

# Multiple eco-regions contribute to the seasonal cycle of Antarctic aerosol size distributions

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## Abstract

In order to reduce the uncertainty of aerosol radiative forcing in global climate models, we need to better understand natural aerosol sources which are important to constrain the current and pre-industrial climate. Here, we analyze Particle Number Size Distributions (PNSD) collected during a year (2015) across four coastal and inland Antarctic research bases (Halley, Marambio, Dome C and King Sejong). We utilise k-means cluster analysis to separate the PNSD data into six main categories. Nucleation and Bursting PNSDs occur 28-48% of the time between sites, most commonly at coastal sites Marambio and King Sejong where air masses mostly come from the west and travel over extensive regions of sea ice, marginal ice, and open ocean, and likely arise from new particle formation. Aitken high, Aitken low, and bimodal PNSDs occur 37-68% of the time, most commonly at Dome C on the Antarctic Plateau, and likely arise from atmospheric transport and aging from aerosol originating likely in both coastal boundary layer and free troposphere. Pristine PNSDs with low aerosol concentrations occur 12-45% of the time, most common at Halley located at low altitudes and far from the coastal melting ice and influenced by air masses from the west. Not only the sea spray primary aerosols and gas to particle secondary aerosols sources, but also the different air masses impacting the research stations should be kept in mind when deliberating upon different aerosol precursors sources across research stations. We infer that both primary and secondary components from pelagic and sympagic regions strongly contribute to the annual seasonal cycle of Antarctic aerosols. Our simultaneous aerosols measurements stress the importance of the variation in atmospheric biogeochemistry across the Antarctic region.

## Introduction

The pristine region of the Southern Ocean plays a major role in modulating Earth's climate (Carslaw et al., 2013), and natural aerosols are one of the greatest sources of uncertainty in estimates of global radiative forcing (Reddington et al., 2017). This uncertainty becomes greater in polar regions (Carslaw et al., 2013). Aerosols modulate the climate both directly, by absorbing and reflecting radiation, and indirectly, by acting as cloud condensation nuclei (CCN), modulating cloud properties. The surface of the Southern Ocean near the Antarctic continent goes through an annual freezing cycle; this large frozen area harbours one of the globe's largest ecosystems providing a stable habitat for diverse microbial assemblages (Arrigo et al., 2009; 2015). Understanding these processes is key in polar environments where warming processes are rapidly affecting delicate ecosystems (Clem et al., 2020). For example, the

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**Eliminado:** In order to reduce the uncertainty of aerosol radiative forcing in global climate models, we need to better understand natural aerosol sources which are important to constrain the current and pre-industrial climate. Here, we analyze Particle Number Size Distributions (PNSD) collected during a year (2015) across four coastal and inland Antarctic research bases (Halley, Marambio, Concordia/Dome C and King Sejong). ¶

We find that the four Antarctic locations have striking differences in PNSD, stressing multiple aerosol sources and processes likely exist. We utilise k-means cluster analysis to separate the PNSD data into six main categories. Nucleation and Bursting PNSDs occur 28-48% of the time between sites, most commonly at coastal sites Marambio and King Sejong where air masses mostly come from the west and travel over extensive regions of sea ice, marginal ice, and open ocean, and likely arise from new particle formation. Aitken high, Aitken low, and bimodal PNSDs occur 37-68% of the time, most commonly at Concordia/Dome C on the Antarctic Plateau, and likely arise from atmospheric transport and aging from aerosol originating likely in both coastal boundary layer and free troposphere. Pristine PNSDs with low aerosol concentrations occur 12-45% of the time, most common at Halley located at low altitudes and far from the coastal melting ice, and influenced by air masses from the west. ¶

The Antarctic Atmospheric circulation has a strong control on the air mass origin type. Most of the time Marambio and King Sejong stations are affected by Easterly air masses, whereas Halley gets air masses mainly from the Weddell sea marginal and consolidated pack ice. Not only the sea spray primary aerosols and gas to particle secondary aerosols sources, but also the different air masses impacting the research stations should be kept in mind when deliberating upon different aerosol precursors sources across research stations. ¶

We provide evidence that both primary and secondary components from pelagic and sympagic regions strongly contribute to the annual seasonal cycle of Antarctic aerosols which add insight on the possible sources of aerosol production/activity in the whole Antarctic region. Our simultaneous aerosol measurements stress the importance of the variation in atmospheric biogeochemistry across the Antarctic region.

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Antarctic Peninsula has shown some of the largest increases in near-surface air temperature measured globally across the last 50 years (Turner et al., 2005). Study of the Antarctic environment can also provide an insight into natural preindustrial aerosol processes, and allows us to further our understanding of the preindustrial baseline (Hamilton et al., 2014).

Antarctic sea ice covers between 1% (summer) and 5% (winter) of the global ocean. Antarctic terrestrial productivity and biodiversity occur almost exclusively in ice-free areas that cover less than 1% of the continent. In Antarctica the coastline extends for 17,968 km and comprises about 34.8-36.4x10<sup>6</sup> km<sup>2</sup> where 80% of this surface is covered by ice, even in summer (Peck et al., 2018; Ronowicz et al., 2019). Overall, there is about 390,071 km<sup>2</sup> of coast shallower than 200 m (Peck et al., 2018). Antarctic coastal systems harbour a high diversity of marine and terrestrial ecosystems including Antarctic seaweeds (benthonic macroalgae) and bird colonies (mainly penguins). Antarctic seaweeds (often called macroalgae) are found in shallow (<200m) coastal, rocky shores and can cover more than 80% of the benthic surface (Amsler et al., 2005; Wiencke and Amsler, 2012). The Antarctic and Sub-Antarctic region is home for about half of the total worldwide seabird population (Otero et al., 2018). Penguins represents a high proportion of the avian biomass, and their fecal material is one of the main source of phosphorus and nitrogen, representing about 80% of these elements in the Antarctic marine environment. Seabird colonies also represent a significant source of atmospheric ammonia (NH<sub>3</sub>) in remote maritime systems (Riddick et al., 2012, Schmale et al., 2013). These emissions are environmentally relevant as they primarily occur as "hot-spots" in otherwise pristine environments.

The role of aerosols in the Antarctic is poorly understood as their sources are many and varied in a relatively understudied region. Emissions from the Southern Ocean and the Antarctic region are characterized by multiple environmental systems including open ocean water, sea ice regions, and land, snow covered and snow-free. These regions are being affected by our changing climate (Chen et al., 2009). It has been long known that the seasonal cycle of Antarctic particle number concentrations has a strong seasonal cycle (Shaw 1988), leading to the inference that most of the Antarctic aerosol concentrations is linked with biological processes occurring in the surrounding oceans. It has been proposed that a large "pulse" over summer months arises from the upper Antarctic plateau (Ito, 1993; James, 1989), although particle number concentrations are much higher in coastal Antarctica. In other words, it is still debatable if the aerosols originate in the upper troposphere, being transported by the Antarctic

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drainage flow (James, 1989) to the coast by katabatic winds (Ito, 1993; Koponen et al., 2002; Fiebig et al., 2014; Hara et al., 2011; Järvinen et al., 2013; Humphries et al., 2015) or the marine pelagic (ice-free) and sympagic (ice-containing) boundary layer are a source of ultrafine particles (Herenz et al., 2019; Weller et al., 2011, 2015, 2018; Dall'Osto et al. 2017; Heintzenberg et al., 2000, 2023).

The Antarctic aerosol summer maximum concentrations (mainly ultrafine particles, <100 nm) may be largely explained by new particle formation (NPF) events, as reviewed by Kerminen et al. (2018). Super-micrometre aerosols (>1 µm) will mostly arise from primary sea spray (O'Dowd et al., 1997a, 1997b; Rankin and Wolff, 2003). The accumulation mode (100 nm – 1 µm) is a complex intermediary region with both primary and secondary sources related to their multitude of sources and long atmospheric lifetimes (Fossum et al., 2018; Gras and Keywood, 2017). The relative importance of these sources undergoes seasonal cycles related to meteorology and biological productivity.

During the austral summer the concentration of DMS in the water of the Southern Ocean is the highest of the planet (Lana et al., 2011), with high fluxes into the atmosphere, potentially producing high concentrations of sulphuric acid and methanesulphonic acid. However, under typical boundary layer conditions the concentration of sulphuric acid is too low to form particles alone, and another molecule, such as ammonia, is required to stabilize the nucleating clusters (Kirkby et al., 2011). Jokinen et al. (2018) reported the first molecular characterization of NPF from the Aboa station in Antarctica, showing that new particles are formed via clustering of sulfuric acid and ammonia (Kirby et al., 2011). Sources of ammonia and alkylamines are related to animals, mainly birds and seal colonies and melting sea ice, but the relative importance of each source is unknown (Riddick et al., 2012; Schmale et al., 2013, Brean et al., 2021; Quelever et al., 2022). Iodine oxoacids are also capable of efficiently forming new particles (Baccarini et al., 2020; He et al., 2021; Sipilä et al., 2016), with iodous acid being able to efficiently stabilise clusters of iodine and sulphuric acid, even in the absence of ammonia (He et al., 2023), with modelling studies showing high gas-phase iodine concentrations across Antarctica (Saiz-Lopez et al., 2007). Further, the formation of marine primary aerosol particles and of gas-to-particles precursors is influenced by the uppermost layer of the ocean, the so called sea surface microlayer (SML) (Cunliffe et al., 2013). The SML - with physicochemical characteristics different from those of subsurface waters (SSW) - results in dense and active viral and microbial communities (Vaque et al., 2021, Heinrichs et

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175 al., 2024). The microlayer of the ocean is a source of oxygenated VOCs in the Arctic leading to secondary aerosol formation (Mungall et al., 2017), and it is also likely a source in the Southern Ocean too although no measurements of this type are available. Therefore, while the limited number of measurement studies implicate sulphuric acid and nitrogenous bases as the drivers of NPF, iodine and VOCs may play a substantial role.

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180 The roles of secondary aerosols produced from biogenic sulfur mainly derived from the atmospheric oxidation of dimethylsulfide (DMS, a trace gas produced by marine plankton) and primary sea-spray aerosols in regulating cloudiness above the Southern Ocean is still a matter of debate (Meskhidze et al., 2006; Korhonen 2008, Asmi et al., 2018; Quin and Bates, 2011, Fossun 2018, Lachlan-Cope et al., 2020). The seasonal cycle of primary and secondary aerosol emissions from the Southern Ocean affect also light scattering by aerosols over all Antarctica, including the upper plateau (Virkkula et al., 2022). A recent intensification in Antarctic aerosol measurement field campaigns is revealing that aerosol chemistry in the higher latitudes of Antarctica can be much more complex than two broad natural sources governing the aerosol populations: sea spray (primary, mostly composed of sea salt) and non-sea salt sulfate (nssSO<sub>4</sub><sup>2-</sup>; secondary). For example, Paglione et al. (2024) stressed that various, not yet fully understood sources and aerosol processes are controlling the Antarctic aerosol primary and secondary population, with the emissions from sympagic and pelagic ecosystems affecting the variability of submicron aerosol composition both in maritime areas as in inner Antarctic regions.

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195 McCoy et al. (2015) suggested that primary marine organic ultrafine aerosols are important in the Southern Ocean region. Saliba et al. (2021) found that the large total organic fraction of particles <0.1 µm diameter may have important implications for CCN number concentrations and indirect radiative forcing over the Southern Ocean. Recently, Humphries et al., (2021) identified three main aerosol sources in the Southern Ocean: northern (40-45 S), mid-latitude (45-65 S) and Southern sector (65-70 S), with different mixture of continental and anthropogenic, primary and secondary aerosols depending on the studied region.

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200 Recently, several long-term measurements (in the order of a year or longer) of aerosol particle number size distributions (PNSD) at a high time resolution allow investigation of the aerosol sources around the Antarctic continent. The PNSD is typically measured from ~10 – 1,000 nm, and in Antarctica is usually comprised of particles from secondary (NPF) and primary (blowing snow - BS, Sea Spray Aerosols - SSA) sources.

At King Sejong Station on the Antarctic Peninsula, (Kim et al., 2017; Park et al., 2023; Kim et al., 2019), Halley on the mainland coast (Lachlan-Cope et al., 2020) and Concordia/Dome C in the centre of the continent (Järvinen et al., 2013), PNSD measurements indicated NPF as a mostly summertime phenomenon. At Concordia/Dome C weak and rare but real NPF has been observed even in June, the darkest time of the year (Järvinen et al., 2013). Particle counts and CCN numbers at Macquarie Island and throughout the Southern Ocean undergo a summertime maximum (Humpries et al., 2023). The CCN concentration at the onset of NPF showed an average instantaneous increase of 44% or 11% across the whole period compared with the background concentration (Kim et al., 2019; Park et al., 2023). NPF therefore seems to modulate particle and CCN counts in the Antarctic.

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At Halley, NPF events were shown to arise both from the sea ice marginal zone and the Antarctic plateau, indicating a marine and free tropospheric source (Lachlan-Cope et al., 2020). NPF events at King Sejong Station were found more frequently associated with air masses originating from the Bellingshausen Sea than those from the Weddell Sea, with a fraction of events being associated with sea ice (Park et al. 2023). This suggests the phytoplankton composition of the Bellingshausen Sea may be a source of NPF precursors (Jang et al., 2019).

In Marambio, Quéléver et al. (2022) reported neutral and charged aerosol precursor molecules and clusters, as well as charged and neutral PNSDs. NPF precursors were inferred to be related with local fauna (mainly penguins), and oceanic emissions from the Bellingshausen sea.

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Hara et al. (2021) hypothesized that NPF mainly occurred in the Antarctic free troposphere during spring and fall, and in both the free troposphere and boundary layer during summer, based on PNSD measurements and their effect on cloud properties observed at Syowa Station.

Concordia/Dome C's observations reported the background PNSD had its largest mode below 30 nm, suggesting these particles were produced in the upper atmosphere before entering continental Antarctica (Järvinen et al., 2013). However, also ground-level particle formation takes place on the upper plateau. Chen et al. (2017) reported NPF events that were observed at Concordia/Dome C by using a combination of an Air Ion Spectrometer (AIS) and a Differential

Mobility Particle Sizer (DMPS). In several events particle formation and growth started from the cluster ion size range of 0.9 – 1.9 nm which means that in these cases the formation took place at the ground level, however, NPF precursor emissions from the plateau itself are believed to be negligible, implying they are transported from elsewhere. NPF in Antarctica is therefore dependent on biogenic emissions from the ice-covered and ice-free regions of the

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ocean, with NPF occurring both at ground-level and in the colder upper atmosphere where particle formation rates can proceed more rapidly.

265 Previous analyses of PNSDs at Halley using k-means cluster analysis has shown that wintertime PNSDs were characterised by extremely low particle concentrations, with a bimodal PNSD appearing with a blowing snow or sea spray origin (Lachlan-Cope et al., 2020).

