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New particle formation events measured on board the ATR-42 aircraft during the EUCAARI campaign

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Abstract

Aerosol properties were studied during an intensive airborne measurement campaign that took place at Rotterdam in Netherlands in May 2008 within the framework of the European Aerosol Cloud Climate and Air Quality Interactions project (EUCAARI). The objective of this study is to illustrate seven events of new particle formation (NPF) observed with two Condensation Particle Counters (CPCs) operated on board the ATR-42 research aircraft in airsectors around Rotterdam, and to provide information on the spatial extent of the new particle formation phenomenon based on 1-s resolution measurements of ultra-fine particle (in the size range 3–10 nm diameter, denoted N_{3-10} hereafter) concentrations. The results show that particle production occurred under the influence of different air mass origins, at different day times and over the North Sea as well as over the continent. The number concentration of freshly nucleated particles (N_{3-10}) varied between 5000 and 100 000 cm^{-3} within the boundary layer (BL). Furthermore the vertical extension for all nucleation events observed on the ATR-42 never exceeded the upper limit of the BL. The horizontal extent of N_{3-10} could not be delimited due to inflexible flight plans which could not be modified to accommodate real-time results. However, the NPF events were observed over geographically large areas; typically the horizontal extension was about 100 km and larger.

1 Introduction

The atmospheric aerosol particle formation and processing results from complex sets of homogeneous and heterogeneous physicochemical processes. The smallest particles are either formed from supersaturated vapour in absence of any seed nuclei acting as a condensation surface, the so-called homogeneous nucleation (Kulmala, 2003), or due to the process of ion-induced nucleation (Yu and Turco, 2000; Curtius, 2006). The formation and growth of these initial clusters to detectable sizes ($D_p > 2-3$ nm) is counteracted by their simultaneous removal from the ultra fine particle mode by coagu-

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lation with the pre-existing aerosol particle population (Kerminen et al., 2001; Kulmala, 2003). Because of the two competitive processes of homogeneous/heterogeneous particle nucleation plus growth and condensation of gas-phase species onto the pre-existing particle populations, the number concentration of particles smaller than 20 nm has been observed to be anti-correlated with the aerosol volume and mass (Rodriguez et al., 2005). Hence, an increase of the nucleation mode aerosol number concentration during nucleation events would mean that the total aerosol volume is rather small. However, a more profound understanding of processes modifying the nucleation mode particle number and size distributions is clearly needed. Indeed nucleation particles received increasing attention as a potentially important source of aerosol particles affecting climate and human health.

Homogeneous nucleation events in the atmosphere have been observed around the world (Kulmala et al. 2004a; Kulmala and Kerminen, 2008) in various environments from the boundary layer (BL) at urban locations (Wehner and Wiedensohler, 2003; Ganguli et al. 2006; Lonati et al, 2006; Tuch et al. 2006) to remote polar background areas (Komppula et al. 2006, Virkkula et al. 2007) and in the free troposphere (Venzac et al., 2008). Brock et al. (1995) suggest the homogeneous nucleation to form aerosol of a binary solution of sulphuric acid and water in the Tropical Tropopause Layer (TTL) providing a source of the background stratospheric aerosol, i.e. for the Junge Layer aerosol (Junge et al., 1961). Current knowledge of submicron aerosol has generally identified the number size distribution to be composed of a Nucleation mode (3–10 nm diameter) representative of recent new particle formation, an Aitken mode (40–80 nm diameter) and an accumulation mode (100–300 nm diameter) representing more aged particles (Kulmala et al., 2004b). The aerosol size distributions have been classically described as a function of the air mass type (marine, continental, mixed) (Birmili et al., 2001; O’Dowd et al., 2001; Tunved et al., 2005; Venzac et al., 2009). However, even within the same air mass type, the aerosol may be rapidly transformed by input of additional reactant sources (particles or gases), dilution, or deposition and coagulation processes within few tens of hours, leading to major differences in aerosol particle

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size distributions, chemical composition, state of mixture (internal or external) and thus their physico-chemical behaviour, even when comparing locations separated by a few hundreds of kilometres (Tunved et al., 2005).

The occurrence of new particle formation episodes depends on various factors including emissions strength of precursors, composition and number concentration of pre-existing aerosol population, meteorological parameters (in particular solar radiation, temperature and relative humidity) and photo-chemical processes (Kulmala et al., 2001, 2003). Air masses of different origins have different meteorological, physical and chemical characteristics, and therefore the probability of occurrence of new particle formation (NPF) events at a given location and time depends not only on local emissions, but also on long range transport (Sogacheva et al., 2005, 2007; Tunved et al., 2006). Moreover, regarding the spatial extension of NPF events, some important questions arise: does nucleation occur in the free troposphere or in the BL? What is the horizontal extend of NPF events, and are they homogeneously distributed over this spatial scale?

The European Aerosol Cloud Climate and Air Quality Interactions (EUCAARI, Kulmala et al., 2009) project has been created to investigate the effects of aerosol particles on climate and air quality. Quantifying the effect of aerosols on the planet's radiative balance is one of the most urgent tasks in our effort to understand future climate change (IPCC, 2007). As a whole, the contribution of the various aerosol sources, the role of long-range transport, and the contribution of primary and secondary particulate matter to the atmospheric aerosol concentrations over Europe are still not well known. Particularly uncertain remain the aerosol parameters influencing the radiative balance and the properties of clouds (Adams and Seinfeld, 2002; Kaufman et al., 2002; Spichtinger and Cziczo 2008), such as the number size distribution, chemical composition and particle mixing state. The EUCAARI observational system consists of long-term and spatially extensive surface-based measurements, including the European network of supersites for aerosol research (EUSAAR, Philippin et al., 2009). In addition, the use of satellites contributes to understanding regional aerosol variations

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and provides the spatial distribution of aerosol and cloud properties (Meskhidze et al., 2009).

Within the frame of the EUCAARI project an intensive measurement campaign including the deployment of European instrumented research aircraft (DLR Falcon-20, British Bae-146, NERC Dornier, French ATR-42 and French Falcon-20) have been performed in May 2008. Some of the results from these flights have already been reported for example by Mirme et al. (2010). The French ATR-42 has been equipped with state-of-the-art aerosol and cloud instrumentation. The study presented here focuses on the measurements of nucleation mode particles (concentrations, vertical and horizontal extension) on board the ATR-42 aircraft. Lagrangian back-trajectory calculations were used to interpret the impact of the air masses origin on the frequency and intensity of observed nucleation mode particles and thus, related to a NPF event.

