak at 15 nm) and mostly occur when air masses blow from westerly sectors, which are compatible with the location of motorways and Tunnel Rd, the main roadway linking LHR to the M4 motorway. In summer, the hourly count pattern exhibits two maxima (6-8 am and 4-8 pm) related to morning and evening rush hours; this pattern is compatible with fresh road traffic emissions. However, the diurnal pattern in winter also presents a high number of counts at 3-5 am, i.e. not directly compatible with rush hours. A possible explanation involves the stronger effect of the winter mixing layer dynamics on the air quality due to the presence of more frequent low level thermal inversions, which may build up the pollutants at ground-level especially overnight. This may increase the signal of the less intense, but still significant, nighttime traffic emissions present in the study area.

3.3 PMF Analysis

The interpretation of PMF results was then attempted by considering: (i) the knowledge of sources impacting the study area; (ii) the comparison with the results reported by Vu et al. (2016), who performed a PMF analysis of SMPS data collected in North Kensington (London urban background); (iii) the shape of resulting profiles for both the particle number and volume concentrations; (iv) the analysis of diurnal patterns; (v) the directional analysis using the polar plot and polar annuli; (vi) the correlations between the source contributions and the other air pollutants monitored at the site or with weather variables, and (vii) the analysis of possible remote source areas by applying the CWT model.

Six-factor solutions were extracted for both the seasons. The resulting factor profiles are presented in Figures 5 and 6 for the warm and cold season, respectively. The factor profiles are expressed as: (i) particle number concentrations and their DISP ranges; (ii) particle volume concentrations, and (iii) explained variations showing how much of the variance (from 0 to 1) in the original dataset is accounted for by each extracted factor. The Figures 5 and 6 also show the diurnal patterns and the polar plots computed from the hourly-averaged contributions. Table 1 summarises the PMF results and spectral characteristics, while Table 2 shows the Pearson correlation matrices with weather and air quality variables. Selected PMF solutions were very stable: no errors or unmapped factors and few swaps (none in summer and <7% in winter) were found in BS; no swaps or errors even at dQ_{max}=25 were found for DISP, i.e. solutions were affected by small rotational ambiguity and, therefore, their interpretation can be considered robust.

DISP analysis is designed to explore the realistic bounds on the optimal (base run) PMF solutions that do not result in appreciable increases in the *Q* values (Brown et al., 2015). In this study, the ranges calculated by DISP for the dQ=4 were used to assess the uncertainty boundaries associated with the final PMF profiles, as suggested in Zikova et al. (2016) and Masiol et al. (2017). This

strategy is useful to better interpret the results, as the regions of spectra affected by high rotational ambiguity are disclosed in the resulting profiles.

3.3.1 Warm season

Factor 1 includes most of the particles in the nucleation range (<20 nm), exhibits a sharp mode in the number distribution below the SMPS detection limit (14 nm) and makes the largest contribution to the total PNC (31.6%, DISP range 31-36%) (Figure 5). However, its contribution to the volume distribution is ~1%. Several studies report that particles in the nucleation range are emitted from aircraft engines (e.g., Anderson et al., 2005; Herndon et al., 2008; Kinsey et al., 2010; Mazaheri et al., 2009;2013; Masiol and Harrison, 2014; Lobo et al., 2015) as well as from other anthropogenic (e.g., Schneider et al., 2005; Chen et al., 2011; Cheung et al., 2012; Stevens et al., 2012; Kumar et al., 2013;2014; Vu et al., 2015b) and natural (e.g., Kulmala et al., 1998; O'Dowd et al., 1998;1999; Kulmala and Kerminen, 2008; Riccobono et al., 2014) sources. This factor does not show any significant ($p < 0.05$) and strong ($r \geq |0.6|$) correlation with other measured species, but shows a weak ($|0.4| \leq r < |0.6|$) correlation with Factor 2 (Table 2). Its diurnal variation (Figure 5) shows higher concentrations between 6 am and 10 pm, and well agrees with the airport flight movements (Figure 2). The polar plot analysis also indicates enhanced levels when winds > 2 m s^{-1} blow from the airfield sectors (SW). All these insights are consistent with the location of Heathrow, i.e. the most plausible interpretation is related to the aircraft engine exhaust emissions. This interpretation is also supported by Keuken et al. (2015), which shows that the PNSD in an area affected by emissions from Schiphol airport (The Netherlands) is dominated by ultrafine (10-20 nm) particles. The large contribution of this factor to the total PNC is not surprising if compared to the results reported for the Los Angeles international airport by Hudda et al. (2014) (emissions from the airport increased PNC 4- to 5-fold at 8–10 km downwind the airfield). Since the airport of Los Angeles and LHR have comparable aircraft traffic, the quite high concentrations found in this study (on annual average nucleation particles are ~10 times higher than those measured in North Kensington urban

543 background by Vu et al. (2016)) are consistent with the sampling location chosen in this study (~1.2
544 km to the airfield). In addition, this result also agrees with previous studies on the impacts of LHR
545 on local air quality; Carslaw et al. (2006) and Masiol and Harrison (2015) found comparable
546 percent contributions of LHR emissions on NO_2 levels in the study area (approx. 25-30%).
547 However, the lack of correlations with NO and NO_2 (tracers for aircraft emissions) is probably due
548 to the presence of several other sources of nitrogen oxides in the area, such as the heavy traffic
549 generated from the airport and from the nearby motorways.
550
551 *Factor 2* is made up of ultrafine particles in the nucleation-Aitken range (one main peak at 20-35
552 nm) and accounts for 28% (DISP 25-30%) of PNC; its contribution to the volume distribution is
553 low (~2%) and peaks at 22-45 nm and at 140-220 nm (Figure 5; Table 1). Several insights seem to
554 link this factor to road traffic emissions: (i) the modal structure; (ii) the strong association with
555 morning and evening rush hours, and (iii) the significant increase for winds in the west and south-
556 westerly sectors consistent with emissions generated from local busy roads close to LHR, Tunnel
557 Rd. and M25 motorway. A similar mode in the nucleation range has been extensively attributed to
558 the size distribution from road traffic (e.g., Vogt et al., 2003; Zhang et al., 2004; Ntziachristos et al.,
559 2007; Vu et al., 2015b) and the growth of nucleation particles from diesel vehicles (Mayer and
560 Ristovski, 2007; Wehner et al., 2009). For example, Charron and Harrison (2003) reported that
561 particles in the range 30–60 nm show a stronger association with light-duty traffic at a traffic
562 hotspot in central London (Marylebone Rd.); Janhäll et al. (2004) reported an average particle size
563 distribution peaking at 15-30 nm during morning peak high traffic intensity in the city of Göteborg
564 (Sweden), which has a car fleet comparable to the UK; Ntziachristos et al. (2007) found a sharp
565 mode at 20-30 nm in sampling from engine exhausts. In addition, PMF factors with similar modal
566 structures were found in other studies and were attributed to road traffic emissions: among others,
567 Harrison et al. (2011) linked a factor peaking at 20 nm to primary road traffic emissions near a
568 major UK highway; Masiol et al. (2016) measured PNSD in an international airport in Northern

Italy during summer and interpreted a factor with a clear mode at 35-40 nm as road traffic from the nearby city; Beddows et al. (2015) and Vu et al. (2016) found traffic factors with modal diameter at around 30 nm in an urban background site in London (North Kensington); Sowlat et al. (2016) reported a factor peaking at 20–40 nm in number concentration and at around 30–40 nm in volume concentration in Los Angeles (US) and interpreted it as traffic tailpipe emissions. However, this factor lacks significant positive correlations with primary road traffic tracers (nitrogen oxides, eBC; Table 2), while other studies have reported weak positive correlations with such species (Harrison et al., 2011; Masiol et al., 2016; Vu et al., 2016; Sowlat et al., 2016). Similarly to factor 1, this latter result may be due to the difference in the time resolution between chemical species and PNSD and the presence of several sources of nitrogen oxides in the area.