The Antarctic wintertime at this site is therefore mostly devoid of secondary aerosol sources and is instead dominated by primary sources. Some of these primary aerosols will be organic,

270 and NMR analyses of ambient aerosol samples show that aerosols arising from the ice-free Southern Ocean are rich in lipids and sugars, and aerosols arising from coastal areas are rich in sugars associated with plant vegetation (Decesari et al., 2020). These sources are likely primary, and the primary aerosol sources are therefore many and varied across Antarctica.

Here, we apply k-means cluster analysis to simultaneous measurements in the year 2015 at four

275 Antarctic research sites, extending the study of Lachlan-Cope et al., (2020). We also compare the yearly data with field studies, including the PEGASO cruise in 2015 (Dall'Osto et al., 2017a, 2019b; Rinaldi et al., 2019; Decesari et al., 2020) and the continental Antarctic site Kohlen (Weller et al., 2018). We show a prevalence of new particle formation at the coastal sites, and associate this new particle formation with air masses flowing over regions of sea ice and ocean. At the more southerly and inland sites, primary particles dominate the particle number concentrations, while air masses primarily travel over regions of land. Ambiguity remains in this analysis, as some PNSD clusters likely contains a substantial contribution from primary and secondary processes. Nonetheless, we provide further evidence for the roles of emissions from sympagic and pelagic ocean regions in new particle formation, and highlight

285 the many and varied sources of particles across Antarctica.

## 2 Methodology

### 2.1 Sampling sites and measurements

Simultaneous PNSD measurements at four sites across Antarctica were collected for analysis. Their locations are stated in Table 1, and shown in Figure 1. Data coverage shown in Figure

290 S1. PNSD measurements were aggregated to 1 hour for this study.

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300 The South Korean King Sejong Station, the highest latitude site, is located in the Antarctic Peninsula. The PNSD from 10 to 300 nm were measured every 3 min with a mobility particle size spectrometer (MPSS) consisting of a differential mobility analyzer (DMA; HCT Inc., LDMA 4210) and a CPC (TSI 3772). Details of the site can be found in (Kim et al., 2017, 2019, Park et al., 2023).

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305 The Argentinian Marambio Station is located in the Antarctic Peninsula, approximately 3 km from the coast. The PNSD from 12 – 800 nm were measured every 6 min with a differential MPSS (Asmi et al., 2018; Quelever et al., 2022).

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The French & Italian operated Concordia/Dome C station is operated on the East Antarctic Plateau at an elevation >3000 m a.s.l, and 900 km from the nearest coast. The Concordia/Dome C sampling site is located 1 km southwest of Concordia/Dome C's main building. The PNSD from 10 – 620 nm were measured every 10 min with a differential MPSS system (Järvinen et al., 2013).

315 The British Halley VI station is located in coastal mainland Antarctica on the floating Brunt Ice Shelf ~ 20 km from the coast of the Weddell Sea. The The Clean Air Sector Laboratory where PNSD measurements are taken is located 1 km south-east of the station. The PNSD from 6 to 209 nm is measured every 1 min using a TSI Inc. MPSS, comprising an Electrostatic Classifier (model 3082), a Condensation Particle Counter (CPC; model 3775) and a long Differential Mobility Analyser (DMA; model 3081) (Lachlan-Cope et al., 2020).

## 2.2 Data processing

320 K-means Cluster Analysis was applied to the PNSD data to apportion the PNSDs according to their shape (Beddows et al., 2009), and is routinely applied in pristine environments to PNSD data (Dall'Osto et al., 2017b, 2019b; Lachlan-Cope et al., 2020), including the Halley dataset included in this paper, (Lachlan-Cope et al., 2020). k-means apportions data into k clusters such as the sum of squares of distances of data to the cluster centre is minimised. In the case of MPSS data, this produces well-separated clusters (Beddows et al., 2009). First, data were normalised so the Euclidian length of each PNSD was 1, so the clustering is done solely on the merit of the shape of the PNSDs. Here, a degree of distance between clusters was achieved with 16 MPSS clusters. These were assigned into 6 categories typical of Antarctic PNSDs (Lachlan-Cope et al., 2020). Compared to previous work, we combined the three “pristine”

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335 [clusters identified by Lachlan-Cope et al. \(2020\) into one, as they follow the same seasonal trend, producing 6 clusters instead of 8.](#)

The condensation sink (CS, s<sup>-1</sup>) represents the rate at which a vapour phase molecule will collide with pre-existing particle surface, and was calculated from the PNSD data as follows (Kulmala et al., 2012):

$$CS = 2\pi D \sum_{d_p} \beta_{m,d_p} d_p N_{d_p} \quad (1)$$

340 [Here,  \$D\$  represents the diffusion coefficient of the condensing vapor, which is assumed to be sulfuric acid. The transitional regime correction factor is denoted by  \$\beta\_{m,d\_p}\$ ,  \$d\_p\$  is the diameter of a measurement bin, and  \$N\_{d\_p}\$  is the number of particles in size bin  \$d\_p\$ .](#)

### 2.3 Air mass trajectories

345 Air mass back trajectories were calculated using the HYSPLIT4 (Hybrid Single Particle Lagrangian Integrated Trajectory) trajectory model (Draxler and Rolph, 2010) arriving at the sampling site every one hour. Each back-trajectory data point was assigned to a surface-type (land, sea, ice, or snow over land. A cell is considered ice-covered if more than 40% of the cell is covered with ice) on a 24 km grid from the daily Interactive Multisensor Snow and Ice Mapping System (IMS) (Ice, 2008). To investigate air masses concurrent with high particle count, these 72-hour back-trajectories were gridded to 1x1 grid cells of 1 degree each, and linked back to the total integrated particle number concentration from the PNSD measurements by the following equation:

$$\ln(\bar{C}_{ij}) = \frac{1}{\sum_{k=1}^N \tau_{ijk}} \sum_{k=1}^N \ln(c_k) \tau_{ijk} \quad (2)$$

355 Where  $\bar{C}_{ij}$  is the concentrated weighted trajectory (CWT) at cell  $i, j$ ,  $N$  is the total number of trajectories,  $c_k$  is the value of particle number ( $N$ ) associated with the arrival of trajectory  $k$ , and  $\tau_{ijk}$  is the residence time of trajectory  $k$  in grid cell  $i, j$ .  $\bar{C}_{ij}$  therefore describes the source strength of condensable vapour that drives particle growth from any particular grid cell (Hsu et al., 2003; Lupu and Maenhaut, 2002). This was done using the trajLevel function in the  
360 Openair package in R 3.6.3.

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### 3 Results

#### 365 3.1 Trends in aerosol PNSDs

The mean PNSD for the measured periods are shown in Figure 2a. PNSDs at the two Antarctic Peninsula sites, Marambio and King Sejong show large nucleation modes. Aitken mode peaks are present at all sites with maximum concentrations between 30 and 50 nm. Accumulation mode peaks are present at all sites with maximum concentrations >100 nm. The prominent nucleation modes at Marambio and King Sejong are also visible in the daily contour plots, with the visible signature of a new nucleation mode in the late and early afternoons (local times) at the two sites, respectively (Figure S3). Total average particle counts are highest at King Sejong & Marambio, ( $312\pm 601$  and  $270\pm 541$   $\text{cm}^{-3}$ ), particle counts at Halley are lower ( $223\pm 245$   $\text{cm}^{-3}$ ), while those at Concordia/Dome C are  $44\pm 67$   $\text{cm}^{-3}$ .

375 The monthly variation in total particle number ( $N_{\text{tot}}$ ) is similar between all sites, highest in the austral summer, with minimum values in July & August.  $N_{\text{tot}}$  at Concordia/Dome C is substantially lower in near all months compared to all other sites, with the monthly average only exceeding  $100$   $\text{cm}^{-3}$  in the Austral Summer months (Nov, Dec). Relative contributions of the modes <30 nm, 30-100 nm, and >100 nm to  $N_{\text{tot}}$  at each site is shown in Figure 2d. The relative contribution of <30 nm particles is highest at Marambio and King Sejong, exceeding 50% of the total  $N_{\text{tot}}$  at Marambio, consistent with the prominence of the presence of the nucleation mode (Fig. 2a). The contribution of <30 nm particles to total  $N_{\text{tot}}$  is lowest at Halley, where ~50% of the  $N_{\text{tot}}$  comes from the mode 30-100 nm, and ~30% from the mode >100 nm. At Concordia/Dome C, ~65% of  $N_{\text{tot}}$  comes from the mode 30-100 nm, with small contributions from particles above or below this size range.

#### 3.2 Cluster analysis

##### 3.2.1 Categorising PNSD K-means clusters

390 K-means cluster analysis was used to partition the PNSD data into days with similarly shaped PNSD data. In k-means cluster analysis, starting with a higher number of clusters and then merging them based on similarity can enhance the separation and interpretability of the final clusters by capturing finer details. These 16 daily K-means clusters (Figure S2, referred to

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395 onwards as the initial PNSD clusters) were therefore aggregated into 6 daily categories typical  
of Antarctic PNSD data (referred to as simply the PNSD clusters) for further analysis based on  
the shape of the PNSD and their temporal variation (Figure 3a and 3b). These are Nucleation,  
Bursting, Aitken [high](#), Aitken [low](#), [Pristine](#), and [Bimodal](#), following previous classification  
described in previous studies (Dall'Osto et al., 2017b; Dall'Osto et al., 2019, Lachlan-Cope et  
400 al., 2020). The diurnal cycle of the PNSD belonging to each of the initial 16 K-means clusters  
at each of these sites is shown in Figure S4, while the diurnal cycle of the merged clusters is  
shown in Figure S5, separated by site, and the PNSD per cluster separate by season is shown  
in Figure S6.

Two clusters relate to NPF: Nucleation (mean particle count  $502 \pm 1087 \text{ cm}^{-3}$ ) has a mean PNSD  
with a peak at 12 nm averaged between all sites. This corresponds to fresh new particles  
405 generated during NPF events and will correspond to days of measurements dominated by a  
signature of new particles appearing in the PNSD, typically with signs of growth. Bursting  
(mean particle count  $177 \pm 350 \text{ cm}^{-3}$ ) has a mean PNSD with a peak at 22 nm averaged between  
all sites, and typically corresponds to days where new particles form, either fail to grow, are  
lost to coagulation, or are diluted and lost at the receptor site. [Note, the name "bursting" refers  
410 to "bursts" in particle number concentrations due to secondary formation, rather than bubble  
bursting.](#) These likely still do correspond to secondary formation processes. We presume here  
that there is no primary contribution to these PNSDs

Aitken is the most common PNSD category separated into Aitken high (mean particle count  
 $401.6 \pm 45.5 \text{ cm}^{-3}$ ), and Aitken low (mean particle count  $208.8 \pm 216.7 \text{ cm}^{-3}$ ). The mean PNSD  
415 shows peak concentrations at about 20-35 nm and 35-55 nm respectively. These two clusters  
are separated based on their aerosol total concentrations. The name of this category emerges  
from growing ultrafine aerosol resulting from the processing of local and regional marine  
aerosols, a phenomenon previously described to occur mainly in summer (Dall'Osto et al.,  
2017b).

420 Pristine (mean particle count  $142 \pm 134 \text{ cm}^{-3}$ ) has a mean PNSD with a peak at 88 nm. The name  
of this category emerges from the extremely low "pristine" particle number concentrations. The  
three K-means clusters belonging to this categories have some specific aerosol modes, peaking  
at 65 nm, 85 nm and a much larger accumulation mode at 160 nm (Figure S2) likely associated  
at pristine conditions with high loadings of blowing snow, as previously discussed in Lachlan-  
425 Cope et al. (2020).

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430 Bimodal (mean particle count  $107 \pm 133 \text{ cm}^{-3}$ ) has two peaks in the PNSD at 39 and 113 nm. Both are characterised by low particle counts. The name of this category is associated with the bimodal PNSD. The two initial clusters differ slightly because of the Hoppel minimum at 55 nm and 75 nm, respectively: [the Hoppel minimum refers to a specific dip in the number concentration of aerosol particles at these sizes, suggesting variations in particle stability, growth, or origin \(Hoppel et al., 1994\)](#).

**Eliminado:** , possibly indicating different chemistry and particle origin

435 The condensation sink represents the main loss process for many low-volatility vapours which contribute to new particle formation. The condensation sinks for the Bursting K-means cluster is lowest ( $2.3 \cdot 10^{-4} \text{ s}^{-1}$ ), followed by Bimodal ( $2.4 \cdot 10^{-4} \text{ s}^{-1}$ ), Pristine ( $3.4 \cdot 10^{-4} \text{ s}^{-1}$ ), Nucleation ( $3.7 \cdot 10^{-4} \text{ s}^{-1}$ ), and Aitken low ( $3.4 \cdot 10^{-4} \text{ s}^{-1}$ ), and Aitken high ( $4.0 \cdot 10^{-4} \text{ s}^{-1}$ ). Particle counts change  
440 by around 1 order of magnitude between sites, but the shapes of the PNSD are similar (Figure S2, S5). All Nucleation PNSDs show formation of and growth of particles at midday and afternoon times (local time), while Bursting show particles at or near the smallest sizes, sometimes with indication of particle growth (Figures S4, S5).

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### 3.2.2 Trends in PNSD K-means clusters

445 The frequency of occurrence of these different PNSD clusters can indicate the dominant sources of aerosol particles at each of these time periods. All sites have a large contribution from Aitken low and Aitken high PNSD clusters, with 37% of PNSDs falling into these categories across all four sites. Marambio is also dominated by PNSD clusters falling into the Nucleation and Bursting category, (25% and 23% of PNSDs, respectively). There is a smaller  
450 contribution of Nucleation and Bursting to the PNSDs at King Sejong (8% and 20% respectively). These PNSD clusters contribute little at Halley (3% and 3%, respectively), and at Concordia/Dome C (2% and 5%, respectively). Halley instead sees a large contribution of Pristine PNSDs (45% of PNSDs), and Concordia/Dome C sees a large contribution of Bimodal PNSDs (34% of PNSDs) (Figure 3a).

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455 The seasonal evolution of these K-means clusters is shown in Figure 3c. At Marambio, the contribution of Nucleation and Bursting PNSDs occurs in nearly all months. At King Sejong, Nucleation and Bursting PNSDs are most frequent in the spring and autumn months, while Pristine air masses are most common in the Austral winter. At Halley, Aitken PNSDs dominate the warmer months, while Pristine air masses dominate the colder winter months. At  
460 Concordia/Dome C, where data are available, Bimodal PNSDs dominate the winter, while

465 Aitken dominates the warmer months. There is a substantial contribution of Nucleation and  
Bursting in the month of September also. When all sites are averaged, a clear seasonal cycle is  
seen, with pristine and bimodal PNSDs dominating in the coldest months, and a substantial  
contribution of Aitken in the warmer months. Nucleation is slightly higher in the warmer  
months (particularly January through March), while bursting PNSDs are higher outside of these  
months.