The following section (Sect. 2), describes briefly the airborne sampling strategy as well as the data set of measured aerosol properties, whereas the basic findings of this study are presented in section 3, by presenting the air mass analysis via back-trajectories correlated to nucleation mode particle observations.

2 Methods

Within the European integrated project EUCAARI on aerosol-cloud-climate research, west-east and north-south station-to-station networks including the Cabauw (CBW) supersite have been used together with quasi-Lagrangian and Eulerian airborne measurements in order to quantify the effects of emissions on regional aerosol properties, and to study aerosol formation, processing, transport and deposition. To improve our understanding on processes related to atmospheric aerosol physics and chemistry, from nanometer scale to the overall aerosol-cloud-climate scale, the EUCAARI intensive observation campaign, including deployment of research aircraft, has been performed in May 2008.

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2.1 Ground based measurements

The Cabauw (CBW) field site is located in the western part of the Netherlands (51.971° N, 4.927° E, 0.7 m below sea level), 35 km east of Rotterdam. The air masses arriving to this rural site are typically originating from clean marine sector from the north or from polluted continental Europe. The speciality in CBW is the 213 m high mast which allows for sampling at different altitudes and hence is used to establish relations between the state of the lowermost atmospheric BL, aerosol particles, clouds, the weather situation, and land surface conditions (Van Ulden and Wieringa, 1996).

The Neutral cluster and Air Ion Spectrometer (NAIS, Kulmala et al., 2007) was measuring aerosol sampled through an inlet installed at 60 m height on the CBW mast. The NAIS measures simultaneously electrical mobility distributions of both negative and positive air ions and total particles in the range from 2.39 to 0.001 cm² V⁻¹ s⁻¹, which corresponds to a mobility diameter range of 0.8–40 nm for ions and in case of particles ~2–40 nm. Controlled corona charging together with electrical filtering also enables measurements of the neutral aerosol particles down to ~2 nm. The instrument consists of two identical cylindrical differential mobility analyzers where the sample is collected to insulated electrometer rings at the outer electrode. A more detailed description of the instrument and examples of its atmospheric applications can be found in Manninen et al. (2009).

2.2 Instrumented ATR-42 aircraft

During May 2008 the French ATR-42 research aircraft operated by SAFIRE (Service des Avions Français Instrumenté pour la Recherche en Environnement) was based at Rotterdam airport in the Netherlands for the duration of the EUCAARI intensive observation period. A total of 22 research flights was performed between 2 and 30 May 2008 (Table 1). For the present study these 22 flights are grouped into six different types of Research Flights (RF_type_n, Fig. 1). The 9 flights corresponding to type RF_type_1 show flight patterns which were centred at and around the ground-based measure-

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ment area of CBW supersite. The objectives of these flights were to specify origins and regional characteristics of the air masses sampled at Cabauw, thus correlating aircraft characterization of the vertical structure of the atmosphere above Cabauw with the ground-based measurements at the supersite and the IfT-helicopter platform (Wehner et al., 2010).

Since the occurrence of the NPF phenomenon in the proximity of the oceans and seas is still not well understood (O'Dowd et al., 2007), nucleation particle concentration observations near the coastline of the North Sea are of great interest in our study. This type of research flight close to the coastline is denoted type RF_type_2 with a total of four corresponding flights. The goals of RF_type_3/RF_type_4/RF_type_5/RF_type_6 (number of flights is 2, 2, 4, and 1 respectively) aimed at studies related to aerosol properties along “quasi-Lagrangian” flight tracks: West-East and North-South transects, most of the time in coordination with the British Bae-146, DLR -Falcon 20 flights and ground sites.

The deployment of the French research aircraft ATR-42 has been particularly dedicated to aerosol-cloud interaction research as already made in previous recent studies (CrumeYrolle et al., 2008; Matsuki et al., 2010). The aerosol instrumentation has been connected to the ATR-42 community aerosol inlet (CAI) and switched to the Counterflow Virtual Impactor (CVI, Schwarzenboeck et al., 2000) during cloud measurements. The isokinetic and isoaxial CAI, based on the University of Hawai shrouded solid diffuser designed by A. Clarke and modified by Meteo France, has a 50% sampling efficiency for particle size higher than 5 μm (McNaughton et al., 2007; Gomes et al., in preparation).

The relevant aerosol instrumental setup (for the presented aerosol nucleation study) consisted of two condensation particle counters (CPC, McMurry, 2000), one TSI model 3025 with measured nominal 50% lower cut-off diameter of 3 nm and one TSI model 3010 with a cut off diameter of 10 nm (Mertes et al., 1995; Stolzenburg and McMurry, 1991), and a SMPS (Scanning Mobility Particle Sizer) measuring particle size distribution between 20–500 nm. The SMPS system consisted of a TSI model 3010 conden-

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sation particle counter, a DMA (Differential Mobility Analyzer) as described by Villani et al. (2007) and a krypton aerosol neutralizer (Kr-85). The sheath flow rate was controlled with a critical orifice in a closed loop arrangement (Jokinen and Mäkelä, 1996). Typically the scan time was chosen to be 90 s. Thus, within the entire instrumental setup on the ATR-42 aircraft, we were able to record nucleation mode particle concentrations by subtracting the data readings of TSI-3025, TSI-3010 (i.e. using the 50% cut-size for respective CPCs), and SMPS integral concentrations. To take into account the coincidence effect, particle concentrations were corrected using the equation given by the producer specifications. In this way we deduced estimates of the ultrafine particle concentrations between 3–10 nm (N_{3-10}), 3–20 nm (N_{3-20}) and 10–20 nm (N_{10-20}). The time resolution of CPC counters was set to 1 s, with relative uncertainties in concentration measurements of about 5% according to the producers specifications.

Besides, the ATR-42 was equipped with state of the art aerosol instrumentation (PSAP, Bond et al., 1999), nephelometer (TSI 3563, TSI Inc., St. Paul, MN), Aerosol Mass Spectrometer (AMS, Jayne et al., 2000; Drewnick et al., 2005; Canagaratna et al., 2007), particle counting/sizing instruments, and cloud instrumentation (Fast Forward Scattering Spectrometer Probe, Fast-FSSP, Brenguier et al., 1998).

