

Measurement report: Analysis of aerosol optical depth variation at Zhongshan Station in Antarctica

Lijing Chen^{1,2}, Lei Zhang¹, Yong She², Zhaoliang Zeng¹, Yu Zheng¹, Biao Tian¹, Wenqian Zhang¹, Zhaohui Liu³, Huizheng Che¹, Minghu Ding^{*1}

¹ State Key Laboratory of Severe Weather, Chinese Academy of Meteorological Sciences, Beijing, 100081, China.

² Chengdu University of Information Technology, Chengdu, 610103, China.

³ Polar Surveying and Mapping Engineering Center of Heilongjiang Administration of Surveying, Mapping and Geoinformation, Harbin 150081, China

Correspondence to: Minghu Ding (dingminghu@foxmail.com)

Three key findings:

- The AOD level over Zhongshan Station in Antarctica is low in summer and high in winter. AE indicates the dominance of fine (coarse) aerosols in summer (winter).
- In winter and spring, high AOD values are related to the increase of coarse mode particles, while in summer and autumn, high AOD values may be related to the growth of fine mode particles.
- AOD varied inversely with wind speed and showed an insignificant positive correlation with temperature but a significant negative correlation with relative humidity.

Abstract: Our understanding of aerosol optical depth (AOD) in Antarctica remains limited due to the scarcity of ground observation stations and limited daylight days. Utilizing data from the CE318-T photometer spanning from January 2020 to April 2023 at Zhongshan Station, we analysed the seasonal, monthly, and diurnal variations in AOD and Ångström exponent (AE). AOD median values increased from spring (0.033) to winter (0.115), while AE peaked during summer (1.010) and autumn (1.034), declining in winter (0.381), indicating a transition in dominant aerosol particle size from fine to coarse mode between summer and winter. Monthly mean AOD variation closely paralleled the proportion of $AE < 1$, suggesting fluctuations in coarse mode particle proportions drive AOD variation. The high AOD values during winter and spring were associated with increased contribution of coarse mode particles, while high AOD values during summer and autumn were associated with the growth of fine mode particles. We observed a peak in AOD (~ 0.06) at 14:00 local time at Zhongshan Station, possibly

30 associated with a slight decrease in boundary layer height (BLH). Additionally, higher (lower) wind
31 speeds corresponded to lower (higher) AOD values, indicating the diffusion (accumulation) effect. The
32 temperature and AOD showed an insignificant positive correlation ($R = 0.22$, $p = 0.40$), relative humidity
33 exhibited a significant negative correlation with AOD ($R = -0.59$, $p = 0.02$). Backward trajectory analysis
34 revealed that coarse particles from the ocean predominantly contributed to high AOD daily mean values,
35 while fine particles on low AOD days originated mainly from the air mass over the Antarctic Plateau.
36 This study enhances the understanding of the optical properties and seasonal behaviors of aerosols in the
37 coastal Antarctic. Specifically, AOD measurements during the polar night address the lack of validation
38 data for winter AOD simulations. Additionally, we revealed that lower wind speeds, higher temperatures,
39 and lower relative humidity contribute to increased AOD at Zhongshan Station, and air masses from the
40 ocean significantly impact local AOD levels. These findings help us infer AOD variation patterns in the
41 coastal Antarctic based on meteorological changes, providing valuable insights for climate modeling in
42 the context of global climate change.

43 **1 Introduction**

44 Aerosols play an important role in impacting the climate system by absorbing and scattering solar
45 radiation (Li et al., 2022). Antarctica, considered one of the most pristine lands, serves as an ideal
46 background area for evaluating the climate effects of aerosols (Kamra, 2022). Marine aerosols emitted
47 from the Southern Ocean are a primary source contributing to the aerosol load in Antarctica (Thakur,
48 2019). The retreat of sea ice in Antarctica is expected to escalate the release of sea salt and secondary
49 biogenic aerosols (Yan et al., 2020). Sea salt particles with strong scattering may produce negative
50 effective radiative forcing or indirect radiative effect by influencing cloud condensation nuclei within
51 the marine boundary layer over Antarctica (Thornhill et al., 2021; Udisti et al., 2012). However, the
52 heating effect of absorbent aerosols, such as black carbon (BC), may be amplified by the high surface
53 albedo in Antarctica (Kang et al., 2020). In recent years, there has been a notable increase in BC
54 concentrations in Antarctica, with BC deposition on snow and ice surfaces contributing to reduced
55 surface albedo and increased solar radiation absorption, subsequently accelerating snow and ice melt
56 (Kannemadugu et al., 2023). Given the close connection between aerosol radiation effects and their

57 optical properties (Che et al., 2024), it is necessary to investigate the optical parameters of Antarctica
58 aerosols.

59 Aerosol optical depth (AOD), as a key parameters of aerosol optical properties, serves as an effective
60 measure of aerosol load and can influence the solar radiation components (Alghoul et al., 2009). AOD
61 observation records from Antarctica sites indicate that the values range from 0.006 to 0.220 in coastal
62 regions and from 0.007 to 0.034 in inland regions (Kannemadugu et al., 2023; Tomasi et al., 2007, 2012;
63 Yang et al., 2021). Typically, coastal aerosols consist primarily of coarse-mode sea salt particles during
64 austral winter, while fine-mode particles (such as dimethyl sulfide and its oxidation product mesylate,
65 DMS, and MSA) lead to elevated particle number concentrations in summer (20-100 times higher than
66 in winter) (Lachlan-Cope et al., 2020; Shaw, 1979). Conversely, aerosols over the Antarctic Plateau
67 predominantly comprise fine-mode particles of non-sea-salt sulfate (NSS) and DMS (Harder et al., 2000;
68 Walters et al., 2019).

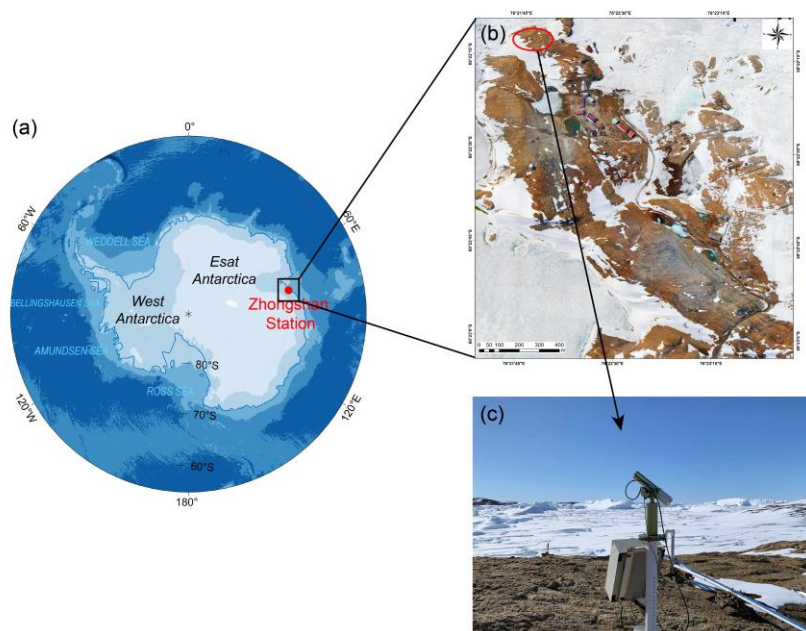
69 Additionally, particle size plays a significant role in aerosol extinction. The Ångström exponent (AE)
70 serves as an important indicator of aerosol size, with value greater (less) than 1 indicating a predominance
71 of fine (coarse) mode particles (Schuster et al., 2006). Weller and Lampert report that the mean AE at
72 Neumayer Station was 1.5 ± 0.6 and 1.2 ± 0.5 during summer and winter, respectively, suggesting an
73 increased contribution of fine-mode biological sulfate particles in summer (Weller and Lampert, 2008).
74 Virkkula et al. observed higher scattering AE estimate values during summer (~ 1.9) and lower values
75 during winter (~ 0.8) at Dome C on the Antarctic Plateau, indicating a prevalence of fine particles in
76 summer (Virkkula et al., 2022).

77 Currently, the challenging environment and the limited number of daylight days per year restrict the
78 availability of ground sites capable of obtaining adequate AOD and AE observations. Consequently, the
79 optical properties of aerosols across large parts of Antarctica remain unexplored. To improve our
80 comprehension of aerosol properties in Antarctica, we analyse the seasonal, monthly, and diurnal
81 variations of AOD and AE using data obtained from the recently installed sun-sky-lunar CE318-T
82 photometer at Zhongshan Station.