Factor 3 is mostly represented by 25–90 nm particles and contributes about 19% (17-21%) to the total number concentration (Figure 5; Table 1). It also shows a second mode below the SMPS detection limit (14 nm), however, the DISP range clearly indicates that this part of the profile is affected by a large amount of rotational ambiguity, so that the presence of this second mode should be interpreted with caution. The volume concentration peaks at around 40–100 nm and 250–450 nm. The factor contribution is higher during rush hours, but the morning peak occurs 1 h later than in factor 2. The wind directionality shows increases for air masses blowing gently (<4 m s^{-1}) from W and for calm wind periods, suggesting a quite local source; however, also an increase of concentrations is found for higher wind regimes (>6 m s^{-1}) from the East (London). Factor 3 also shows significant positive correlations with NO (0.43) and NO_2 (0.61) (Table 2). All these insights seem to point to an aged road traffic source. This interpretation is also supported by Vu et al. (2016), who found a similar factor in London (North Kensington) peaking at ~20–100 nm. In this context, several source apportionment studies on PNSDs have attributed more than one factor to road traffic (e.g. Kasumba et al., 2009; Thimmaiah et al., 2009; Harrison et al., 2011; Liu et al., 2014; Al-Dabbous and Kumar, 2015; Vu et al., 2016; Sowlat et al., 2016). This result is not

surprising in areas where heavy traffic is widespread, as particles may undergo condensation, agglomeration, evaporation and dilution processes and, consequently, they may change modal characteristics in time and space. Such atmospheric processes are the main mechanisms reshaping PNSDs after primary exhaust is emitted into the atmosphere and have been discussed in several studies (Shi et al., 1999; Kim et al., 2004; Zhang et al., 2005; Zhou et al., 2005; Kulmala and Kerminen, 2008; Zhang et al., 2011; Harrison et al., 2016).

Factor 4 is made up of particles over a wide range (50-200 nm with a clear mode at ~80 nm for PNC and 60-300 nm for PVC). The factor contributes 14% of PNC, but accounts for the main percentage of the volume concentration (33%). This factor correlates well with gaseous pollutants linked to combustion sources (mostly road traffic), i.e. NO (0.6), NO_2 (0.76), and non-volatile primary pollutants, such as eBC (0.62), $NVPM_{2.5}$ (0.62) and EC (0.75) (Table 2). The factor also strongly correlates with OC (0.84) and sulphate (0.75). The diurnal pattern shows two main peaks in the morning and evening rush hours (Figure 5), but the concentrations recorded between the two maxima are higher overnight than during daytime. This pattern suggests that both local emission sources and the dynamics of the mixing layer may play a key role in shaping its diurnal cycle, i.e. emitted pollutants undergo a wide dispersion within the expanded mixing layer during the daytime, while the drop of the mixing layer top occurring overnight restricts those pollutants to a layer close to ground level. The polar plot indicates increased levels for calm wind conditions or winds blowing from London (East sectors); in addition, the factor is strongly negatively correlated with wind speed (-0.64) (Table 2).

All these insights suggest that Factor 4 represents the fingerprint of the London pollution. Several studies carried out in London (Beddows et al., 2009;2015; Vu et al., 2016) and other megacities (e.g., New York: Masiol et al., 2017) have reported similar results, all interpreting this source profile as urban background (or urban accumulation mode). This source comprises both the solid

particle mode from traffic emissions (Harrison et al., 2011; Pant and Harrison, 2013; Dall'Osto et al., 2012) and secondary species condensed upon pre-existing particles acting as condensation nuclei, including secondary sulphate, nitrate and organic aerosols. Secondary sulphate is formed through the atmospheric processing of local or distant SO_2 emissions (Kerminen et al., 2000) and neutralisation with ammonia (Benson et al., 2011). Nitrate aerosol is formed through the oxidation of NO_2 to nitrate and the consequent neutralization with ammonia (Seinfeld and Pandis, 2006) and occurs during both daytime and night-time; however the semivolatile nature of ammonium nitrate, makes its partitioning to the condensed-phase very weak. This behaviour also favours the occurrence of negative artefacts in filter-based sampling, which may explain the lack of significant correlations between the factor and the $PM_{2.5}$-bound nitrate (Table 2). On the contrary, the increase of the intensity of factor 4 during the night-time and the significant association with NO_2 are highly consistent with the chemistry driving the heterogeneous reactions of N_2O_5 and NO_3 on aerosol surfaces (Bertram and Thornton, 2009; Brown and Stutz, 2012). In view of this, Dall'Osto et al. (2009) reported that most nitrate particles in London are: (i) locally produced in urban locations during nighttime; (ii) mainly present in particles smaller than 300 nm and (iii) internally mixed with sulphate, ammonium, EC and OC.

Factors 5 and 6 make small contributions to PNC (4-7% and 1-4%, respectively), but are relevant for the volume concentration (37% and 21%, respectively). Factor 5 shows a main accumulation mode in number concentration at 110-250 nm and two more modes at ~30-70 nm and below 14 nm (Figure 5; Table 1); however, the latter two modes suffer of large rotational ambiguity and should be interpreted with care. On the contrary, it exhibits a wide mode in volume concentration ranging from ~100 to ~500 nm. Factor 6 has two relevant modes in number concentration at 55-120 nm and 230-400 nm, and two modes in volume concentration at 260-500 nm and 75-140 nm.

These factors still present two peaks corresponding to the rush hours, but the morning peak occurs 1-2 h earlier than in the road traffic-related factors, i.e. when ambient temperature reaches its daily minimum. Both factors correlate well with secondary aerosol tracers (nitrate, sulphate, OC) and non-volatile components (eBC, EC, $NVPM_{2.5}$), but Factor 6 exhibits much higher correlation coefficients (Table 2). Despite the polar plots indicating the main wind directionality toward N-E sectors, the analysis of air mass histories though the CWT model (Figure 7) clearly indicates likely continental origin areas rather than local sources.

Vu et al. (2016) observed two factors in North Kensington with very similar modal structures, daily patterns, correlations with $PM_{2.5}$-bound species and external source areas maps. Therefore, their interpretation is confirmed also in this study, i.e. mixed secondary aerosol (Factor 5) and inorganic secondary aerosol (Factor 6). Both factors are clearly originated from continental Europe and are consistent with a previous receptor modelling study carried out in a rural background site representative of the southern UK (Charron et al., 2013). Similar origin and formation mechanisms also explain their strong correlation (0.75). Although it is not reasonable to extract much more information from these data due to the short period of sampling and the large uncertainty associated with back-trajectory analysis, it can be observed that Factor 5 shows a wide source area all over Central Europe, while Factor 6 exhibits two distinct hotspots (Central and North-eastern Europe).

3.3.2 Cold season

The 6 factors identified during the cold period (Figure 6) are similar to those for the warm season. *Factor 1* is composed of a high proportion of particles in the nucleation range with a sharp mode at ~15 nm. It accounts for 33% (32-35%) of PNC and less than 2% of PVC. The polar plot reveals increased concentrations for moderate winds blowing from the airport sector and the diurnal pattern is also compatible with the aircraft traffic. No statistically significant correlations are found with

any other monitored species (Table 3). Therefore, Factor 1 may be attributed to the airport emissions related to aircraft engine exhaust. As in the warm season, factor 1 is moderately correlated with factor 2 (fresh road traffic, r=0.55), indicating a quite clear relationship between the two sources.

Factor 2 represents particles in the 15-35 nm range of number concentration, accounting for 35% (33-37%) of total PNC (Figure 6; Table 1). Its importance for volume concentration is minimal (3%) with two modes at 30 and 200 nm. The diurnal pattern and the wind directionality are compatible with LHR as a source and it shows a weak positive correlation with NO_2 (0.42) and a strong correlation with nitrate (0.63) (Table 3). Despite its similarity and relationship with Factor 1 and the consequent similar potential origin, Factor 2 may represent a different source: Factors 1 and 2 remain clearly separated even at solutions down to 4 factors, demonstrating their structural robustness and the lack of potential artefacts affecting the PMF solution. Consequently, it can be concluded that they to not represent over-resolved solutions (i.e. factor splitting). The most plausible interpretation for Factor 2 is therefore the same as for the warm season, i.e. fresh road traffic emissions. Furthermore, this factor can be attributed to the road traffic generated by the airport and nearby major roads.

Factor 3 includes most of the particles in the Aitken range and accounts for 19% (18-20%) of PNC. It contribution to particle volume concentration is relevant (9%) with a main peak at around 100 nm and a secondary peak at 400 nm (Table 1). It presents two rush hours peaks and the polar plot reveals an origin from the SW quadrant. However, as with the warm period, the wind directionality suggests increases for slower wind regimes than the fresh road traffic factor and for more westerly sectors, which are not compatible with the airfield location. Since factor 3 correlates well (Table 3) with a number of other pollutants linked to primary emissions from road traffic (NO (0.51), NO_2

(0.81), eBC (0.52), $PM_{2.5}$ (0.53), OC (0.79) and EC (0.83)), it represents a second road traffic factor, more affected by aging in the atmosphere than factor 2.