3 Results

3.1 Observation of atmospheric nucleation

New particle formation (NPF) events are frequently monitored on single measurement sites, where contour plots of the temporary evolution of nucleation particle size spectra are utilised to illustrate distinct features of the new particle formation and growth. Within EUCAARI a NPF event classification was performed visually from the NAIS and SMPS data measured at the CBW ground based site and ATR-42, respectively. NPF event classification follows methods introduced earlier by Dal Maso et al. (2005). In addition to the NPF observation at the CBW site, considerable amounts of nucleation mode

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particles, thus related to NPF events, have been observed during 10 out of 22 research flights performed with the ATR-42.

The NPF observations could be associated to all types of research flights from RF_type_1 to RF_type_6. Most of the time, when NPF events were observed with the aircraft CPCs and SMPS, the phenomenon was simultaneously observed with the NAIS and SMPS at the CBW supersite. As a consequence, the new particle formation was not limited to a local small scale nucleation source, but was a spatially widespread phenomenon. The overview of the nucleation observations at the CBW supersite and ATR-42 measurements is given in Table 1. Only in two cases (9 and 29 May 2008), the nucleation mode particle event observations differ when comparing the CBW ground site measurements with the aircraft observations. Indeed, on 9 May, the NPF event was observed over the CBW ground site while the ATR-42 (Flight 47) flew from Oberpfaffenhofen (Germany) to CBW. For the case of 29 May, the NPF event is observed with the ATR-42 in an air sector 150 km east of CBW, but not at CBW supersite.

The horizontal and vertical extension of the nucleation mode particles observed during four flights (RF44, RF55, RF56 and RF62) are on the scale of only a few kilometres, and not subject of further study here. Only seven flights showing nucleation mode particles (3–10 nm) on a large scale will be considered in this study. All the seven events with observed nucleation mode particles happened during the daytime (10:00–16:00), which is consistent with earlier observations (Dal Maso et al., 2005; Kuang et al., 2008; Laaksonen et al., 2008; Yu et al., 2008), where NPF events are reported to be observed frequently in very different locations (Mäkelä et al., 1997; Birmili and Wiedensohler, 2000, O'Dowd et al. 2007). The nucleation mode particles have been observed in the BL during different day times: (a) during the morning (before 11:00, flight 53), (b) during the early afternoon (before 16:00, flight 61), but mostly (c) at midday (before 14:00, RF43, RF45, RF48, RF50, RF54). Then, 45% of NPF events were detected at noon, 33% were observed in the afternoon and only 11% in the morning. These frequencies are consistent with continuous measurements at different ground sites (Dal Maso et al., 2004; Vehkamäki et al., 2004; Dal Maso et al., 2008; Hussein et al., 2008; Kuang

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3.2 Air mass origins

For all ATR-42 flights performed during EUCAARI IOP, air mass back trajectories have been calculated from HYSPLIT (Draxler et al., 2003) along the ATR flight path regardless of occurrence of the nucleation mode particles. For all ATR-42 flights, the prevailing atmospheric flows in the BL are given in Table 1. Back-trajectories have been calculated 24 h, and 48 h backward in time. Thus, the respective air mass origin, once calculated, has been attributed to the corresponding segment of the aircraft path. Figure 2 represents the frequencies (colour coded from 0–1) of observed nucleation mode particles as a function of air mass origin (squares) determined for two chosen back-trajectory calculation times of 24h (Fig. 2a), and 48 h (Fig. 2b). The numbers associated to each square denote the total number of calculated air mass origins assigned from back trajectory calculations for the complete set of EUCAARI flights (flight segments were selected in a horizontally equidistant way, then 138 back-trajectories were calculated subsequently). To represent only statistically significant results, only the NPF probability calculated when the total number of air masses is higher than five are represented in Fig. 2a and b.

As a consequence of the air mass dispersion, the frequencies of detected NPF events decrease with the duration of backward calculations (from 24 to 48 h). For trajectory calculation of 24 h (Fig. 2a), the majority of the “air mass origins” sampled by the ATR-42 stemmed from the South of Scandinavia (115 back-trajectory origins). The probability of NPF is highest for air masses originating from the western Scandinavia and the South-West of Europe and it is lowest when air masses are coming from South-East of Europe (<20%). These observations are confirmed with the 48-hours back-trajectory analysis, which shows that the probability of NPF events occurrence when the air mass originates from the South-West of Europe and from North of Scandinavia is higher than 90% (Fig. 2b). One hypothesis to explain the observation may be the transport of air masses with low background particle concentration and high con-

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centration of gaseous sulphate and/or organic species (Clarke et al., 1998, O'Dowd et al., 1998, 1999; Komppula et al., 2006; Metzger et al., 2010). Since measurements of these gases have not been performed on-board the ATR-42, the hypothesis that those (clean) air masses with lower particle concentrations contain high concentrations of gaseous aerosol precursors and which have been transported to the air sectors of NPF observations cannot be proven. Besides, air masses with initially high particle contain such as air masses originating from eastern Europe are not favourable to NPF events.

3.3 Horizontal and vertical extension of NPF events

While in the preceding sections, all flights (including NPF events and nonevents) have been treated in common, in this section, only results from flights during which nucleation mode particles have been observed will be used to determine the vertical and the horizontal extension of these events. To highlight the atmospheric structure more in detail, Figs. 3 to 6 present horizontal cross sections of the aircraft flight path with measured N_{3-10} observed during six flights at a constant level static pressure. The grey line corresponds to the flight track when aerosol particles were analysed in clear sky conditions. During some of the research flights in the vicinity of the Rotterdam airport, the N_{3-10} particle concentration has been high, probably as a consequence of fine mode aerosol production or of high gas phase concentrations due to aircraft activity or local urban pollution. Therefore, observations made around the airport will not be taken into account here.

3.3.1 Horizontal extension

While for one of the six ATR-42 flights (flight 61) we can give the horizontal extension at least along the flight path of observed nucleation mode particles, this is not possible for the other five flights. The ATR-42 flight path for those flights (flights 43, 45, 48, 53 and 54) could not reach the limit of the event in its horizontal extension. In the next

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paragraphs, the six flights with NPF events are discussed in further detail. To study the horizontal extension of NPF events, flight legs are classified into four groups: Group I events occurred under continental South-East flow and were observed over the sea, Group II events occurred under continental South-East flow but were observed over the continent, Group III events occurred under marine flow and were observed over the continent and Group IV events occurred under continental South-West flow and were observed over the continent.