83 2 Site, Instrument, and Data

84 2.1 Site Introduction

85 Zhongshan Station ($69^{\circ}22'12''\text{S}$, $76^{\circ}21'49''\text{E}$, 18 m a.s.l.) is located at the Larsemann Hills of Prydz Bay
86 on the east Antarctic continent. The sun-sky-lunar CE318-T photometer is installed at Swan Ridge,
87 northwest of the Nella fjord (Fig. 1) (Tian et al., 2022). This location experiences 54 polar days and 58
88 polar nights annually, with snow covering the surrounding surface during winter and revealing bare rock
89 in summer. In this study, the austral spring, summer, autumn, and winter are referred to the season from
90 September to November (SON), December to February of next year (DJF), March to May (MAM), and
91 June to August (JJA), respectively. The average annual air temperature is -10°C , with a relative humidity
92 of 58% and prevailing wind speeds of 6.9 m s^{-1} , primarily from the east or east-southeast direction
93 (Ding et al., 2022).



94
95 **Figure 1 (a) The location of Zhongshan Station in Antarctica, (b) the aerial view of Zhongshan Station, and**
96 **(c) the sun-sky-lunar photometer CE318-T at Zhongshan Station.**

97 2.2 Instrument and Data

98 The AOD measurement data utilized in this study were obtained from the sun-sky-lunar CE318-T
99 photometer, manufactured by CIMEL Electronique, France. The CE318-T is a ground-based multiband
100 radiometer capable of inverting aerosol optical parameters by measuring the spectral data of direct solar
101 and lunar radiation extinction and the angular distribution of sky radiances (Barreto et al., 2016).

102 We collected AOD level 1.5 (cloud-screened) data across various wavelengths spanning from January
103 2020 to April 2023 (Fig. S1). However, the operation of CE318-T in polar environment is impeded by
104 solar radiation and weather conditions, leading to a significant number of missing measurements.
105 Consequently, we categorize daily observations with less than 20 measurements and the coefficient of
106 dispersion (CV) exceeding 1 as invalid data, which are systematically eliminated from our analysis.
107 Typically, these invalid data manifest with exceedingly high AOD values, often attributed to instrument
108 downtime caused by factors such as precipitation or cloudy weather. Moreover, to ensure the accuracy
109 of AOD measurement at Zhongshan Station, we refine our data by cross-referencing station operation
110 records and the time series of black carbon (BC) concentrations. This process allows us to exclude AOD
111 data associated with significant station activities and periods of elevated BC concentrations, thereby
112 enhancing the reliability of our analysis. It should be noted that there are uncertainties in the AOD
113 measurements of CE318-T during field observations due to atmospheric conditions, instrument noise,
114 and calibration. It is estimated that during daytime measurements, the AOD uncertainty ranges from
115 0.010 to 0.021. For night-time measurements, the AOD uncertainty depends on the calibration technique
116 used. Specifically, when calibrated using the Moon Ratio technique, the uncertainty ranges from 0.011
117 to 0.019. With the application of the new Sun Ratio technique, the uncertainty for the 440 nm channel is
118 between 0.012 and 0.015 (0.017), while for longer wavelengths, it ranges from 0.015 to 0.021. By
119 employing the new Sun-Moon gain factor technique and using the Langley-calibrated instrument for
120 calculation of the amplification between daytime and night-time measurements, the uncertainty range is
121 from 0.016 to 0.019 (Barreto et al., 2016).

122 The meteorology data, including temperature, relative humidity, wind direction, and wind speed, were
123 obtained from the Zhongshan Station meteorology observatory, with the temporal resolution of 1 hour.
124 BLH data was obtained from ERA5 reanalysis provided by European Centre for Medium Range Weather
125 Forecasts (ECMWF) with the temporal and spatial resolution of 1 hour and 0.25 (latitude) \times 0.25
126 (longitude).

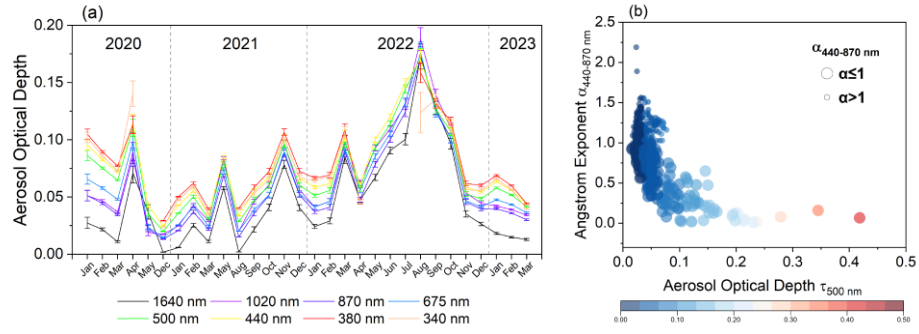
127 The Hybrid Single-Particle Lagrangian Integrated Trajectory (HYSPLIT) model, is a comprehensive
128 model developed by the National Oceanic and Atmospheric Administration (NOAA) and the Air
129 Resources Laboratory (ARL) to calculate and analyse the source, transport, and diffusion trajectories of

130 atmospheric pollutants. The meteorological data used in the HYSPLIT model comes from the National
131 Center for Environmental Prediction (NCEP) Global Data Assimilation System (GDAS). In this study,
132 the HYSPLIT model is utilized to calculate the 168h backward air mass trajectory from 3 altitudes of
133 50,500 and 1000 m (amsl) to Zhongshan Station.

134 **3 Results**

135 **3.1 Variation Characteristics of AOD**

136 From January 2020 to March 2023, the monthly mean AOD values at various wavelengths varied from
137 0.00 to 0.20, with the lowest values in December 2020 and the highest values in August 2022 (Fig. 2a).
138 The monthly mean AOD values at 500 nm ($AOD_{500\text{ nm}}$) generally remained below 0.10, consistent with
139 findings by Gadhavi and Achuthan at the Maitri Station, where AOD variation fell within the range of
140 0.01 to 0.10 (Gadhavi and Achuthan, 2004). The annual mean \pm SD (standard deviation) values of the
141 $AOD_{500\text{ nm}}$ were 0.074 ± 0.090 , 0.051 ± 0.066 , 0.071 ± 0.117 , and 0.053 ± 0.031 in 2020, 2021, 2022, and
142 2023, respectively (Table 1). Similarly, the annual mean \pm SD values of the $AE_{440-870\text{ nm}}$ were
143 1.134 ± 0.411 , 0.953 ± 0.338 , 0.883 ± 0.374 , 0.753 ± 0.206 for the same years, respectively, suggesting that
144 the aerosols over Zhongshan Station were mainly dominated by fine mode particles in 2020, and coarse
145 mode particles in 2021, 2022, and 2023. The relationship between multi-year $AOD_{500\text{ nm}}$ and $AE_{440-870\text{ nm}}$
146 illustrates that fine mode particles are primarily concentrated in the range of $AOD_{500\text{ nm}} < 0.1$, while high
147 $AOD_{500\text{ nm}}$ values, which occur occasionally, are caused by coarse mode particles (Fig. 2b). Although
148 fine mode particles have a longer suspension time in the atmosphere and can efficiently scatter and absorb
149 sunlight, leading to lower AOD ranges, it is worth mentioning that in the coastal regions of Antarctica,
150 the dominant role in AOD is sometimes played by coarse mode particles. These particles, with larger
151 radii and higher volume concentrations, originate mainly from abundant sea salt sources. Their presence
152 results in increased scattering and absorption of sunlight, emphasizing the significance of coarse mode
153 particles in determining AOD levels in the Antarctic coastal areas (Su et al., 2022)



154
 155 **Figure 2 (a) Monthly variation of mean aerosol optical depth at different wavelengths measured over**
 156 **Zhongshan Station in Antarctica from 2020 to 2023. (b) Relationship between AOD_{500 nm} and AE_{440-870 nm} over**
 157 **Zhongshan Station from 2020 to 2023.**

158 **Table 1 Annual mean and standard deviation of aerosol optical depth at different wavelengths and Angstrom**
 159 **Exponent at 440-870 nm at Zhongshan Station from 2020 to 2023.**

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	2020	2021	2022	2023
AOD _{1640 nm}	0.028±0.102	0.026±0.079	0.050±0.141	0.016±0.036
AOD _{1020 nm}	0.049±0.095	0.045±0.073	0.067±0.131	0.040±0.034
AOD _{870 nm}	0.047±0.093	0.039±0.070	0.060±0.126	0.037±0.032
AOD _{675 nm}	0.059±0.091	0.042±0.068	0.063±0.122	0.044±0.031
AOD _{500 nm}	0.074±0.090	0.051±0.066	0.071±0.117	0.053±0.031
AOD _{440 nm}	0.081±0.089	0.057±0.065	0.077±0.116	0.057±0.031
AOD _{380 nm}	0.089±0.091	0.063±0.065	0.077±0.117	0.061±0.032
AOD _{340 nm}	0.088±0.095	0.059±0.064	0.073±0.118	0.058±0.032
AE _{440-870 nm}	1.134±0.411	0.953±0.338	0.883±0.374	0.753±0.206

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162 3.2 Seasonal and Monthly Variations in AOD and Ångström Exponent

