

Formation and growth of atmospheric nanoparticles in the eastern Mediterranean: Results from long-term measurements and process simulations

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1 Abstract

2 Atmospheric New Particle Formation (NPF) is a common phenomenon all over the world. In
3 this study we present the longest time series of NPF records in the eastern Mediterranean
4 region by analyzing ten years of aerosol number size distribution data obtained with a mobility
5 particle sizer. The measurements were performed at the Finokalia environmental research
6 station on Crete, Greece during the period June 2008-June 2018. We found that NPF took
7 place 27% of the available days, undefined days were 23% and non-event days 50%. NPF is
8 more frequent in April and May probably due to the terrestrial biogenic activity and is less
9 frequent in August. Throughout the period under study, nucleation was observed also during
10 the night. Nucleation mode particles had the highest concentration in winter and early spring,
11 mainly because of the minimum sinks, and their average contribution to the total particle
12 number concentration was 8%. Nucleation mode particle concentrations were low outside
13 periods of active NPF and growth, so there are hardly any other local sources of sub-25 nm
14 particles. Additional atmospheric ion size distribution data simultaneously collected for more
15 than two years period were also analyzed. Classification of NPF events based on ion
16 spectrometer measurements differed from the corresponding classification based on a
17 mobility spectrometer, possibly indicating a different representation of local and regional NPF
18 events between these two measurement data sets. We used the MALTE-box model for
19 simulating a case study of NPF in the eastern Mediterranean region. Monoterpenes
20 contributing to NPF can explain a large fraction of the observed NPF events according to our
21 model simulations. However the adjusted parameterization resulting from our sensitivity tests
22 was significantly different from the initial one that had been determined for the boreal
23 environment.

24 1) Introduction

25 Most of the atmospheric aerosol particles, and a substantial fraction of particles able to act as
26 cloud condensation nuclei (CCN), have been estimated to originate from new particle
27 formation (NPF) taking place in the atmosphere (Spracklen et al. 2006; Kerminen et al., 2012;
28 Gordon et al., 2017). The exact mechanisms driving atmospheric NPF and subsequent particle
29 growth processes are still not fully understood, nor are the roles of different vapors and ions
30 in these processes (Kulmala et al., 2014; Lehtipalo et al., 2016; Tröstl et al., 2016). In order to
31 understand how aerosol particles affect regional and global climate and air quality, it is
32 necessary to quantify the factors that determine the occurrence of NPF and characterize the

1 parameters that describe the strength of NPF, such as the new particle formation and growth
2 rates, in various environments.

3 While NPF has been reported to take place worldwide (Kulmala et al., 2004a; Wang et al.,
4 2017), observational studies on this subject are scarce in rural sub-tropical environments. It
5 has been shown that the processes responsible for particle formation and growth differ
6 substantially across the European continent (Dall'Osto et al., 2018).

7 Several studies have investigated NPF in eastern Mediterranean and found it to be a frequent
8 phenomenon. Lazaridis et al. (2006) first reported NPF at the area and correlated these events
9 to polluted air masses. Petäjä et al. (2007) presented NPF in Athens metropolitan area and
10 showed that under the influence of urban pollution, condensing species leading to growth of
11 the new particles are far more hygroscopic than under cleaner conditions. NPF events have
12 also been reported to be frequent at the urban environment of Thessaloniki (Siakavaras et al.,
13 2016). Kalivitis et al. (2008) showed that precursors and nucleation mode particles experience
14 strong scavenging on Crete island during summer. Pikridas et al. (2012) suggested that
15 nucleation events occurred only when accumulation mode particles were neutral, being
16 consistent with the hypothesis that a lack of NH_3 , during periods when particles are acidic,
17 may limit nucleation in sulfate-rich environments such as the eastern Mediterranean.
18 Additionally, based on ion observations, Pikridas et al. (2012) showed that NPF is more
19 frequent in winter. By using the same data set from eastern Mediterranean, Kalivitis et al.
20 (2012) reported night-time enhancements in ion concentrations with a plausible association
21 with NPF, being among the very few locations where such observations have been made.
22 Manninen et al. (2010) presented an analysis of a full year of observations of NPF with
23 atmospheric ion spectrometers at various locations across Europe during the EUCAARI project
24 and showed that NPF is less frequent in the eastern Mediterranean site than in other, mostly
25 continental, European sites. On the other hand, Berland et al. (2017) showed that similar
26 patterns are being observed throughout the Mediterranean when comparing observations
27 from the island of Crete to a western Mediterranean site in terms of the frequency of
28 occurrence, seasonality, and particle formation and growth rates. Kalivitis et al. (2015) studied
29 for the first time the NPF-CCN link using observations of particle number size distributions,
30 CCN and high-resolution aerosol chemical composition for the eastern Mediterranean
31 atmosphere. From the hygroscopicity of the particles in different size fractions, it was
32 concluded that smaller particles during active NPF periods tend to be less hygroscopic (and
33 richer in organics) than larger ones. Finally, Kalkavouras et al. (2017) reported that NPF may

1 result in higher CCN number concentrations, but the effect on cloud droplet number is limited
2 by the prevailing meteorology.

3 In this work, we present results from the analysis of ten years of aerosol particles number size
4 distributions and more than two years of atmospheric ion size distributions, representing one
5 of the longest published NPF data set in the Mediterranean atmosphere. The main questions
6 we wanted to address were: 1) How often does NPF take place in eastern Mediterranean,
7 what are the characteristics of this phenomenon and to what extent has it changed over the
8 period under study? 2) Are there features in NPF observed at the study area that are not
9 common in other locations? and 3) How well can numerical models, used in different
10 environmental conditions, represent NPF in this subtropical environment?

11 2) Materials and methods

12 2.1) Measurements

13 Measurements presented in this work were carried out at the atmospheric observation
14 station of the University of Crete at Finokalia, Crete, Greece (35°20'N, 25°40'E, 250m a.s.l.)
15 over ten years, between June 2008 and June 2018. The Finokalia station
16 (<http://finokalia.chemistry.uoc.gr/>) is a European supersite for aerosol research, part of the
17 ACTRIS (Aerosols, Clouds, and Trace gases Research Infrastructure) Network. The station is
18 located at the top of a hill over the coastline, in the north east part of the island of Crete
19 (Mihalopoulos et al., 1997). The station is representative for the marine background
20 conditions of eastern Mediterranean (Lelieveld et al., 2002), with negligible influence by local
21 anthropogenic sources. The nearest major urban center in the area is Heraklion with
22 approximately 200 000 inhabitants, located about 50 km to the west of the station.

23 In order to monitor the NPF events, a TROPOS type custom-built Scanning Mobility Particle
24 Sizer (SMPS), similar to IFT-SMPS in Wiedensohler et al. (2012), was used at Finokalia. Particle
25 number size distributions were measured in the diameter range of 9–848 nm every five
26 minutes. The system was a closed-loop, with a 5:1 ratio between the aerosol and sheath flow
27 and it consisted of a Kr-85 aerosol neutralizer (TSI 3077), a Hauke medium Differential Mobility
28 Analyzer (DMA) and a TSI-3772 Condensation Particle Counter (CPC). The sampling was made
29 through a PM_{10} sampling head and the sample humidity was regulated below the relative
30 humidity of 40% with the use of Nafion® dryers in both the aerosol and sheath flow. The
31 measured number size distributions were corrected for particle losses by diffusion on the
32 various parts of the SMPS according to the methodology described in Wiedensohler et al.

(2012). Three different types of calibration were performed for the SMPS, DMA voltage supply calibration, aerosol and sheath flows calibrations and size calibrations. These measurements have been performed at Finokalia on a continuous basis since 2008. The instrument used at Finokalia was audited on-site with good results in the framework of EUSAAR (European Supersites for Atmospheric Aerosol Research) project (<http://www.wmo-gaw-wcc-aerosol-physics.org/audits.html>) and has successfully passed twice laboratory intercomparison workshops (2013 and 2016, reports available at <http://www.wmo-gaw-wcc-aerosol-physics.org/instrumental-workshops.html>) in the framework of ACTRIS project. The instrument has been operated following the recommendations described in Wiedensohler et al. (2012). Additional information for newly formed particles were obtained with the use of an Air Ion Spectrometer (AIS- AIREL Ltd., Institute of Environmental Physics, University of Tartu, Estonia, Mirme et al., 2007). AIS is a cluster ion air spectrometer used to simultaneously measure electrical mobility distribution of positive and negative air ions (mobilities in the range of 2.4 to $1.3 \cdot 10^{-3} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$). The mobility distributions were then transformed to size distributions in the size range 0.8-42 nm. The number counting threshold was approximately 10 cm^{-3} and the uncertainties of the AIS measurements were $\sim 10\%$ for negative and positive ion concentrations and $\sim 0.5 \text{ nm}$ in size (Manninen et al., 2010). The diameter of the AIS inlet tube was 35 mm and the sample flow rate was 60 L m^{-1} . The time step of the measurements was five minutes.