### 470 3.3 Air mass analysis

72 hour HYSPLIT back trajectories arriving at the four different measurement sites for every  
hour with PNSD data were calculated. The amount of time each of these air masses spent  
flowing over land, open ocean, marginal ice, or sea ice per K-means cluster across all sites is  
plotted in Figure 4a, and per site is plotted in Figure 4b. On average, the air masses arriving at  
475 the sites had flown over mostly the land of the Antarctic continent (54%), with a 24%  
contribution from sea ice, 15% contribution from open water, and a small (6%) contribution  
from marginal ice. Nucleation & Bursting PNSDs occur when air masses have travelled under  
substantially less land and over more sea ice, marginal ice, and ocean than average (40%, 8%,  
and 26% of air mass hours for Nucleation, and 34%, 10%, and 25% of air mass hours for  
480 Bursting, respectively). At Marambio, the main contribution is air masses arising from sea-ice  
regions (50%) which is highest for Bimodal PNSDs (76%). The contribution of open water to  
Aitken, Bursting & Nucleation is higher than average (23%), while the contribution of sea ice  
to these air masses is lower (44%). At King Sejong, air masses spend the majority of their time  
travelling over regions of open water (41%) and sea ice (33%). The contribution of open water  
485 is higher for bimodal (51%) and Aitken PNSDs (58%). At Halley, air masses spend the majority  
of their time flowing over land regions (79%), with a small contribution from sea ice (15%).  
The contribution of sea ice to Nucleation PNSDs is substantially higher (30%). At  
Concordia/Dome C, air masses arriving at the receptor site mostly flow over land (99%). The  
contribution of sea ice, marginal ice, and open water are higher for bursting PNSDs (9% in  
490 total).

Our CWT analysis grids back trajectories to 1x1 degree squares and weighs each segment of  
the back trajectory with the corresponding  $N_{tot}$  observed upon the air mass's arrival, performed  
individually for each PNSD cluster. These are plotted in Figure 5. A map highlighting source  
regions for  $N_{tot}$  unseparated by cluster per site is shown in Figure S8. The CWTs aggregated

Eliminado: (concentrated weighted trajectory)

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[together for each site are shown in](#) Figure S9. Mean heights of these trajectories are shown in Figure S10.

The sources of particles by number (or,  $N_{\text{tot}}$ ) across all PNSD data at the two Antarctic Peninsula sites Marambio and King Sejong are to the west of the peninsula, across the iced and open water regions of the Bellingshausen Sea and to a lesser extent the Weddell sea. Those at Halley are concentrated at the coastal and land-based regions, with some influence from the Weddell sea, while the highest aerosol concentrations at Concordia/Dome C arise from mostly land regions, with some trajectories extending past the southern tip of the continent over ocean regions (Figure S8).

Air masses arriving at the Antarctic sites corresponding to Nucleation & Bursting K-means clusters pass mostly over the Bellingshausen sea at Marambio and King Sejong. Air masses passing over the regions just west of the peninsula have a high source contribution to high number concentrations of particles especially, as do the more northerly regions off the southern tip of south America. At Halley, these PNSD clusters are mostly associated with coastal regions, consistent with the higher contribution of sea ice. The source contributions at Concordia/Dome C correspond to a small number of trajectories. All trajectories corresponding to Nucleation PNSDs have a lower-than-average trajectory height, [although the total number of trajectories this corresponds to is relatively small](#) (Figure S10).

Aitken PNSDs have strong source contributions from the entire Bellingshausen sea region at Marambio and King Sejong. At Halley, there is a strong contribution from land mass regions, while at Concordia/Dome C, land and regions mostly contribute to this PNSD. Bimodal and Pristine air masses at Marambio and King Sejong have strong source contributions from the iced regions either side of the Antarctic peninsula, covering the Weddel and Bellingshausen seas. This is consistent with the higher contribution from sea-ice regions to these K-means clusters. At Halley, the Bimodal PNSDs arise when air masses flow over coastal and land-based regions, while Pristine PNSDs, which are extremely frequent at this site, arise largely from the Antarctic continent itself. At Concordia/Dome C, there is a strong source contribution to Bimodal & Pristine PNSDs from air masses flowing over the coastal region of southern Antarctica, as well as some land regions. Trajectories corresponding to Pristine PNSDs have, on average, a higher trajectory height than average.

### 3.4 Intercomparison with other existing data across the year 2015

#### 3.4.1 PEGASO cruise

The PEGASO (Plankton-derived Emissions of trace Gases and Aerosols in the Southern Ocean) cruise was conducted on board the RV Hesperides in the regions of Antarctic Peninsula, South Orkney and South Georgia Islands from 2 January to 11 February 2015. It was found that the microbiota of sea ice and the sea ice-influenced ocean can be a source of atmospheric primary and secondary organic nitrogen (ON), specifically low molecular weight alkylamines (Dall'Osto et al., 2017; 2019). Other follow up studies also claim that the potential impact of the sea ice (sympagic) planktonic ecosystem on aerosol composition were overlooked in past studies, and multiple eco-regions act as distinct aerosol sources around Antarctica (Decesari et al., 2020; Rinaldi et al., 2020).

Figure 6a shows the intercomparison of MPSS PNSDs for the PEGASO cruise. These are separated into times when the air masses flowed over the Pacific ocean, and over the Weddel sea. This is compared with stations used for this study where overlapping data are available (Marambio, King Sejong, and Halley). Air masses from the Southern Pacific Ocean with anthropogenic and terrestrial influence from Patagonia are likely responsible for the higher Aitken mode seen for the "Pacific" PNSD during the PEGASO cruise. By contrast, cruise measurements influenced by the Weddel sea show a PNSD similar to the three stations intercompared, suggesting little anthropogenic influence of the latter. Dall'Osto et al., (2017) showed that sea ice regions are a strong source of sub-3 nm particles in Antarctica relative to open water regions. The present study, considering a much broader dataset, also shows that the Open Water (OW) regions sampled at Marambio and at King Sejong stations are also associated with enhanced NPF.

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#### 3.4.2 Kohnen station

Weller et al., (2018) reported PNSDs and conducted bulk and size-segregated aerosol sampling during two summer campaigns in January 2015 and January 2016 at the continental Antarctic station Kohnen (Dronning Maud Land). The transport of aerosols was associated with two main weather conditions: (1) the sporadic impact of cyclones causing high loadings of marine aerosol concentrations and NPF and (2) clear sky conditions causing long range transport of aged aerosols. Figure 6b shows similar aerosol PNSDs among Kohnen, Halley and King Sejong

stations for the same sampling period in 2015. By contrast, much higher concentrations of ultrafine particles - likely due to NPF - were seen occurring at Marambio. This may be a combination of local sources (nearby penguin colonies) and melting sea ice during the summer period, given Marambio is the station among all closest to marginal sea ice zones (Quelever et al., 2022).

## 4 Discussion

### 4.1 Possible primary sources

Of our six aerosol categories, nucleation and bursting are related to secondary aerosols, while the two Aitken K-means clusters likely have significant contributions from primary and secondary processes (Figure 2a), although the latter may dominate. Bimodal PNSDs are likely associated with aged and processed marine aerosols (the Hoppel minimum mode is seen at about 70 nm, Hoppel et al., 1994). We cannot apportion primary aerosols from our PNSD measurements without any chemical composition information, but can hypothesise some. Sea salt aerosol particles are formed from open ocean regions when sea spray is produced by wave breaking and bubble bursting generating film and jet drops (De Leeuw et al., 2011). Sea salt aerosol is also formed from blowing snow above sea ice (Frey et al., 2020). These mechanisms are thought to contribute equally to sea salt aerosol loads (Legrand et al., 2016), while blowing snow over land is a source from continental Antarctica (Giordano et al., 2018).

In the periods where NPF is negligible, there are still other primary aerosol sources. We show that pristine and bimodal PNSD clusters dominate through the winter (Figure 3c, S6). It is therefore important to characterize these pristine conditions. The three K-means clusters of the pristine categories have similar low particle number concentrations, but one of the three has a distinct PNSD. The cluster (pristine\_160) seen with a bimodal PNSD (75 and 160 nm, respectively), was strongly associated with high wind speed and possibly associated with blowing snow and sea spray sea salt, dominating the winter aerosol population (Lachlan-Cope et al., 2020). When the four stations are compared in this study, it can clearly be seen that Halley has the most frequent pristine conditions (45%, relative to the 12-20% of the other three), also relative to Concordia/Dome C where we propose that free troposphere transport and aged aerosols populate the PNSD with Aitken modes and bimodal PNSDs.



The pristine initial PNSD cluster with a strong peak at 160nm (table S1) is seen mainly at Halley (12%) relative to the other stations (2-3%). A study by Yang et al. (2019) proposes a source for ultrafine sea salt aerosol particles from blowing snow, dependent on snow salinity.

590 This mechanism could account for the small particles seen during Antarctic winter at coastal stations (Giordano et al., 2018; Frey et al., 2020). Recently, similar PNSD were reported by Gong et al., (2023) in the central Arctic over an entire year from September 2019 to October 2020 during the Multidisciplinary drifting Observatory for the Study of Arctic Climate (MOSAIC) expedition, showing that blowing snow was observed more than 20% of the time  
605 from November to April. The sublimation of blowing snow generates high concentrations of fine-mode sea salt aerosol (diameter below 300 nm), enhancing cloud condensation nuclei concentrations up to tenfold above background levels (Gong et al., 2023).

Primary marine organic aerosol in the submicron regime is likely produced separately from sea salt (Gantt and Meskhidze, 2013). At the coastal sites of Marambio and Halley, Bimodal and  
600 Pristine PNSDs with the highest particle number concentration arise mostly from the Bellingshausen Sea, where DMS(P)-rich phytoplankton concentrations are very high. A dominant biological oceanic mechanism of primary particle origin at these sites is therefore likely. Given the vastly different source characteristics between these sites, it is likely that the dominant mechanisms of origin of these primary particles, and therefore our bimodal and  
605 pristine clusters also differ between sites, but it is clear that primary emissions dominate the PNSD in the Antarctic winter.

#### 4.2 New particle formation and possible secondary sources

We identify two PNSD clusters which we classify as arising from secondary processes, these are Nucleation and Bursting PNSDs. The Aitken PNSD clusters cannot be definitively said to  
610 contain solely particles from secondary processes, although they have similar source regions (Figure S5) and could consist of grown particles formed either elsewhere in the boundary layer or in the free troposphere, with average growth rates around  $\sim 0.1 - 1 \text{ nm h}^{-1}$  (Brean et al., 2021; Järvinen et al., 2013; Jokinen et al., 2018; Kim et al., 2019), particles formed through NPF would grow to the sizes observed in this PNSD cluster on the order of  $\sim 2$  days. The contribution  
615 of Aitken PNSDs is greatest outside of the winter months (Figure S7). However, below, we will purely discuss the contribution of the Bursting and Nucleation PNSDs.

NPF in Antarctica is a summertime phenomenon largely responsible for the seasonal cycle in particle number concentrations (Figure 2c). This cycle has been observed at at King Sejong Station (Kim et al., 2017; Park et al., 2023; Kim et al., 2019), Halley (Lachlan-Cope et al., 2020) and Concordia/Dome C (Järvinen et al., 2013; Chen et al., 2017). This corresponds with periods of high biological and photochemical activity (Jang et al., 2019; Kim et al., 2019). These observations are consistent with our observed higher frequency of Nucleation PNSD clusters in January through March (Figure 3c, S6). However, we consistently observe Nucleation and Bursting PNSDs year-round, even at sites where manual inspection of these PNSDs has found little to no NPF in the winter seasons (Kim et al., 2019).

We frequently observe PNDs with a large nucleation mode in the Austral summer (Figure S7). NPF is mainly influenced by the source rates of vapours, including emissions of dimethylsulfide (DMS), volatile organic compounds (VOCs), bases such as NH<sub>3</sub>, and iodine, as well as OH<sup>-</sup> and O<sub>3</sub> production rates, loss rates of vapours and new particles (CS), ion pair production rates, and temperature (Lee et al., 2019). Vapour source rates will be lowest in winter, however, ion pair production from cosmic rays is likely constant, while loss rates of vapours and temperatures have a wintertime maximum. Some number of particles may therefore form and be identified by Cluster analysis, even when they do not give the visual signature of an NPF event.

Our analyses show that coastal Antarctic sites nearest to the melting sea ice are most influenced by NPF (Figure 3c) where a large fraction of the PNSDs are classified as Nucleation and Bursting (48 and 28% at Marambio and King Sejong respectively). Further, we show that NPF is frequent when air masses flow over the ice-influenced oceanic Antarctic regions (Figure 4,5). NPF events have been shown to be frequent and strong at coastal Antarctic sites (Brean et al., 2021; Jokinen et al., 2018), but most rapid at those nearest to melting ice (Brean et al., 2021) as well as near local sources associated with penguin colonies (Quelever et al., 2022). Recent observations shows that this does not extend across the whole of the Southern Ocean (Baccarini et al., 2021). Our PNSD cluster analysis is consistent with this. Melting sea is therefore a likely source of NPF precursors in the Antarctic Peninsula region.

High aerosol number concentrations arising from the Bellingshausen sea is consistent with a previous analysis of NPF at King Sejong (Jang et al., 2019). Bursting and nucleation aerosol categories in particular exhibit high number concentrations when air masses pass over the regions just west of the peninsula (Figure 5). These regions have been shown to be rich in

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alkylamines (Dall'Osto et al., 2019a) that lead to rapid and efficient particle formation (Brean et al., 2021). There is evidence that these melting sea-ice regions are sources of numerous NPF precursors (Atkinson et al., 2012; Dall'Osto et al., 2017a, 2019a). This is also consistent in the sympagic Arctic environment (Dall'Osto et al., 2017b, 2018). Consistent with this, the results of our K-means cluster analysis show that the sites with the highest exposure to air masses that pass over regions of sea ice and marginal ice have by far the greatest contribution from nucleation and bursting PNSD types (Figure 2, 4). The sites where air masses primarily travel over snow-covered land regions have, by contrast, very little contribution of nucleation and bursting (Figure 4a).

The role of new particles formed in the free-troposphere and transported down to our receptor sites is unknown, but likely. Particle formation processes are highly efficient in the free troposphere given the lower temperatures and higher ion pair production rates. A wider number of potential nucleation mechanisms are therefore likely (Kirkby et al., 2011; Wang et al., 2022). Modelling studies predict a substantial fraction of CCN in all seasons (Korhonen et al., 2008). These particles, if arriving at the receptor site long enough after formation, will contribute either to Nucleation, Bursting, or Aitken PNSDs. While we therefore highlight source regions leading to boundary layer NPF processes, we cannot state what fraction of PNSDs arise from free tropospheric particle formation processes.