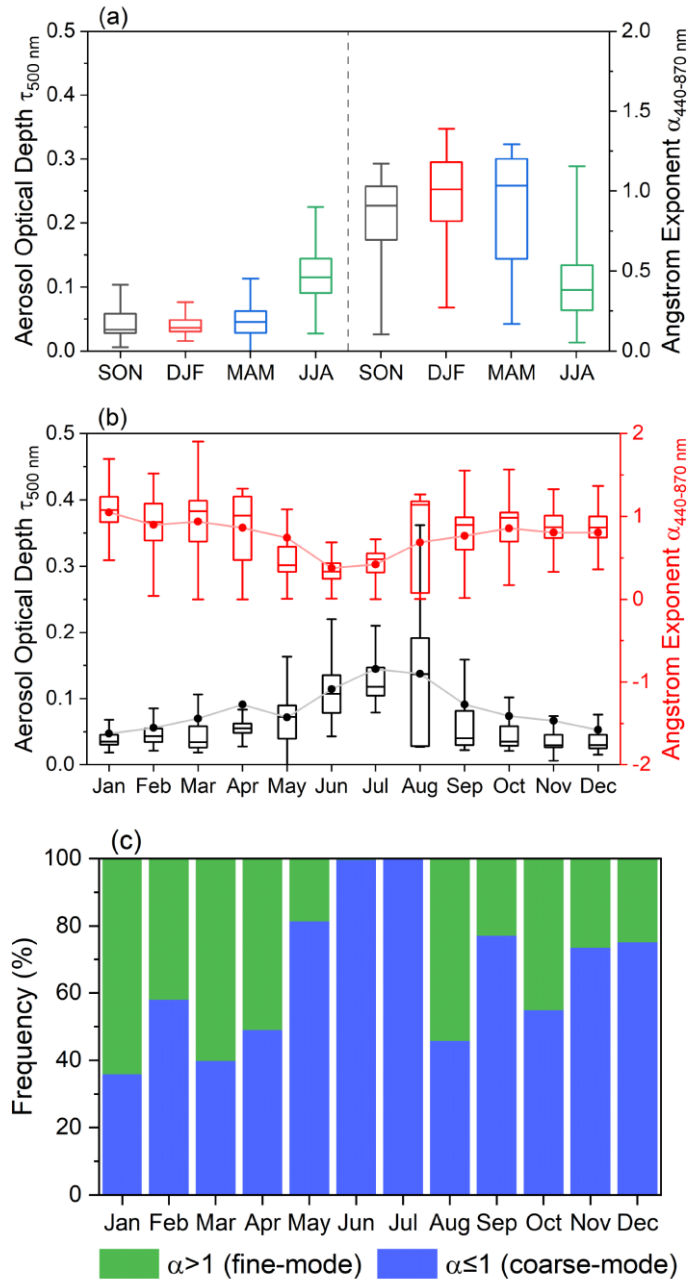
163 The seasonal variation of AOD_{500 nm} and AE_{440-870 nm} over Zhongshan Station suggests the median AOD₅₀₀
 164 nm values are lower in spring (0.033), summer (0.036), and autumn (0.045), but higher in winter (0.115),
 165 while the AE_{440-870 nm} values are 0.908, 1.010, 1.036, and 0.381, respectively (Fig. 3a). The frequency
 166 histograms show that the highest frequency range of AOD_{500 nm} is 0.02 to 0.04 in spring, summer, and
 167 autumn, while 0.08 to 0.12 in winter (Fig. S2). The normal fitting curves of the frequency histograms of

168 $AE_{440-870\text{ nm}}$ indicate that the peak in winter is in the low-value range (0.3~0.4), while the peaks in spring,
169 summer, and autumn are in the high-value range (1.0~1.2).

170 The seasonal variations in AOD and AE are consistent with previous findings on sea salt aerosol
171 concentrations, although the mechanism behind this seasonal variation is multifaceted. Wang and Huang
172 et al. have indicated that higher winter wind speeds at Zhongshan Station can elevate marine source
173 aerosol concentrations, primarily composed of sea salt, potentially explaining the winter peak in sea salt
174 concentration (Hong et al., 2009; Huang et al., 2005). However, Hall and Wolff propose that the high
175 sea salt load correlates more with moderate wind speeds and shifts in wind direction, rather than high
176 wind speeds, with concentrated brine on freshly formed ice surfaces acting as a source of winter sea salt
177 (Hall and Wolff, 1998). Moreover, blowing snow over sea ice generates aerosols primarily made of sea
178 salt, contributing to the winter peak in sea salt aerosols (Frey et al., 2020). In summer, lower sea salt
179 concentrations lead to lower background levels of AOD, but the effect of enhanced marine biogenic
180 emissions on AOD may increase. In the marine boundary layer over the eastern Southern Ocean sector,
181 $nssSO_4^{2-}$ and MSA contribute approximately 40% of the total mass of fine aerosols (particle size < 0.56
182 μm) (Xu et al., 2021). Xu et al. reported the annual mean concentrations of $nssSO_4^{2-}$ and MSA at
183 Zhongshan Station were 0-79 $ng\ m^{-3}$ and 19-41 $ng\ m^{-3}$, respectively, with the maximum
184 concentrations were observed in summer (Xu et al., 2019). This increase in summer concentrations is
185 attributed to enhanced solar radiation, phytoplankton blooms in the polynyas releasing DMS (Zhang et
186 al., 2015), and the DMS in the atmosphere is oxidized by radicals such as O_3 (significant at high latitudes),
187 OH, and BrO in the gas phase (Boucher et al., 2003), resulting in elevated concentrations of MSA and
188 $nssSO_4^{2-}$. The positive correlation between mean surface chlorophyll and AOD in the Southern Ocean
189 confirmed the contribution of DMS flux to aerosol load during summer (Gabric et al., 2005).

190 The monthly variations in $AOD_{500\text{ nm}}$ and $AE_{440-870\text{ nm}}$ at Zhongshan Station suggest an opposite trend,
191 with the mean values of $AOD_{500\text{ nm}}$ peaking in July and $AE_{440-870\text{ nm}}$ reaching its lowest in June (Fig. 3b).
192 Median $AOD_{500\text{ nm}}$ values increase slightly from January to February, followed by a decrease in March
193 and increase continuously from March to August, reach the maximum value, then gradually decrease,
194 and reach the minimum in November and December. The percentages of $AE_{440-870\text{ nm}} > 1.0$ and $AE_{440-8870}$
195 $nm < 1.0$ represent the proportion of the monthly occurrence frequency of fine and coarse mode particles

196 (Fig. 3c). The monthly mean and median AOD_{500 nm} values are consistent with the proportion of coarse
 197 mode particles ($AE_{440-870 \text{ nm}} > 1.0$), suggesting that the variation characteristics of AOD_{500 nm} at
 198 Zhongshan Station are primarily influenced by coarse mode particles. Given that Zhongshan Station is
 199 located in the coastal area of Antarctica, it is suspected that these coarse particles may be sea salt aerosols.

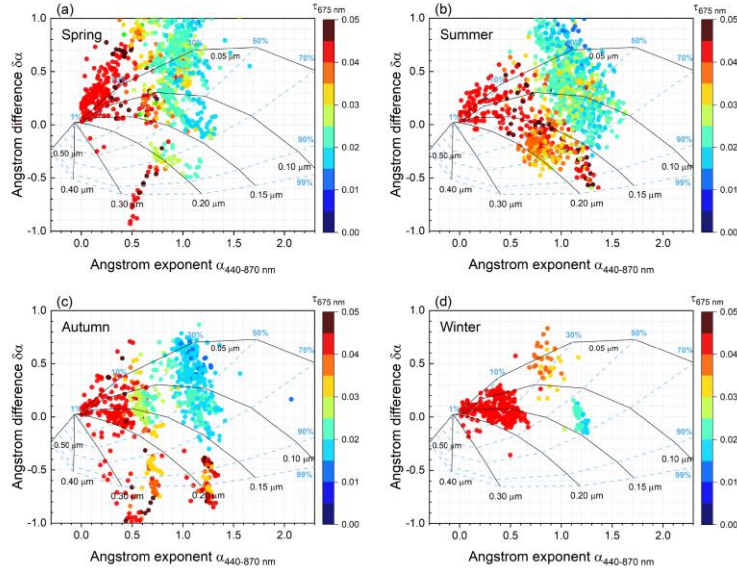


200
 201 **Figure 3 (a) Seasonal variation of aerosol optical depth at 500 nm and Angstrom exponent at 440-870 nm over**
 202 **Zhongshan Station. For each monthly box, the central line indicates the median; and the bottom and top edges**
 203 **of the box indicate the 25th and 75th percentiles, respectively. (b) Variations in monthly AOD_{500 nm} and AE₄₄₀₋**
 204 **870 nm at Zhongshan Station. For each monthly box, the central line indicates the median; the dot represents**
 205 **the mean; and the bottom and top edges of the box indicate the 25th and 75th percentiles, respectively. (c)**

206 **Monthly percentages of Ångström exponent >1.0 (green) and Ångström exponent ≤ 1.0 (blue) at Zhongshan**
207 **Station from 2020 to 2023.**