Events of Group I events were observed during three flights: RF43 on 5 May 2008 (Fig. 3a), RF45 on 8 May 2008 (Fig. 3b) and RF48 on 11 May 2008 (Fig. 3c). For these three flight cases, air masses were transported from the continent to the North Sea in a south-eastern flow. Recent studies (O'Dowd et al., 2002a, b) showed that NPF events in a coastal environment are linked with biogenic vapours which are greatly enhanced during the low tide period. Then the tide conditions have been determined (Table 2) using data from the French "Service Hydrographique et Océanographique de la Marine" (<http://www.shom.fr/>). The NPF events of RF43 and RF45 occurred when the sea was at its highest level, meaning that these events are probably more influenced by transport of pollution from cities close to the coastline while the NPF event of the RF48 might probably be linked to emission of biogenic gas, such as halocarbon species (Carpenter et al., 1999, 2000; Laturnus et al., 2000), from stressed biota during low tide conditions. The NPF event observed during RF43 was horizontally spread out over spatially very limited area (40 km) suggesting a NPF burst occurred here compared to the NPF events observed over the sea during research flights RF45 and RF48 with horizontal extensions higher than 100 km.

Furthermore, comparing both ATR-42 legs parallel to the coastline (RF43, RF45), the N_{3-10} is homogeneously distributed on RF43 (Fig. 3a) and shows more inhomogeneous character during the RF45 (Fig. 3c). During the RF43, the N_{3-10} reached high concentration ($80\,000\text{ cm}^{-3}$), whereas during RF45, the N_{3-10} was lower than $18\,000\text{ cm}^{-3}$. This inhomogeneity in N_{3-10} concentrations during RF45 seems to be a consequence of surface wind speed inhomogeneities. Indeed, along the flight track

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the concentration of particles with diameter higher than 50 nm (N_{50}) fluctuate between 1000 and 12 000 cm^{-3} . The highest N_{3-10} values are associated to air masses with enhanced N_{50} . This effect of inhomogeneous air mass dispersion implies that sources of gas and particles might be local and close to the observation area. As the above NPF events (RF43, RF45) were localized upwind Rotterdam (150 km) and Amsterdam (133 km), the metropolises along the coast might be the sources of NPF events.

Moreover, N_{3-10} gradient and fluctuation decrease when the distance from the Netherland coastline increases (Fig. 3b and c). At 50 km away from the coastline of the Netherlands the N_{3-10} particle concentration averages 18 000 cm^{-3} and at more than 130 km away from the coastline the N_{3-10} is lower than 10 000 cm^{-3} (Fig. 3b). The gradient of N_{3-10} is more pronounced during the RF48. Indeed the N_{3-10} particle concentration reached 100 000 cm^{-3} close to the coastline (<30 km) and was lower than 15 000 cm^{-3} at more than 130 km from the coastline (Fig. 3c). The horizontal N_{3-10} gradient is a result of the air mass dispersion, as well as nucleation particle growth and coagulation of nucleation particles during the transport from their source to the observation area. The particle concentrations (N_{3-10}) were higher during the RF48 which might be due to the environmental differences (low tide and higher solar radiation). Besides these favourable NPF event conditions, during the RF48, the freshly nucleated particles were propagating deeply over the sea due to high surface wind speeds ($>10 \text{ ms}^{-1}$). Then the gradient intensity depends on the distance between the localization of the ultra-fine particle precursor sources and the observation area, and on the surface wind speed.

The Group II events were observed during two flights: RF43 on 5 May 2008 (Fig. 4a) and RF48 on 11 May 2008 (Fig. 4b). The uppermost limit of Group II events horizontal extension can not be determined with the observations ($>50 \text{ km}$). The N_{3-10} were homogeneous ($\sim 80\,000 \text{ cm}^{-3}$) over the flight track at a constant level of pressure, but one can note that close or upwind to the big cities (Rotterdam and Eindhoven) the N_{3-10} decreased drastically to reach values lower than $20\,000 \text{ cm}^{-3}$. This results is likely a consequence of highly polluted air masses above and downwind of cities enriched in

particles (N_{50}) on which condensable vapours could condense. For both cases the N_{50} is higher than 4000 cm^{-3} in the vicinity of cities.

The Group III events were observed during two flights that have been performed on a single day: RF53 on 18 May 2008 during the morning (Fig. 5a) and RF54 on 18 May 2008 during the afternoon (Fig. 5b). Both research flights were performed nearby the Netherlands coastline. The air mass trajectories were similar all day long. Air masses originated from the Norway coast and, during their transport, never crossed a pollution plume. Thus, these air masses should have been enriched in organic vapours originating from volatile organic compounds (VOC) photo-oxidation from the Boreal forest and impoverished in particles ($<1000 \text{ cm}^{-3}$). These conditions are highly favourable for the occurrence of NPF events (Clarke et al., 1998; O'Dowd et al., 1998, 1999; Komppula et al., 2006; Metzger et al., 2010). In the morning, the NPF event was observed at lower altitudes of the BL (600 m) between 5.2 and 6.7° E . N_{3-10} concentrations have been around $52\,000 \text{ cm}^{-3}$ over the entire leg (longitude: 5.2 – 6.2° E). During the afternoon, in the BL (850 m), the averaged N_{3-10} particle concentration was about $43\,000 \text{ cm}^{-3}$ (longitude: 5.2 – 6.8° E). The tide conditions were not the same during both flights, low tide in the morning and high tide in the early afternoon (Table 2). Thus, for these cases, the tide conditions may influence the intensity of the NPF event but not the fact that these events occurred. Upwind of Amsterdam and Rotterdam, the N_{50} concentration reached maximum values of 2000 cm^{-3} . During both flights, the horizontal extension of the NPF events could not be delineated. Indeed, NPF events were spread over a distance larger than 200 km. Despite of this large horizontal extension, the N_{3-10} was homogeneous during both flight segments (RF53 and RF54) close to the coastline. As there are many environmental differences between both flight (flight altitude, vertical dynamics), the comparison of N_{3-10} is not straightforward, however, the time evolution of the NPF event is demonstrated. Indeed 3 h after the NPF event (flight 53) sampled with the ATR-42, the phenomena is still persisting in the lower layer (flight 54). The SMPS measurements onboard the ATR-42 demonstrate nicely the temporal growth of nucleation particles (i) within each of the two flights and (ii) particularly when compar-

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ing flights. The observed modal diameter reached 30–35 nm in diameter towards the end of the afternoon flight.