Despite the wind regimes from NE sectors being poorly represented during the cold campaign, *Factor 4* is the only one showing a possible origin from London and for calm wind periods. As with the warm season, it is composed of a wide range of particles encompassing the Aitken and accumulation modes (50 to 150 nm), while the peak in volume concentration is at 170 nm (Table 1). The diurnal pattern (Figure 6) is clearly related to the mixing layer dynamics and the correlation analysis reveals strong relationships with many species (NO, NO_2, eBC, Delta-C, $NVPM_{2.5}$, OC, EC, nitrate, ammonium and potassium; Table 3). Consequently, it is concluded that it represents the urban accumulation mode, whose contribution to the total volume concentration is also similar to the warm season (33%). It is interesting to note the large similarity with the urban accumulation mode found in the warm season, from which it differs slightly only in the diurnal pattern (higher overnight) and in the presence of a strong correlation with nitrate (r=0.88), possibly due to the lesser extent of negative artefacts on $PM_{2.5}$ filter samples.

The last two factors are interpreted as due to secondary aerosols. Their modal structures, their contributions to total PNC and PVC, and their correlations with $PM_{2.5}$-bound species (Table 3; Figure 6) largely reflect the results obtained for the warm period. However, the CWT maps (Figure 7) highlight different source areas, i.e. the origin of the secondary aerosols is regional (UK and Northern Europe). In addition, the presence of strong positive correlations with chloride may also indicate a contribution from the transport of sea-salt aerosol.

3.3 Comparison of *k*-means and PMF

The cluster analysis revealed the presence of 5 characteristic PNSD shapes during both the seasons. These spectra have been linked to potential sources in the study area, i.e. road traffic, airport

activities, and secondary aerosol formation processes. However, the cluster analysis is mostly driven by the spectral size regions with higher particle number concentrations, i.e. it has the disadvantage of partitioning the single observations predominantly according to the finest region of the size distribution. This limitation is well illustrated by the poor (almost null) separation of clusters based on the particle volume distributions (all clusters showed quite similar particle volume spectra). In addition, cluster analysis also has the disadvantage of linking each cluster to a single source and does not easily account for PNSD resulting from the mix of two or more different sources.

In contrast, the PMF analysis computed over the PNSD also accounts well for the sources with a small impact on the number distribution, but having a larger influence on the particle volume size distributions and, therefore, on the particle mass concentration. Despite the differences in the two methods, some further information can be extracted by combining the results of cluster and PMF analysis. Figure 8 shows the statistics of normalised PMF source contributions relating to each single cluster.

For the warm period, significantly higher (0.05 significance) PMF contributions of the airport factor (F1) are measured for cluster 1 (average normalised contribution ~3.5). This result indicates that the airport fingerprint was well captured by both source apportionment methods. During the cold season, the airport factor (F1) is significantly higher for both clusters 1 and 5 (average normalised contributions of ~2 and ~3, respectively). While cluster 5 presents significant high PMF contributions only for factor 1, cluster 1 also shows high contributions of factor 2 (fresh road traffic). This result indicates that cluster 5 may be linked as the typical PNSD spectra for airport emissions, while cluster 2 likely represents mixed emissions from aircraft and airport-related traffic. A close analysis of wind roses for the two clusters in the cold season (Figure 4) reveals that cluster 5 occurs at significantly higher wind speed regimes than cluster 1 (Mann-Whitney-Wilcoxon test at

749 0.05 significance level), i.e. average wind speeds of 8.3 and 5.9 m s^{-1}, respectively. As a
750 consequence, the different wind regimes may well be responsible for the split between the two
751 clusters.
752
753 Results for fresh traffic emissions also agree between the two methods. Factors 2 exhibit the higher
754 normalised contributions to clusters 5 (normalised contribution 2.5) and 1 (normalised contribution
755 ~3) for the warm and cold period, respectively (Figure 8). However, in winter it is evident that
756 PNSDs grouped on cluster 1 are also strongly influenced by airport emissions, probably due to the
757 lower mixing layer height and, thus, a lesser dispersion in the atmosphere.
758
759 Clusters 4 for both the periods show enrichments in the contributions for 4 PMF sources (aged road
760 traffic, urban accumulation and the two secondary aerosols) (Figure 8). This further emphasises that
761 cluster 4 represents the typical PNSD during daytime resulting from the mixing of different sources.
762 In a similar way, clusters 3 and 2 in the warm and cold periods, respectively, represent the typical
763 nighttime spectra (Figures 3 and 4), i.e. they exhibit similar partitioning over the PMF sources and
764 similar daily cycles.
765

766 **3.4 Analysis of a Large Regional Nucleation Event**

767 Regional photochemical nucleation episodes are regularly recorded in the Southern and Eastern
768 UK. Their general characteristics have been reported in a number of studies (e.g., Alam et al., 2003;
769 Charron et al., 2007; 2008; Beddows et al., 2015; Vu et al., 2016) and can be summarised as
770 follows: (i) particle modality at around 20 nm; (ii) higher frequency around noon in association with
771 the peak in actinic flux intensities; (iii) clear seasonal cycles (higher average contribution levels in
772 the summer, from June to September); (iv) marked directionality from the westerly sectors,
773 reflecting maritime atmospheric circulation regimes, with high wind speed and low $PM_{2.5}$
774 concentrations.

A strong regional nucleation event occurred during the warm period sampling campaign (starting on 7th September at 1 pm UTC and lasting for about 12 h). Increases of PNC were almost simultaneously recorded at Harlington and at Harwell, a national network rural background site located approx. 60 km WNW of LHR and representative of the regional background levels of air pollution across the Southern UK. The comparison of PNC time series at the two sites is provided as Figure SI10. Figure 9 shows the contour plots of SMPS data recorded at Harlington between 7th and 8th September as well as the hourly averaged concentrations of nucleation, Aitken and accumulation particles, TEOM-FDMS $PM_{2.5}$ mass and the contributions of Factors 1 to 4 extracted by the PMF. Figure 9 also reports the hourly counts of number of clusters extracted by the *k*-means analysis. The contour plot shows a typical "banana" shape with particle mode growing from ~20 nm (1 pm) to ~100 nm (overnight). The episode strongly influenced the PNSDs until around midnight; however its effect is also visible over the first half of 8th September. The time series (Figure 9) exhibits a clear peak in nucleation particles between 1 pm and 3 pm followed by peaks of Aitken (3-11 pm) and accumulation mode (8 pm-2 am) particles. The back-trajectory analysis (Figure SI11) revealed that the event occurred when north-westerly fresh (and clean) maritime air masses were advected from the Atlantic. This is also supported by the $PM_{2.5}$ mass, which exhibited a fast drop of concentrations just a few hours before the event (-30 µg m^{-3} in 3 hours, i.e. from 40 µg m^{-3} at 6 am to 10 µg m^{-3} at 9 am, Figure 9), probably reducing the condensation sink and facilitating nucleation.

Both atmospheric nucleation and aircraft engines are recognised to produce particles in the nucleation range. The analysis of this single –but strong– episode gives insights into how much the source apportionment results can potentially be affected by regional nucleation. This latter analysis is possible because the wind directionality during the entire episode was from N sectors, i.e. the contribution of LHR can be considered negligible.

The results of cluster analysis were affected by the event. Before the episode, the PNSD spectra were mostly categorised as clusters 3 and 4 (urban background and daytime pollution, respectively), i.e. the clusters mostly recorded under north-easterly wind regimes (Figure 3). About 50% and 30% of the clusters were then categorised as "airport" in the first and second hour of the episode, respectively (Figure 9). Since the wind directionality is inconsistent with an origin from the airfield, this categorisation is likely the result of the nucleation event. The growing of particles in the hours after the beginning of the event has further driven the cluster results: (i) about 60-80% of PNSDs were categorised as "fresh road traffic" (cluster 5) after 2-3 hours, and (ii) 80-100% of PNSDs were clustered as "nighttime regional pollution" (cluster 2) after 4-6 hours. In a similar way, PMF results were affected by the event (Figure 9), with a sharp increase of contribution levels for: (i) factor 1 (airport) from 1.5 x 10^3 particles cm^{-3} at noon to 13.3 x 10^3 particles cm^{-3} at 2 pm; (ii) factor 2 (fresh road traffic) from 0.5 x 10^3 particles cm^{-3} at 1 pm to 21 x 10^3 particles cm^{-3} at 3 pm; and (iii) factor 3 (aged road traffic) from 2.1 x 10^3 particles cm^{-3} at 2 pm to approx. 15 x 10^3 particles cm^{-3} at 5-6 pm.