208 Additionally, we used a graphical method proposed by Gobbi et al (Gobbi et al., 2007), which is based
209 on Mie calculation and correlates Ångström exponent (α) and Ångström exponent spectral difference
210 ($\delta\alpha$) with fine mode aerosol effective radius (R_{eff}) and fine mode fraction to investigate the aerosol
211 modification processes at Zhongshan Station in different seasons. Figure 4 presents a schematic diagram
212 of the classification of aerosol types using the α and $\delta\alpha$ functions of a dual-mode, lognormal
213 distribution with refractive index = $1.4 - 0.001i$ as reference. It is known from Jurányi and Weller'
214 research that the refractive index of Antarctic coastal aerosol is about 1.4, so it seems reasonable to use
215 this reference (Jurányi and Weller, 2019). We utilized AOD_{440nm} , AOD_{675nm} , and AOD_{870nm} to calculate
216 $\alpha_{440-675nm}$, $\alpha_{440-870nm}$, and $\alpha_{675-870nm}$, and then get the $\delta\alpha = \alpha_{440-675nm} - \alpha_{675-870nm}$. The
217 negative values of $\delta\alpha$ indicate the dominance of fine mode aerosol, while positive values indicate the
218 effect of two separate particle modes (Kaufman, 1993). The solid black line represents the size of fine
219 mode particles (R_{eff}), and the dashed blue line represents the proportion of the contribution of fine mode
220 particles to AOD (η). In Fig. 4, increasing AOD_{675nm} is associated with the declining η (spring and
221 winter) and increasing R_{eff} (summer and autumn). This indicates that higher aerosol loads in spring and
222 winter are attributed to increased coarse-mode particle fractions, whereas in summer and autumn are
223 primarily associated with the increase of fine-mode particle size. Previous studies have indicated that sea
224 salt dominates winter aerosols in the coastal areas of Antarctica (Hall and Wolff, 1998; Weller et al.,
225 2008), and Xu et al observed that the highest mean concentration of sea salt in September at Zhongshan
226 Station, these can explain the $\delta\alpha$ values are mainly positive in spring and winter, and η is concentrated
227 within the range of less than 50% (Xu et al., 2019). In summer and autumn, apart from common sea salt
228 aerosols ($\delta\alpha > 0$, $\eta < 50$), the high AOD is mainly related to the particle growth such as hygroscopic
229 growth or condensation of fine mode aerosols ($R_{eff}: 0.10\mu m \sim 0.20\mu m$). This may be linked to the
230 atmospheric oxidation of (DMS) emitted by biological sources in coastal regions, or the aging process
231 of aerosols originating from other sources, as the rate of new particle formation and particulate matter
232 growth in summer is much greater than in winter in the Antarctica (Davison et al., 1996; Lachlan-Cope
233 et al., 2020; Weller et al., 2015).

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235

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Figure 4 Ångström exponent difference ($\delta\alpha = \alpha_{440-675 \text{ nm}} - \alpha_{675-870 \text{ nm}}$) as a function of the $\alpha_{440-870 \text{ nm}}$ and $\text{AOD}_{675 \text{ nm}}$ (colour scale) during (a) spring, (b) summer, (c) autumn, and (d) winter at Zhongshan Station. The black lines indicate the R_{eff} of fine-mode aerosols, while the blue lines correspond to fine-mode fraction (η).

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3.3 Relationship between AOD, Ångström Exponent and Meteorological Conditions

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In this section, we analyse the diurnal variation characteristics of $\text{AOD}_{500 \text{ nm}}$ and $\text{AE}_{440-870 \text{ nm}}$ during

242

summer and explore their correlation with meteorological variables within the planetary boundary layer

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(PBL), such as wind directions and speeds, temperature, and relative humidity. We calculated the diurnal

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variations of $\text{AOD}_{500 \text{ nm}}$ and $\text{AE}_{440-870 \text{ nm}}$ based on observations collected at Zhongshan Station during

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summer (December-February, 2020-2023), with each hourly mean containing at least one thousand

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individual observations (Fig. 5). The mean $\text{AOD}_{500 \text{ nm}}$ exhibited an increase from 5:00 to 14:00 (local

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time of Zhongshan Station), reaching a maximum value (0.06 ± 0.04), and then decreased. The mean

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$\text{AE}_{440-870 \text{ nm}}$ decreased from 5:00 to 12:00, reaching the lowest value (0.85 ± 0.25), and then increased.

249

These results indicate that the highest aerosol load occurs at 14:00, accompanied by a larger aerosol

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particle size during this period. The diurnal variation of boundary layer height (BLH) is almost consistent

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with the variation of $\text{AOD}_{500 \text{ nm}}$, which is inconsistent with the general conclusion that the negative

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correlation between BLH and particulate matter concentration in the mid-latitudes (Lou et al., 2019;

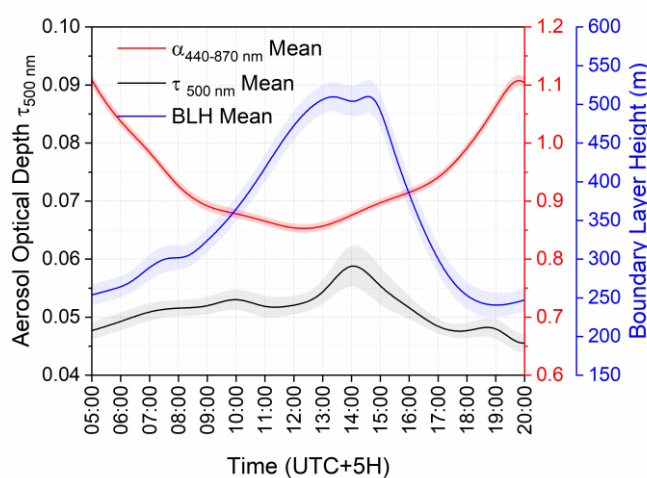
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Miao and Liu, 2019). However, a minor decline in BLH is noticeable when the $\text{AOD}_{500 \text{ nm}}$ value reaches

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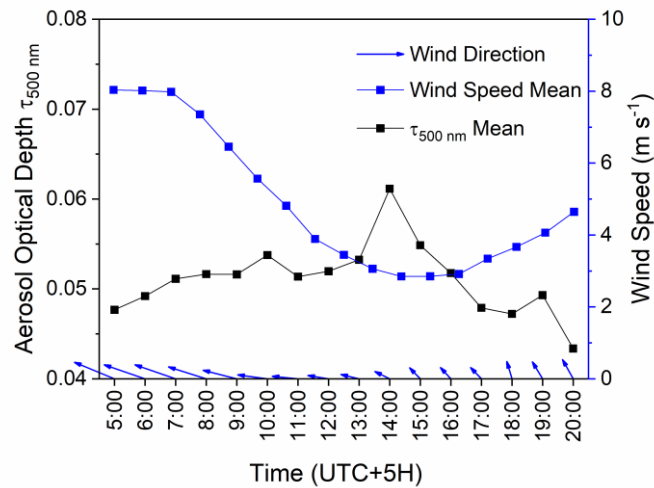
its peak at 14:00. Consequently, we suspect that the weak absorption and low content of Antarctic

255 aerosols typically do not suffice to form an “aerosol-boundary layer” positive feedback mechanism, but
 256 may contribute to reducing the BLH when AOD is high (Lou et al., 2019; Petäjä et al., 2016).



257
 258 **Figure 5 Diurnal variation of AOD_{500 nm} and AE_{440-870 nm} at Zhongshan Station. The black line indicates the**
 259 **mean of AOD_{500 nm}; the red line represents the mean of AE_{440-870 nm}; the blue line represents the mean of BLH.**
 260 **The shadow represents the standard deviation of the mean.**

261 Moreover, the diurnal variation of the 2-minute wind at Zhongshan Station reveals prevailing southeast
 262 direction, with average speeds ranging from 2 to 9 m s⁻¹. There is a noticeable decline in wind speeds
 263 between 5:00 and 14:00, followed by a gradual increase thereafter (Fig. 6). Given that the CE318-T is
 264 positioned westward of the main Zhongshan Station building, the eastward winds may carry emissions
 265 originating from western stations such as Zhongshan and Progress Station. The relationship between the
 266 diurnal variation of AOD_{500 nm} and wind speed is more obvious: AOD_{500 nm} exhibits a decline (increase)
 267 concurrent with decreasing (increasing) wind speeds. This correlation stems from the fact that higher
 268 wind speeds facilitate the dispersion of pollutants, leading to a reduction in AOD, and vice versa (Coccia,
 269 2021; Liu et al., 2020; Wang et al., 2022).