## 5 Implication and conclusion

Figure 7 shows a schematic illustration of the sea ice, microbiota, sea-to-air emissions, and primary and secondary aerosols in Antarctica. Figure 7 highlights the dominance of NPF in summertime PNSDs, and a dominance of primary aerosols during the wintertime, with these primary aerosols being more prevalent inland than at the coast, a key finding of this study. It also highlights the retreat of sea ice in the summer, leading to increased marine emissions, alongside a reduction in terrestrial biological activity and sunlight intensity during winter months.

To gain insight into the influence of marine Antarctic biogeochemistry on atmospheric aerosol, we report simultaneous aerosol PNSDs collected across an entire year (2015) at four research stations (Marambio, King Sejong, Halley, and Concordia/Dome C), as well as a brief intercomparison with a cruise around the regions of Antarctic Peninsula, South Orkney, and South Georgia Islands (Dall'Osto et al., 2017) and a field campaign in Kohnen station (Weller

**Movido hacia arriba[1]:** During the austral summer the concentration of DMS in the water of the Southern Ocean is the highest of the planet (Lana et al., 2011), with high fluxes into the atmosphere, potentially producing high concentrations of sulphuric acid and methanesulphonic acid. However, under typical boundary layer conditions the concentration of sulphuric acid is too low to form particles alone, and another molecule, such as ammonia, is required to stabilize the nucleating clusters (Kirkby et al., 2011). Jokinen et al. (2018) reported the first molecular characterization of NPF from the Aboa station in Antarctica, showing that new particles are formed via clustering of sulfuric acid and ammonia (Kirby et al., 2011). Sources of ammonia and alkylamines are related to animals, mainly birds and seal colonies and melting sea ice, but the relative importance of each source is unknown (Riddick et al., 2012; Schmale et al., 2013; Brean et al., 2021; Quelever et al., 2022). Iodine oxoacids are also capable of efficiently forming new particles (Baccarini et al., 2020; He et al., 2021; Sipilä et al., 2016), with iodosous acid being able to efficiently stabilise clusters of iodic and sulphuric acid, even in the absence of ammonia (He et al., 2023), with modelling studies showing high gas-phase iodine concentrations across Antarctica (Saiz-Lopez et al., 2007). Further, the formation of marine primary aerosol particles and of gas-to-particles precursors is influenced by the uppermost layer of the ocean, the so called sea surface microlayer (SML) (Cunliffe et al., 2013). The SML - with physicochemical characteristics different from those of subsurface waters (SSW) - results in dense and active viral and microbial communities (Vaquer et al., 2021; Heinrichs et al., 2024). The microlayer of the ocean is a source of oxygenated VOCs in the Arctic leading to secondary aerosol formation (Mungall et al., 2017), and it is also likely a source in the Southern Ocean too although no measurements of this type are available. ¶ Therefore, while the limited number of measurement studies implicate sulphuric acid and nitrogenous bases as the drivers of NPF, iodine and VOCs may play a substantial role. ¶

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et al., 2018). Our study shows that the aerosol PNSDs across the Antarctic have striking differences, likely due to multiple eco-regions and subsequent atmospheric chemical and physical processes act as multiple aerosol sources around Antarctica. These analyses suggest that the PNSD of Antarctic sub-micrometre aerosols may have been oversimplified in the past (Ito, 1993) and that complex interactions between multiple ecosystems, coupled with different atmospheric circulation, result in very different PNSDs populating Antarctica. Our knowledge on aerosol sources of primary and secondary origin is limited.

The Southern Ocean is among the largest source of Sea Spray Aerosols (SSA) on planet Earth. Current aerosol models have a large uncertainty in the SSA abundance (Lapere et al., 2023) and the relative importance of the sublimation of blowing snow is not yet quantified (Giordano et al., 2018; Frey et al., 2020). The biogenic organic component of SSA in Antarctica is thought to be important (Mc Coy et al., 2015) but again not fully quantified. Sea ice may also modulates SSA production, with potentially significant climate impacts (Dall'Osto et al., 2022a). Other leached material from Antarctic media including seaweeds and penguin guano may also affect cloud-relevant aerosol SSA production (Dall'Osto et al., 2022b).

Recently, Brean et al. (2023) emphasized how understanding the geographical variation in surface types across the Arctic is key to understanding secondary aerosol sources, highlighting that particle formation and growth rate vary tremendously likely due to different regions producing different precursors source rates. The same complexity applies to Antarctica - an ensemble of regions with substantial spatial heterogeneity across marine, terrestrial, and freshwater biomes, with productivity and biodiversity patchiness super-imposed on strong environmental gradients. Antarctica, one of the world's eight major biogeographical realms, is split into 16 Antarctic Conservation Biogeographic Regions (also known as ecoregions or bioregions, Terauds et al., 2012, 2016), but is made up almost entirely of the ice-covered land mass, coastal tundra, and sea ice surrounding the main continent. It contains two additional marine bioregions - the Antarctic Peninsula & Scotia Sea and the Subantarctic Indian Ocean Islands (Chown et al., 2007). A bioregion is smaller in scale than a biogeographical realm but larger than an eco-region or an ecosystem, it allows the integrations of multiple eco-regions, including terrestrial, freshwater and marine into a cohesive system. Whilst these definitions may be challenging in the context of atmospheric biogeochemistry, we argue a better understanding of the interactions between the biosphere and the geosphere are needed to better understand aerosol sources in Antarctica.

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**Eliminado:** Recently, Brean et al., (2023) emphasized how understanding the geographical variation in surface types across the Arctic is key to understanding secondary aerosol sources, highlighting that particle formation and growth rate vary tremendously likely due to different regions producing different precursors source rates. We argue that the same complexity applies to Antarctica - an ensemble of regions with substantial spatial heterogeneity across marine, terrestrial, and freshwater biomes, with productivity and biodiversity patchiness super-imposed on strong environmental gradients. Antarctica, one of the world's eight major biogeographical realms, is made up almost entirely of one single bioregion - the ice-covered land mass, coastal tundra, and sea ice of the main continent. It contains two additional marine bioregions - the Antarctic Peninsula & Scotia Sea and the Subantarctic Indian Ocean Islands (Chown et al., 2007; Terauds et al., 2012, 2016). A bioregion is smaller in scale than a biogeographical realm but larger than an eco-region or an ecosystem, it allows the integrations of multiple eco-regions, including terrestrial, freshwater and marine into a cohesive system. Whilst these definitions may be challenging in the context of atmospheric biogeochemistry, we argue a better understanding of the interactions between the biosphere and the geosphere are needed to better understand aerosol sources in Antarctica. ¶

**Movido hacia arriba[2]:** Antarctic sea ice covers between 1% (summer) and 5% (winter) of the global ocean. Antarctic terrestrial productivity and biodiversity occur almost exclusively in ice-free areas that cover less than 1% of the continent. In Antarctica the coastline extends for 17,968 km and comprises about 34.8-36.4x10<sup>6</sup> km<sup>2</sup> where 80% of this surface is covered by ice, even in summer (Peck et al., 2018; Ronowicz et al., 2019). Overall, there is about 390,071 km<sup>2</sup> of coast shallower than 200 m (Peck et al., 2018). Antarctic coastal systems harbour a high diversity of marine and terrestrial ecosystems including Antarctic seaweeds (benthonic macroalgae) and bird colonies (mainly penguins). Antarctic seaweeds (often called macroalgae) are found in shallow (<200m) coastal, rocky shores and can cover more than 80% of the benthic surface (Amsler et al., 2005; Wiencke and Amsler, 2012). The Antarctic and Sub-Antarctic region is home for about half of the total worldwide seabird population (Otero et al., 2018). Penguins represent a high proportion of the avian biomass, and their fecal material is one of the main source of phosphorus and nitrogen, representing about 80% of these elements in the Antarctic marine environment. Seabird colonies also represent a significant source of atmospheric ammonia (NH<sub>3</sub>) in remote maritime systems (Riddick et al., 2012; Schmale et al., 2013). These emissions are environmentally relevant as they primarily occur as "hot-spots" in otherwise pristine environments.¶

**Eliminado:** Figure 7 (a, b) shows a schematic illustration of the sea ice, microbiota, sea-to-air emissions, and primary and secondary aerosols in Antarctica.

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The changes occurring in the Antarctic environment will modify the climate with feedbacks and exchanges between the biosphere and cryosphere with the atmosphere. Changes in marine and terrestrial life - including adaptation of ecophysiology, food and nutrient availability - will affect the emissions of primary and secondary aerosol precursors. Interdisciplinary studies and international cooperation in Antarctica are reducing the gap in our knowledge of these key environmental factors.

#### **Code and data availability.**

The code and data used to produce all non-illustrative figures are available from the corresponding authors under reasonable request.

#### **Supplement.**

The supplement related to this article is available online at: X

#### **Author contributions.**

Conceptualization: MDO conceptualized the paper following the study of Lachlan-Cope et al (2020).

Data Curation: EA provided data for Marambio; TLC and AJ provided data for Halley, AL, AV, PA provided data for Concordia/Dome C, and JP and YJY provided data for King Sejong. Software: DCSB and JB.

Formal Analysis: DCSB, JB, MDO.

Funding Acquisition: See acknowledgement.

Methodology: DCBS, JB.

Project Administration: MDO.

Resources: See acknowledgement.

Validation: Validated by all co-authors.

Visualization: JB.

Writing: JB, MDO

Review & Editing: all authors.

#### **Competing interests.**

At least one of the (co-)authors is a member of the editorial board of Atmospheric Chemistry and Physics.

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850

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#### **Review statement.**

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855

#### **References**

Amsler et al., 2005 C.D. Amsler, K. Iken, J.B. McClintock, M.O. Amsler, K.J. Peters, J.M. Hubbard, F.B. Furrow, B.J. Baker Comprehensive evaluation of the palatability and chemical defenses of subtidal macroalgae from the Antarctic peninsula *Mar. Ecol. Prog. Ser.*, 294 (2005), pp. 141-159

860

Arrigo, K. R., Lizotte, M. P., Mock, T. Primary producers and sea ice. *Science* 2009 pp 283–326.

Arrigo, K. R.; van Dijken, G. L.; Strong, A. L. Environmental controls of marine productivity hot spots around Antarctica. *J. Geophys Res. - Oceans* 2015, 120, 5545–5565.

865

[Asmi, E., Neitola, K., Teinilä, K., Rodriguez, E., Virkkula, A., Backman, J., Bloss, M., Jokela, J., Lihavainen, H., De Leeuw, G., Paatero, J., Aaltonen, V., Mei, M., Gambarte, G., Copes, G., Albertini, M., Perez Fogwill, G., Ferrara, J., Elena Barlasina, M. and Sanchez, R.: Primary sources control the variability of aerosol optical properties in the Antarctic Peninsula, \*Tellus B: Chemical and Physical Meteorology\*, vol. 70 , 2018. <https://doi.org/10.1080/16000889.2017.1414571>](#)

870

Atkinson, H. M., Huang, R. J., Chance, R., Roscoe, H. K., Hughes, C., Davison, B., Schönhardt, A., Mahajan, A. S., Saiz-Lopez, A., Hoffmann, T. and Liss, P. S.: Iodine emissions from the sea ice of the Weddell Sea, *Atmos. Chem. Phys.*, 12(22), 11229–11244, doi:10.5194/acp-12-11229-2012, 2012.

875

Baccarini, A., Karlsson, L., Dommen, J., Duplessis, P., Vüllers, J., Brooks, I. M., Saiz-lopez, A., Salter, M., Tjernström, M., Baltensperger, U., Zieger, P. and Schmale, J.: Frequent new particle formation over the high Arctic pack ice by enhanced iodine emissions, *Nat. Commun.*, (2020), 1–11, doi:10.1038/s41467-020-18551-0, 2020.

880

Baccarini, A., Dommen, J., Lehtipalo, K., Henning, S., Modini, R. L., Geysel-Ber, M., Baltensperger, U. and Schmale, J.: Low-Volatility Vapors and New Particle Formation Over the Southern Ocean During the Antarctic Circumnavigation Expedition, *J. Geophys. Research Atmos.*, (126), 2021.

885

Beddows, D. C. S., Dall'Osto, M., Harrison, R. M., Dall'Osto, M., Harrison, R. M., Dall'Osto, M. and Harrison, R. M.: Cluster Analysis of Rural, Urban, and Curbside Atmospheric Particle Size Data, *Environ. Sci. Technol.*, 43(13), 4694–4700, doi:10.1021/es803121t, 2009.

890

Brean, J., Dall'Osto, M., Simó, R., Shi, Z., Beddows, D. C. S. and Harrison, R. M.: Open ocean and coastal new particle formation from sulfuric acid and amines around the Antarctic Peninsula, *Nat. Geosci.*, doi:10.1038/s41561-021-00751-y, 2021.

895

Brean, J., Beddows, D. C. S., Harrison, R. M., Song, C., Tunved, P., Ström, J., Krejci, R., Freud, E., Massling, A., Skov, H., Asmi, E., Lupi, A., and Dall'Osto, M.: Collective geographical ecoregions and precursor sources driving Arctic new particle formation, *Atmos. Chem. Phys.*, 23, 2183–2198, <https://doi.org/10.5194/acp-23-2183-2023>, 2023.