These measurements have been used to identify NPF for the whole period and provide a historical perspective for the frequency and the characteristics of NPF phenomena in the eastern Mediterranean. Calculations for formation rates of new particles (J), growth rates (GR) in various size ranges and condensation sink (CS) were made according to Kulmala et al. (2012). Formation rates of particles with diameter D_p (in this study $D_p=9 \text{ nm}$) were calculated as:

$$J_{D_p} = \frac{\Delta N_{D_p}}{\Delta t} + CoagS \cdot N_{D_p} + \frac{GR}{\Delta D_p} \cdot N_{D_p} + S_{losses} \quad (1)$$

ΔN_{D_p} is the increase in nucleation mode particles' number concentration ($D_p < 25 \text{ nm}$), $CoagS$ is the coagulation of 9-nm particles on larger particles, GR is the growth rate in the size range 9-25 nm. S_{losses} takes into account additional losses and was neglected in this study. GR was calculated using the mode-fitting method (Dal Maso et al., 2005). The aerosol size distributions were fitted with lognormal distributions and the nucleation mode geometric

mean diameter was plotted as a function of time. *GR* was calculated as the slope of the linear fit so that:

$$GR = \frac{dD_p}{dt} (2)$$

CS is the condensation sink caused by the pre-existing aerosol population and was calculated using the properties of sulfuric acid as condensing vapor.

All important meteorological parameters were monitored every five minutes using an automated meteorological station, including the temperature, wind velocity and direction, relative humidity, solar irradiance and precipitation. Ozone concentrations were measured with a TEI 49C instrument and nitrogen oxides with a TEI 42CTL, both commercially available, with a time step of five minutes.

2.2) NPF simulations with the MALTE-Box model

The simulations of NPF events in the eastern Mediterranean atmosphere were here performed using the MALTE-box model of the University of Helsinki. This 0-d model able to simulate aerosol dynamics and chemical processes has successfully reproduced observations of aerosol formation and growth in the boreal environment (Boy et al., 2006) as well as in highly polluted areas (Huang et al., 2016). For the present study, chemical reactions relevant to the production of condensing species from the Master Chemical Mechanism (MCM) were incorporated in the MALTE-box chemical mechanism, as described in Boy et al. (2013). These include the full MCM degradation scheme of the following volatile organic compounds (described in more detail in Tzitzikalaki et al., 2017): C₁-C₄ alkanes, C₂-C₃ alkenes, acetylene, isoprene, α - and β -pinene, aromatics, methanol, dimethyl sulfide, formaldehyde, formic and acetic acids, acetaldehyde, glycoaldehyde, glyoxal, methylglyoxal, acetone, hydroxyacetone, butanone and marine amines. The Kinetic PreProcessor (KPP) was used to produce the Fortran code for the calculations of the concentrations of each individual compound (Damian et al., 2002), except for those species whose concentrations were manually input from large scale model simulations.

The major aerosol dynamical processes for clear sky atmosphere were simulated by the size-segregated aerosol model UHMA (University Helsinki Multicomponent Aerosol Model, Korhonen et al., 2004) impended in the MALTE-Box model. Measured aerosol number size distributions were used to initialize UHMA daily, which simulates NPF, coagulation, growth and dry deposition of particles. UHMA simulated new cluster formation using the activation

1 nucleation parameterization, so that the nucleation rate has a linear relationship with sulfuric
2 acid concentration, depending on the nucleation coefficient K_{act} .

3 Apart from sulfuric acid, about 20 extremely low-volatility organic compounds (ELVOCs) and
4 7 selected semi-volatile organic compounds (SVOCs) were treated as condensing vapors,
5 following the simplified chemical mechanism presented in Huang et al. (2016). All condensing
6 compounds were treated either as sulfuric acid or organic compounds and the condensation
7 of organic vapors was determined by the nano-Kohler theory (Kulmala et al., 2004b).

8 As input to the MALTE-Box model were used the observations at Finokalia station and when
9 such observations were not available, the results from numerical simulations with the global
10 3-dimensional chemistry transport model (CTM) TM4-ECPL (Daskalakis et al., 2015, 2016;
11 Myriokefalitakis et al., 2010, 2016) for Finokalia. Observational data include temperature,
12 relative humidity, total radiation (meteorological input), ozone (O_3) and nitrogen oxides (NO_x)
13 concentrations as well as aerosol number size distributions. The aerosol number size
14 distribution measured by the SMPS was used to calculate the condensation sink for H_2SO_4
15 vapors. Due to the lack of detailed measurements of VOC at Finokalia, as a first approximation,
16 biogenic and anthropogenic concentrations of all the above mentioned VOCs resolved every
17 3 hours were taken from the TM4-ECPL model.

18 The global TM4-ECPL model was run driven for this study by ECMWF (European Centre for
19 Medium – Range Weather Forecasts) Interim re-analysis project (ERA – Interim) meteorology
20 (Dee et al., 2011) of the year 2012 at an horizontal resolution of 3° in longitude x 2° in latitude
21 with 34 vertical layers up to 0.1 hPa. The model used year-specific meteorology and emissions
22 of trace gases and aerosols. For this study, that of the year 2012 was used, except for soil NO_x
23 and oceanic CO and VOCs emissions which were taken from POET inventory database for the
24 year 2000 (Granier et al., 2005). TM4-ECPL simulations for this work were performed with a
25 model time-step of 30 min, and the simulated VOC concentrations every 3-hours were used
26 as input to MALTE box model; SO_2 surface levels at Finokalia were taken from Monitoring
27 Atmospheric Composition and Climate (MACC) data assimilation system (Inness et al., 2013).

28 For the calculations of the photo-dissociation rate coefficient by the MALTE-Box model, the
29 solar actinic flux (AF) is needed. Unfortunately, AF was not measured at Finokalia in 2012,
30 therefore AF levels were calculated by the Tropospheric Ultraviolet and visible Radiation
31 Model (TUV, Madronich, 1993) version v.5 for cloud free conditions. The ability of TUV to
32 calculate the AF at Finokalia was investigated by comparing observations of photo dissociation

1 rates of O_3 (JO^1D) and NO_2 (JNO_2) and model calculations. The measurements of these photo
2 dissociation rates were performed by filter radiometers (Meteorologie Consult, Germany).
3 The JO^1D was measured at wavelengths <325 nm, while for JNO_2 wavelengths <420 nm were
4 used.

5 A series of sensitivity tests of AF to different input parameters was also performed to optimize
6 the calculations. The model uses extra-terrestrial solar spectral irradiance (200-1000 nm by
7 0.01nm steps) and computes its propagation through the atmosphere taking into account
8 multiple scattering and the absorption and scattering due to gases and particles. TUV inputs
9 of interest were surface reflectivity (albedo), O_3 column, Aerosol Optical Depth at 500 nm
10 (AOD), Single Scattering Albedo of aerosol (SSA), NO_2 column, air density. Total O_3 column
11 values were taken from the Ozone Monitoring Instrument (OMI) on the Aura spacecraft of
12 NASA (Levelt et al., 2006). Aerosol columnar optical properties were obtained from the
13 Aerosol Robotic Network (AERONET). AOD data were measured at the FORTH_Crete station
14 which is located 35 km west of Finokalia (Fotiadi et al., 2006). Data level 1.5 from Version 2
15 were used (cloud-screened). Total NO_2 column values were taken from GOME2 and OMI
16 satellites. The calculations were carried out at wavelength from 280 to 650 nm with a
17 resolution of 5 nm. Simulations using surface reflectivity of 0.075 and simulation using O_3
18 column taken from OMI had the best correlation with measurements. However, the TUV
19 model still significantly overestimated JO^1D and JNO_2 data. Thus, a parameterisation took
20 place following a simple empirical approach, according to Mogensen et al. (2015) and the
21 ratios between the measured and modelled (from TUV) photolysis rate were calculated and
22 used in the model.