270

271 **Figure 6 Diurnal variations of 2-minute wind direction and speed, and AOD_{500 nm} in summer at Zhongshan**
 272 **Station.**

273 The influence of temperature and relative humidity on aerosol parameters is relatively complex.

274 Temperature affects aerosol particle concentration by influencing the air convection and influencing the

275 formation and optical properties of secondary by controlling chemical transformation (Han et al., 2007;

276 Li et al., 2020). Relative humidity affects the chemical composition, size distribution, and optical

277 properties of aerosol particles by affecting their aqueous-phase reactions and gas-particle partitioning

278 (Altieri et al., 2008; Ding et al., 2021; Hennigan et al., 2008; Sun et al., 2013). The diurnal variations of

279 AOD_{500 nm}, temperature, and relative humidity in summer at Zhongshan Station show that AOD_{500 nm} is

280 positively correlated with temperature with a correlation coefficient of 0.22, and AOD_{500 nm} is negatively

281 correlated with relative humidity with a correlation coefficient of -0.59 (Fig. 7). This indicates that rising

282 (declining) temperature and declining (rising) relative humidity during the day may contribute to an

283 increase (declining) in aerosol load. Previous studies have shown a positive correlation between

284 temperature and AOD (Basharat et al., 2023). During the summer at Zhongshan Station, high

285 temperatures may destroy the physical properties of bare rocks and promote the formation and diffusion

286 of particulate matter, thereby increasing the aerosol load (Zhang, 2024). However, there is a study

287 showing that higher temperatures may reduce methane sulfinic acid (MSIA) yield (Cecilia Arsene et al.,

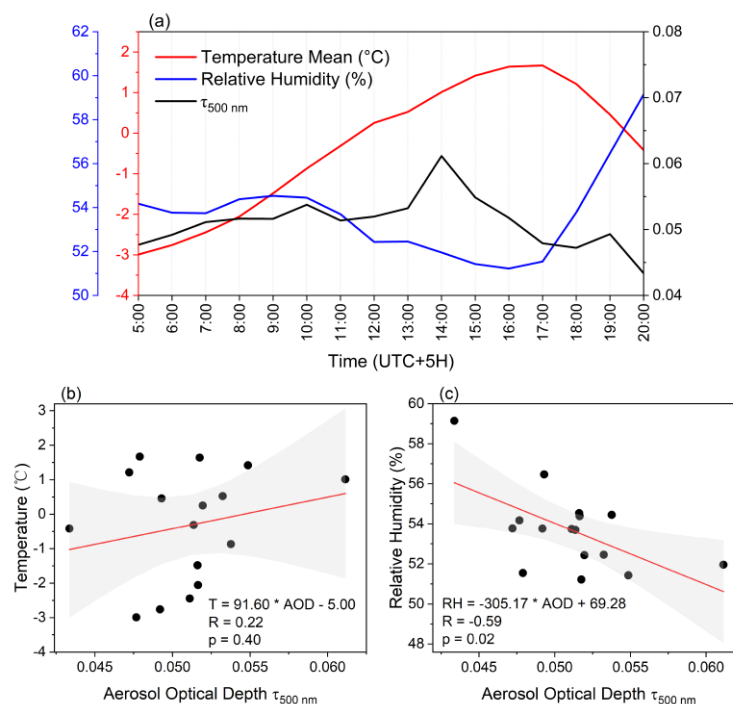
288 1999). Therefore, the effect of temperature on the AOD at Zhongshan Station is complex, resulting in an

289 insignificant positive correlation. The relationship between relative humidity and AOD is inconclusive

290 (Gautam et al., 2022), as high relative humidity may contribute to the increase of aerosol hygroscopic

291 properties leading to an increase in AOD (Meng et al., 2024), or it may contribute to a decrease in AOD

292 by reducing dust particles in the air (Zhang, 2024). Therefore, the influence of temperature and relative
 293 humidity on AOD may be related to the physicochemical properties of local aerosols and their sourcing
 294 and sink processes.



295
 296 **Figure 7 (a) Diurnal variations of AOD_{500 nm} (black), temperature (red), and relative humidity (blue) in**
 297 **summer at Zhongshan Station; (b) relationship between AOD_{500 nm} and temperature; and (c) relationship**
 298 **between AOD_{500 nm} and relative humidity. The red line indicates the regression line obtained by fitting a linear**
 299 **regression, and the grey bands indicate the confidence intervals for the linear regression.**

300 4 Discussion

301 4.1 Potential effects of aerosol sources on AOD levels

302 Besides meteorological conditions, aerosol sources may also influence the diurnal variation
 303 characteristics of AOD. We classified days with mean AOD below the 5th percentile as low AOD day
 304 and those above the 95th percentile as high AOD day (Fig. S3 and Table S1). Using the HYSPLIT
 305 backward trajectory model, we found that air masses on high AOD days primarily originated from the
 306 ocean, whereas those on low AOD days mostly came from the interior of Antarctica (Fig. S4). The
 307 altitudes of the backward trajectories show that during low AOD days, the air mass originating from the
 308 ocean usually starts at a lower altitude (<1000 m), rises to a higher altitude (~2000 m) and then descends
 309 to Zhongshan Station (2020-05-15 and 2020-12-25), while the air mass originating from the interior of

310 Antarctica usually starts at a higher altitude (~3000 m) and then descends to Zhongshan Station. This
311 indicates that particles from the Antarctic plateau or the free troposphere above the Antarctic interior are
312 transported to Zhongshan Station by katabatic winds. Researches show that the katabatic winds driven
313 by latent cooling occurring in the high-wind East Antarctic can rush the dense air from the interior plateau
314 to the coast (Simmons et al., 2021; Yu et al., 2020). Combined with the AE values, we can find that the
315 AE values of low AOD days are usually greater than 1, indicating the small particle size, thus, we suspect
316 that these fine particles may be $nssSO_4^{2-}$ from the Antarctic interior (Pei et al., 2021). In contrast, in
317 high AOD days, the air mass all originates in the ocean and usually starts at a lower altitude. The AE
318 values corresponding to high AOD moment on high AOD days are extremely low (<0.5), indicating that
319 the particle size is large, thus, we suspect that these aerosols may consist of coarse sea salt particles.

320 **4.2 Potential effects of aerosol particles on cloud and radiative forcing**

321 The optical properties of aerosols play a crucial role in their impact on radiative forcing, cloud formation,
322 and local climate. In our analysis of the variations in AOD and AE, we provided insights into the aerosol
323 loading, particle sizes, and possible formation and growth mechanisms in the atmosphere over
324 Zhongshan Station. During winter and spring, coarse mode particles are predominantly derived from sea
325 salt. Studies have shown that aerosols larger than 0.13 μm in the marine boundary layer contain sea salt,
326 contributing to most of the aerosol scattering and inducing cooling effects (Murphy et al., 1998).
327 Additionally, the size and inhomogeneity of sea salt particles are often associated with relative humidity.
328 Compared to remote oceans, the low relative humidity in coastal Antarctica may introduce more
329 inhomogeneous sea salt particles, resulting in up to a 12% change in direct radiative forcing due to
330 inhomogeneity (Wang et al., 2019).

331 However, we are particularly interested in the behaviour of aerosol particles during summer since solar
332 radiation is limited in winter. In summer and autumn, the increase in fine mode particles is closely related
333 to the release of biogenic aerosols, such as DMS, emitted by phytoplankton in the marginal ice zone.
334 When particles grow to a size suitable for cloud condensation nuclei or ice nucleating particles, they can
335 affect the formation of low-level mixed-phase clouds in coastal areas, contributing to the formation of
336 low-level ice clouds. At the same time, the increased number density of cloud droplets enhances cloud
337 reflectivity, resulting in negative radiative forcing (Satheesh and Krishna Moorthy, 2005). A recent study

338 revealed that in the shallow mixed-phase clouds over Antarctica, the concentrations of cloud-relevant
339 aerosol particles match the concentrations of ice crystals and cloud droplets (Radenz et al., 2024). the
340 number of particles plays a crucial role in cloud growth. Increasing particle concentration results in a
341 higher abundance of liquid droplets and ice crystals within clouds, which can impact cloud lifespan and
342 potentially influence local weather and climate. Therefore, continuous monitoring of aerosol optical
343 properties in coastal Antarctica is vital to improve our comprehension of aerosol radiative forcing
344 variations caused by changes in aerosol loading and particle size.

345 **5 Summary**

346 This study analysed the AOD and AE variations retrieved from CE318-T sun photometer data spanning
347 from January 2020 to April 2023 at Zhongshan Station in Antarctica. The main conclusions we draw are
348 as follows:

349 At Zhongshan Station, AOD varied from 0.00 to 0.20. Fine mode particles were predominantly found in
350 the lower AOD range, while higher AOD values were mainly attributed to coarse mode particles.
351 Seasonally, AOD exhibited a pattern of lower values in summer and higher values in winter, and the AE
352 displayed an opposite trend. The increases in AOD during summer and autumn may be linked to particle
353 growth, whereas the increases during spring and winter are associated with a decline in the fraction of
354 fine mode particles.