Finally, the Group IV events were observed during RF61 on 28 May 2008 during the early afternoon (Fig. 6). As mentioned before, this event is different from all the others because of its localized character. Indeed, the longitudinal and vertical bounds of the NPF event are well determined during the flight (5.1–5.8° E and 620–1350 m). The N_{3-10} heterogeneity along the flight segment was striking as compared to the Group I–III events. The N_{3-10} maximum values (40 000 cm⁻³) were reached at 5.6° E and 51.8° N. Over the leg the terrain has been similar, consisting of a succession of smallest towns/villages and arable farmland. Thus, localised sources of NPF precursors over the area of observed NPF events may have been insignificant, rather longer range transport into the area is the most likely source for NPF observations. Generally, the environmental conditions were totally different during the last five flights (RF59, RF60, RF61, RF62, RF63) as compared to the previous ones (particularly Groups I–III). Indeed, the air mass trajectories were coming from the south and then crossed many pollution plumes from industrialized regions of France and Germany. Along the whole flight track at 600 m of RF61, the N_{50} concentration was higher than 5000 cm⁻³ and reached 45 000 cm⁻³ where the NPF event has been detected. Thus, this inhomogeneous case occurred under highly polluted conditions.

3.3.2 Vertical extension

During all presented ATR-42 flights, where nucleated particles have been observed, at least one vertical profiling has been made, either at the beginning or at the end of the flight between 0.3 km and 3.5 km (highest flight level), providing a rapid characterization of the vertical profiles of thermodynamic conditions (inversion level, etc) and microphysical parameters. For each flight the height of the BL, determined by the vertical temperature and humidity progress, and the height of the NPF event, estimated by the N_{3-10} , have been reported in Table 3. This table shows that, in our cases, NPF

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events appear to be initiated within the BL and to be limited by the top of the BL. The lower limit (600 m) of a NPF event has been observed only during the sounding S13 on 18 May. The comparison of soundings performed over the sea (S4, S5, S6, S7 and S8) and over the continent (all the others) shows no clear N_{3-10} vertical trends related to the surface differences. The NPF event observed over the sea during RF43 (S4 and S5) was only detected within the first 50 m of the soundings and the vertical exploration during S11 was not sufficient, thus, these three soundings will not be discussed hereafter.

Independently from the horizontal extension study, three different groups of soundings are defined: Group SI soundings (soundings S1, S3, and S9) corresponds to homogenous events throughout the PBL, Group SII soundings (S2, S6, S8 and S10) corresponds to positive vertical gradients of N_{3-10} with the altitude and Group SIII soundings (S12 and S13) corresponds to negative vertical gradients of N_{3-10} as a function of altitude. During the sounding S7, a special NPF event has been observed, thus S7 will be discussed separately.

The observed meteorological and environmental conditions (air mass origins, temperature, pressure, wind speed, solar radiation) were somewhat similar during Group SI and Group SII soundings. In contrast, both observed soundings with negative vertical gradients of N_{3-10} (Group SIII) were observed after rain events (>0.8 mm), thus related to potential scavenging of aerosol particles and to potential entrainment of pollutant into the lower atmospheric layers. Moreover, the prevailing air mass trajectories, in the BL, of the Group I and Group II soundings were similar and came from South-East while Group III trajectories came from North-East (clean marine air, S12) and South-West (polluted air, S13). Thus, after rainy events, a nucleation event was observed whatever the pollution level of the air mass. Moreover, one can note that the negative vertical gradients were not well-marked compare to the positive ones.

Within the frame of the small number of aircraft soundings performed during the EUCAARI campaign, when nucleation particles have been observed, the classification in different groups yields, that the frequency of positive vertical gradient of N_{3-10} is higher

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than the frequency of observed negative gradients, which would mean that nucleated particles are likely to be increased more frequently in the upper part of the BL. Two processes may explain the presence of ultra-fine particles in this altitude range: (1) particles have been nucleated at the ground and then uplifted before being sampled with the ATR-42 and (2) particles have been nucleated in the upper part of the BL.

To understand which process has been involved in our cases, we first assume that particles have been formed at the ground, calculate the mean diameter that particles should have after being uplifted into the upper part of the BL and compare their mean diameter with the observations. Following the work of Hirsikko et al. (2005), the temporal evolution of ion size distributions, measured with the NAIS instrument at the CBW station, has been utilized to estimate growth rates of newly formed particle populations. To accomplish this estimation, lognormal distributions were fitted to the NAIS size distributions (including the ultra-fine particle mode) as a function of time. After determining the mean diameters corresponding to the peak concentrations in the lognormally fitted ultra-fine mode, the growth rates were obtained by a linear fit through the data points of peak concentrations in that mode with time (Hirsikko et al., 2005). We calculated the growth rates in nm per hour, applying linear fits within following size ranges: 1.5 to 3, 3 to 7 and 7 to 20 nm (Table 3).

The three resulting growth rates for the above size ranges vary from day to day. The total diameter increase ($\Delta D\rho$) is then calculated during vertical transport from the ground to the upper limit of the NPF event, assuming that particle growth rates are similar to the growth rates determined for the measurements at the Cabauw site (Table 4). The time which is needed for freshly formed particles to reach altitudes that has been found by this study to be the upper altitude limit of NPF events has been determined using the observed vertical wind velocity data (W). The mean diameter of uplifted particles have been calculated with the total diameter increase plus initial diameter estimated at 1nm. For three cases (RF50 RF53 and RF54), particles assumed having been newly formed at the ground and vertically transported to the upper limit of the NPF event, have a mean diameter lower than 10nm at this upper limit which is

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consistent with the observed nucleation particles diameter. Thus, for these two cases we cannot rule out that N_{3-10} particle concentrations observed at 1650 m and 1100 m, respectively, might have been solely due to transport of NPF events originating at the ground. However, for at least the other four cases the exclusive transport of freshly formed particles from the surface to higher altitudes cannot be inferred from these considerations. Indeed, the mean diameter of uplifted particles is higher than 10nm thus higher than the observed particle diameter. These latter NPF events have not been limited to the ground, but should have originated from elsewhere in the BL.

During sounding 7, the N_{3-10} concentration abruptly decreased ($20\,000\text{ cm}^{-3}$) in a sub-layer of the BL between 950 m and 1100 m. This particular stratification is related to the presence of two different layers in terms of pre-existing particle concentrations (N_{50}). Whereas in the lower part of the BL ($200\text{m} < z < 950\text{m}$), N_{50} is homogeneous and lower than 1000 cm^{-3} , in the upper part of the BL ($950\text{m} < z < 1800\text{m}$) the N_{50} concentration reached 5000 cm^{-3} . Thus, this sounding illustrates two different NPF events: the first one under almost clean conditions and the second one under polluted conditions.