23 3) Results and discussion

24 3.1) Particle size distribution and its connection with NPF

25 We analyzed all available measurements of number size distributions of atmospheric aerosol
26 particles measured at Finokalia in order to identify and analyze the NPF phenomenon in the
27 eastern Mediterranean. The data coverage for the period 2008-2018 was 82 %, providing one
28 of the longest time series of size distributions not only in this region but also in the southern
29 Europe and a unique data base for aerosol physical properties.

30 First, we calculated the total particle number concentration (median concentration was
31 2202 cm^{-3} , standard deviation (SD) 528 cm^{-3}) and corresponding number concentration in the
32 nucleation mode ($D_p < 25\text{ nm}$, median 80 cm^{-3} , SD 528 cm^{-3}), Aitken mode ($25\text{ nm} < D_p < 100\text{ nm}$,

median 1028 cm^{-3} , SD 894 cm^{-3}) and accumulation mode ($D_p > 100 \text{ nm}$, median 898 cm^{-3} , SD 605 cm^{-3}). We found that Aitken mode accounted for 50% and accumulation mode 42% of the total particle number concentration, while the nucleation mode accounted only for 8%. The standard deviation of the nucleation particle number concentration was 528 cm^{-3} , indicating that the abundance of these smallest particles is of episodic nature. The highest monthly average concentrations of nucleation mode particles were observed during winter and early spring and the lowest ones during summer (Fig. 1a). Calculating the median diurnal variability of the nucleation mode, we can see that there is a clear pattern for all seasons of the year (Fig. 2a) with a sudden burst in the number concentration around noon that is most pronounced in winter and least in summer. Such an observation suggests that the nucleation particle number concentration is controlled by NPF episodes rather than other sources such as combustion processes. As can be seen in Fig. 2b where a typical “banana shaped” pattern of an NPF event at Finokalia is presented, the sudden burst at noon is typical for a NPF event. In summer, nucleation mode particles have the highest concentrations during the night, yet another concentration relative maximum at noon can be attributed to NPF (Fig. 2a). The shift in the average time of the daytime burst of nucleation mode particles can be attributed to the annual variation of the daylight length. Similar observations to ours have been reported in Cusack et al. (2013) for the western Mediterranean where the diurnal variation of nucleation mode particles presents a clear maximum at noon under both polluted and clean conditions.

It is worth noticing that during night-time the median nucleation mode particle number concentrations were similar in all the seasons. This suggests that there is some new particle production mechanism at night, especially in summer and autumn, that operates separately from daytime NPF. Frequently during the night-time, we observed a pronounced appearance of new nucleation mode particles over several hours as illustrated by Fig. 3. While nocturnal NPF has been reported in the literature (see Salimi et al. (2017) and references therein), this phenomenon seems to be rare and it remains unclear what are the exact mechanisms leading to it. Given that we observed no or little growth during nighttime NPF, we may assume that the sources leading to the formation of new particles are local rather than regional and that the lack of photochemistry during night limits the abundance of condensable vapors driving particle growth. Observations of very localized NPF have been reported in Mace Head, Ireland, where intense NPF frequently takes place under low tide conditions when algae are exposed to the atmosphere (O’Dowd et al., 2002). Henceforth, we will exclude the nighttime NPF events from our further analysis. We refer the interested reader to Kalivitis et al. (2012) for a more detailed description of this phenomenon.

Overall, we observed atmospheric NPF to take place during both day and night at Finokalia, but no sign of any other source of nucleation mode particles in measured air masses. We therefore hypothesize that atmospheric NPF is the dominant source of nucleation mode particles in this Mediterranean environment.

3.2) Characteristics of NPF in the eastern Mediterranean

We analyzed the dataset of aerosol size distributions following the approach of Dal Maso et al. (2005) in order to mark the available days as 1) NPF event days when a clear new nucleation mode and subsequent growth of newly-formed particles to larger diameters can be observed, 2) non-event days and 3) undefined days when either new particles appear into the Aitken mode or nucleation mode particles do not show a clear growth. The available days were manually inspected and classified.

We used the Statistica software package for Windows to carry out factor analyses, including meteorological parameters, ozone concentrations (as an important oxidant in the atmosphere) and PM_{10} mass concentration (as an index of particulate pollutant levels), in order to examine whether any of these factors were associated with the formation of new particles, represented by the nucleation mode number concentration. Furthermore, we divided our data to night and day time periods in order to separate daytime NPF from that taking place during nighttime. The only parameter that had some effect on the nucleation mode particle number concentration was the wind velocity: when strong winds were prevailing at Finokalia, it was more unlikely to observe nucleation particles. On the other hand, the lack of correlation to any other parameter may indicate that the NPF is not sensitive to local meteorological conditions, preexisting particulate matter and ozone levels in this environment. Air mass back trajectories calculated using the HYSPLIT model showed little difference during NPF events from air masses typical for the prevailing situation at Finokalia: air masses arriving at Finokalia from the northeast were the most frequent during NPF events (30% against 24% of all days), followed by northern directions (20% against 21%) and northwestern air masses that were more frequent than the average (19% against 17%).

Next, we focused on determining the main characteristics of daytime NPF at Finokalia. Overall, 837 NPF events were identified. This is one of the longest time series of the NPF phenomenon recorded in the Mediterranean atmosphere, providing a representative climatology of NPF events in this region. NPF took place 27% of the 3057 available measurement days whereas no event occurred on 50% of those days. It is worth noting that 23% of the days were

1 characterized as undefined, which means that while no clear NPF event could be observed,
2 there was some evidence of secondary particle formation although not at the immediate
3 vicinity of the station (Table 1). We found that NPF is most frequent in April and May, probably
4 due to the biogenic activity and the onset of intense photochemistry, and least frequent in
5 August (Fig. 4) probably due to high wind speeds occurring these month (not shown) and
6 additionally the high Condensational Sink (Fig. 1b). Rain season in southeastern Europe in
7 early autumn leads to gradual CS decrease, and as a result a local maximum in NPF frequency
8 is observed in October. NPF at Finokalia takes place throughout the year.

9 As a next step, we classified the NPF events into Class I or Class II events depending on whether
10 the particle formation rate at 9 nm (J_9) and growth rates from 9 to 25 nm diameter (GR_{9-25})
11 could be calculated with a good confidence or not respectively. Overall, Class I events
12 corresponded to 8% of the available measuring days and 28% of the event days, and they were
13 observed throughout the year, providing enough data for a statistical analysis of particle
14 formation and growth rates during NPF events (Fig. 5).

15 The average value of J_9 during the Class I NPF events in Finokalia was $0.9 \text{ cm}^{-3} \text{ s}^{-1}$ (median 0.5
16 $\text{cm}^{-3} \text{ s}^{-1}$, SD $1.2 \text{ cm}^{-3} \text{ s}^{-1}$). This is well in the range of values reported for J_{10} in other locations
17 (Kulmala et al., 2004a), higher though than J_{16} reported by Berland et al. (2017) at the Finokalia
18 site in 2013 ($0.26 \text{ cm}^{-3} \text{ s}^{-1}$), but substantially lower than the values found by Kopanakis et al.
19 (2013) in western Crete ($13.1 \pm 9.9 \text{ cm}^{-3} \text{ s}^{-1}$). The monthly variation of J_9 (Fig. 6a) shows that
20 the highest average formation rates were observed in December and January, probably as a
21 result of the low CS values observed in winter, although it is difficult to say which factors
22 determine the monthly variability of J_9 at Finokalia. Seasonal averages of J_9 , GR_{9-25} and CS are
23 summarized in Table 2. Moreover, we found that J_9 and N_{9-25} have a clear linear relation (Fig.
24 7), which supports our earlier hypothesis that at Finokalia the main source of nucleation mode
25 particles is their secondary formation in the atmosphere.

26 We calculated the average growth rate of the newly formed particles to be 5.4 nm hr^{-1} (median
27 4.5 nm hr^{-1} , SD 3.9 nm hr^{-1}). We found that GR_{9-25} is highest in summer until September and
28 lowest in winter and early spring, probably in line with the seasonal cycle of photochemical
29 activity and biogenic emission patterns, producing condensable species that are driving the
30 growth process (Fig. 6b). Additionally, transported pollution in summer at Finokalia may
31 contribute except of CS to GR as well, since transported anthropogenic SO_2 is a precursor for
32 condensable sulfuric acid.