This episode was the main nucleation event recorded during the two sampling campaigns. Other possible episodes also occurred (mostly during the warm season), but they were much less significant and often hard to detect. This qualitative analysis points to some conclusions: (i) regional photochemical nucleation events may have an effect on clustering and PMF results; (ii) the effect may lead to an "additive" bias, mostly over the "airport" and "road traffic" factors and clusters; (iii) the effect of regional nucleation events in the study area is largely overwhelmed by the strength of local sources, but in other locations with more frequent nucleation events it may be more important to identify and separate them.

4 CONCLUSIONS

The effect of airport emissions upon the particle number concentration and size distribution was assessed at a site close to a major European airport (Heathrow) serving a megacity (London). The conclusions to be drawn are:

- High particle number concentrations were recorded for the finest sizes (nucleation <30 nm and Aitken nuclei 30-100 nm) if compared to an urban background site in London (N. Kensington).
- Polar plot analysis indicates that Heathrow is a strong potential source for NO_2, nucleation and Aitken particles, but its contribution to the mass concentration of $PM_{2.5}$ and eBC is very small. On the contrary, the urban area of London appears to be the main source for PM and eBC.
- The *k*-means cluster analysis has revealed that 20% of PNSDs are mostly shaped by airport direct emissions, but particle size spectra are also strongly affected by other local sources (mostly fresh and aged road traffic during daytime) and the reduction of mixing layer depth (during nighttime). Typical PNSD spectra have been identified for nighttime and daytime pollution as well. Such spectra are likely the result of multiple source mixtures.
- PMF analysis revealed that the fingerprint of Heathrow has a peculiar modal structure peaking at <20 nm. The direct airport emissions account for 30-35% of total particles in both the seasons. Such results are in line with percent estimations for NO_2 reported in previous studies.
- Other major contributors to PNC are fresh (24-36%) and aged (16-21%) road traffic emissions. Despite both applied source apportionment methods failing to fully disaggregate the emissions from the local traffic (including motorway) and traffic generated by the airport, results suggest that road traffic sources may contribute to the total PNC more than Heathrow (40-56%). However, making a clear distinction between the influence of traffic generated by the airport from other road traffic is not feasible from this analysis.
- An urban accumulation mode was found. This source presents a wide mode between 50-150 nm and accounts for around 10% of PNC. The wind directionality is consistent with the advection of air masses from London. It is more evident overnight due to the drop of the

852 mixing layer top, the subsequent increase in air pollutants at ground level and the generation of
853 nighttime secondary nitrate aerosols.
854 • Secondary sources accounted for less than 6% in number concentrations but for more than 50%
855 in volume concentration. Long-range transport has a key role in advecting polluted air masses
856 from mainland Europe.
857
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873
874

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TABLE LEGENDS:

Table 1. Summary of PMF results for both seasons.

Table 2. Results of Pearson's correlation analysis among extracted factor contributions and other measured variables recorded at different time resolutions. Only correlations significant at $p<0.05$ are reported, strong correlations ($\rho>|0.6|$) are highlighted in bold.

FIGURE LEGENDS:

Figure 1. Statistics of size distribution spectra for particle number (red) and volume (blue) concentrations categorised by sampling periods and time of the day (daytime= 7am-7pm and nighttime=7pm- 7am local time). For the particle number spectra, solid lines represent the median concentrations, while shaded areas report the 1st-3rd quartile intervals (interquartile range, IQR). For the particle volume spectra, only medians are reported (dotted lines).

Figure 2. Diurnal patterns of PNC, LHR traffic, solar irradiance and eBC. Plots report the average levels as a filled line and the associated 95th confidence interval calculated by bootstrapping the data (n= 200). Outliers (data >99.5th percentile) were removed for computing the diurnal patterns. Hours are given in UTC. LHR traffic movements (bottom right plot) are reported as arrivals (dotted lines) and departures (solid lines). The offset between the seasons is largely due to daylight saving time (BST = UTC + 1) in the summer data. The diurnal patterns of all the measured variables in reported in Figure SI4.

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1482 **Table 1**. Summary of PMF results for both seasons.

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Factor number and interpretation	Particle Number Concentration		Particle Volume Concentration	
Warm season (Aug-Sep 2014)	No. modes[a] (peak ranges[b])	Percent contribution (DISP range)	No. modes[a] (peak ranges[b])	Percent contribution
Factor 1: Airport	1 (<20 nm)	31.6 (30.8–36.2)	2 (60–160 nm; <25 nm)	1.2
Factor 2: Fresh road traffic	1 (20–35 nm)	27.9 (24.7–30.2)	2 (22–45 nm; 140–220 nm)	1.7
Factor 3: Aged road traffic	1 (30–60 nm)	18.9 (16.6–21.1)	2 (40–100 nm; 250–450 nm)	5.6
Factor 4: Urban accumulation	1 (50–150 nm)	14.4 (13.8–18)	1 (80–250 nm)	33.2
Factor 5: Mixed SA[c]	1 (110–250 nm)	5.2 (3.6–6.9)	1 (160–350 nm)	37.4
Factor 6: Inorganic SA	2 (55–120 nm; 230–400 nm)	2.1 (1.1–3.5)	2 (260–500 nm; 75–140 nm)	20.8
Cold season (Dec 2014-Jan 2015)				
Factor 1: Airport	1 (<20 nm)	33.1 (31.7–34.8)	2 (160–350 nm; 15–25 nm)	1.7
Factor 2: Fresh road traffic	1 (18–35 nm)	35.2 (33.4–36.9)	2 (22–45 nm; 150–300 nm)	3.1
Factor 3: Aged road traffic	1 (28–60 nm)	18.9 (17.9–19.7)	2 (40–150 nm; 330–450 nm)	8.7
Factor 4: Urban accumulation	1 (55–170 nm)	7.6 (7.3–8.3)	1 (100–250 nm)	32.5
Factor 5: Mixed SA	2 (130–280 nm, <17 nm)	2.3 (2.1–3.3)	1 (170–400 nm)	30.8
Factor 6: Inorganic SA	3 (17–28 nm; 55–100 nm, 250–400 nm)	2.9 (2.4–3.9)	2 (280–550 nm; 90–140 nm)	23.3

(a) Only modes above the DISP ranges are shown; (b) Range endpoints are taken at approx. half the mode height; (c) SA = secondary aerosol

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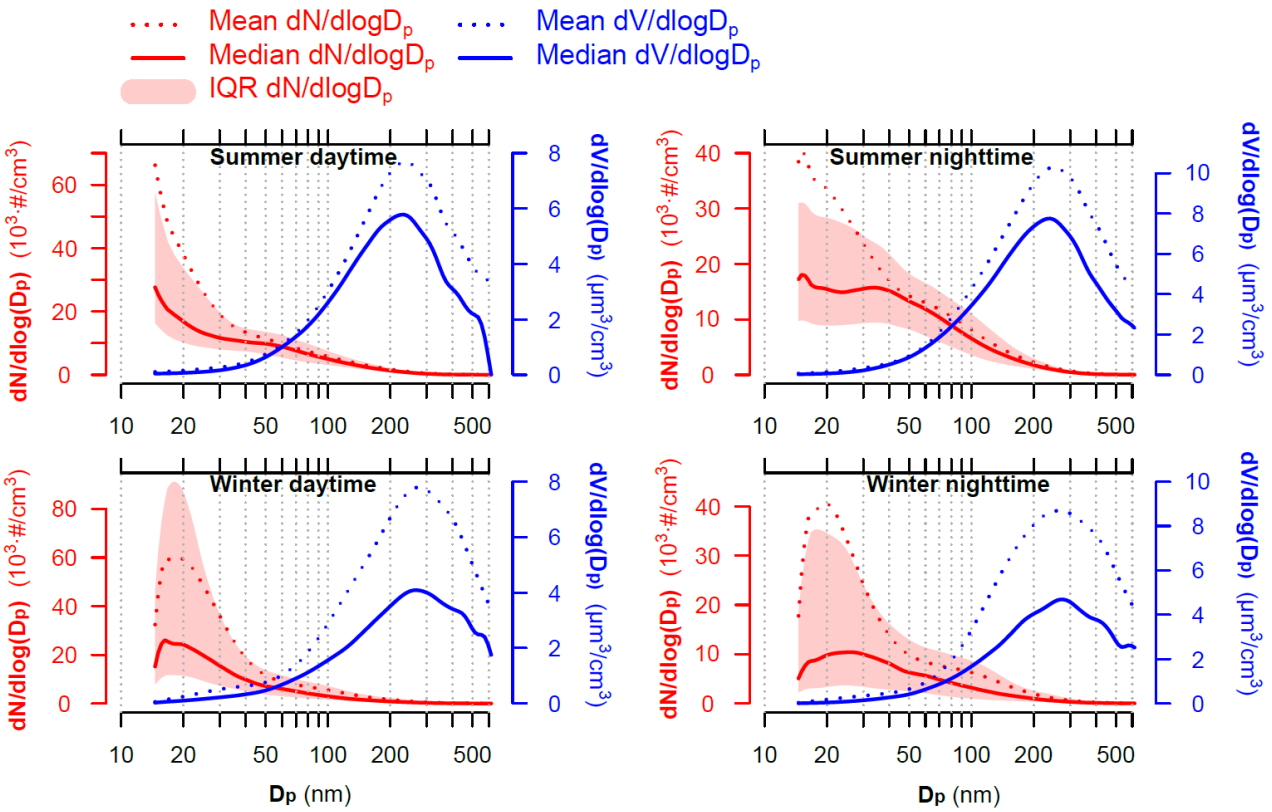
Table 2. Results of Pearson's correlation analysis among extracted factor contributions and other measured variables recorded at different time resolutions. Only correlations significant at $p<0.05$ are reported, strong correlations ($\rho>|0.6|$) are highlighted in bold.