355 Low aerosol load over Zhongshan Station was not enough to form an “aerosol-boundary layer” positive
356 feedback mechanism, but the slight decrease in BLH may be related to AOD diurnal peak at 14:00.
357 Moreover, high (low) wind speeds facilitated pollutant dispersion (accumulation), leading to reduced
358 (increased) AOD. A weak positive correlation was noted between temperature and AOD ($R = 0.22$, $p =$
359 0.40), and a negative correlation between relative humidity and AOD ($R = -0.59$, $p = 0.02$). The
360 mechanisms underlying temperature and humidity's influence on aerosols remain unclear, possibly
361 linked to local aerosol properties at Zhongshan Station. In addition, we discuss the influence of aerosol
362 sources on AOD. The backward trajectories show that the air masses on high AOD days come from the
363 ocean, and the low AE values indicate that the particle size is larger, we speculate that the main
364 composition of the aerosols is sea salt. The air masses on the low AOD days mainly come from the

365 interior of Antarctica, and the high AE values indicate that the particle size is small. We speculate that
366 the katabatic winds rush the air from the Antarctic plateau to Zhongshan Station.

367 **Data availability**

368 The data included in this study can be accessed via <https://zenodo.org/records/10983098>. Boundary layer
369 height data downloaded from ECMWF ERA5 ([https://www.ecmwf.int/en/forecasts/dataset/ecmwf-](https://www.ecmwf.int/en/forecasts/dataset/ecmwf-reanalysis-v5)
370 [reanalysis-v5](https://www.ecmwf.int/en/forecasts/dataset/ecmwf-reanalysis-v5)). Backward trajectory of air mass and the meteorological data are obtained from NOAA
371 Air Resources Laboratory (https://www.ready.noaa.gov/HYSPLIT_traj.php).

372 **Author contributions**

373 The paper is a result of the lead author's research work under the supervision of MD, LZ, YS. ZZ and
374 YZ provided constructive comments. MD, QW and BT provided experimental data. ZL provided aerial
375 photos of Zhongshan Station. LC wrote the first draft of the paper with the help and support of all the
376 authors. HC provided guidance for the manuscript revisions.

377 **Competing interests**

378 The contact author has declared that none of the authors has any competing interests.

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383 **Reference**

384 Alghoul, M., Khamies, H., Assadeg, J., Yahya, M., Alfegi, E., and Sopian, K.:
385 Impact of Aerosol Optical Depth on Solar Radiation Budget, in: Proceedings of the 3rd
386 World Scientific and Engineering Academy and Society Int., Conference on renewable
387 energy sources, 2009.

388 Altieri, K. E., Seitzinger, S. P., Carlton, A. G., Turpin, B. J., Klein, G. C., and

389 Marshall, A. G.: Oligomers formed through in-cloud methylglyoxal reactions:
390 Chemical composition, properties, and mechanisms investigated by ultra-high
391 resolution FT-ICR mass spectrometry, *Atmospheric Environment*, 42, 1476–1490,
392 <https://doi.org/10.1016/j.atmosenv.2007.11.015>, 2008.

393 Barreto, Á., Cuevas, E., Granados-Muñoz, M.-J., Alados-Arboledas, L., Romero,
394 P. M., Gröbner, J., Kouremeti, N., Almansa, A. F., Stone, T., Toledano, C., Román, R.,
395 Sorokin, M., Holben, B., Canini, M., and Yela, M.: The new sun-sky-lunar Cimel
396 CE318-T multiband photometer - a comprehensive performance evaluation,
397 *Atmospheric Measurement Techniques*, 9, 631–654, [https://doi.org/10.5194/amt-9-](https://doi.org/10.5194/amt-9-631-2016)
398 631-2016, 2016.

399 Basharat, U., Tariq, S., Chaudhry, M. N., Khan, M., Bonah Agyekum, E., Fendzi
400 Mbasso, W., and Kamel, S.: Seasonal correlation of aerosols with soil moisture,
401 evapotranspiration, and vegetation over Pakistan using remote sensing, *Heliyon*, 9,
402 e20635, <https://doi.org/10.1016/j.heliyon.2023.e20635>, 2023.

403 Boucher, O., Moulin, C., Belviso, S., Aumont, O., Bopp, L., Cosme, E., von
404 Kuhlmann, R., Lawrence, M. G., Pham, M., Reddy, M. S., Sciare, J., and Venkataraman,
405 C.: DMS atmospheric concentrations and sulphate aerosol indirect radiative forcing: a
406 sensitivity study to the DMS source representation and oxidation, *Atmospheric*
407 *Chemistry and Physics*, 3, 49–65, <https://doi.org/10.5194/acp-3-49-2003>, 2003.

408 Cecilia Arsene, Barnes, I., and Becker, K. H.: FT-IR product study of the photo-
409 oxidation of dimethyl sulfide: Temperature and O₂ partial pressure dependence,
410 *Physical Chemistry Chemical Physics*, 1, 5463–5470,
411 <https://doi.org/10.1039/A907211J>, 1999.

412 Che, H., Xia, X., Zhao, H., Li, L., Gui, K., Zheng, Y., Song, J., Qi, B., Zhu, J.,
413 Miao, Y., Wang, Y., Wang, Z., Wang, H., Dubovik, O., Holben, B., Chen, H., Shi, G.,
414 and Zhang, X.: Aerosol optical and radiative properties and their environmental effects
415 in China: A review, *Earth-Science Reviews*, 248, 104634,
416 <https://doi.org/10.1016/j.earscirev.2023.104634>, 2024.

417 Coccia, M.: How do low wind speeds and high levels of air pollution support the
418 spread of COVID-19?, *Atmos Pollut Res*, 12, 437–445,
419 <https://doi.org/10.1016/j.apr.2020.10.002>, 2021.

420 Davison, B., O’ Dowd, C., Hewitt, C. N., Smith, M. H., Harrison, R. M., Peel, D.
421 A., Wolf, E., Mulvaney, R., Schwikowski, M., and Baltensperger, U.: Dimethyl sulfide
422 and its oxidation products in the atmosphere of the Atlantic and Southern Oceans,
423 *Atmospheric Environment*, 30, 1895–1906, [https://doi.org/10.1016/1352-](https://doi.org/10.1016/1352-2310(95)00428-9)
424 2310(95)00428-9, 1996.

425 Ding, J., Dai, Q., Zhang, Y., Xu, J., Huangfu, Y., and Feng, Y.: Air humidity affects

426 secondary aerosol formation in different pathways, *Science of The Total Environment*,
427 759, 143540, <https://doi.org/10.1016/j.scitotenv.2020.143540>, 2021.

428 Ding, M., Zou, X., Sun, Q., Yang, D., Zhang, W., Bian, L., Lu, C., Allison, I., Heil,
429 P., and Xiao, C.: The PANDA automatic weather station network between the coast and
430 Dome A, East Antarctica, *Earth System Science Data*, 14, 5019–5035,
431 <https://doi.org/10.5194/essd-14-5019-2022>, 2022.

432 Frey, M. M., Norris, S. J., Brooks, I. M., Anderson, P. S., Nishimura, K., Yang, X.,
433 Jones, A. E., Nerentorp Mastromonaco, M. G., Jones, D. H., and Wolff, E. W.: First
434 direct observation of sea salt aerosol production from blowing snow above sea ice,
435 *Atmospheric Chemistry and Physics*, 20, 2549–2578, [https://doi.org/10.5194/acp-20-](https://doi.org/10.5194/acp-20-2549-2020)
436 2549-2020, 2020.

437 Gabric, A. J., Shephard, J. M., Knight, J. M., Jones, G., and Trevena, A. J.:
438 Correlations between the satellite-derived seasonal cycles of phytoplankton biomass
439 and aerosol optical depth in the Southern Ocean: Evidence for the influence of sea ice,
440 *Global Biogeochemical Cycles*, 19, <https://doi.org/10.1029/2005GB002546>, 2005.

441 Gadhavi, H. and Achuthan, J.: Aerosol characteristics and aerosol radiative forcing
442 over Maitri, Antarctica, *Current Science*, 86, 296, 2004.

443 Gautam, S., Elizabeth, J., Gautam, A. S., Singh, K., and Abhilash, P.: Impact
444 Assessment of Aerosol Optical Depth on Rainfall in Indian Rural Areas, *Aerosol*
445 *Science and Engineering*, 6, 186–196, <https://doi.org/10.1007/s41810-022-00134-9>,
446 2022.

447 Gobbi, G. P., Kaufman, Y. J., Koren, I., and Eck, T. F.: Classification of aerosol
448 properties derived from AERONET direct sun data, *Atmospheric Chemistry and*
449 *Physics*, 7, 453–458, <https://doi.org/10.5194/acp-7-453-2007>, 2007.

450 Hall, J. S. and Wolff, E. W.: Causes of seasonal and daily variations in aerosol sea-
451 salt concentrations at a coastal Antarctic station, *Atmospheric Environment*, 32, 3669–
452 3677, [https://doi.org/10.1016/S1352-2310\(98\)00090-9](https://doi.org/10.1016/S1352-2310(98)00090-9), 1998.