**Eliminado:** Asmi, E., Frey, A., Virkkula, A., Ehn, M., Manninen, H. E., Ti- monen, H., Tolonen-Kivimäki, O., Aurela, M., Hillamo, R., and Kulmala, M.: Hygroscopicity and chemical composition of Antarctic sub-micrometre aerosol particles and observations of new particle formation, *Atmos. Chem. Phys.*, 10, 4253–4271, <https://doi.org/10.5194/acp-10-4253-2010>, 2010

- 905 Carslaw, K. S., Lee, L. A., Reddington, C. L., Pringle, K. J., Rap, A., Forster, P. M., Mann, G. W., Spracklen, D. V., Woodhouse, M. T., Regayre, L. A. and Pierce, J. R.: Large contribution of natural aerosols to uncertainty in indirect forcing, *Nature*, 503(7474), 67–71, doi:10.1038/nature12674, 2013.
- 910 Clem, K. R., Fogt, R. L., Turner, J., Lintner, B. R., Marshall, G. J., Miller, J. R. and Renwick, J. A.: Record warming at the South Pole during the past three decades, *Nat. Clim. Chang.*, 10(8), 762–770, doi:10.1038/s41558-020-0815-z, 2020.
- Chen, J. L., Wilson, C. R., Blankenship, D., and Tapley, B. D.: Accelerated Antarctic ice loss from satellite gravity measurements, *Nat. Geosci.*, 2, 859–862, <https://doi.org/10.1038/ngeo694>, 2009.
- 915 Chen, X., Virkkula, A., Kerminen, V.-M., Manninen, H. E., Busetto, M., Lanconelli, C., Lupi, A., Vitale, V., Del Guasta, M., Grigioni, P., Väänänen, R., Duplissy, E.-M., Petäjä, T., and Kulmala, M.: Features in air ions measured by an air ion spectrometer (AIS) at Dome C, *Atmos. Chem. Phys.*, 17, 13783–13800, <https://doi.org/10.5194/acp-17-13783-2017>, 2017.
- 920 Chown, S.L. & Convey, P. (2007) Spatial and temporal variability across life's hierarchies in the terrestrial Antarctic. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 362: 2307–2331
- Cunliffe, M.; Engel, A.; Frka, S.; Gasparovic, B.; Guitart, C.; Murrell, J.C.; Salter, M.; Stolle, C. Sea surface microlayers: A unified physicochemical and biological perspective of the air-ocean interface. *Prog. Oceanogr.* 2013, 1109, 104–116.
- 925 Dall'Osto, M., Ovadnevaite, J., Paglione, M., Beddows, D. C. S., Ceburnis, D., Cree, C., Cortés, P., Zamanillo, M., Nunes, S. O., Pérez, G. L., Ortega-Retuerta, E., Emelianov, M., Vaqué, D., Marrasé, C., Estrada, M., Sala, M. M., Vidal, M., Fitzsimons, M. F., Beale, R., Airs, R., Rinaldi, M., Decesari, S., Facchini, M. C., Harrison, R. M., O'Dowd, C. and Simó, R.: Antarctic sea ice region as a source of biogenic organic nitrogen in aerosols, *Sci. Rep.*, 7(1), 1–10, doi:10.1038/s41598-017-06188-x, 2017a.
- 930 Dall'Osto, M., Beddows, D. C. S., Tunved, P., Krejci, R., Ström, J., Hansson, H. C., Yoon, Y. J., Park, K. T., Becagli, S., Udristi, R., Onasch, T., O'Dowd, C. D., Simó, R. and Harrison, R. M.: Arctic sea ice melt leads to atmospheric new particle formation, *Sci. Rep.*, 7(1), 1–10, doi:10.1038/s41598-017-03328-1, 2017b.
- 935 Dall'Osto, M., Geels, C., Beddows, D. C. S., Boertmann, D., Lange, R., Nøjgaard, J. K., Harrison, R. M., Simo, R., Skov, H. and Massling, & A.: Regions of open water and melting sea ice drive new particle formation in North East Greenland OPEN, *Sci. Rep.*, 8, 6109, doi:10.1038/s41598-018-24426-8, 2018.
- 940 Dall'Osto, M., Airs, R. L., Beale, R., Cree, C., Fitzsimons, M. F., Beddows, D., Harrison, R. M., Ceburnis, D., O'Dowd, C., Rinaldi, M., Paglione, M., Nenes, A., Decesari, S. and Simó, R.: Simultaneous Detection of Alkylamines in the Surface Ocean and Atmosphere of the Antarctic Sympagic Environment, *ACS Earth Sp. Chem.*, 3(5), 854–862, doi:10.1021/acsearthspacechem.9b00028, 2019a.
- 945 Dall'Osto, M., Beddows, D. C. S., Tunved, P., Harrison, R. M., Lupi, A., Vitale, V., Becagli, S., Traversi, R., Park, K. T., Jun Yoon, Y., Massling, A., Skov, H., Lange, R., Strom, J. and Krejci, R.:



- 950 Simultaneous measurements of aerosol size distributions at three sites in the European high Arctic, *Atmos. Chem. Phys.*, 19(11), 7377–7395, doi:10.5194/acp-19-7377-2019, 2019b.
- Dall'Osto, Manuel, Ana Sotomayor-Garcia, Miguel Cabrera-Brufau, Elisa Berdalet, Dolors Vaqué, Sebastian Zeppenfeld, Manuela van Pinxteren, Hartmut Herrmann, Heike Wex, Matteo Rinaldi, Marco Paglione, David Beddows, Roy Harrison, Conxita Avila, Rafael P. Martin-Martin, Jiyeon Park, Andrés Barbosa, Leaching material from Antarctic seaweeds and penguin guano affects cloud-relevant aerosol production, *Science of The Total Environment*, Volume 831, 2022, 154772, ISSN 0048-9697, <https://doi.org/10.1016/j.scitotenv.2022.154772>.
- 955
- Decesari, S., Paglione, M., Rinaldi, M., Dall'Osto, M., Simó, R., Zanca, N., Volpi, F., Facchini, M. C., Hoffmann, T., Götz, S., Kampf, C. J., O'Dowd, C., Ceburnis, D., Ovadnevaite, J., and Tagliavini, E.: Shipborne measurements of Antarctic submicron organic aerosols: an NMR perspective linking multiple sources and bioregions, *Atmos. Chem. Phys.*, 2020, 20, 4193–4207, <https://doi.org/10.5194/acp-20-4193-2020>
- 960
- De Leeuw, G., Andreas, E. L., Anguelova, M. D., Fairall, C. W., Lewis, E. R., O'Dowd, C., Schulz, M. and Schwartz, S. E.: Production flux of sea spray aerosol, *Rev. Geophys.*, 49(2), 1–39, doi:10.1029/2010RG000349, 2011.
- 965
- Draxler, R. R. and Rolph, G. D.: HYSPLIT (HYbrid Single-Particle Lagrangian Integrated Trajectory) Model access via NOAA ARL READY Website, [online] Available from: <https://ready.arl.noaa.gov/HYSPLIT.php>, 2010.
- 970
- Fetterer, F., Knowles, K., Meier, W. N., Savoie, M., and Wind-nagel, A. K.: Sea Ice Index, Version 3. [Indicate subset used]. Boulder, Colorado USA, NSIDC: National Snow and Ice Data Center, <https://doi.org/https://doi.org/10.7265/N5K072F8.2019>, 2017, updated daily.
- 975
- Fiebig, M., Hirdman, D., Lunder, C. R., Ogren, J. A., Solberg, S., Stohl, A., and Thompson, R. L.: Annual cycle of Antarctic baseline aerosol: controlled by photooxidation-limited aerosol formation, *Atmos. Chem. Phys.*, 14, 3083–3093, <https://doi.org/10.5194/acp-14-3083-2014>, 2014.
- 980
- Fossum, K. N., Ovadnevaite, J., Ceburnis, D., Dall'Osto, M., Marullo, S., Bellacicco, M., Simó, R., Liu, D., Flynn, M., Zuend, A. and O'Dowd, C.: Summertime Primary and Secondary Contributions to Southern Ocean Cloud Condensation Nuclei, *Sci. Rep.*, 8(1), 1–14, doi:10.1038/s41598-018-32047-4, 2018.
- 985
- Frey, M. M., Norris, S. J., Brooks, I. M., Anderson, P. S., Nishimura, K., Yang, X., Jones, A. E., Nerentorp Mastromonaco, M. G., Jones, D. H. and Wolff, E. W.: First direct observation of sea salt aerosol production from blowing snow above sea ice, *Atmos. Chem. Phys.*, 20(4), 2549–2578, doi:10.5194/acp-20-2549-2020, 2020.
- 990
- Gantt, B. and Meskhidze, N.: The physical and chemical characteristics of marine primary organic aerosol: A review, *Atmos. Chem. Phys.*, 13(8), 3979–3996, doi:10.5194/acp-13-3979-2013, 2013.

- 995 Giordano, M. R., Kalnajs, L. E., Douglas Goetz, J., Avery, A. M., Katz, E., May, N. W., Leemon, A., Mattson, C., Pratt, K. A. and DeCarlo, P. F.: The importance of blowing snow to halogen-containing aerosol in coastal Antarctica: Influence of source region versus wind speed, *Atmos. Chem. Phys.*, 18(22), 16689–16711, doi:10.5194/acp-18-16689-2018, 2018.
- 1000 Gras, L. J. and Keywood, M.: Cloud condensation nuclei over the Southern Ocean: Wind dependence and seasonal cycles, *Atmos. Chem. Phys.*, 17(7), 4419–4432, doi:10.5194/acp-17-4419-2017, 2017.
- Gong, X., Zhang, J., Croft, B., Yang, X., Frey, M.M., Bergner, N., Chang, R.Y.-W., Creamean, J.M., Kuang, C., Martin, R.V., Ranjithkumar, A., Sedlacek, A.J., Uin, J., Willmes, S., Zawadowicz, M.A., Pierce, J.R., Shupe, M.D., Schmale, J., Wang, J., 2023. Arctic warming by abundant fine sea salt aerosols from blowing snow. *Nat. Geosci.* 16, 768–774. <https://doi.org/10.1038/s41561-023-01254-8>
- 1005 Hamilton, D. S., Lee, L. A., Pringle, K. J., Reddington, C. L., Spracklen, D. V. and Carslaw, K. S.: Occurrence of pristine aerosol environments on a polluted planet, *Proc. Natl. Acad. Sci. U. S. A.*, 111(52), 18466–18471, doi:10.1073/pnas.1415440111, 2014.
- 1010 Hara, K., Osada, K., Nishita-Hara, C., and Yamanouchi, T.: Seasonal variations and vertical features of aerosol particles in the Antarctic troposphere, *Atmos. Chem. Phys.*, 11, 5471–5484, <https://doi.org/10.5194/acp-11-5471-2011>, 2011
- 1015 Hara, K., Nishita-Hara, C., Osada, K., Yabuki, M., and Yamanouchi, T.: Characterization of aerosol number size distributions and their effect on cloud properties at Syowa Station, Antarctica, *Atmos. Chem. Phys.*, 21, 12155–12172, <https://doi.org/10.5194/acp-21-12155-2021>, 2021.
- 1020 Heinrichs, M.E. et al. (2024). Breaking the Ice: A Review of Phages in Polar Ecosystems. In: Tumban, E. (eds) Bacteriophages. Methods in Molecular Biology, vol 2738. Humana, New York, NY. [https://doi.org/10.1007/978-1-0716-3549-0\\_3](https://doi.org/10.1007/978-1-0716-3549-0_3)
- 1025 [Heintzenberg, J., Covert, DS and Van Dingenen, R. 2000. Size distribution and chemical composition of marine aerosols: A compilation and review. Tellus, 52B: 1104–1122. DOI: <https://doi.org/10.1034/j.1600-0889.2000.00136.x>](#)
- 1030 Heintzenberg, J, Legrand, M, Gao, Y, Hara, K, Huang, S, Humphries, RS, Kamra, AK, Keywood, MD and Sakerin, SM. 2023. Spatio-Temporal Distributions of the Natural Non-Sea-Salt Aerosol Over the Southern Ocean and Coastal Antarctica and Its Potential Source Regions. *Tellus B: Chemical and Physical Meteorology*, 75(1): 47–64. DOI: <https://doi.org/10.16993/tellusb.1869>
- 1035 Herenz, P., Wex, H., Mangold, A., Laffineur, Q., Gorodetskaya, I. V., Fleming, Z. L., Panagi, M., and Stratmann, F.: CCN measurements at the Princess Elisabeth Antarctica research station during three austral summers, *Atmos. Chem. Phys.*, 19, 275–294, <https://doi.org/10.5194/acp-19-275-2019>, 2019.

#### Eliminado: ¶

Humphries, R. S., Schofield, R., Keywood, M. D., Ward, J., Pierce, J. R., Gionfriddo, C. M., Tate, M. T., Krabbenhoft, D. P., Galbally, I. E., Molloy, S. B., Klekociuk, A. R., Johnston, P. V., Kreher, K., Thomas, A. J., Robinson, A. D., Harris, N. R. P., Johnson, R., and Wilson, S. R.: Boundary layer new particle formation over East Antarctic sea ice – possible Hg-driven nucleation?, *Atmos. Chem. Phys.*, 15, 13339–13364, <https://doi.org/10.5194/acp-15-13339-2015>, 2015¶

¶

Humphries, R. S., Keywood, M. D., Ward, J. P., Hamwell, J., Alexander, S. P., Klekociuk, A. R., Hara, K., McRobert, I. M., Protat, A., Alroe, J., Cravigan, L. T., Miljevic, B., Ristovski, Z. D., Schofield, R., Wilson, S. R., Flynn, C. J., Kulkarni, G. R., Mace, G. G., McFarquhar, G. M., Chambers, S. D., Williams, A. G., and Griffiths, A. D.: Measurement report: Understanding the seasonal cycle of Southern Ocean aerosols, *Atmos. Chem. Phys.*, 23, 3749–3777, <https://doi.org/10.5194/acp-23-3749-2023>, 2023.¶

**Movido hacia abajo[3]:** He, X.-C., Simon, M., Iyer, S., Xie, H.-B., Rörup, B., Shen, J., Finkenzeller, H., Stolzenburg, D., Zhang, R., Baccarini, A., Tham, Y. J., Wang, M., Amanatidis, S., Piedehierro, A. A., Amorim, A., Baalbaki, R., Brasseur, Z., Caudillo, L., Chu, B., Dada, L., Duplissy, J., El Haddad, I., Flagan, R. C., Granzin, M., Hansel, A., Heinritzi, M., Hofbauer, V., Jokinen, T., Kempainen, D., Kong, W., Krechmer, J., Kürten, A., Lamkaddam, H., Lopez, B., Ma, F., Mahfouz, N. G. A., Makhmutov, V., Manninen, H. E., Marie, G., Marten, R., Massabò, D., Mauldin, R. L., Mentler, B., Onnela, A., Petäjä, T., Pfeifer, J., Philippov, M., Ranjithkumar, A., Rissanen, M. P., Schobesberger, S., Scholz, W., Schulze, B., Surdu, M., Thakur, R. C., Tomé, A., Wagner, A. C., Wang, D., Wang, Y., Weber, S. K., Welti, A., Winkler, P. M., Zauner-Wieczorek, M., Baltensperger, U., Curtius, J., Kurtén, T., Worsnop, D. R., Volkamer, R., Lehtipalo, K., Kirby, J., Donahue, N. M., Sipilä, M., and Kulmala, M.: Iodine oxoacids enhance nucleation of sulfuric acid particles in the atmosphere, *Science*, 382, 1308–1314, [10.1126/science.adh2526](https://doi.org/10.1126/science.adh2526), 2023.¶

#### Eliminado: ¶

1075 [He, X.-C., Simon, M., Iyer, S., Xie, H.-B., Rörup, B., Shen, J., Finkenzeller, H., Stolzenburg, D., Zhang, R., Baccarini, A., Tham, Y. J., Wang, M., Amanatidis, S., Piedehierro, A. A., Amorim, A., Baalbaki, R., Brasseur, Z., Caudillo, L., Chu, B., Dada, L., Duplissy, J., El Haddad, I., Flagan, R. C., Granzin, M., Hansel, A., Heinritzi, M., Hofbauer, V., Jokinen, T., Kempainen, D., Kong, W., Krechmer, J., Kürten, A., Lamkaddam, H., Lopez, B., Ma, F., Mahfouz, N. G. A., Makhmutov, V., Manninen, H. E., Marie, G., Marten, R., Massabò, D., Mauldin, R. L., Mentler, B., Onnela, A., Petäjä, T., Pfeifer, J., Philippov, M., Ranjithkumar, A., Rissanen, M. P., Schobesberger, S., Scholz, W., Schulze, B., Surdu, M., Thakur, R. C., Tomé, A., Wagner, A. C., Wang, D., Wang, Y., Weber, S. K., Welti, A., Winkler, P. M., Zauner-Wieczorek, M., Baltensperger, U., Curtius, J., Kurtén, T., Worsnop, D. R., Volkamer, R., Lehtipalo, K., Kirkby, J., Donahue, N. M., Sipilä, M., and Kulmala, M.: Iodine oxoacids enhance nucleation of sulfuric acid particles in the atmosphere, \*Science\*, 382, 1308-1314, \[10.1126/science.adh2526\]\(https://doi.org/10.1126/science.adh2526\), 2023.](#)

Movido (inserción)[3]

1090 Hoffmann, E. H., Tilgner, A., Schrödner, R., Bräuer, P., Wolke, R. and Herrmann, H.: An advanced modeling study on the impacts and atmospheric implications of multiphase dimethyl sulfide chemistry, *Proc. Natl. Acad. Sci. U. S. A.*, 113(42), 11776–11781, doi:10.1073/pnas.1606320113, 2016.