	Warm period					
	Factor 1	**Factor 2**	**Factor 3**	**Factor 4**	**Factor 5**	**Factor 6**
Variables	Airport	Fresh road traffic	Aged road traffic	Urban accumulation	Mixed SA	Inorganic SA
Weather parameters (1 h-resolution time)						
Solar irr.	0.12	-0.15	-0.24	-0.26	-0.24	-0.28
Air temp.	0.25	-0.21	-0.37	-0.1	0.1	
RH		0.1	0.32	0.22	0.26	0.33
Wind speed	0.38		-0.47	**-0.64**	-0.45	-0.49
5 min-resolution time						
Factor 1	–					
Factor 2	0.46	–				
Factor 3	0.03	0.28	–			
Factor 4	-0.17	-0.04	0.47	–		
Factor 5	-0.15	-0.06	0.21	0.56	–	
Factor 6	-0.17	-0.14	0.15	0.56	**0.75**	–
eBC	-0.1	-0.03	0.33	**0.62**	0.52	0.53
Delta-C			0.13	-0.07		-0.06
1 h-resolution time						
NO			0.43	**0.6**	0.32	0.33
NO_2		0.18	**0.61**	**0.76**	0.52	0.52
NO_x		0.11	0.58	**0.77**	0.48	0.48
O_3	0.14	-0.19	-0.57	-0.54	-0.37	-0.43
$PM_{2.5}$	-0.23	-0.24	0.13	**0.61**	**0.63**	**0.77**
$NVPM_{2.5}$	-0.22	-0.22	0.17	**0.62**	**0.61**	**0.75**
$VPM_{2.5}$	-0.17	-0.24		0.42	0.54	**0.65**
1 day-resolution time $PM_{2.5}$-bound species						
OC				**0.84**	**0.74**	**0.83**
EC	-0.47	-0.54		**0.75**	0.51	**0.67**
TC	-0.45	-0.44		**0.85**	**0.69**	**0.82**
Chloride						
Nitrate		-0.45			**0.83**	**0.85**
Sulphate		-0.57		**0.75**	0.5	**0.67**
Oxalate		-0.47		0.59	**0.89**	**0.93**
Sodium						
Ammonium	-0.44	-0.52		0.57	0.54	**0.71**
Potassium		-0.47		0.46	0.5	**0.66**
Magnesium	0.5			-0.53		
Calcium						

Table 2. Continued.

	Cold period					
	Factor 1	**Factor 2**	**Factor 3**	**Factor 4**	**Factor 5**	**Factor 6**
Variables	Airport	Fresh road traffic	Aged road traffic	Urban accumulation	Mixed SA	Inorganic SA
Weather parameters (1 h-resolution time)						
Solar irr.				-0.11		
Air temp.	0.38		-0.43	**-0.67**	-0.5	-0.59
RH			0.23	0.38	0.46	0.46
Wind speed	0.3		-0.49	**-0.67**	-0.54	**-0.61**
5 min-resolution time						
Factor 1	–					
Factor 2	0.55	–				
Factor 3	0.24	0.54	–			
Factor 4	-0.11	0.08	0.53	–		
Factor 5	-0.05	0.15	0.38	**0.65**	–	
Factor 6	-0.09	0.08	0.39	**0.7**	**0.81**	–
eBC		0.16	0.52	**0.77**	**0.60**	**0.63**
Delta-C			0.35	**0.62**	0.55	0.52
1 h-resolution time						
NO	-0.14		0.51	**0.81**	**0.62**	**0.63**
NO_2	0.13	0.42	**0.81**	**0.82**	**0.61**	**0.66**
NO_x		0.17	**0.63**	**0.85**	**0.64**	**0.68**
O_3		-0.29	**-0.71**	**-0.78**	**-0.65**	**-0.7**
$PM_{2.5}$	-0.1	0.16	0.53	**0.82**	**0.88**	**0.88**
$NVPM_{2.5}$	-0.11	0.16	0.53	**0.82**	**0.85**	**0.85**
$VPM_{2.5}$			0.19	0.39	0.49	0.48
1 day-resolution time $PM_{2.5}$-bound species						
OC			**0.79**	**0.79**	**0.76**	**0.8**
EC			**0.83**	**0.8**	**0.64**	**0.66**
TC			**0.81**	**0.8**	**0.73**	**0.77**
Chloride				0.58	**0.82**	**0.85**
Nitrate		**0.63**	**0.73**	**0.88**	**0.93**	**0.9**
Sulphate					**0.92**	**0.88**
Oxalate					**0.87**	**0.81**
Sodium		-0.58	**-0.74**	**-0.64**		
Ammonium			**0.63**	**0.78**	**0.99**	**0.97**
Potassium				**0.71**	**0.98**	**0.97**
Magnesium						
Calcium						



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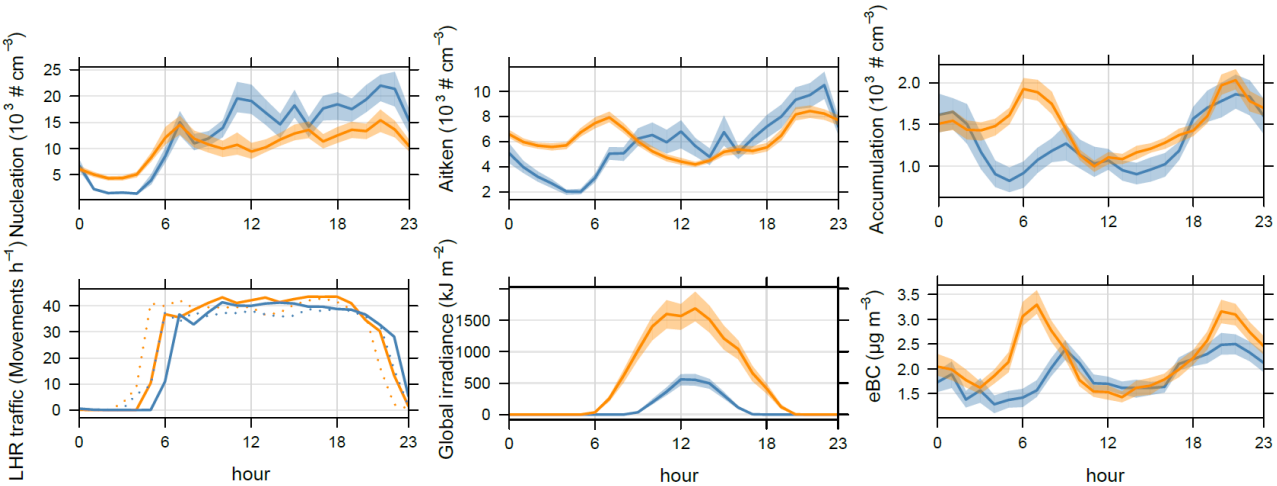