453 Han, D., Liu, W., Zhang, Y., Lu, Y., Liu, J., and Zhao, N.: Influence of temperature
454 and relative humidity upon aerosol mass concentrations vertical distributions, *Journal*
455 *of University of Chinese Academy of Sciences*, 24, 619,
456 <https://doi.org/10.7523/j.issn.2095-6134.2007.5.011>, 2007.

457 Harder, S., Warren, S. G., and Charlson, R. J.: Sulfate in air and snow at the South
458 Pole: Implications for transport and deposition at sites with low snow accumulation,
459 *Journal of Geophysical Research: Atmospheres*, 105, 22825–22832,
460 <https://doi.org/10.1029/2000JD900351>, 2000.

461 Hennigan, C. J., Bergin, M. H., Dibb, J. E., and Weber, R. J.: Enhanced secondary
462 organic aerosol formation due to water uptake by fine particles, *Geophysical Research*
463 *Letters*, 35, <https://doi.org/10.1029/2008GL035046>, 2008.

464 Hong, J., Chen, L., and Yang, X.: Characteristics of the aerosols in Zhongshan
465 Station, Antarctica (in Chinese), *Chinese Journal of Polar Research*, 21, 1, 2009.

466 Huang, Z., Ji, W., Yang, X., Huang, R., Tang, R., Yu, T., and Zhang, G.: The
467 chemical composition of marine aerosol over Zhongshan Station in Antarctica and its
468 sources discrimination in 1998 (in Chinese), *Acta Oceanologica Sinica*, 27, 59–66,
469 2005.

470 Jurányi, Z. and Weller, R.: One year of aerosol refractive index measurement from
471 a coastal Antarctic site, *Atmospheric Chemistry and Physics*, 19, 14417–14430,
472 <https://doi.org/10.5194/acp-19-14417-2019>, 2019.

473 Kamra, V. P., Devendraa Siingh, A. K.: Antarctic Aerosols and Climate:
474 Measurements at a Coastal Antarctic Station, in: *Climate Variability of Southern High*
475 *Latitude Regions*, CRC Press, 2022.

476 Kang, S., Zhang, Y., Qian, Y., and Wang, H.: A review of black carbon in snow
477 and ice and its impact on the cryosphere, *Earth-Science Reviews*, 210, 103346,
478 <https://doi.org/10.1016/j.earscirev.2020.103346>, 2020.

479 Kannemadugu, H. B. S., Sudhakaran Syamala, P., Taori, A., Bothale, R. V., and
480 Chauhan, P.: Atmospheric aerosol optical properties and trends over Antarctica using
481 in-situ measurements and MERRA-2 aerosol products, *Polar Science*, 38, 101011,
482 <https://doi.org/10.1016/j.polar.2023.101011>, 2023.

483 Kaufman, Y. J.: Aerosol optical thickness and atmospheric path radiance, *Journal*
484 *of Geophysical Research: Atmospheres*, 98, 2677–2692,
485 <https://doi.org/10.1029/92JD02427>, 1993.

486 Lachlan-Cope, T., Beddows, D. C. S., Brough, N., Jones, A. E., Harrison, R. M.,
487 Lupi, A., Yoon, Y. J., Virkkula, A., and Dall’Osto, M.: On the annual variability of
488 Antarctic aerosol size distributions at Halley Research Station, *Atmospheric Chemistry*
489 *and Physics*, 20, 4461–4476, <https://doi.org/10.5194/acp-20-4461-2020>, 2020.

490 Li, J., Wang, W., Li, K., Zhang, W., Peng, C., Zhou, L., Shi, B., Chen, Y., Liu, M.,
491 Li, H., and Ge, M.: Temperature effects on optical properties and chemical composition
492 of secondary organic aerosol derived from *n*-dodecane, *Atmospheric Chemistry and*
493 *Physics*, 20, 8123–8137, <https://doi.org/10.5194/acp-20-8123-2020>, 2020.

494 Li, J., Carlson, B. E., Yung, Y. L., Lv, D., Hansen, J., Penner, J. E., Liao, H.,
495 Ramaswamy, V., Kahn, R. A., Zhang, P., Dubovik, O., Ding, A., Lacis, A. A., Zhang,

496 L., and Dong, Y.: Scattering and absorbing aerosols in the climate system, *Nat Rev*
497 *Earth Environ*, 3, 363–379, <https://doi.org/10.1038/s43017-022-00296-7>, 2022.

498 Liu, Y., Zhou, Y., and Lu, J.: Exploring the relationship between air pollution and
499 meteorological conditions in China under environmental governance, *Sci Rep*, 10,
500 14518, <https://doi.org/10.1038/s41598-020-71338-7>, 2020.

501 Lou, M., Guo, J., Wang, L., Xu, H., Chen, D., Miao, Y., Lv, Y., Li, Y., Guo, X., Ma,
502 S., and Li, J.: On the Relationship Between Aerosol and Boundary Layer Height in
503 Summer in China Under Different Thermodynamic Conditions, *Earth and Space*
504 *Science*, 6, 887–901, <https://doi.org/10.1029/2019EA000620>, 2019.

505 Meng, H., Bai, G., and Wang, L.: Analysis of the spatial and temporal distribution
506 characteristics of AOD in typical industrial cities in northwest China and the influence
507 of meteorological factors, *Atmospheric Pollution Research*, 15, 101957,
508 <https://doi.org/10.1016/j.apr.2023.101957>, 2024.

509 Miao, Y. and Liu, S.: Linkages between aerosol pollution and planetary boundary
510 layer structure in China, *Science of The Total Environment*, 650, 288–296,
511 <https://doi.org/10.1016/j.scitotenv.2018.09.032>, 2019.

512 Murphy, D. M., Anderson, J. R., Quinn, P. K., McInnes, L. M., Brechtel, F. J.,
513 Kreidenweis, S. M., Middlebrook, A. M., Pósfai, M., Thomson, D. S., and Buseck, P.
514 R.: Influence of sea-salt on aerosol radiative properties in the Southern Ocean marine
515 boundary layer, *Nature*, 392, 62–65, <https://doi.org/10.1038/32138>, 1998.

516 Pei, Q., Saikawa, E., Kaspari, S., Widory, D., Zhao, C., Wu, G., Loewen, M., Wan,
517 X., Kang, S., Wang, X., Zhang, Y.-L., and Cong, Z.: Sulfur aerosols in the Arctic,
518 Antarctic, and Tibetan Plateau: Current knowledge and future perspectives, *Earth-*
519 *Science Reviews*, 220, 103753, <https://doi.org/10.1016/j.earscirev.2021.103753>, 2021.

520 Petäjä, T., Järvi, L., Kerminen, V.-M., Ding, A. J., Sun, J. N., Nie, W., Kujansuu,
521 J., Virkkula, A., Yang, X., Fu, C. B., Zilitinkevich, S., and Kulmala, M.: Enhanced air
522 pollution via aerosol-boundary layer feedback in China, *Sci Rep*, 6, 18998,
523 <https://doi.org/10.1038/srep18998>, 2016.

524 Radenz, M., Engelmann, R., Henning, S., Schmithüsen, H., Baars, H., Frey, M. M.,
525 Weller, R., Bühl, J., Jimenez, C., Roschke, J., Muser, L. O., Wullenweber, N.,
526 Zeppenfeld, S., Griesche, H., Wandinger, U., and Seifert, P.: Ground-based Remote
527 Sensing of Aerosol, Clouds, Dynamics, and Precipitation in Antarctica —First results
528 from the one-year COALA campaign at Neumayer Station III in 2023,
529 <https://doi.org/10.1175/BAMS-D-22-0285.1>, 2024.

530 Satheesh, S. K. and Krishna Moorthy, K.: Radiative effects of natural aerosols: A
531 review, *Atmospheric Environment*, 39, 2089–2110,

532 <https://doi.org/10.1016/j.atmosenv.2004.12.029>, 2005.

533 Schuster, G. L., Dubovik, O., and Holben, B. N.: Angstrom exponent and bimodal
534 aerosol size distributions, *Journal of Geophysical Research: Atmospheres*, 111,
535 <https://doi.org/10.1029/2005JD006328>, 2006.

536 Shaw, G. E.: Considerations on the origin and properties of the Antarctic aerosol,
537 *Reviews of Geophysics*, 17, 1983–1998, <https://doi.org/10.1029/RG017i008p01983>,
538 1979.

539 Simmons, J. B., Humphries, R. S., Wilson, S. R., Chambers, S. D., Williams, A.
540 G., Griffiths, A. D., McRobert, I. M., Ward, J. P., Keywood, M. D., and Gribben, S.:
541 Summer aerosol measurements over the East Antarctic seasonal ice zone, *Atmospheric*
542 *Chemistry and Physics*, 21, 9497–9513, <https://doi.org/10.5194/acp-21-9497-2021>,
543 2021.