The survival probability of newly-formed particles is closely related to the ratio of *CS* to *GR*, at least for cluster sizes (Kerminen and Kulmala, 2002; Kulmala et al., 2017) and at Finokalia they present the same annual cycle. The survival probability for nucleation mode particles for Class I events was calculated based on the formula in Kulmala et al. (2017). It was found that on a seasonal basis the median survival probability is higher in summer and winter, however varies between the seasons only within 5%. The concentrations of nucleation mode particles are lower during summer and the average duration of the NPF in summer seems to be shorter as shown in Fig. 1 and 2a respectively. These observations may be explained by the higher *CS* and *GR* during summer. The *CS* (and hence *CoagS*) may directly affect the maximum concentrations observed. The slightly higher survival probability in summer perhaps explains that given high *CS* values, new particles need to grow fast in order to survive. On the other hand, one would expect NPF to be most frequent in winter when the highest concentrations of nucleation particles are observed and *CS* is the lowest, however this was not the case. A possible explanation for the high nucleation mode particle number concentrations in winter could be that the survival probability is higher than in spring or autumn.

3.3) NPF trends during the 2008-2018 period

During the period under study no statistically significant trends in NPF events were observed at Finokalia for the 120 available months. It should be noted though, that since 2010 a decreasing trend is observed, which is statistically significant with a p-value of 0.005. During the measurement period under study, no trend in J_9 was observed (Fig. 8c). Although no statistically significant trend was observed for GR_{9-25} as well (Fig. 8d), we observed a decreasing trend during the period 2008-2015 of about $0.3 \text{ nm hr}^{-1} \text{ yr}^{-1}$. This trend can be considered statistically significant (p-value of 0.03). In order to explain this trend, we need to emphasize the regional characteristics of the observations at Finokalia, as this site is greatly affected by long-range transported pollutants of marine, desert dust and polluted continental origin (Lelieveld et al., 2002). Non-sea salt sulfate (nss-SO_4^{2-}) can be considered as an indicator of regional pollution from anthropogenic activities (SO_2 emissions), and since the beginning of the economic crisis in Europe, especially in Greece, we observed a clear decline in its concentration since 2008 (Paraskevopoulou et al., 2015) which however has stopped after 2015. We can therefore assume also a regional decrease in SO_2 emissions, since the main source of SO_2 at Finokalia is attributed to transported pollution (Sciare et al., 2003). This could

1 result in a decrease in the availability of sulfuric acid, a major condensable species responsible
2 for the particle growth (Bzdek et al., 2012).

3 Hamed et al. (2010) studied the effect of the reduction in anthropogenic SO₂ emissions in
4 Germany between the years 1996-97 and 2003-06 as a result of the socio-economic changes
5 in East Germany after the reunification. They observed a notable decrease in the NPF event
6 frequency but an increase in the growth rate of nucleated particles. A decrease in the NPF
7 frequency due to the reduction of anthropogenic SO₂ emissions in eastern Lapland was also
8 reported by Kyrö et al. (2014), and this decrease was most pronounced for the Class I NPF
9 events. Nieminen et al. (2014) analyzed the longest data set reported in literature from
10 Finland and found that, despite major decreases in ambient SO₂ concentrations observed all
11 over Europe as a result of overall air quality improvements, there was a slight upward trend
12 in the particle formation and growth rates. This feature was attributed partly to increased
13 biogenic emissions over the same period.

14 In our case the reasons for the variations in the NPF frequency, J_9 and GR_{9-25} remain unclear,
15 even though factors like meteorological conditions and organic vapor abundance have
16 probably played some role in this respect.

17 3.4) Atmospheric ion observations related to new particle formation

18 At the Finokalia station, atmospheric ion observations relevant to new particle formation were
19 performed during two separate periods, 2008-2009 during the EUCAARI project (Manninen et
20 al., 2010) and 2012-2014 during the FRONT (Formation and growth of atmospheric
21 nanoparticles) project. Here we will focus only on FRONT data, since the EUCAARI dataset is
22 discussed in detail in Manninen et al. (2010) and Pikridas et al., (2012). A typical nucleation
23 event is presented in Fig. 9 as recorded by both the AIS and SMPS. AIS observations may
24 provide information about the initial stages of new particle formation as particles can be
25 observed emerging in the intermediate ion diameter range 1.6-7.4 nm. Intermediate ions
26 appear only under certain circumstances, such as during precipitation, at high wind speeds,
27 and when NPF is taking place (Hörrak et al. 1998; Tamm et al., 2014; Leino et al., 2016; Chen
28 et al., 2017). In the following we will focus on NPF and use only the observations from the
29 negative polarity due to the better representation of NPF events in those data compared with
30 corresponding positive ions in our dataset (Kalivitis et al., 2012).

31 We classified all of the available AIS measurement days into event, non-event and undefined
32 days, once again according to methods introduced by Dal Maso et al. (2005), and subsequently

1 compared the findings from AIS data to those from the SMPS. In Fig. 9 an NPF event is
2 presented observed with both the AIS and the SMPS at Finokalia. Surprisingly, the two data
3 sets for the same time period gave quite different results in terms of the NPF event frequency:
4 in the AIS data the NPF event frequency peaked earlier during the year than in the SMPS data
5 (Fig. 10). This feature was evident in both periods of AIS measurements and has been also
6 reported at a rural site in Hungary (Yli-Juuti et al., 2009), probably because AIS detects only
7 naturally charged particles while SMPS detects all particles. Additionally, it is possible that AIS
8 data are more representative of local NPF events with limited particle growth, and such events
9 may not be seen in the SMPS data. On the other hand, the SMPS measures neutral particles
10 but has a much higher detection limit (9 nm), so its data may be more representative of
11 regional NPF that takes place over distances of hundreds of kilometers (Kalkavouras et al.,
12 2017).

13 We calculated the growth rates at three different size ranges for the FRONT project similarly
14 to Manninen et al. (2010) and Pikridas et al (2012) for the EUCAARI project data. The particle
15 growth rates in the size ranges 1.5-3 nm, 3-7 nm and 7-20 nm were $1.6 \pm 1.8 \text{ nm hr}^{-1}$, 5.4 ± 4.9
16 nm hr^{-1} and $9.1 \pm 9.5 \text{ nm hr}^{-1}$, respectively. These values are lower than those in Pikridas et al.
17 (2012) but comparable to those observed during the EUCAARI project for the first two size
18 ranges, and higher than those observed during the EUCAARI project for the last size range
19 (Manninen et al., 2010). Overall, we observed much faster growth of newly-formed charged
20 particles in the eastern Mediterranean atmosphere after their first growth steps beyond 3 nm
21 in diameter, reflecting probably the strong Kelvin effect at small particle sizes preventing
22 condensation and hence growth, and the abundance of precursors leading to nucleation and
23 condensing species contributing to each growth stage.

24 3.5) Simulations of NPF using the zero-dimensional model MALTE-box

25 In order to evaluate our understanding of the observed NPF events in the eastern
26 Mediterranean we chose to simulate two distinct cases of one week duration each, during
27 which NPF events have been observed (event week) or not (no event week). The selection was
28 done from the summer of the year 2012, when JO^1D and JNO_2 photodissociation
29 measurements were also available at Finokalia. Two weeks in August 2012 were chosen,
30 28/08– 03/09 as event week and 09/08– 15/08 as non-event week. The “event week” was
31 described in detail by Kalivitis et al. (2015). Applying the MALTE-Box model the aerosol size
32 distribution and its evolution over the week has been simulated for these two cases.

During the “event week” the simulated formation of new particles successfully coincided with the observations. The NPF events simulated using the nucleation rates as parameterized for the boreal environment overestimated the observations while the simulated growth of newly-formed particles was greatly underestimated as shown in Tzitzikalaki et al. (2017). The most likely reason for this is the very low concentration of monoterpenes, calculated by TM4-ECPL global model for the Finokalia model grid box, on which the ELVOC and SVOC chemistry was built on. Indeed, the TM4-ECPL model results for Finokalia were too low compared to monoterpenes observations in 2014 (not shown). Therefore, we performed a number of sensitivity tests to improve the simulations by adjusting the nucleation coefficient and the monoterpene concentrations until we simulated efficiently the nucleation and growth rates observed during the second day of the “event week” when the most pronounced NPF event was observed. The best agreement between model results and observations was reached by decreasing the nucleation coefficient from 10^{-11} s^{-1} (the value commonly used for the boreal environment) to $5 \times 10^{-16} \text{ s}^{-1}$ and increasing by a factor of 10 the α - and β -pinene concentrations. With these modifications the model results improved and the aerosol number size distributions were better simulated, as well as total number and volume concentration of aerosol particles (Fig. 11a and b respectively). This was the first time that we were able to simulate NPF in the eastern Mediterranean environment. The almost five orders of magnitude lower nucleation coefficient used here for the sub-tropical set-up could be related to the contribution of still unknown compounds in the cluster-formation process. Huang et al. (2016) applied different kinetic nucleation coefficients at Nanjing, China, with the lowest value for a “China-clean” day of $6.0 \times 10^{-13} \text{ s}^{-1}$. The higher monoterpene concentrations used are comparable to the findings at Finokalia but also at another location in the eastern Mediterranean (Debevec et al., 2018).