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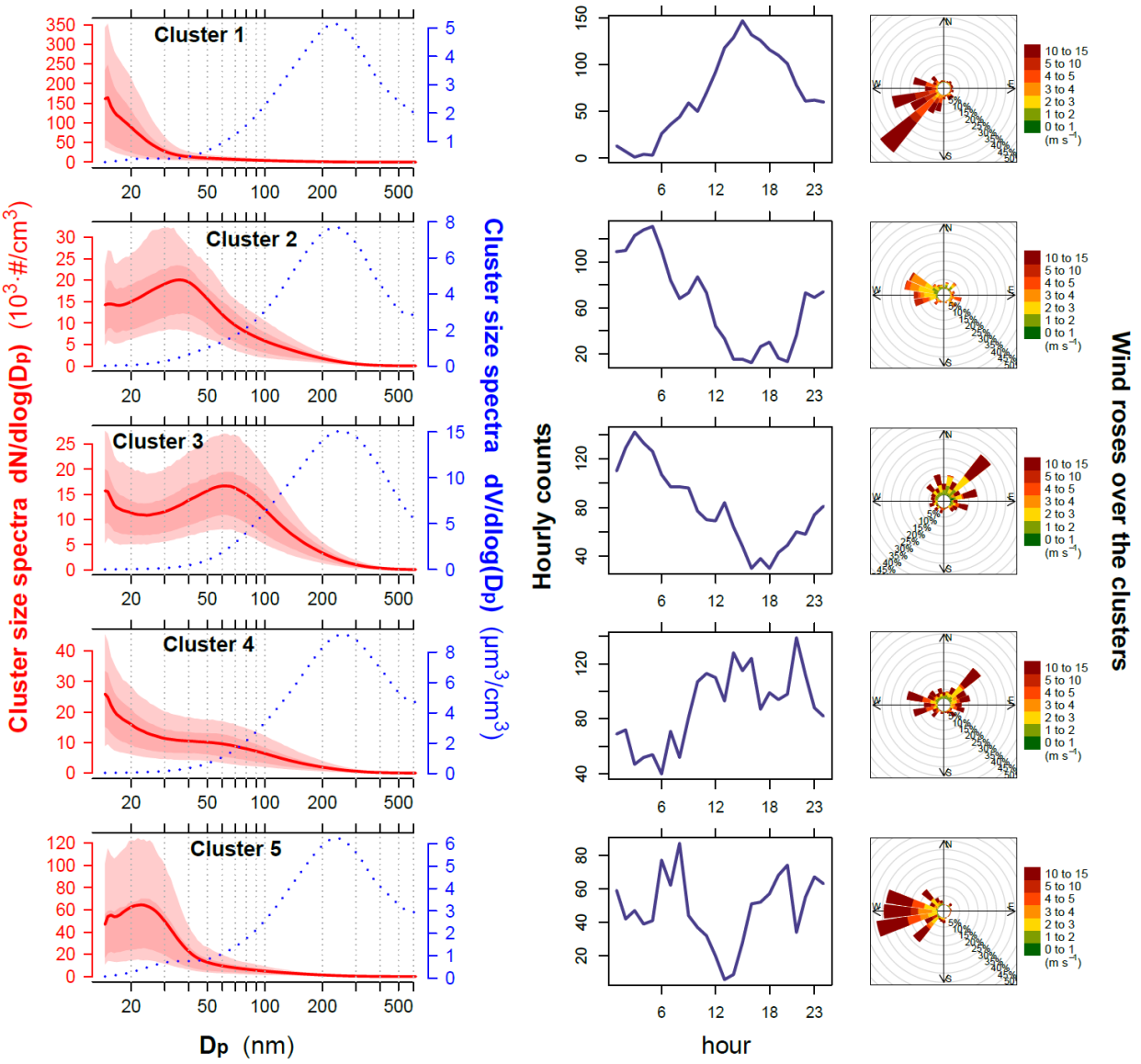

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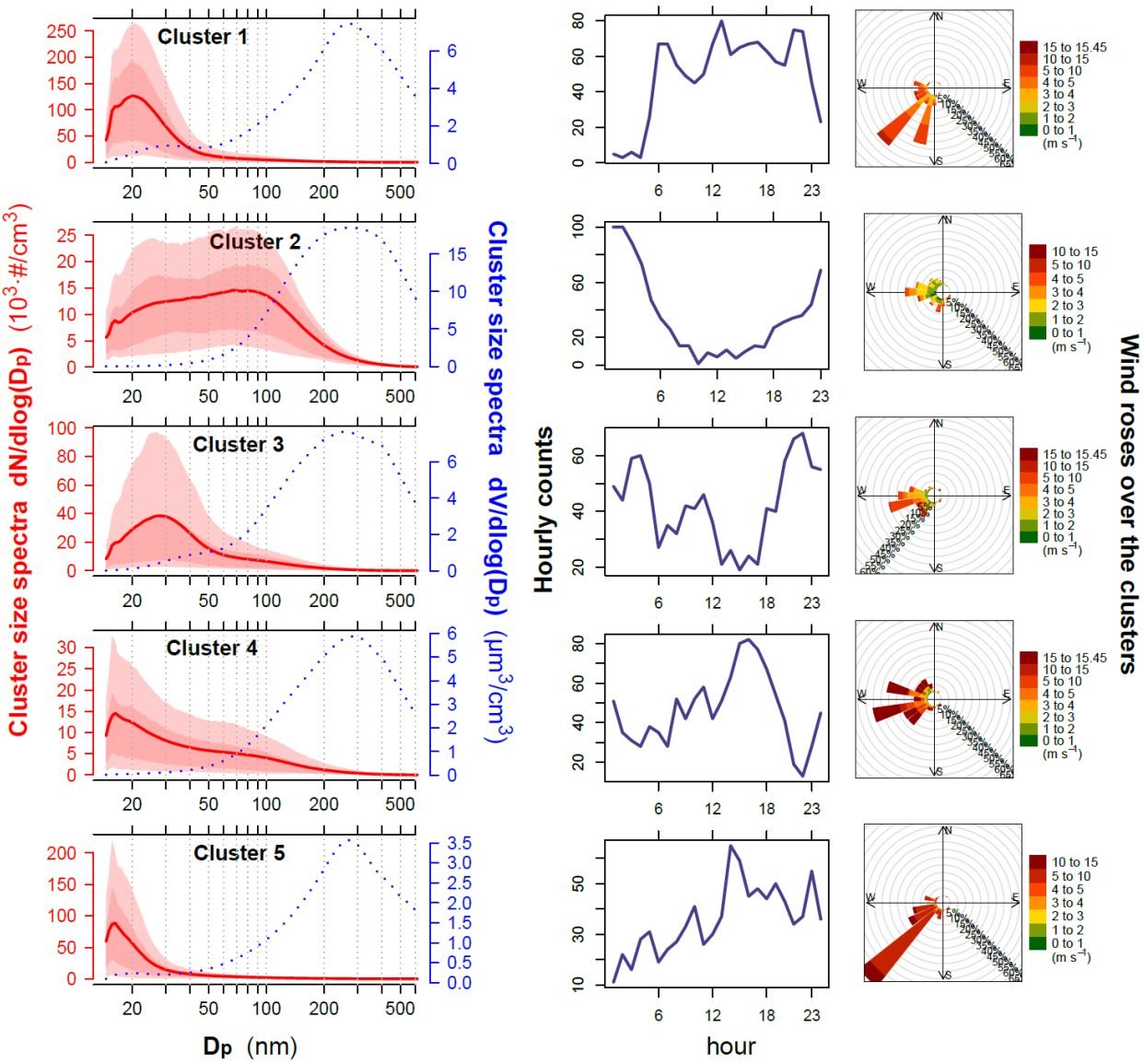