1095 [Hoppel, W. A., Frick, G. M., and Fitzgerald, J. W.: Marine bound- ary layer measurements of new-particle formation and the effects of non-precipitating clouds have on aerosol size distribution, \*J. Geophys. Res.\*, 99, 14443–14459, 1994](#)

Eliminado: ¶

1100 Humphries, R. S., Keywood, M. D., Gribben, S., McRobert, I. M., Ward, J. P., Selleck, P., Taylor, S., Harnwell, J., Flynn, C., Kulkarni, G. R., Mace, G. G., Protat, A., Alexander, S. P., and McFarquhar, G.: Southern Ocean latitudinal gradients of cloud condensation nuclei, *Atmos. Chem. Phys.*, 21, 12757–12782, <https://doi.org/10.5194/acp-21-12757-2021>, 2021

Eliminado: Hoppel, W. A., Frick, G. M., and Fitzgerald, J. W.: Marine bound- ary layer measurements of new-particle formation and the effects of non-precipitating clouds have on aerosol size distribution, *J. Geophys. Res.*, 99, 14443–14459, 1994¶

1105 Humphries, R. S., Keywood, M. D., Ward, J. P., Harnwell, J., Alexander, S. P., Klekociuk, A. R., Hara, K., McRobert, I. M., Protat, A., Alroe, J., Cravigan, L. T., Miljevic, B., Ristovski, Z. D., Schofield, R., Wilson, S. R., Flynn, C. J., Kulkarni, G. R., Mace, G. G., McFarquhar, G. M., Chambers, S. D., Williams, A. G., and Griffiths, A. D.: Measurement report: Understanding the seasonal cycle of Southern Ocean aerosols, *Atmos. Chem. Phys.*, 23, 3749–3777, <https://doi.org/10.5194/acp-23-3749-2023>, 2023.

1110 Hsu, Y., Holsen, T. M. and Hopke, P. K.: Comparison of hybrid receptor models to locate PCB sources in Chicago, *Atmos. Environ.*, 37, 545–562, 2003.

JMS Daily Northern Hemisphere Snow and Ice Analysis at 1 km, 4 km, and 24 km Resolutions, Version 1. Boulder, Colorado USA. NSIDC: National Snow and Ice Data Center, doi:<https://doi.org/10.7265/N52R3PMC>, 2008.

Eliminado: Ice, C. U. N.:

Eliminado: US Natl. Ice Cent.,

Ito, T.: Size distribution of Antarctic submicron aerosols, *Tellus B*, 45, 145–59, 1993.

1115 James, I. M.: The Antarctic drainage flow: implications for hemi- spheric flow on the Southern Hemisphere, *Antarct. Sci.*, 1, 279– 290, 1989.

Jang, E., Park, K. T., Jun Yoon, Y., Kim, T. W., Hong, S. B., Becagli, S., Traversi, R., Kim, J. and Gim, Y.: New particle formation events observed at the King Sejong Station, Antarctic Peninsula -

Part 2: Link with the oceanic biological activities, *Atmos. Chem. Phys.*, 19(11), 7595–7608, doi:10.5194/acp-19-7595-2019, 2019.

- 1130 Järvinen, E., Virkkula, A., Nieminen, T., Aalto, P. P., Asmi, E., Lanconelli, C., Busetto, M., Lupi, A., Schioppa, R., Vitale, V., Mazzola, M., Petäjä, T., Kerminen, V.-M. and Kulmala, M.: Sciences ess Atmospheric Chemistry and Physics Climate of the Past Geoscientific Instrumentation Methods and Data Systems Seasonal cycle and modal structure of particle number size distribution at Dome C, Antarctica, *Atmos. Chem. Phys.*, 13, 7473–7487, doi:10.5194/acp-13-7473-2013, 2013a.
- 1135 Järvinen, E., Virkkula, A., Nieminen, T., Aalto, P. P., Asmi, E., Lanconelli, C., Busetto, M., Lupi, A., Schioppa, R., Vitale, V., Mazzola, M., Petäjä, T., Kerminen, V. M. and Kulmala, M.: Seasonal cycle and modal structure of particle number size distribution at Dome C, Antarctica, *Atmos. Chem. Phys.*, 13(15), 7473–7487, doi:10.5194/acp-13-7473-2013, 2013b.
- 1140 Jokinen, T., Sipilä, M., Kontkanen, J., Vakkari, V., Tisler, P., Duplissy, E.-M., Junninen, H., Kangasluoma, J., Manninen, H. E., Petäjä, T., Kulmala, M., Worsnop, D. R., Kirkby, J., Virkkula, A. and Kerminen, V.-M.: Ion-induced sulfuric acid–ammonia nucleation drives particle formation in coastal Antarctica, *Sci. Adv.*, 4(11), eaat9744, doi:10.1126/sciadv.aat9744, 2018.
- 1145 Kerminen, V. M., Chen, X., Vakkari, V., Petäjä, T., Kulmala, M. and Bianchi, F.: Atmospheric new particle formation and growth: Review of field observations, *Environ. Res. Lett.*, 13(10), doi:10.1088/1748-9326/aadf3c, 2018.
- 1150 Kim, J., Yoon, Y. J., Gim, Y., Kang, H. J., Choi, J. H., Park, K. and Lee, B. Y.: Seasonal variations in physical characteristics of aerosol particles at the King Sejong Station, Antarctic Peninsula, *Atmos. Chem. Phys.*, 17, 12985–12999, 2017.
- 1155 Kim, J., Jun Yoon, Y., Gim, Y., Hee Choi, J., Jin Kang, H., Park, K. T., Park, J. and Yong Lee, B.: New particle formation events observed at King Sejong Station, Antarctic Peninsula - Part 1: Physical characteristics and contribution to cloud condensation nuclei, *Atmos. Chem. Phys.*, 19(11), 7583–7594, doi:10.5194/acp-19-7583-2019, 2019.
- 1160 Kirkby, J., Curtius, J., Almeida, J., Dunne, E., Duplissy, J., Ehrhart, S., Franchin, A., Gagné, S., Ickes, L., Kürten, A., Kupc, A., Metzger, A., Riccobono, F., Rondo, L., Schobesberger, S., Tsagkogeorgas, G., Wimmer, D., Amorim, A., Bianchi, F., Breitenlechner, M., David, A., Dommen, J., Downard, A., Ehn, M., Flagan, R. C., Haider, S., Hansel, A., Hauser, D., Jud, W., Junninen, H., Kreissl, F., Kvashin, A., Laaksonen, A., Lehtipalo, K., Lima, J., Lovejoy, E. R., Makhmutov, V., Mathot, S., Mikkilä, J., Minginette, P., Mogo, S., Nieminen, T., Onnela, A., Pereira, P., Petäjä, T., Schnitzhofer, R., Seinfeld, J. H., Sipilä, M., Stozhkov, Y., Stratmann, F., Tomé, A., Vanhanen, J., Viisanen, Y., Vrtala, A., Wagner, P. E., Walther, H., Weingartner, E., Wex, H., Winkler, P. M., Carslaw, K. S., Worsnop, D. R., Baltensperger, U. and Kulmala, M.: Role of sulphuric acid, ammonia and galactic cosmic rays in atmospheric aerosol nucleation, *Nature*, 476(7361), 429–435, doi:10.1038/nature10343, 2011.
- 1170 Koponen, I. K., Virkkula, A., Hillamo, R., Kerminen, V.-M., and Kulmala, M.: Number size distributions of marine aerosols: observations during a cruise between the English Channel and coast of Antarctica, *J. Geophys. Res.*, 107, 4753, <https://doi.org/10.1029/2002JD002533>, 2002.

- 1175 Korhonen, H., Carslaw, K. S., Spracklen, D. V., Mann, G. W. and Woodhouse, M. T.: Influence of oceanic dimethyl sulfide emissions on cloud condensation nuclei concentrations and seasonality over the remote Southern Hemisphere oceans: A global model study, *J. Geophys. Res. Atmos.*, 113(15), 1–16, doi:10.1029/2007JD009718, 2008.
- 1180 Lachlan-Cope, T., Beddows, D., Brough, N., Jones, A., Harrison, R., Lupi, A., Yoon, Y. J., Virkkula, A. and Dall'Osto, M.: On the annual variability of Antarctic aerosol size distributions at Halley research station, *Atmos. Chem. Phys.*, 20(7), 4461–4476, doi:10.5194/acp-2019-847, 2020.
- 1185 Lana, A., et al. (2011), An updated climatology of surface dimethylsulfide concentrations and emission fluxes in the global ocean, *Global Biogeochem. Cycles*, 25, GB1004, doi:10.1029/2010GB003850.
- Lee, S. H., Gordon, H., Yu, H., Lehtipalo, K., Haley, R., Li, Y., and Zhang, R.: New Particle Formation in the Atmosphere: From Molecular Clusters to Global Climate, *Journal of Geophysical Research: Atmospheres*, 124, 7098–7146, 10.1029/2018jd029356, 2019.
- 1190 Lapere, R., Thomas, J. L., Marelle, L., Ekman, A. M. L., Frey, M. M., Lund, M. T., et al. (2023). The representation of sea salt aerosols and their role in polar climate within CMIP6. *Journal of Geophysical Research: Atmospheres*, 128, e2022JD038235.
- 1195 Legrand, M., Yang, X., Preunkert, S. and Theys, N.: Year-round records of sea salt, gaseous, and particulate inorganic bromine in the atmospheric boundary layer at coastal (Dumont d'Urville) and central (Concordia) East Antarctic sites, *J. Geophys. Res. Atmos.*, (121), 997–1023, 2016.
- 1200 Lupu, A. and Maenhaut, W.: Application and comparison of two statistical trajectory techniques for identification of source regions of atmospheric aerosol species, *Atmospheric Environ.*, 36, 5607–5618, 2002.
- 1205 McCoy, D. T., Burrows, S. M., Wood, R., Grosvenor, D. P., Elliott, S. M., Ma, P. L., Rasch, P. J., & Hartmann, D. L.: Natural aerosols explain seasonal and spatial patterns of Southern Ocean cloud albedo. *Science Advances*, 1(6), e1500157. <https://doi.org/10.1126/sciadv.1500157>, 2015.
- Meskhize, N. and Nenes, A.: Phytoplankton and cloudiness in the Southern Ocean, *Science*, 314, 1419–1423, <https://doi.org/10.1126/science.1131779>, 2006
- 1210 Mungall, E. L., Abbatt, J. P. D., Wentzell, J. J. B., Lee, A. K. Y., Thomas, J. L., Blais, M., Gosselin, M., Miller, L. A., Papakyriakou, T., Willis, M. D., and Liggio, J.: Microlayer source of oxygenated volatile organic compounds in the summertime marine Arctic boundary layer, *P. Natl. Acad. Sci. USA*, 114, 6203–6208, <https://doi.org/10.1073/pnas.1620571114>, 2017.

1215 O'Dowd, C. D., Lowe, J. A., Smith, M. H., Davison, B., Hewitt, C. N. and Harrison, R. M.: Biogenic sulphur emissions and inferred non-sea-salt-sulphate particularly during Events of new particle formation in and around Antarctica, Atlantic, 102(DII), 1997a.

1220 O'Dowd, C. D., Smith, M. H., Consterdine, I. E. and Lowe, J. A.: Marine aerosol, sea-salt, and the marine sulphur cycle: A short review, *Atmos. Environ.*, 31(1), 73–80, doi:10.1016/S1352-2310(96)00106-9, 1997b.

Otero, X. L., De La Peña-Lastra, S., Pérez-Alberti, A., Ferreira, T. O. and Huerta-Diaz, M. A.: Seabird colonies as important global drivers in the nitrogen and phosphorus cycles, *Nat. Commun.*, 9(1), doi:10.1038/s41467-017-02446-8, 2018.

1225  
1230  
Paglione, M., Beddows, D. C. S., Jones, A., Lachlan-Cope, T., Rinaldi, M., Decesari, S., Manarini, F., Russo, M., Mansour, K., Harrison, R. M., Mazzanti, A., Tagliavini, E., and Dall'Osto, M.: Simultaneous organic aerosol source apportionment at two Antarctic sites reveals large-scale and ecoregion-specific components, *Atmos. Chem. Phys.*, 24, 6305–6322, <https://doi.org/10.5194/acp-24-6305-2024>, 2024.

Eliminado: ¶

1235 Park, J., Kang, H., Gim, Y., Jang, E., Park, K.-T., Park, S., Jung, C. H., Ceburnis, D., O'Dowd, C., and Yoon, Y. J.: New particle formation leads to enhanced cloud condensation nuclei concentrations on the Antarctic Peninsula, *Atmos. Chem. Phys.*, 23, 13625–13646, <https://doi.org/10.5194/acp-23-13625-2023>, 2023.

Eliminado: Paglione, M., Beddows, D. C. S., Jones, A., Lachlan-Cope, T., Rinaldi, M., Decesari, S., Manarini, F., Russo, M., Mansour, K., Harrison, R. M., Mazzanti, A., Tagliavini, E., and Dall'Osto, M.: Simultaneous organic aerosol source apportionment at two Antarctic sites reveals large-scale and eco-region specific components, *EGUsphere* [preprint], <https://doi.org/10.5194/egusphere-2023-2275>, 2023.¶

Peck, L.S., 2018. Antarctic marine biodiversity: adaptations, environments and responses to change. *Oceanogr. Mar. Biol. Annu. Rev.* 56, 105–236.

1240 Quinn, P. K. and Bates, T. S.: The case against climate regulation via oceanic phytoplankton sulphur emissions, *Nature*, 480, 51–56, <https://doi.org/10.1038/nature10580>, 2011.