Using the non-event week as our control case, we performed simulations of number size distributions at Finokalia station using the sub-tropical set-up and compared it to our measurements. For the “non-event week”, weak NPF were predicted by the model during the last two days that were not found in the measurements (Tzitzikalaki et al., 2017) but appear to be associated with the rapid drop of CS during day five of the simulations. Nevertheless, even if no NPF took place during the last two days, it was apparent in our measurements that some nucleation particles appeared ($\sim 200 \text{ cm}^{-3}$) and thus the general tendency was captured by the model. Both total number and volume concentrations were adequately simulated by the model (Fig. 12 a, b). These results show the potential of MALTE-box model to simulate the NPF in the eastern Mediterranean and the importance of input data. Therefore, when more

appropriate input data for Malte-box will become available (concurrent detailed measurements of gases and aerosol distributions) at Finokalia, new simulations and VOC measurements will further provide insight in the nucleation mechanisms, the growth process and the factors controlling NPF in the eastern Mediterranean atmosphere.

4) Conclusions

NPF in the atmosphere is a recurrent phenomenon in eastern Mediterranean. In this study, we presented the longest time series of NPF records in the region. We analyzed 3057 days of aerosol number size distribution data from June 2008 until June 2018 and found that NPF took place 27% of the available days, more frequently in spring and less frequently in late summer. Production of nucleation mode particles was common during night-time as well. Nucleation mode particle number concentrations were low outside periods of active NPF and subsequent particle growth indicating absence of local sources. Classification of NPF events based on atmospheric ion measurements differed from the corresponding classification based on mobility spectrometer measurements: the maximum frequency of NPF events was observed earlier in spring from AIS data than from SMPS data, possibly indicating a different representation of local and regional NPF events between these two data sets since SMPS measures new particles after they have grown to diameters larger than 9-nm and hence records only regional events lasting for several hours.

We used the MALTE-box model to simulate NPF observations in the eastern Mediterranean region. Using a “sub-tropical” environment parametrization, we were able to simulate with good agreement the selected time period. The parametrization used was significantly different than the one used for the boreal environment: nucleation rates were much lower, yet monoterpenes seemed to play a key role in the mechanisms governing NPF phenomena.

From the results presented in this work it is evident that the Finokalia site is a unique location in the eastern Mediterranean for studying the processes leading to NPF in the marine environment. As a next step, a more detailed look to the precursors driving these processes is necessary, with special emphasis on VOCs, and the expansion of the available measurements at the site in order to eliminate the uncertainties introduced in our simulations from the use of model outputs instead of observations.

1 5) Author contribution

2 N.K., G.K., I.S., A.B., P.K. and H.E.M. participated in the field measurements and analyzed the
3 data, E.T., N.D., S.M., P.R. and M.B. participated in the modeling study, T.P., V.M.K., M.K.,
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5

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28

1 8) Tables

Day classification	Number of events	%
Total events	837	27.4
Class I	232	7.6
Class II	605	19.8
Undefined	687	22.5
Non-event	1533	50.1
Total days	3057	100.00

2

3 Table 1. Total available measurement days and percentage of NPF events observed at
4 Finokalia during the period June 2008-June 2018

	J_9 ($\text{cm}^{-3} \text{s}^{-1}$)			GR_{9-25} (nm hr^{-1})			$CS \times 10^{-3}$ (s^{-1})		
	Mean	Median	SD	Mean	Median	SD	Mean	Median	SD
Winter	0.9	0.6	1.4	3.3	2.6	2.4	4.3	3.5	2.9
Spring	1.0	0.6	1.0	4.2	3.3	3.1	5.8	5.5	3.0
Summer	0.7	0.5	0.9	7.3	6.8	3.9	9.1	9.0	3.1
Autumn	0.8	0.4	1.0	5.3	4.7	2.9	6.5	6.0	3.4

5

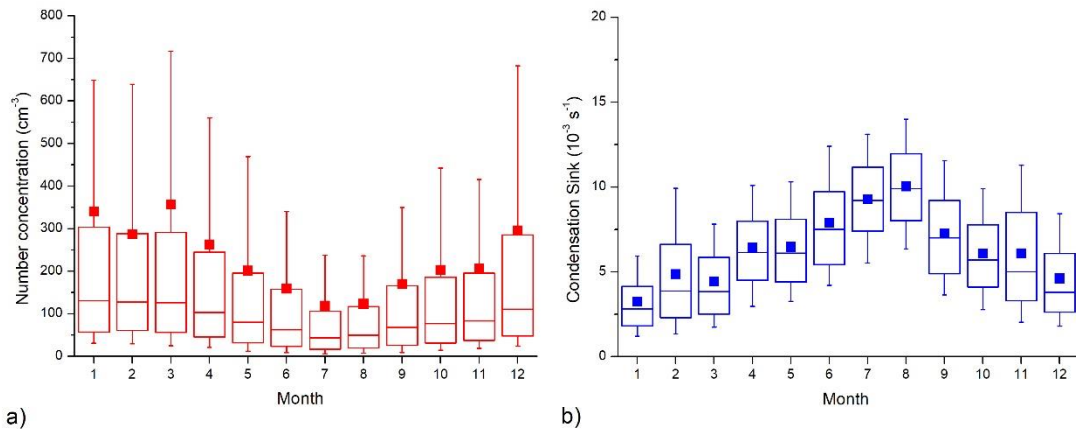
6 Table 2. Formation rates for 9-nm particles (J_9), growth rates in the size range 9-25 nm (GR_{9-25}) for NPF events observed at Finokalia and condensational sink for sulfuric acid (CS) on a
7 seasonal basis during the period June 2008-June 2018 (mean, median and standard deviation).
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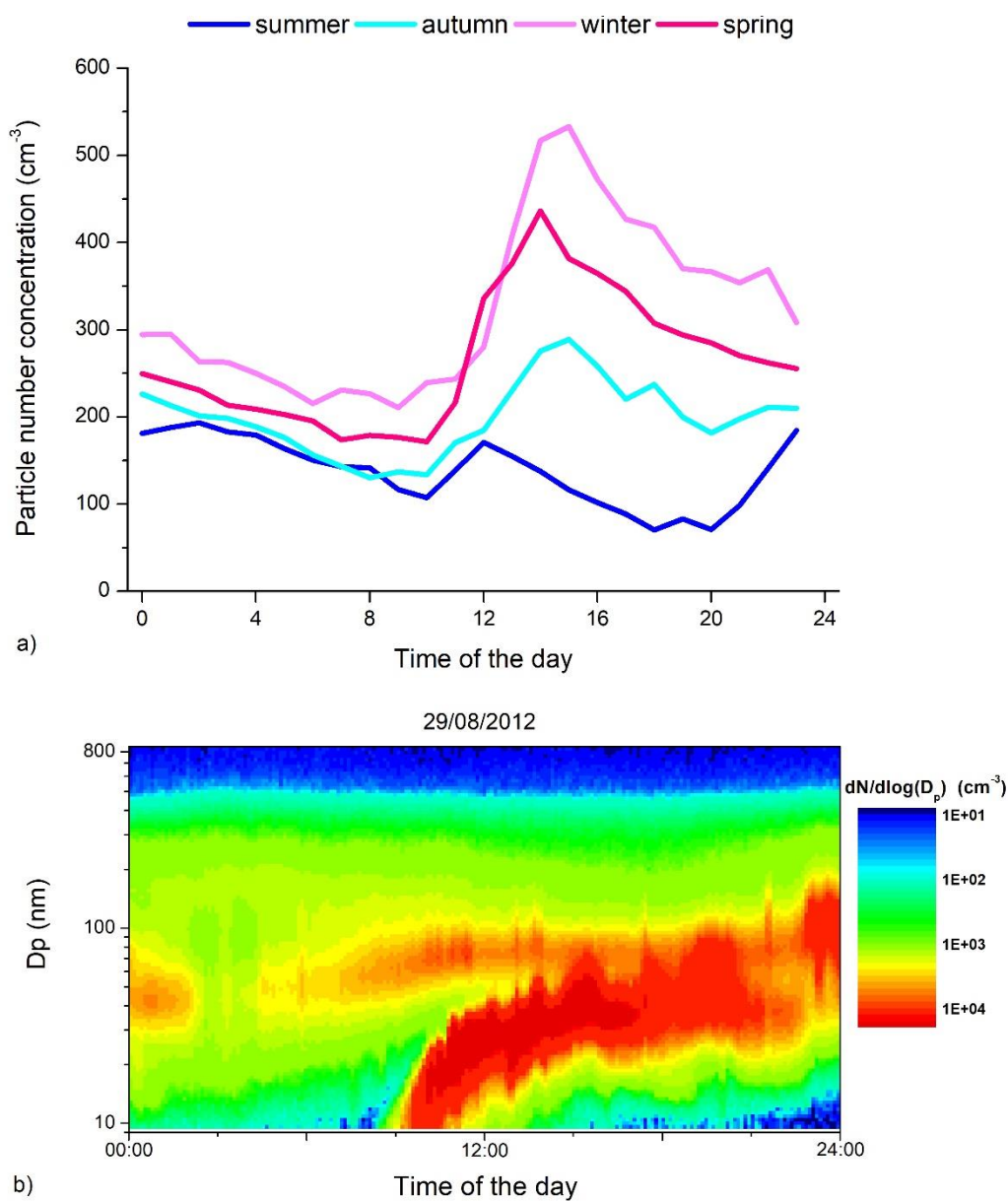
1 9) Figures