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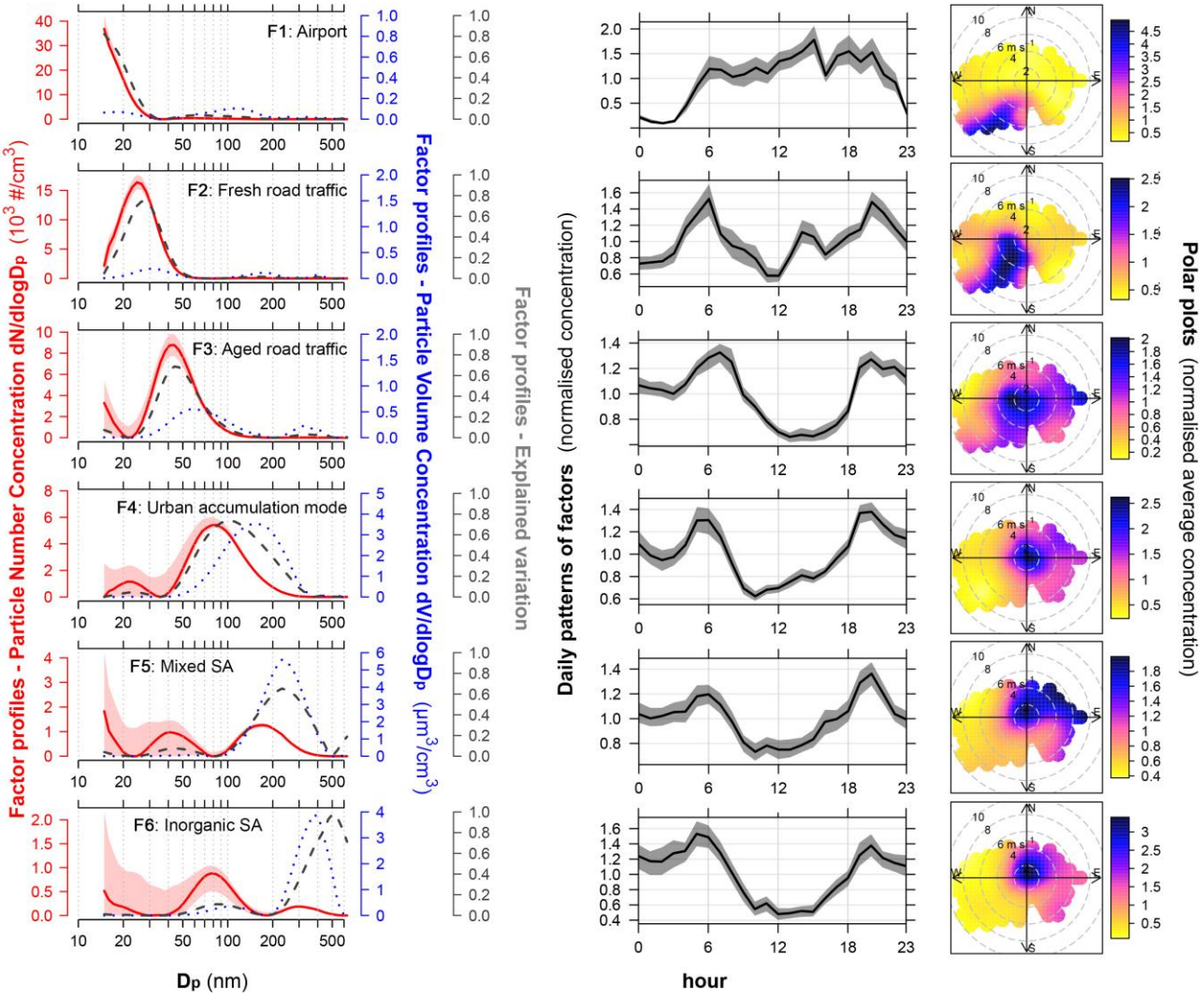

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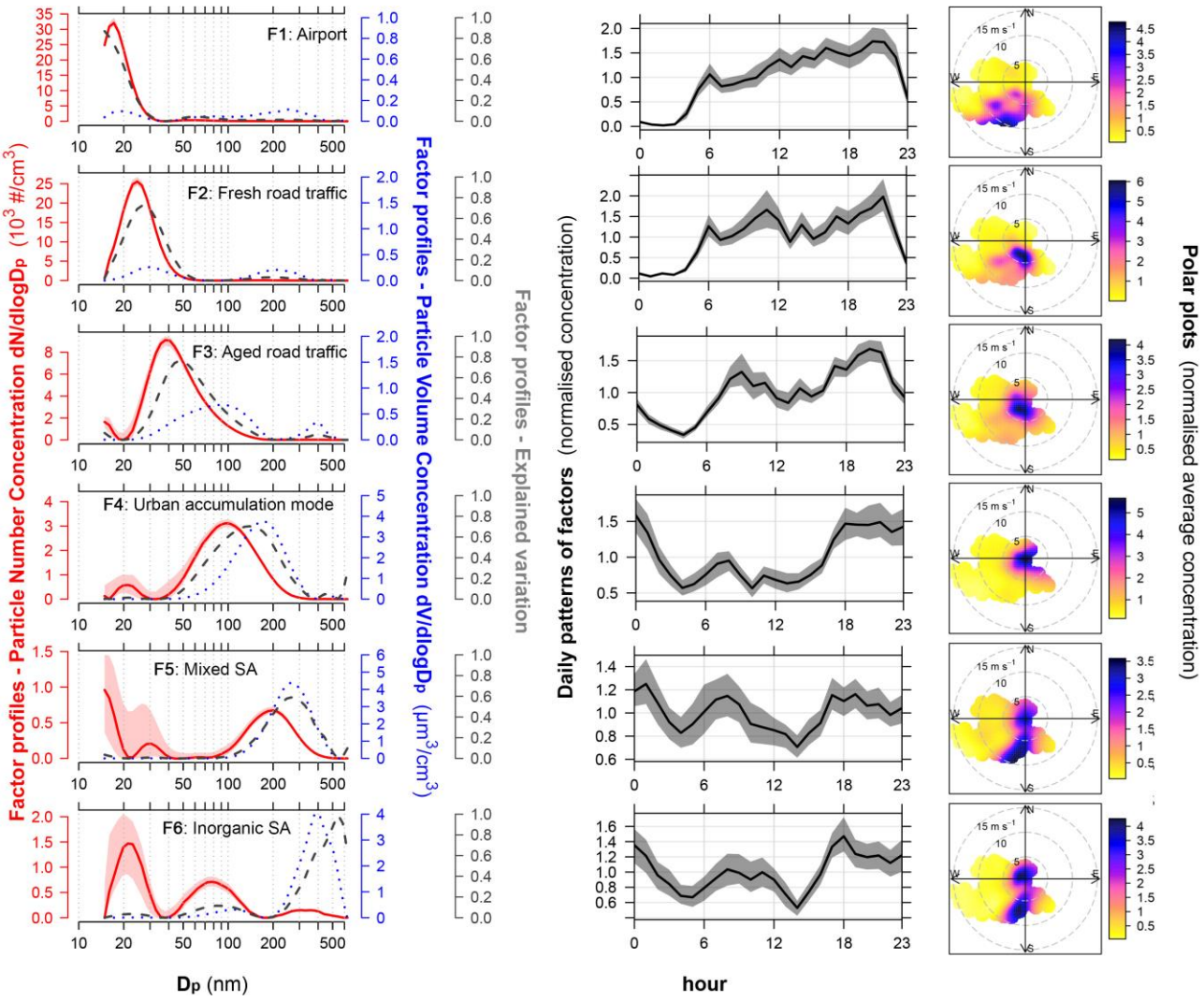

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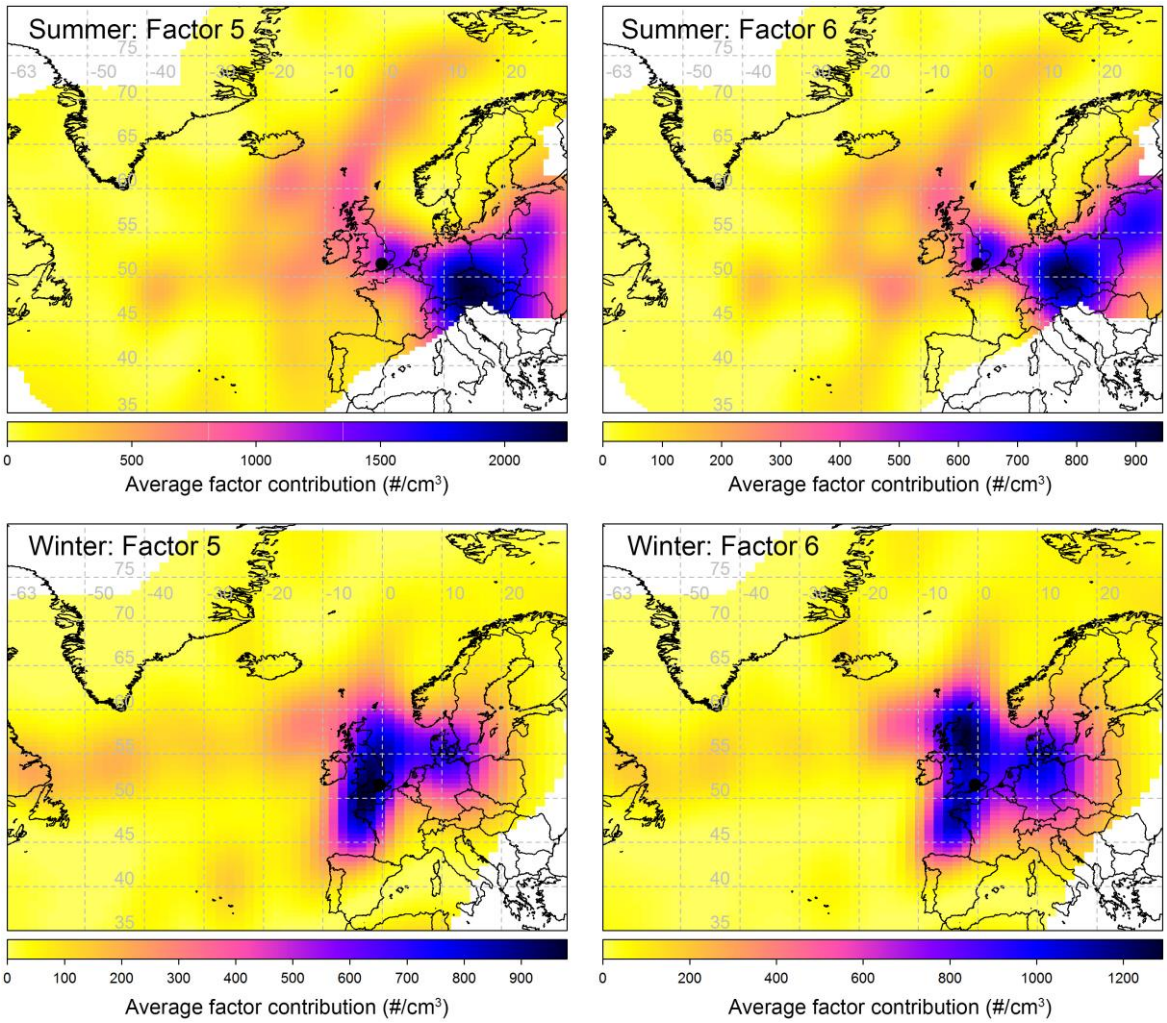