Rankin, A. M. and Wolff, E. W.: A year-long record of size-segregated aerosol composition at Halley, Antarctica, *J. Geophys. Res. Atmos.*, 108(24), 1–12, doi:10.1029/2003jd003993, 2003.

1245  
1250 Reddington, C. L., Carslaw, K. S., Stier, P., Schutgens, N., Coe, H., Liu, D., Allan, J., Browse, J., Pringle, K. J., Lee, L. A., Yoshioka, M., Johnson, J. S., Regayre, L. A., Spracklen, D. V., Mann, G. W., Clarke, A., Hermann, M., Henning, S., Wex, H., Kristensen, T. B., Leaitch, W. R., Pöschl, U., Rose, D., Andreae, M. O., Schmale, J., Kondo, Y., Oshima, N., Schwarz, J. P., Nenes, A., Anderson, B., Roberts, G. C., Snider, J. R., Leck, C., Quinn, P. K., Chi, X., Ding, A., Jimenez, J. L. and Zhang, Q.: The global aerosol synthesis and science project (GASSP): Measurements and modeling to reduce uncertainty, *Bull. Am. Meteorol. Soc.*, 98(9), 1857–1877, doi:10.1175/BAMS-D-15-00317.1, 2017.

1255 Riddick, S. N., Dragosits, U., Blackall, T. D., Daunt, F., Wan-less, S., and Sutton, M. A.: The global distribution of ammonia emissions from seabird colonies, *Atmos. Environ.*, 55, 319–327, doi:10.1016/j.atmosenv.2012.02.052, 2012.

- Rinaldi, M., Paglione, M., Decesari, S., Harrison, R. M., Beddows, D. C., Ovadnevaite, J., Ceburnis, D., O'Dowd, C. D., Simó, R., and Dall'Osto, M.: Contribution of Water-Soluble Organic Matter from Multiple Marine Geographic Eco-Regions to Aerosols around Antarctica, *Environ. Sci. Technol.*, 54, 7807–7817, <https://doi.org/10.1021/acs.est.0c00695>, 2020
- 1270 Ronowicz, M., Peña Cantero, A.L., Mercado Casares, B., Kuklinski, P., Soto Àngel, J.J., 2019. Assessing patterns of diversity, bathymetry and distribution at the poles using hydrozoa (Cnidaria) as a model group. *Hydrobiologia* 833, 25–51. <https://doi.org/10.1007/s10750-018-3876-5>.
- 1275 Saliba, G., Sanchez, K. J., Russell, L. M., Twohy, C. H., Roberts, G. C., Lewis, S., Dedrick, J., McCluskey, C. S., Moore, K., DeMott, P. J., Toohey, D. W.: Organic composition of three different size ranges of aerosol particles over the Southern Ocean, *Aerosol Science and Technology*, 55:3, 268-288, DOI: 10.1080/02786826.2020.1845296, 2021.
- 1280 Schmale, J., Schneider, J., Nemitz, E., Tang, Y. S., Dragosits, U., Blackall, T. D., Trathan, P. N., Phillips, G. J., Sutton, M., and Braban, C. F.: Sub-Antarctic marine aerosol: dominant contributions from biogenic sources, *Atmos. Chem. Phys.*, 13, 8669–8694, <https://doi.org/10.5194/acp-13-8669-2013>, 2013.
- 1285 Shaw, G. E.: Considerations on the Origin and Properties of the Antarctic Aerosol, *Rev. Geophys.*, 17, 1983–1998, 1988
- Sipilä, M., Sarnela, N., Jokinen, T., Henschel, H., Junninen, H., Kontkanen, J., Richters, S., Kangasluoma, J., Franchin, A., Peräkylä, O., Rissanen, M. P., Ehn, M., Vehkamäki, H., Kurten, T., Berndt, T., Petäjä, T., Worsnop, D., Ceburnis, D., Kerminen, V. M., Kulmala, M. and O'Dowd, C.: Molecular-scale evidence of aerosol particle formation via sequential addition of HIO<sub>3</sub>, *Nature*, 537(7621), 532–534, doi:10.1038/nature19314, 2016.
- 1290 Terauds, A., S. L. Chown, F. Morgan, H. J. Peat, D. J. Watts, H. Keys, P. Convey, and D. M. Bergstrom. (2012) Conservation biogeography of the Antarctic. *Diversity and Distributions* 18:726-741.
- Terauds, A., and Lee, J. R. (2016) Antarctic biogeography revisited: updating the Antarctic conservation biogeographic regions. *Diversity and Distributions* 22:836-840.
- 1300 Turner, J., Colwell, S. R., Marshall, G. J., Lachlan-Cope, T. A., Carleton, A. M., Jones, P. D., Lagun, V., Reid, P. A. and Iagovkina, S.: Antarctic climate change during the last 50 years, *Int. J. Climatol.*, 25(3), 279–294, doi:10.1002/joc.1130, 2005.
- 1305 Vaqué, D.; Boras, J.A.; Arrieta, J.M.; Agustí, S.; Duarte, C.M.; Sala, M.M. Enhanced Viral Activity in the Surface Microlayer of the Arctic and Antarctic Oceans. *Microorganisms* 2021, 9, 317. <https://doi.org/10.3390/microorganisms9020317>

Eliminado: ¶

- 1310 Virkkula, A., Grythe, H., Backman, J., Petäjä, T., Busetto, M., Lanconelli, C., Lupi, A., Becagli, S., Traversi, R., Severi, M., Vitale, V., Sheridan, P., and Andrews, E.: Aerosol optical properties calculated from size distributions, filter samples and absorption photometer data at Dome C, Antarctica, and their relationships with seasonal cycles of sources, *Atmos. Chem. Phys.*, 22, 5033–5069, <https://doi.org/10.5194/acp-22-5033-2022>, 2022.
- 1315 Yang, X., Frey, M. M., Rhodes, R. H., Norris, S. J., Brooks, I. M., Anderson, P. S., Nishimura, K., Jones, A. E., and Wolff, E. W.: Sea salt aerosol production via sublimating wind-blown saline snow particles over sea ice: parameterizations and relevant microphysical mechanisms, *Atmos. Chem. Phys.*, 19, 8407–8424, <https://doi.org/10.5194/acp-19-8407-2019>, 2019.
- 1320 Wang, M., Xiao, M., Bertozzi, B., Marie, G., Rörup, B., Schulze, B., Bardakov, R., He, X.-C., Shen, J., Scholz, W., Marten, R., Dada, L., Baalbaki, R., Lopez, B., Lamkaddam, H., Manninen, H. E., Amorim, A., Ataei, F., Bogert, P., Brasseur, Z., Caudillo, L., De Menezes, L.-P., Duplissy, J., Ekman, A. M. L., Finkenzeller, H., Carracedo, L. G., Granzin, M., Guida, R., Heinritzi, M., Hofbauer, V., Höhler, K., Korhonen, K., Krechmer, J. E., Kürten, A., Lehtipalo, K., Mahfouz, N. G. A., Makhmutov, V., Massabò, D., Mathot, S., Mauldin, R. L., Mentler, B., Müller, T., Onnela, A., Petäjä, T., Philippov, M., Piedehierro, A. A., Pozzer, A., Ranjithkumar, A., Schervish, M., Schobesberger, S., Simon, M., Stozhkov, Y., Tomé, A., Umo, N. S., Vogel, F., Wagner, R., Wang, D. S., Weber, S. K., Welti, A., Wu, Y., Zauner-Wieczorek, M., Sipilä, M., Winkler, P. M., Hansel, A., Baltensperger, U., Kulmala, M., Flagan, R. C., Curtius, J., Riipinen, I., Gordon, H., Lelieveld, J., El-Haddad, I., Volkamer, R., Worsnop, D. R., Christoudias, T., Kirkby, J., Möhler, O. and Donahue, N. M.: Synergistic HNO<sub>3</sub>–H<sub>2</sub>SO<sub>4</sub>–NH<sub>3</sub> upper tropospheric particle formation, *Nature*, 605(7910), 483–489, [doi:10.1038/s41586-022-04605-4](https://doi.org/10.1038/s41586-022-04605-4), 2022.
- 1335 Weller, R., Minikin, A., Wagenbach, D., and Dreiling, V.: Characterization of the inter-annual, seasonal, and diurnal variations of condensation particle concentrations at Neumayer, Antarctica, *Atmos. Chem. Phys.*, 11, 13243–13257, <https://doi.org/10.5194/acp-11-13243-2011>, 2011.
- 1340 Weller, R., Schmidt, K., Teinilä, K., and Hillamo, R.: Natural new particle formation at the coastal Antarctic site Neumayer, *Atmos. Chem. Phys.*, 15, 11399–11410, <https://doi.org/10.5194/acp-15-11399-2015>, 2015.
- 1345 Weller, R., Legrand, M., and Preunkert, S.: Size distribution and ionic composition of marine summer aerosol at the continental Antarctic site Kohnen, *Atmos. Chem. Phys.*, 18, 2413–2430, <https://doi.org/10.5194/acp-18-2413-2018>, 2018.
- Wiencke and Amsler, 2012 C. Wiencke, C.D. Amsler *Seaweeds and Their Communities in Polar Regions* Springer, Berlin, Heidelberg (2012), pp. 265-291, [10.1007/978-3-642-28451-9\\_13](https://doi.org/10.1007/978-3-642-28451-9_13)
- 1350 Wohl, C., Li, Q., Cuevas, C. A., Fernandez, R. P., Yang, M., Saiz-Lopez, A., and Simó, R.: Marine biogenic emissions of benzene and toluene and their contribution to secondary organic aerosols over the polar oceans, *Science Advances*, 9, eadd9031, [10.1126/sciadv.add9031](https://doi.org/10.1126/sciadv.add9031),



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Site	Lat	Lon	Elevation a.s.l. (m)
King Sejong	-62.2	-58.8	10
Marambio	-64.2	-56.6	198
Concordia/Dome C	-75.1	123.3	3233
Halley	-75.6	-26.2	30

Table 1: Latitude & longitude of each monitoring site

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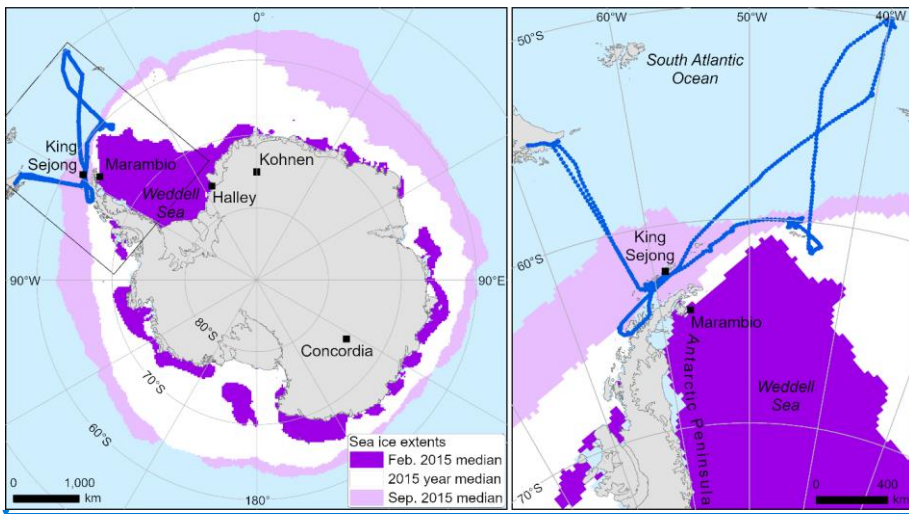
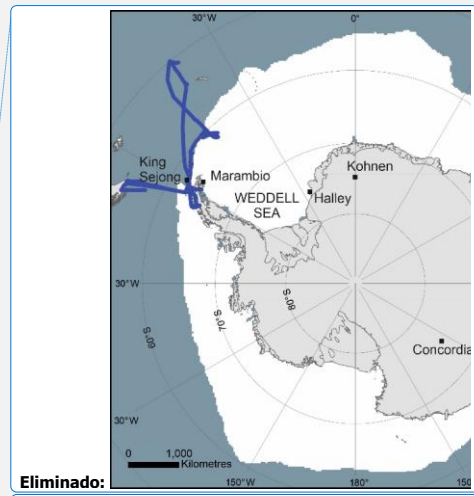


Figure 1: Map of the sampling stations (Halley, Marambio, Concordia/Dome C, King Sejong) for the year dataset collected in 2015. Additional, data for shorter period are intercompared at Kohnen (Weller et al., 2018) and during the PEGASO cruise (Dall'Osto et al., 2017, blue line the PEGASO cruise track). [The February sea ice extent signifies the annual minimum, while the September median signifies the annual maximum](#) (data are from the National Snow and Ice

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1395 | Data Center – NSIDC – at <https://nsidc.org/data/>, last access: 30 July 2024, Fetterer et al., (2017).

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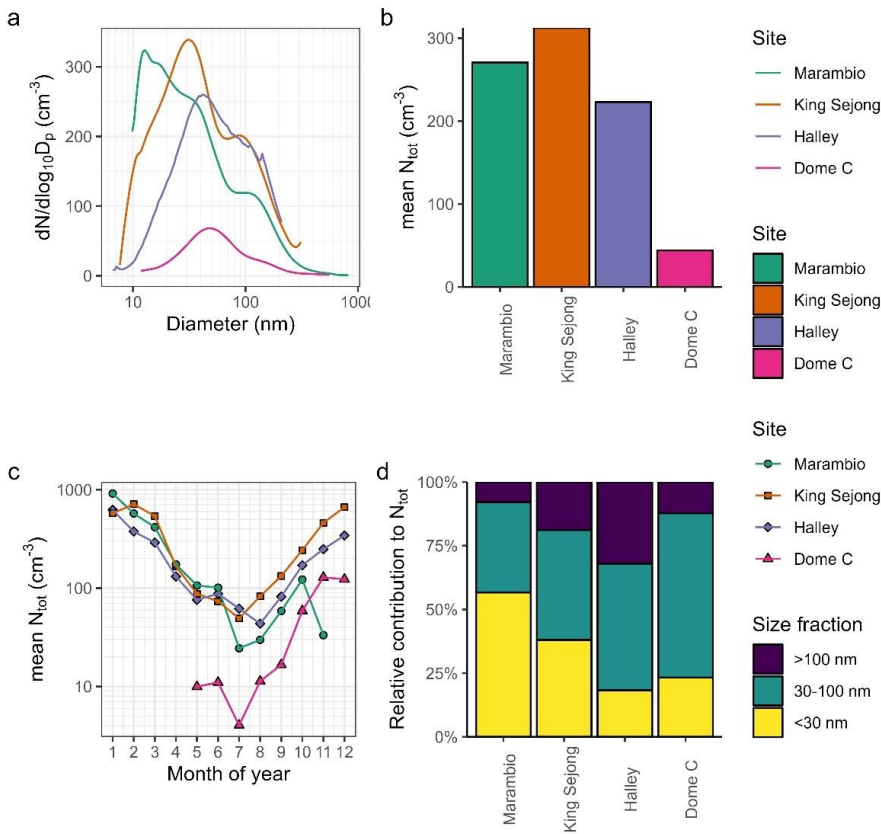


Figure 2: Features of particle PNSD data for four Antarctic sites, showing (a) mean particle PNSD per site, (b) mean particle count per site, (c) seasonal variation in particle count per site, and (d) contribution of different size fractions to particle count per site. Only data from the same time period (April-December) are intercompared and presented in Figure 2a,b,d, whereas all the temporal trend is presented in Figure 2c.