- 2 a)
- 3 1. Monthly average variation of (a) nucleation mode particle number concentration and (b)
- 4 sulfuric acid condensational sink (CS) at Finokalia station over the period June 2008-June 2018.
- 5 Whiskers represent 10th and 90th percentiles, box edges are 75th and 25th percentiles, the
- 6 line in the box is the median, the solid square is the mean.

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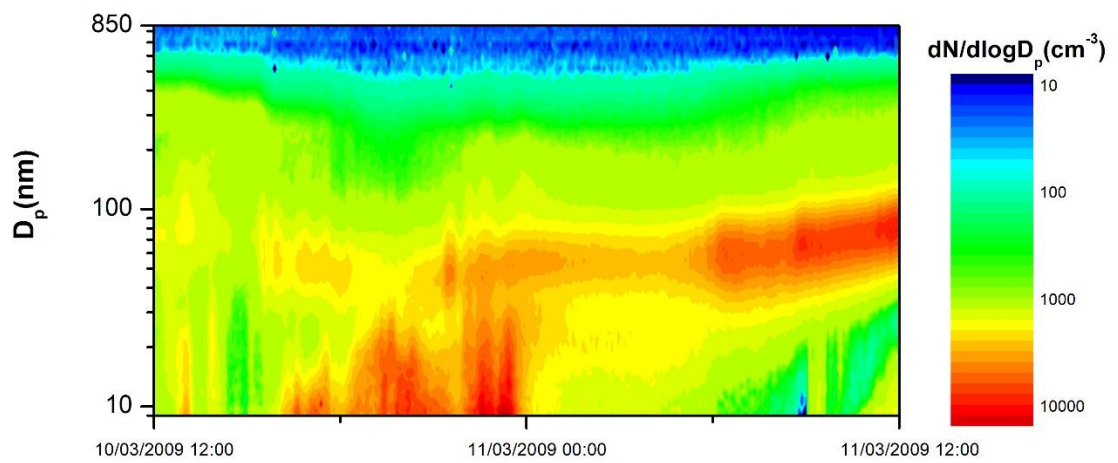
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3 2. (a) Average diurnal variation of nucleation mode particle number concentration (hourly
4 values) at Finokalia over the period June 2008-June 2018. (b) New particle formation event
5 captured at Finokalia on 29/08/2012 (time in UTC+2).

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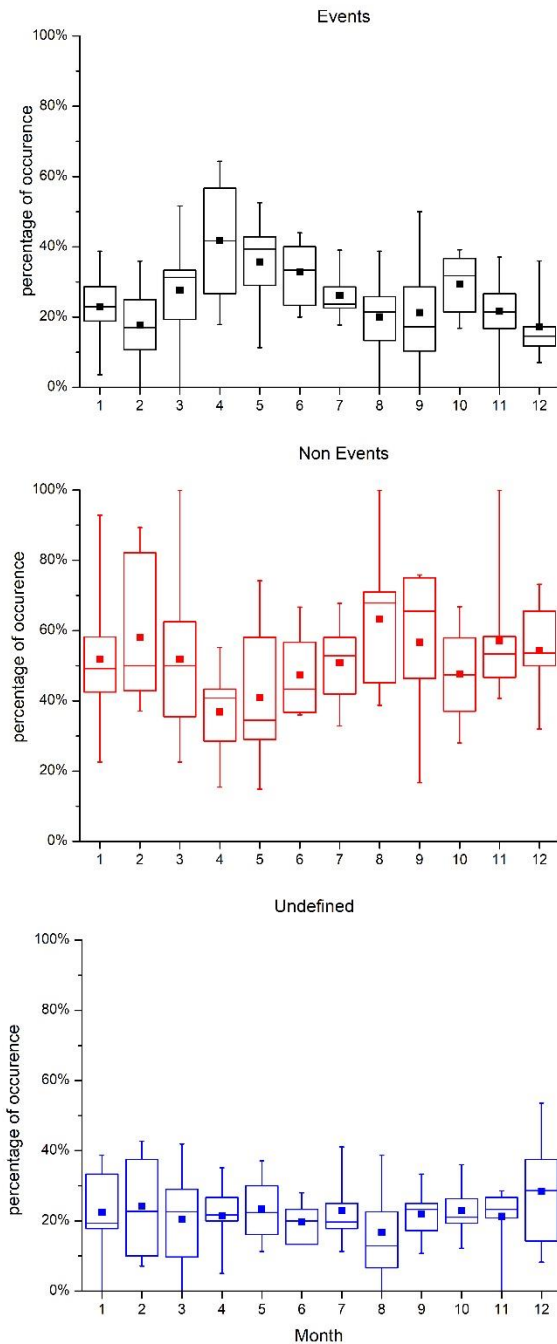
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2 3. Example of appearance of nucleation mode particles during several hours as observed
 3 during the night of 10 to 11/03/2009 (time in UTC+2).

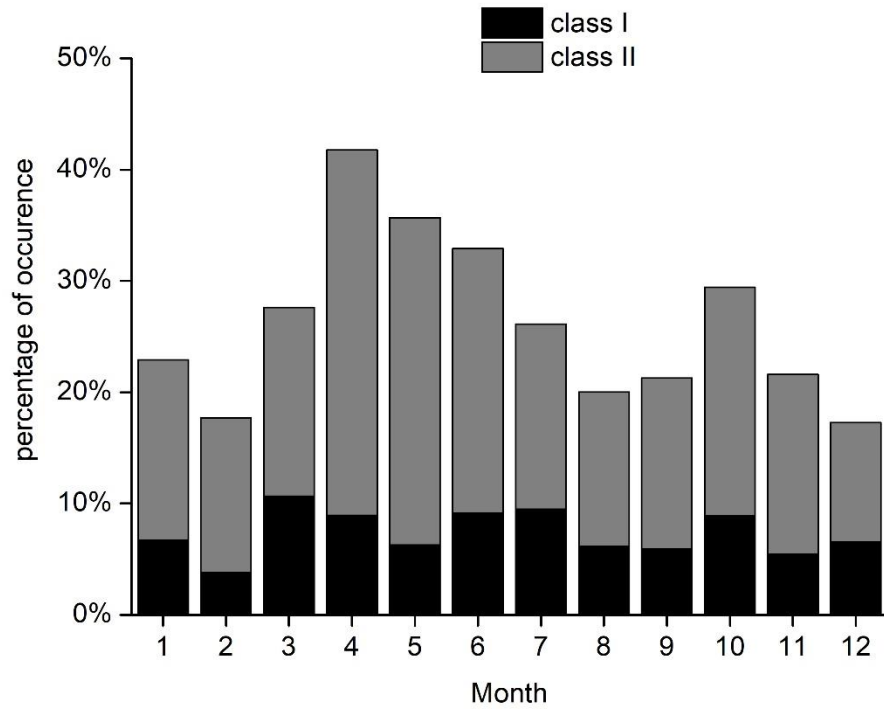
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2 4. Seasonal variation of NPF percentage of occurrence of event, non-event and undefined days
3 relatively to available measurement days at Finokalia for the period June 2008-June2018.
4 Whiskers represent 10th and 90th percentiles, box edges are 75th and 25th percentiles, the
5 horizontal line in the box is the median, square is mean.