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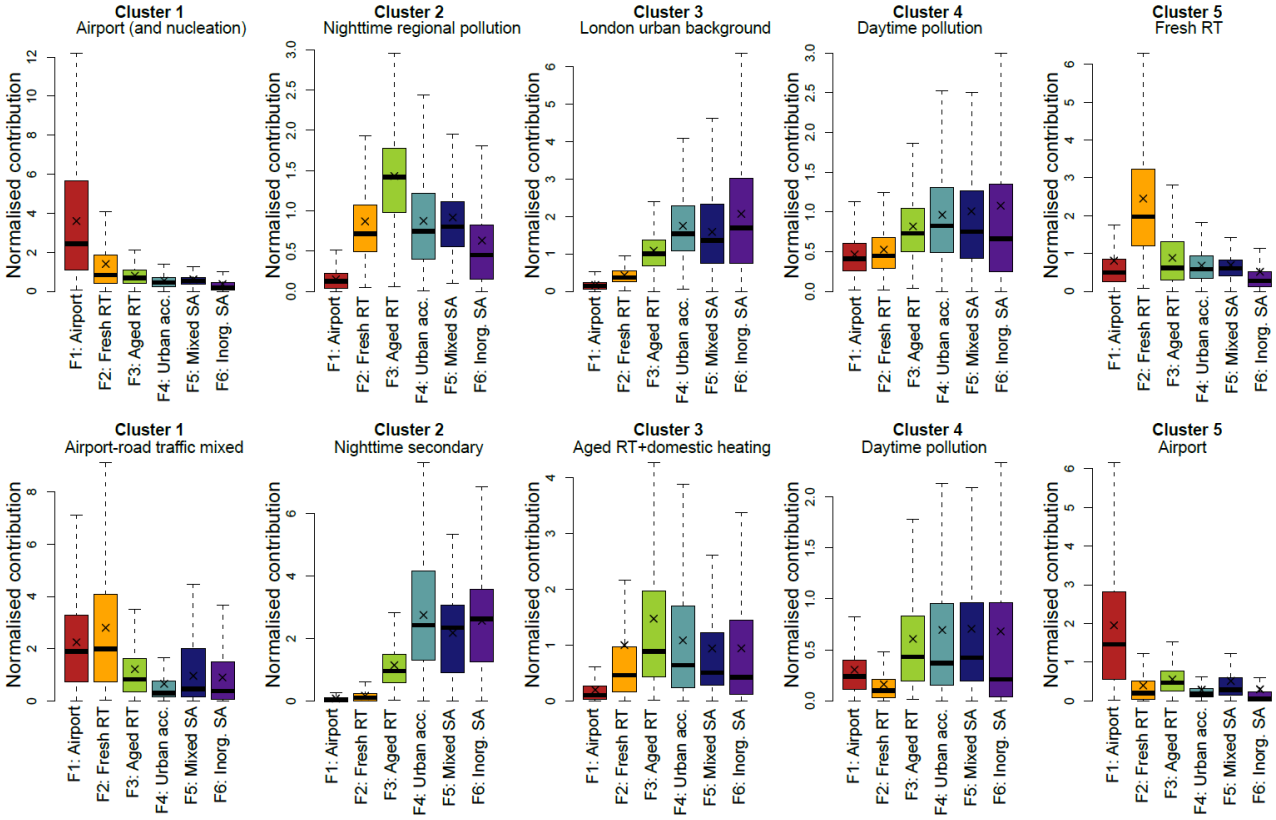

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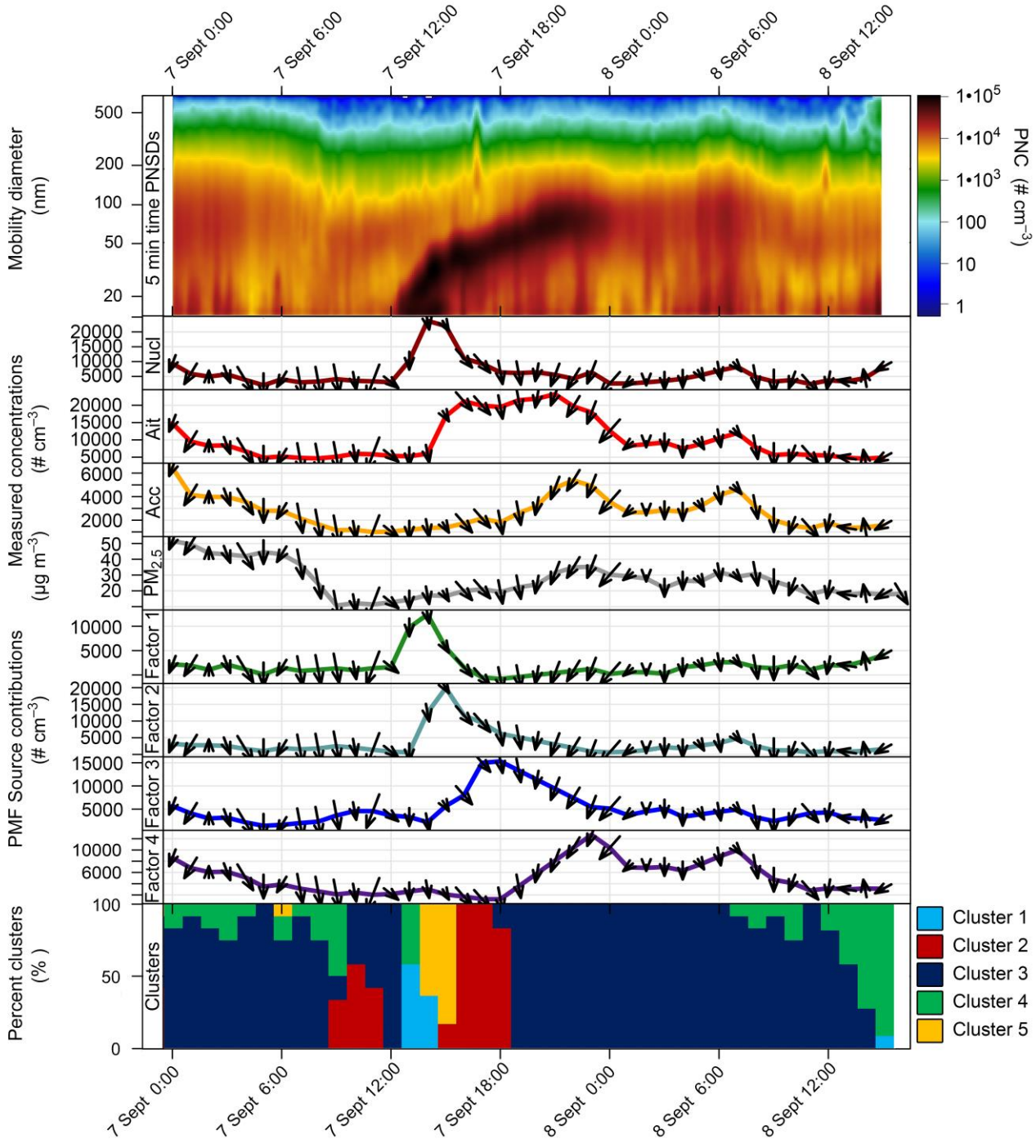

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