544 Su, Y., Han, Y., Luo, H., Zhang, Y., Shao, S., and Xie, X.: Physical-Optical
545 Properties of Marine Aerosols over the South China Sea: Shipboard Measurements and
546 MERRA-2 Reanalysis, *Remote Sensing*, 14, 2453, <https://doi.org/10.3390/rs14102453>,
547 2022.

548 Sun, Y., Wang, Z., Fu, P., Jiang, Q., Yang, T., Li, J., and Ge, X.: The impact of
549 relative humidity on aerosol composition and evolution processes during wintertime in
550 Beijing, China, *Atmospheric Environment*, 77, 927–934,
551 <https://doi.org/10.1016/j.atmosenv.2013.06.019>, 2013.

552 Thakur, R.: Trace elemental variability in aerosols near the two Indian Antarctic
553 research stations during austral summer, No. 26, pp 61–74, 2019.

554 Thornhill, G., Collins, W., Olivie, D., Skeie, R. B., Archibald, A., Bauer, S., Checa-
555 Garcia, R., Fiedler, S., Folberth, G., Gjermundsen, A., Horowitz, L., Lamarque, J.-F.,
556 Michou, M., Mulcahy, J., Nabat, P., Naik, V., O'Connor, F. M., Paulot, F., Schulz, M.,
557 Scott, C. E., Séférian, R., Smith, C., Takemura, T., Tilmes, S., Tsigaridis, K., and Weber,
558 J.: Climate-driven chemistry and aerosol feedbacks in CMIP6 Earth system models,
559 *Atmos. Chem. Phys.*, 21, 1105–1126, <https://doi.org/10.5194/acp-21-1105-2021>, 2021.

560 Tian, B., Ding, M., Putero, D., Li, C., Zhang, D., Tang, J., Zheng, X., Bian, L., and
561 Xiao, C.: Multi-year variation of near-surface ozone at Zhongshan Station, Antarctica,
562 *Environ. Res. Lett.*, 17, 044003, <https://doi.org/10.1088/1748-9326/ac583c>, 2022.

563 Tomasi, C., Vitale, V., Lupi, A., Di Carmine, C., Campanelli, M., Herber, A.,
564 Treffeisen, R., Stone, R. S., Andrews, E., Sharma, S., Radionov, V., von Hoyningen-
565 Huene, W., Stebel, K., Hansen, G. H., Myhre, C. L., Wehrl, C., Aaltonen, V.,
566 Lihavainen, H., Virkkula, A., Hillamo, R., Ström, J., Toledano, C., Cachorro, V. E.,
567 Ortiz, P., de Frutos, A. M., Blindheim, S., Frioud, M., Gausa, M., Zielinski, T., Petelski,

568 T., and Yamanouchi, T.: Aerosols in polar regions: A historical overview based on
569 optical depth and in situ observations, *Journal of Geophysical Research: Atmospheres*,
570 112, <https://doi.org/10.1029/2007JD008432>, 2007.

571 Tomasi, C., Lupi, A., Mazzola, M., Stone, R. S., Dutton, E. G., Herber, A.,
572 Radionov, V. F., Holben, B. N., Sorokin, M. G., Sakerin, S. M., Terpigova, S. A.,
573 Sobolewski, P. S., Lanconelli, C., Petkov, B. H., Busetto, M., and Vitale, V.: An update
574 on polar aerosol optical properties using POLAR-AOD and other measurements
575 performed during the International Polar Year, *Atmospheric Environment*, 52, 29–47,
576 <https://doi.org/10.1016/j.atmosenv.2012.02.055>, 2012.

577 Udisti, R., Dayan, U., Becagli, S., Busetto, M., Frosini, D., Legrand, M., Lucarelli,
578 F., Preunkert, S., Severi, M., Traversi, R., and Vitale, V.: Sea spray aerosol in central
579 Antarctica. Present atmospheric behaviour and implications for paleoclimatic
580 reconstructions, *Atmospheric Environment*, 52, 109–120,
581 <https://doi.org/10.1016/j.atmosenv.2011.10.018>, 2012.

582 Virkkula, A., Grythe, H., Backman, J., Petäjä, T., Busetto, M., Lanconelli, C., Lupi,
583 A., Becagli, S., Traversi, R., Severi, M., Vitale, V., Sheridan, P., and Andrews, E.:
584 Aerosol optical properties calculated from size distributions, filter samples and
585 absorption photometer data at Dome C, Antarctica, and their relationships with seasonal
586 cycles of sources, *Atmospheric Chemistry and Physics*, 22, 5033–5069,
587 <https://doi.org/10.5194/acp-22-5033-2022>, 2022.

588 Walters, W. W., Michalski, G., Böhlke, J. K., Alexander, B., Savarino, J., and
589 Thiemens, M. H.: Assessing the Seasonal Dynamics of Nitrate and Sulfate Aerosols at
590 the South Pole Utilizing Stable Isotopes, *Journal of Geophysical Research:*
591 *Atmospheres*, 124, 8161–8177, <https://doi.org/10.1029/2019JD030517>, 2019.

592 Wang, X., Chen, L., Guo, K., and Liu, B.: Spatio-temporal trajectory evolution
593 and cause analysis of air pollution in Chengdu, China, *Journal of the Air & Waste*
594 *Management Association*, 72, 876–894,
595 <https://doi.org/10.1080/10962247.2022.2058642>, 2022.

596 Wang, Z., Bi, L., Yi, B., and Zhang, X.: How the Inhomogeneity of Wet Sea Salt
597 Aerosols Affects Direct Radiative Forcing, *Geophysical Research Letters*, 46, 1805–
598 1813, <https://doi.org/10.1029/2018GL081193>, 2019.

599 Weller, R. and Lampert, A.: Optical properties and sulfate scattering efficiency of
600 boundary layer aerosol at coastal Neumayer Station, Antarctica, *Journal of Geophysical*
601 *Research: Atmospheres*, 113, <https://doi.org/10.1029/2008JD009962>, 2008.

602 Weller, R., Wöltjen, J., Piel, C., Resenberg, R., Wagenbach, D., König-Langlo, G.,
603 and Kriews, M.: Seasonal variability of crustal and marine trace elements in the aerosol
604 at Neumayer station, Antarctica, *Tellus B*, 60, 742–752, <https://doi.org/10.1111/j.1600->

605 0889.2008.00372.x, 2008.

606 Weller, R., Schmidt, K., Teinilä, K., and Hillamo, R.: Natural new particle
607 formation at the coastal Antarctic site Neumayer, *Atmospheric Chemistry and Physics*,
608 15, 11399–11410, <https://doi.org/10.5194/acp-15-11399-2015>, 2015.

609 Xu, G., Chen, L., Zhang, M., Zhang, Y., Wang, J., and Lin, Q.: Year-round records
610 of bulk aerosol composition over the Zhongshan Station, Coastal East Antarctica, *Air
611 Qual Atmos Hlth*, 12, 271–288, <https://doi.org/10.1007/s11869-018-0642-9>, 2019.

612 Xu, G., Chen, L., Xu, T., He, S., and Gao, Y.: Distributions of water-soluble ions
613 in size-aggregated aerosols over the Southern Ocean and coastal Antarctica, *Environ.
614 Sci.: Processes Impacts*, 23, 1316–1327, <https://doi.org/10.1039/D1EM00089F>, 2021.

615 Yan, J., Jung, J., Lin, Q., Zhang, M., Xu, S., and Zhao, S.: Effect of sea ice retreat
616 on marine aerosol emissions in the Southern Ocean, Antarctica, *Sci Total Environ*, 745,
617 140773, <https://doi.org/10.1016/j.scitotenv.2020.140773>, 2020.

618 Yang, Y., Zhao, C., Wang, Q., Cong, Z., Yang, X., and Fan, H.: Aerosol
619 characteristics at the three poles of the Earth as characterized by Cloud–Aerosol Lidar
620 and Infrared Pathfinder Satellite Observations, *Atmospheric Chemistry and Physics*, 21,
621 4849–4868, <https://doi.org/10.5194/acp-21-4849-2021>, 2021.

622 Yu, L., Zhong, S., and Sun, B.: The Climatology and Trend of Surface Wind Speed
623 over Antarctica and the Southern Ocean and the Implication to Wind Energy
624 Application, *Atmosphere*, 11, 108, <https://doi.org/10.3390/atmos11010108>, 2020.

625 Zhang, F.: Factors Influencing the Spatio–Temporal Variability of Aerosol Optical
626 Depth over the Arid Region of Northwest China, *Atmosphere*, 15, 54,
627 <https://doi.org/10.3390/atmos15010054>, 2024.

628 Zhang, M., Chen, L., Xu, G., Lin, Q., and Liang, M.: Linking Phytoplankton
629 Activity in Polynyas and Sulfur Aerosols over Zhongshan Station, East Antarctica,
630 *Journal of the Atmospheric Sciences*, 72, 4629–4642, [https://doi.org/10.1175/JAS-D-
631 15-0094.1](https://doi.org/10.1175/JAS-D-
631 15-0094.1), 2015.

632