4 Conclusion

This paper describes six cases of new particle formation (NPF) events observed on board the ATR-42 research aircraft during the EUCAARI intensive measurement campaign in May 2008. All these events were observed at different moments of the day, stages of development and locations (both marine and continental). An extended air mass backtrajectory analysis highlights that the most of air masses in the boundary layer that have been observed during the campaign are coming from the North Sea and Scandinavia. The probability, associated with these air masses, to observe a NPF event with the ATR-42 is in average about 40% and 60% respectively. The transport of clean air masses from the North Sea, which are enriched in sulphate and organic components, to Cabauw (CBW) may have an influence on NPF event over The Netherlands (Kulmala et al., 2001, 2004b; Komppula et al., 2006; Tunved et al., 2006).

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As the nucleation phenomenon was not the major focus of the ATR-42 flights during EUCAARI, the ATR-42 flight plans were not performed with the objective to horizontally bound the NPF events, and therefore ATR-42 results presented here provide a best estimate of the horizontal extension (>70 km) of NPF area. Moreover, the highest particle concentrations in the size range 3–10 nm (N_{3-10} , 10^5 cm $^{-3}$) were observed over the continent whereas the lowest nucleation mode particle concentrations (<5000 cm $^{-3}$) were observed over the North Sea. These limiting cases were observed during the same flight (on 5 May). As the N_{3-10} is highly variable during these six NPF events, four types of NPF event groups (Group I–IV) were defined as a function of air mass origin and geographical location of performed flights. All events observed over the sea (Group I) were related to continental flow regimes. These observations highlight a negative gradient of N_{3-10} from the coastline further to the North Sea. The intensity of the gradient appears to be dependant of the tide conditions and low level (<900 mbar) wind speeds. Over the continent, the N_{3-10} are homogeneous along five flight tracks. One inhomogeneous case is observed during research flight 61 (Group IV events) which were observed under highly polluted conditions. Close or upwind to big cities the number concentration of pre-existent particles greater than 50 nm diameter (N_{50}) is important and inhibits the nucleation process.

During the numerous ascents and descents of the ATR-42, the vertical extension of the NPF events did not exceed the top of the BL and no NPF event was detected in the free troposphere. These results have also been observed for boreal nucleation events (Laaksonen et al., 2008; O'Dowd et al. 2009). Over the continent as well as over the North Sea, the vertical extension of NPF events seems to coincide exactly with the top of the BL for most of the observed cases. Three different types of soundings have been observed: homogeneous (Group SI; three soundings), positive vertical (increasing with altitude) N_{3-10} gradient (Group SII; four soundings) and negative vertical (decreasing with altitude) N_{3-10} gradient (Group SIII; two soundings). The frequency of NPF event occurrences in the higher part of the BL is important (number of positive vertical gradients) as already observed by Venzac et al. (2009). Based on the hypothesis that

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particles may have been nucleated solely at the surface, then transported to higher levels within the BL, particle growth rates have been estimated from simultaneous nucleation findings at the CBW ground station. We conclude that most of the nucleated particle events observed with the ATR-42 have not been formed at the surface but must have nucleated elsewhere in the BL.

This work demonstrates that NPF events observed during ATR-42 flights within EU-CAARI have been limited to the top of the boundary layer but showed differences in terms of horizontal and vertical homogeneity and extension. Newly formed particles evolve to larger sizes within hours, which impacts the energy budget and through direct radiative forcing and aerosol-cloud interactions. Airborne measurements of NPF at different locations around the world would help constrain vertical and horizontal scales allowing better implementation of the nucleation phenomenon to be implemented in existing global model studies (Spracklen et al., 2008; Makkonen et al., 2009).

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Table 1. Overview of ATR-42 flights performed during the EUCAARI campaign: day, times, Flight number and plans of all flights. The prevailing atmospheric flows at low level (850 hPa) and observation flag of new particle formation (NPF) event at Cabauw (from SMPS and NAIS measurements) and on-board the ATR-42 (from SMPS measurements) are added.

Day	Take-off – Landing time (UTC)	Flight type	Flight number	Prevailing atmospheric flow	NPF event	
					Cabauw	ATR-42
02/05/2008	08:40–10:09	RF_type_1	42	West	N	N
05/05/2008	11:04–14:20	RF_type_2	43	South-East	Y	Y
06/05/2008	08:57–10:27	RF_type_2	44	South-East	Y	Y
08/05/2008	11:54–15:24	RF_type_2	45	South-East	Y	Y
09/05/2008	06:04–07:53	RF_type_3	46	South-East	N	N
09/05/2008	11:37–14:57	RF_type_3	47	South-East	Y	N
11/05/2008	09:55–12:45	RF_type_2	48	South-East	Y	Y
13/05/2008	12:55–15:55	RF_type_1	49	South-East	N	N
14/05/2008	12:13–15:05	RF_type_1	50	South-East	Y	Y
15/05/2008	06:19–09:50	RF_type_4	51	South-East	N	N
15/05/2008	11:49–15:06	RF_type_4	52	South-East	N	N
18/05/2008	08:50–11:10	RF_type_5	53	North-West	Y	Y
18/05/2008	12:01–14:45	RF_type_5	54	North-West	Y	Y
19/05/2008	11:06–14:24	RF_type_1	55	North-West	Y	Y
20/05/2008	09:02–11:06	RF_type_1	56	North-West	N	N
21/05/2008	10:07–12:06	RF_type_1	57	North-West	N	N
21/05/2008	13:45–15:14	RF_type_1	58	North-West	N	N
26/05/2008	11:59–15:20	RF_type_5	59	South-West	N	N
28/05/2008	07:54–09:55	RF_type_6	60	South-West	N	N
28/05/2008	13:24–15:40	RF_type_1	61	South-West	Y	Y
29/05/2008	13:15–14:36	RF_type_1	62	South-West	N	Y
30/05/2008	11:26–14:42	RF_type_5	63	South-West	N	N

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Table 2. Tide conditions during the three Type I events observed during three flights.