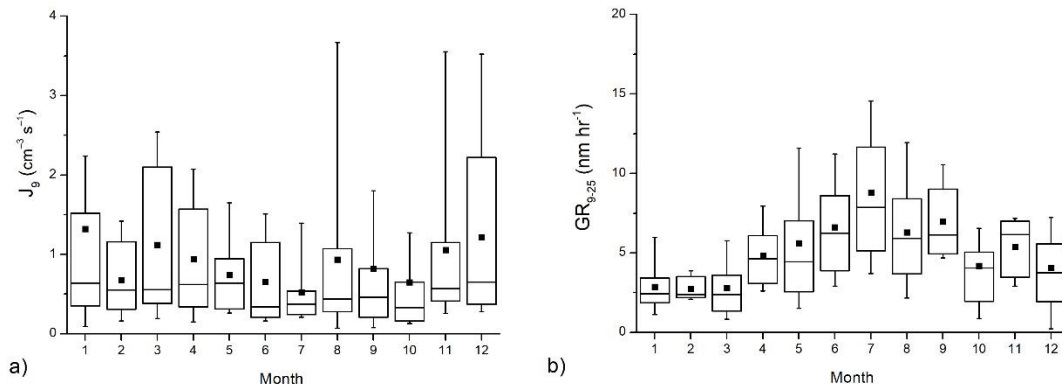
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2 5. Seasonal variation of percentage of occurrence of NPF Class I & II events relatively to
 3 available measurement days at Finokalia in the eastern Mediterranean for the period June
 4 2008-June2018.

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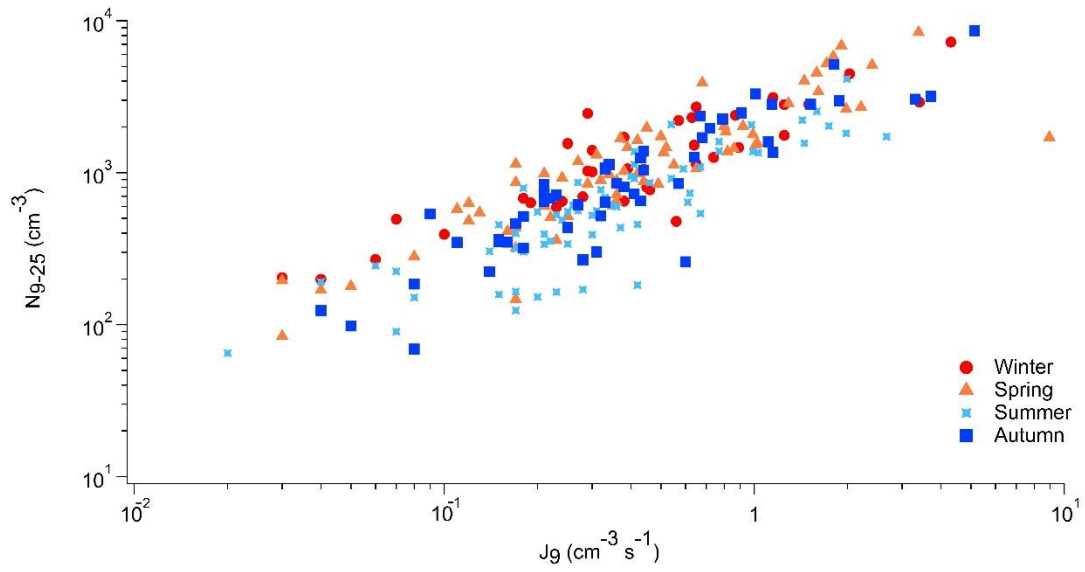
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4 6. Seasonal variation of (a) formation rate of 9-nm particles (J_9) and (b) growth rate in the size
5 range 9-25nm (GR_{9-25}) as calculated during Class I NPF events at Finokalia for the period June
6 2008-June 2018. Whiskers represent 10th and 90th percentiles, box edges are 75th and 25th
7 percentiles, the horizontal line in the box is the median and the solid square is the mean.

8

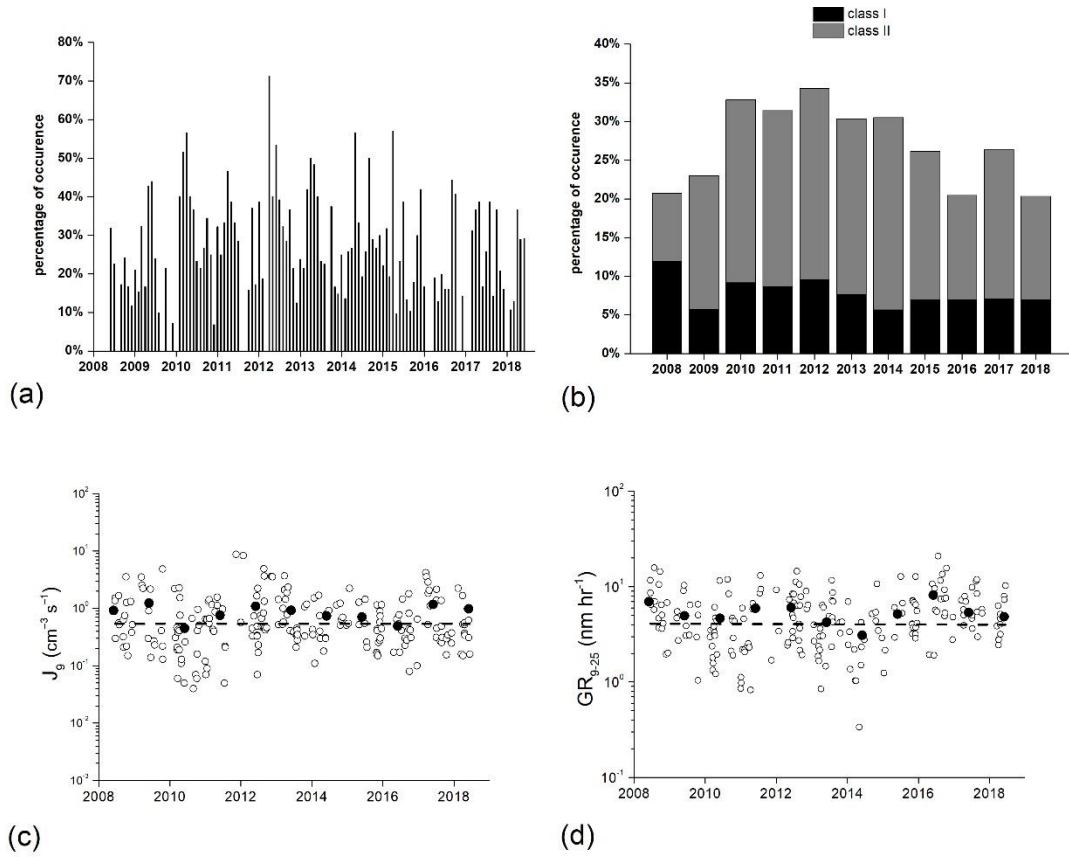


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2 7. Scatter plot of the number concentration of nucleation mode particles (N_{9-25}) (hourly
3 maximum value during the event) versus formation rates of 9-nm particles (J_9) at Finokalia,
4 for events when J_9 could be calculated with a good level of confidence (Class I events).