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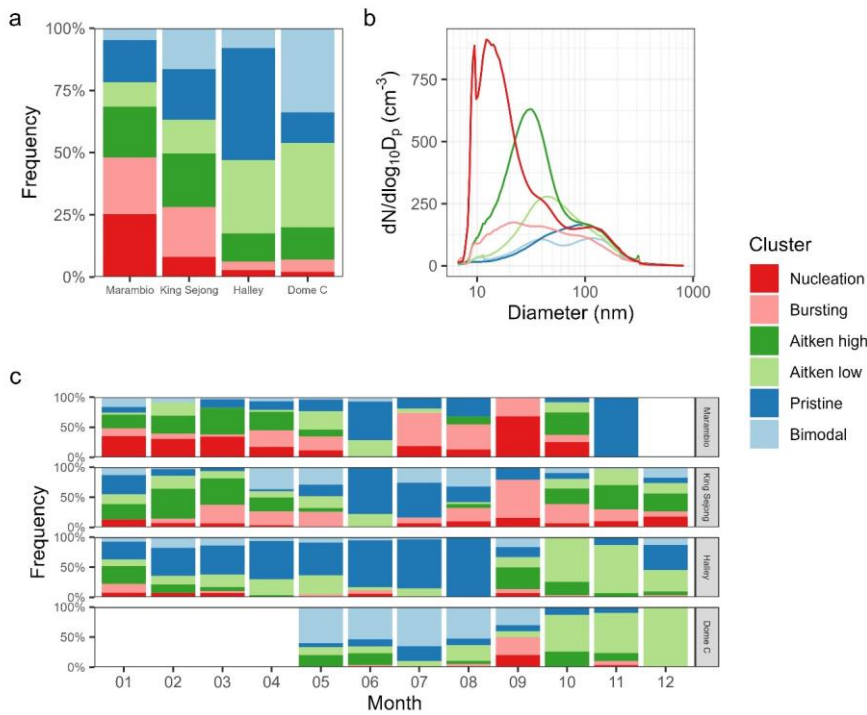
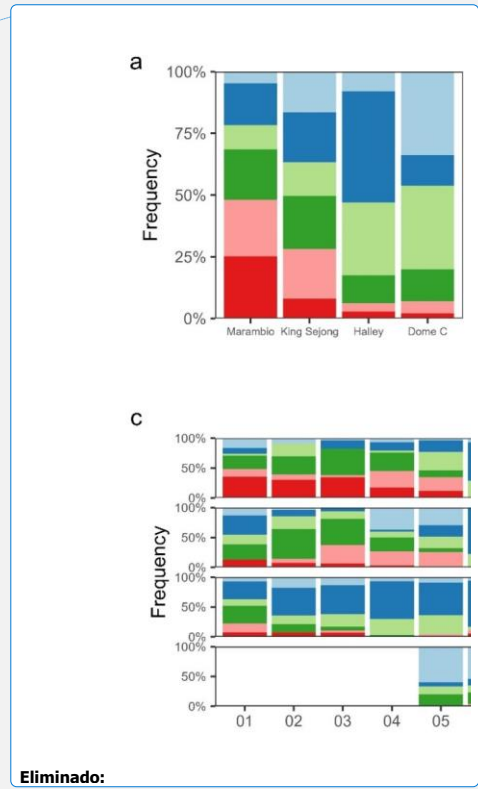


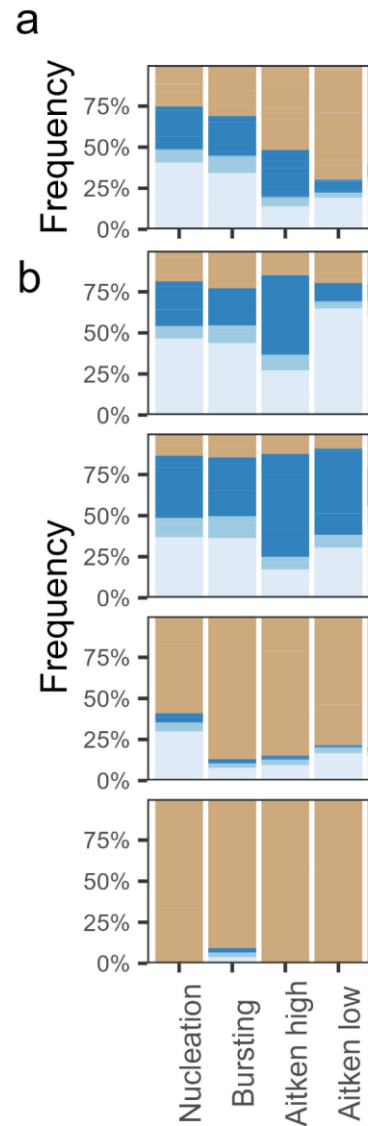
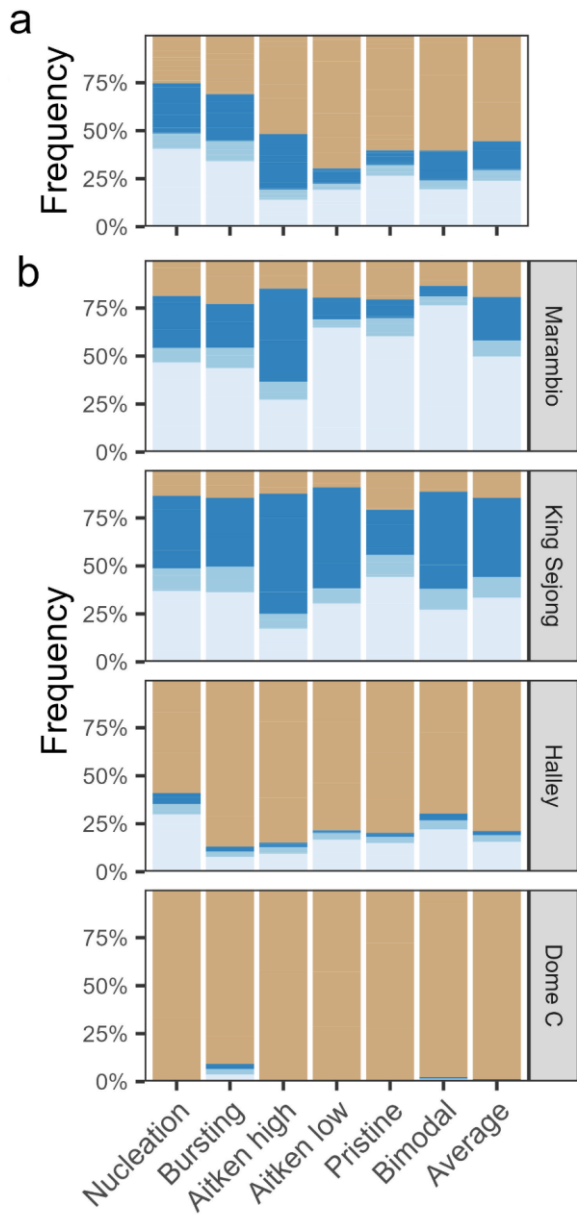
Figure 3: Cluster analysis results, showing (a) average frequency of each cluster per site, (b) mean PNSD per cluster, and (c) seasonal variations in cluster frequency per site

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1430 Figure 4: Land surface types associated with each cluster, showing (a) average association  
 across all sites, and (b) association per site. “Average” is the mean of all clusters.

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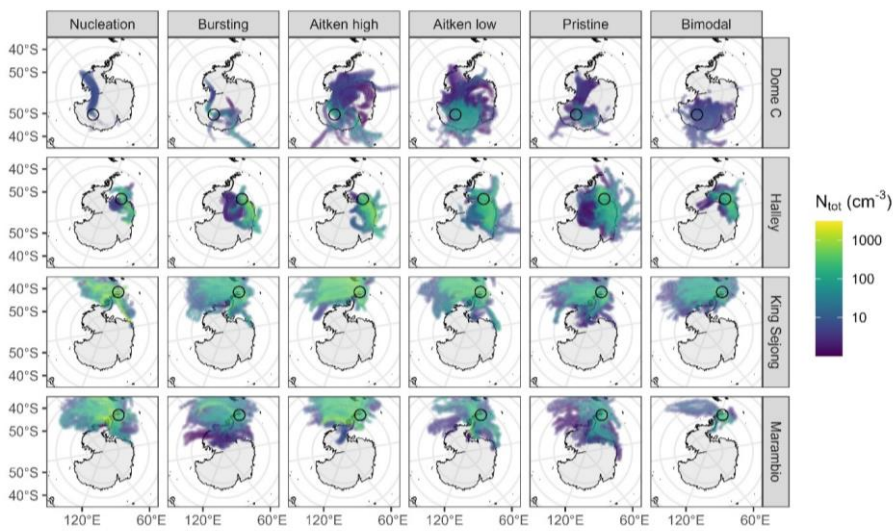
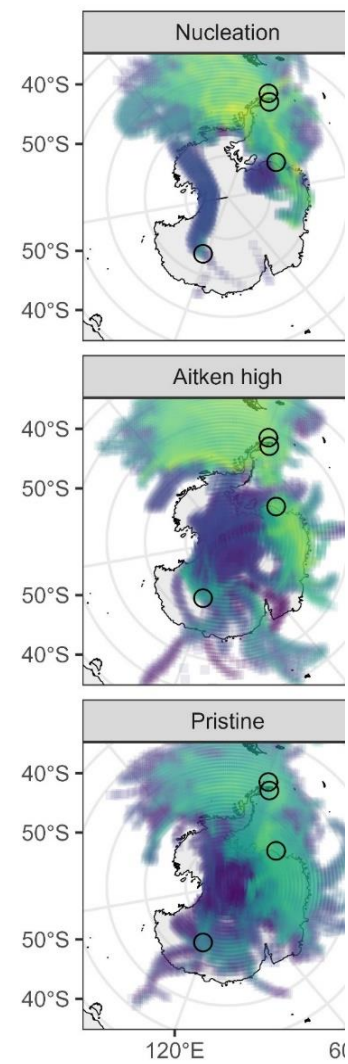


Figure 5: Concentration weighted trajectory maps showing the sources of particles corresponding to each cluster at each site. 72 hour back trajectories calculated with HYSPLIT.

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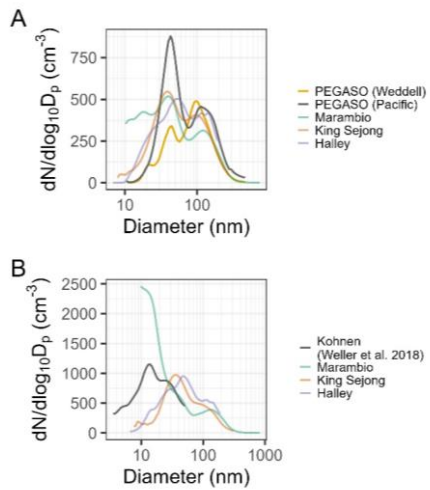
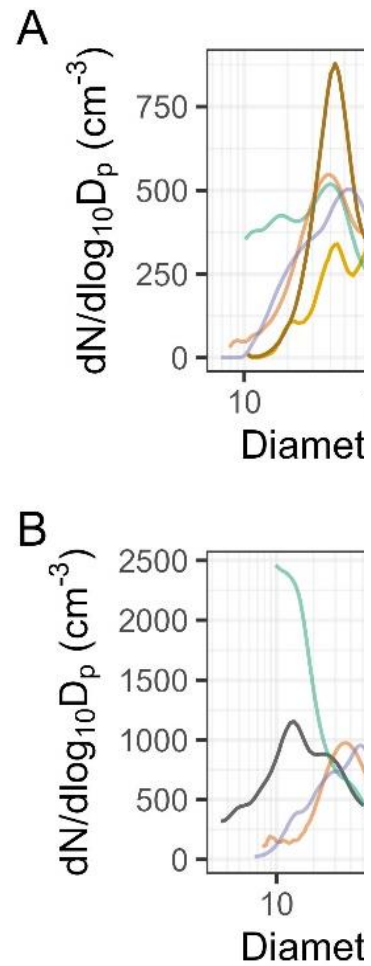


Figure 6: PNSD intercomparisons: (a) PNSD from the PEGASO cruise both when influenced from air masses from the Weddell Sea and Pacific Ocean and the stations used for this study where overlapping data are available (Marambio, Halley and King Sejong stations), and (b) PNSD from the Kohnen station (Weller et al., 2018) and the stations used for this study where overlapping data are available (Marambio, Halley and King Sejong stations)

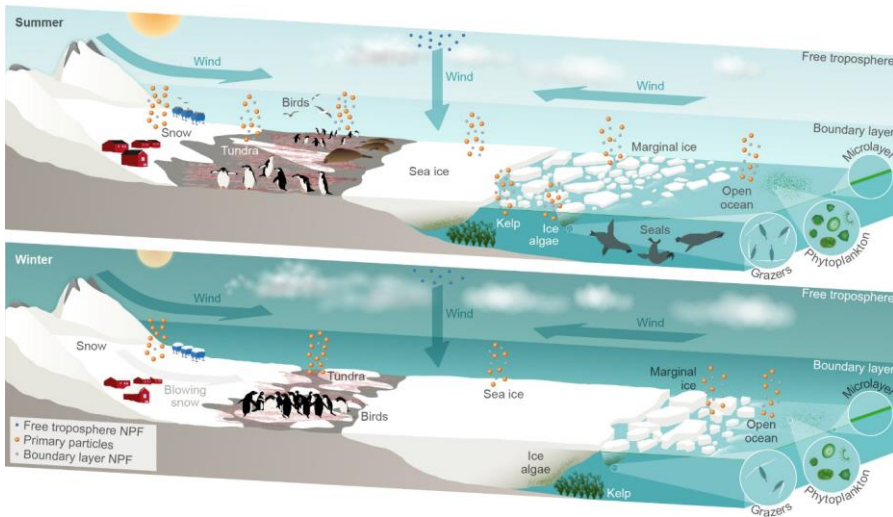


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Figure 7 Schematic illustrations of the sea ice, microbiota, sea-to-air emissions, and primary and secondary aerosols in Antarctica during summer (top) and winter (bottom).