Date	Flight number	ATR-42 leg times (UTC)	Low tide times (UTC)	High tide times (UTC)	Low sea level (m)	High sea level (m)
05-05-2008	43	12:57–13:06	06:46	12:43	0.00	4.88
08-05-2008	45	14:26–14:47	09:05	14:54	0.26	4.50
11-05-2008	48	12:20–12:41	11:36	17:32	0.91	3.74
18-05-2008	53	08:35–09:10	06:17	12:11	0.33	4.40
18-05-2008	54	13:00–13:51	18:31		0.52	

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Table 3. Horizontal and vertical extension of all detected new particle formation (NPF) events. In addition to the vertical extension of NPF events is presented the top of the boundary layer (BL), both derived from numerous ATR-42 ascents and descents.

Flight number	Soundings	Sounding start and end times	Sounding zone	BL height (m) limit (m)	NPF event upper limit (m)	NPF event lower (km)	Horizontal extension
43	S1	11:12–11:31	Continent	1400	1325	–	>100
	S2	11:34–11:51	Continent	1100	1100	–	
	S3	12:03–12:11	Continent	1450	1550	–	
	S4	12:46–12:56	North Sea	1850	300	–	
	S5	13:08–13:24	North Sea	1500	300	–	
	S6	13:48–14:03	North Sea	1100	1100	–	
45	S7	11:54–12:09	North Sea	1750	1800	–	>100
	S8	13:11–13:29	North Sea	1850	1850	–	
48	S9	10:29–10:49	Continent	1500	1450	–	>160
50	S10	12:16–12:55	Continent	1700	1650	–	>70
53	S11	08:30–09:20	Continent	1900	1080	–	>150
54	S12	12:01–12:11	Continent	2100	2100	600	>150
61	S13	13:40–14:30	Continent	1350	1400	–	~100

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Table 4. Mean vertical component of the wind (W , m s^{-1}) observed during all detected new particle formation (NPF) events. The estimated median growth rates for nucleation mode charged particles in size ranges 1.5–3, 3–7 and 7–20 nm. Calculations are based on ion size distributions measured at the Cabauw field site during the flights. Estimation of the minimum diameter increase (ΔD_p) assuming that the particle is formed at the surface site and is transported to the top of the boundary layer. Thus, ΔD_p is calculated using the growth rate, estimated with the NAIS measurements at the Cabauw field site, the mean vertical wind speed (W) observed in the BL and the height of the BL given in Table 2 ($\Delta D_p = \text{GR} \cdot W \cdot \text{BL}$).

Date (dd-mm-yyyy)	Flight number	W (m s^{-1})	GR median 1.5–3 nm [nm/h]	GR median 3–7 nm [nm/h]	GR median 7–20 nm [nm/h]	ΔD_p (nm)
05-05-2008	43	0.0155	3.94	10.64	7.37	204.5 ± 2.5
08-05-2008	45	0.1572	5.92	6.94	5.93	19.9 ± 2
11-05-2008	48	0.0288	3.04	3.32	5.12	68.5 ± 1.5
14-05-2008	50	0.5362	6.54	7.58	6.99	6.2 ± 2.1
18-05-2008	53	0.4314	2.40	7.99	13.06	2.2 ± 1.8
18-05-2008	54	0.4614	2.40	7.99	13.06	7.3 ± 5.2
28-05-2008	61	0.3811	7.17	21.82	14.25	14.5 ± 3.5

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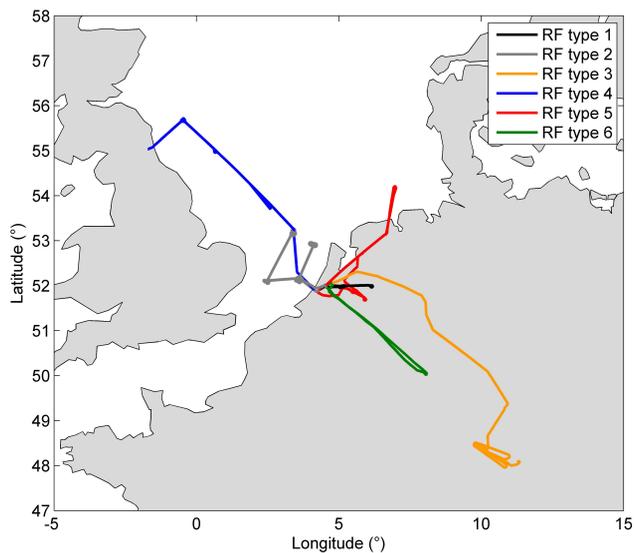


Fig. 1. Different types of research flight (RF) plans performed during the EUCAARI intensive observation campaign in May 2008.

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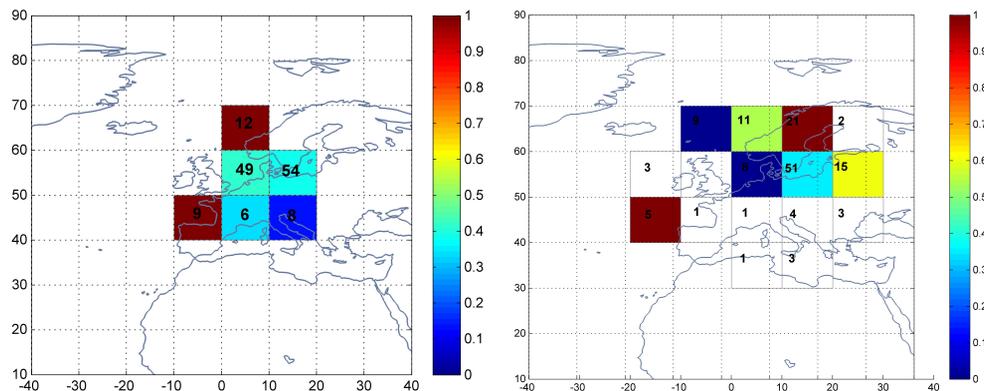


Fig. 2. Frequencies (colour coded from 0-1) of observed particle formation as a function of air mass origins (squares) determined for two chosen back-trajectory calculation times of 24 h **(a)** and 48 h **(b)**. The numbers associated to each square denote the total number of calculated air mass origins assigned to the complete set of EUCAARI flights horizontally divided into equidistant flight segments, for which calculations have been performed.