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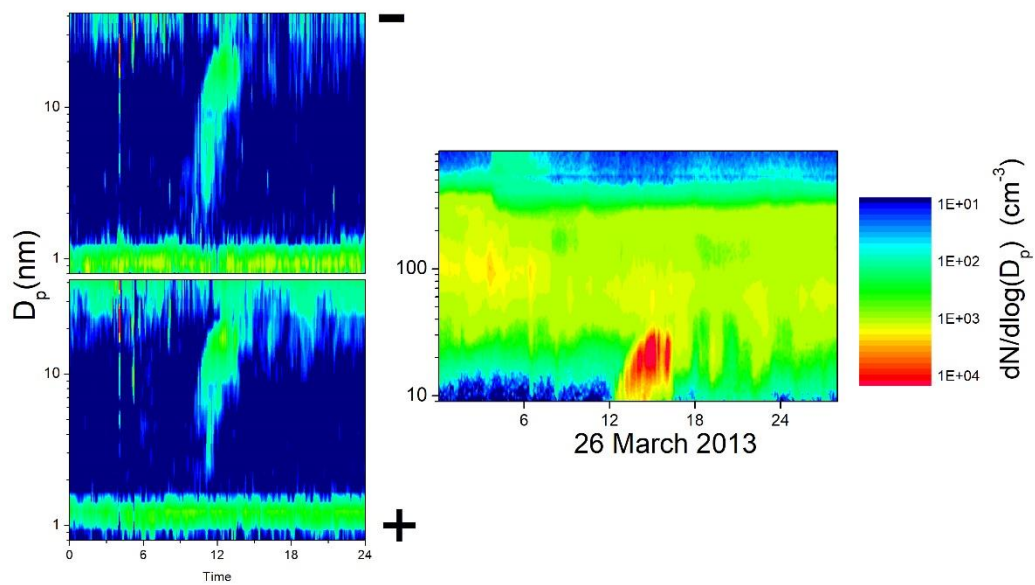
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3 8. (a) Time series of monthly NPF percentage of occurrence at Finokalia for the years 2008-
 4 2018. (b) Annual NPF percentage of occurrence at Finokalia for the period June 2008-June
 5 2018 for Class I&II events. Interannual variation of (c) formation rates of 9-nm particles (J_9)
 6 and (d) growth rate in the size range 9-25 nm (GR_{9-25}) during Class I NPF events at Finokalia for
 7 the period June 2008-June 2015 (the solid circles represent annual averages and the dashed
 8 lines the linear regression).

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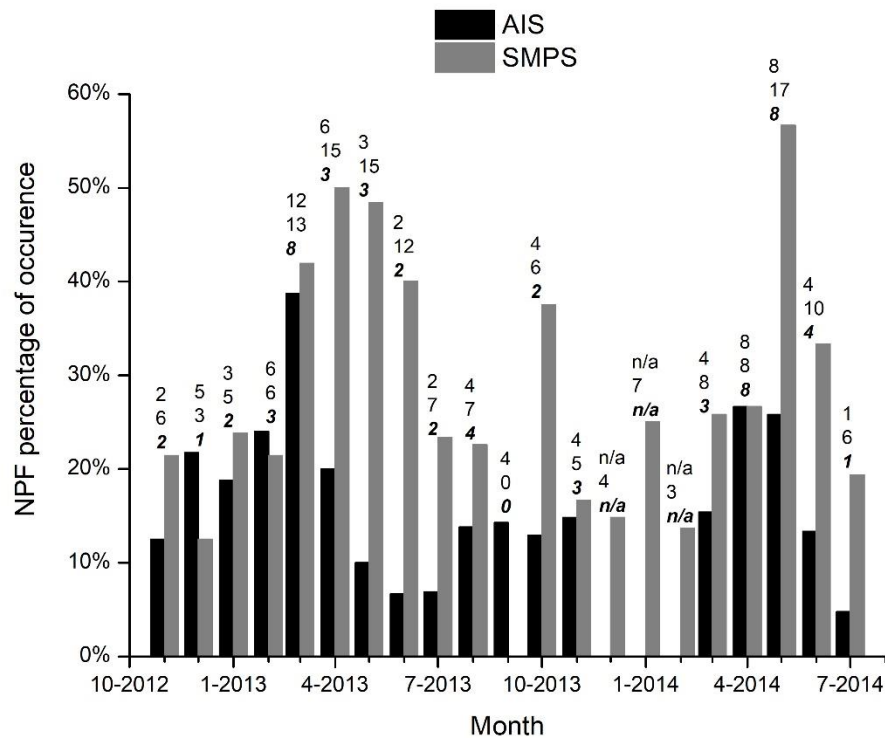
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6 9. Nucleation event observed at Finokalia on 26 March 2013 as captured by AIS (left panels
 7 for negative (up) and positive (bottom) polarity) and SMPS (right panel) (time in UTC+2).

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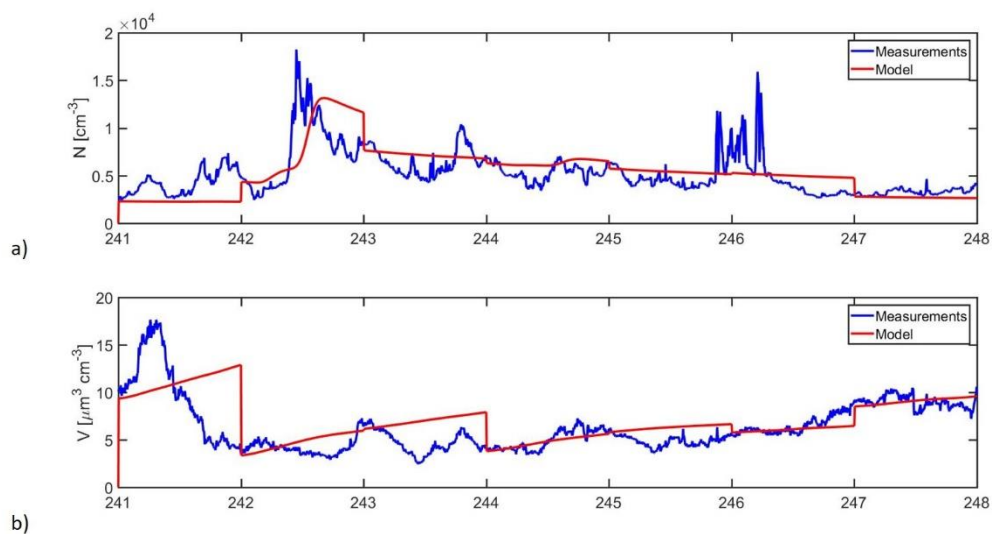


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4 10. Monthly variability of NPF events' percentage of occurrence relatively to available
 5 measurement days at Finokalia as determined by analysis of AIS data during the FRONT
 6 experiment (Nov. 2012-July 2014). For a direct comparison, the monthly variability of NPF
 7 events as obtained from the SMPS measurements for the same period is included. On top of
 8 the columns, the number of NPF events observed for AIS (top), SMPS (middle) and the number
 9 of common events for both instruments (bottom, italic, bold) for each month are presented.

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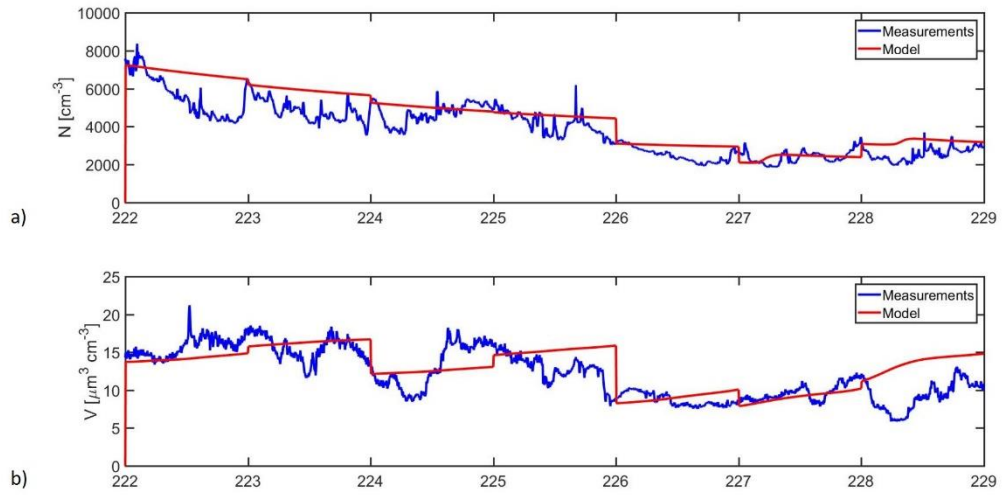


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4 11. Simulations with the MALTE box with the adjusted parameters for the sub-tropical
 5 environment for the “event week” that NPF events were observed at Finokalia. Measured and
 6 modelled (a) total number concentration and (b) total volume concentration for the same
 7 period. The x-axis in both figures is Julian day of 2012.

8



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2 12. Simulations with the MALTE box with the adjusted parameters for the sub-tropical
 3 environment for the “non-event” week that no NPF was observed at Finokalia. Measured
 4 and modelled (a) total number concentration and (b) total volume concentration for the
 5 same period. The x-axis in both figures is Julian day of 2012.

6