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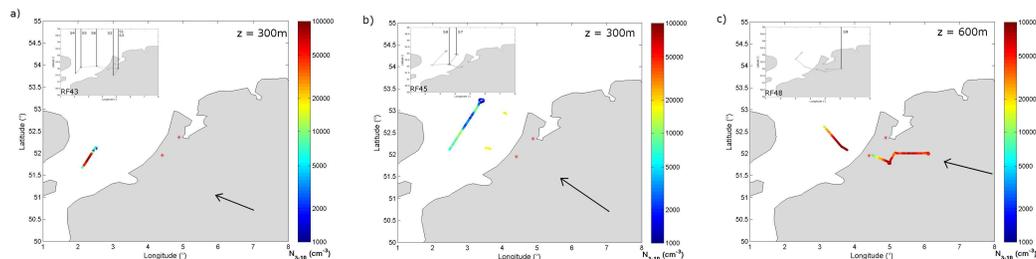


Fig. 3. Horizontal cross sections showing nucleation mode aerosol concentrations of Group I events within the size range of 3–10 nm along the ATR-42 flight trajectories during research flights **(a)** RF43 on 5 May, **(b)** RF45 on 8 May, and **(c)** RF48 on 11 May. Only N_{3-10} at a constant flight leg altitude, indicated on each figure, are represented. The black arrows represent the mean direction of the air mass (deduced from back-trajectories calculated with Hysplit (<http://www.ready.noaa.gov/ready/open/hysplit4.html>)). The vertical black lines refer to the ATR-42 sounding numbers as they appear in Fig. 7.

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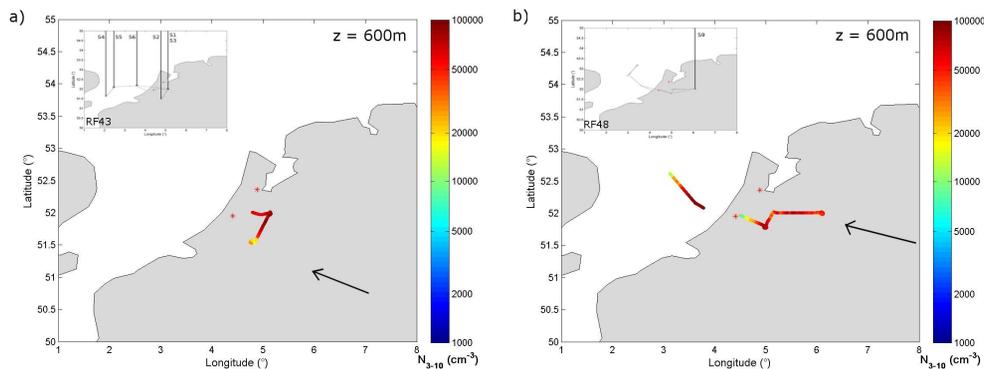


Fig. 4. As for Fig. 3 for Group II events **(a)** research flights RF43 on 5 May, and **(b)** RF48 on 11 May.

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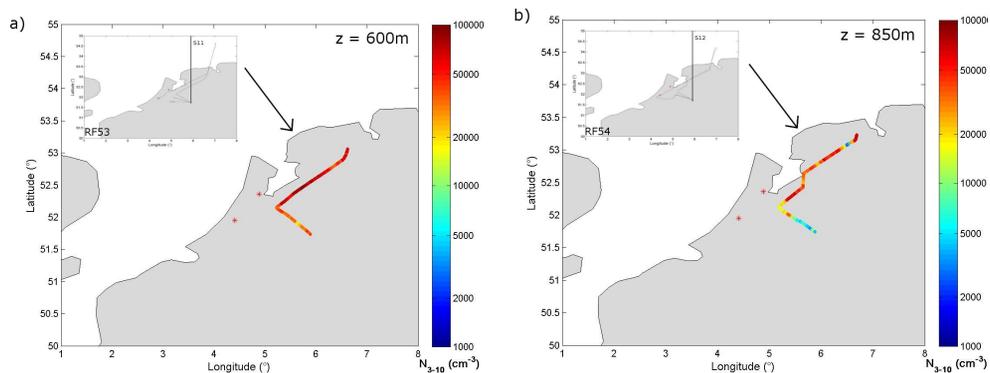


Fig. 5. As for Fig. 3 for Group III events **(a)** RF53 on 18 May and **(b)** RF54 on 18 May.

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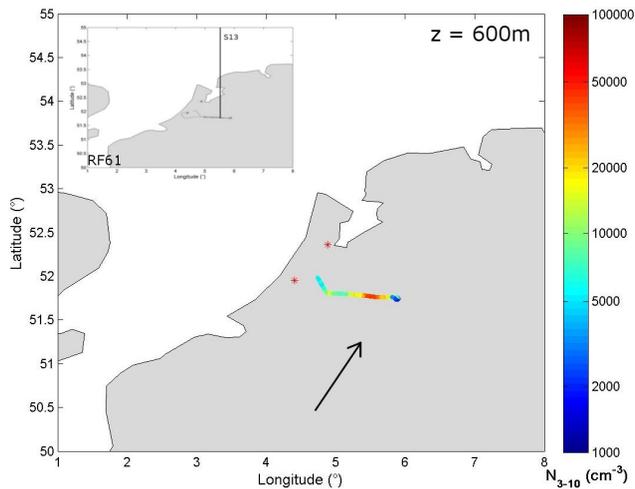


Fig. 6. As for Fig. 3 for research flight RF61 on 28 May (Group IV event).

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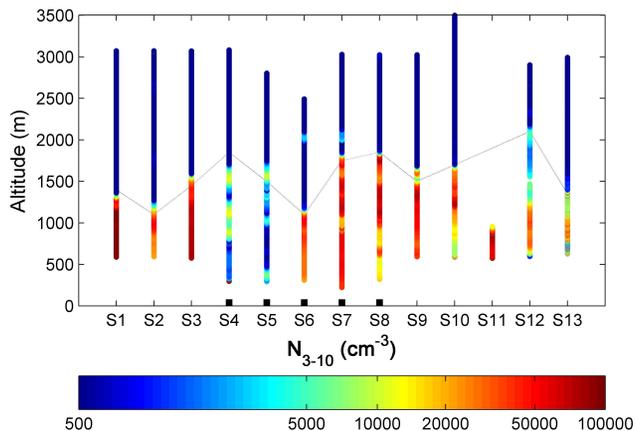


Fig. 7. Profile of the particle concentrations within the size range of 3–10 nm during the ATR-42 soundings for the six flights when NPF events were observed: RF43 (S1, S2, S3, S4, S5, S6) on 5 May, RF45 (S7, S8) on 8 May, RF48 (S9) on 11 May, RF50 (S10) on 14 May, RF53 (S11) on 18 May, RF54 (S12) on 18 May, and RF61 (S13) on 28 May 2008. The black rectangles highlight soundings performed above the sea. The grey line corresponds to the top of the BL as described in Table 3.

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