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Distinct effects of Fine and Coarse Aerosols on Microphysical Processes of Shallow  
Precipitation Systems in Summer over Southern China

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17 **Abstract:** The densely populated South China, adjacent to the South China Sea, which  
18 is associated with shallow precipitation during summer, is an open-air natural  
19 laboratory for studying the impact of aerosols on shallow precipitation events. Using  
20 eight years of GPM DPR, MERRA-2 aerosol and ERA reanalysis data, this study  
21 investigates the potential influence of coarse and fine aerosol modes on the structure of  
22 the precipitation and the microphysical processes of shallow precipitation in South  
23 China. Statistical results indicate that during coarse aerosol-polluted conditions,  
24 shallow precipitation clouds have a lower mean height of the storm top (STH, ~3.2 km),  
25 but a higher mean near-surface rainfall (RR, ~1.78 mm h<sup>-1</sup>), characterized by high  
26 concentrations of large raindrops, driven mainly by significant collision-coalescence  
27 processes (accounting for 74.1%). In contrast, during fine aerosol-polluted conditions,  
28 shallow precipitation clouds develop a deeper median STH ~3.7 km with lower surface  
29 RR characterized by a low concentration of small hydrometeors, resulting from  
30 increased breakup processes (33.1%) and reduced collision-coalescence processes  
31 (69.6%). The coarse (fine) aerosols act as promoters (inhibitors) of radar reflectivity in  
32 the profile of shallow precipitation, regardless of dynamic and humid conditions. The  
33 effect of coarse aerosols in promoting precipitation and the inhibiting effect of fine  
34 aerosols are the most significant under low humidity conditions, mainly attributed to  
35 significantly enhanced collision-coalescence processes, exceeding 22.2%. Furthermore,  
36 the increase in RR above 3 km during coarse aerosol-polluted environments is mainly  
37 driven by the high concentration of hydrometeors in low instability conditions, whereas  
38 by large hydrometeors in high instability environments.  
39

40 **Short Summary:** The [microphysical mechanisms of precipitation](#) responsible for the  
41 varied impacts of aerosols on shallow precipitation remain unclear. This study reveals  
42 that coarse aerosols invigorate shallow rainfall through enhanced coalescence processes,  
43 whereas fine aerosols suppress shallow rainfall [through](#) intensified microphysical  
44 breaks. These impacts are independent of thermodynamic environments, but are more  
45 significant in low-humidity conditions.  
46

## 47 *1 Introduction*

48 Shallow precipitation, [generally identified by](#) storm height, [dominates in marine](#)  
49 [regions such as the](#) ocean and marine continent, potentially accounting for 20% of  
50 rainfall over tropical oceans and 7.5% over tropical land (Liu and Zipser, 2009; Chen  
51 et al., 2016; Short and Nakamura, 2000). This underscores its crucial significance in the  
52 regulation of the global water cycle. However, shallow precipitation is a complex  
53 phenomenon influenced by various factors such as water vapor, thermodynamic  
54 environment, and aerosols (Lang et al., 2021; Chen et al., 2024; Smalley and Rapp,  
55 2020). [Aerosols, as one of these factors, have sparked significant](#) debate due to the  
56 [intricate nature of](#) aerosol-radiation and aerosol-cloud interactions among various  
57 species, [resulting in unanswered questions about whether aerosols will increase or](#)  
58 [decrease shallow precipitation](#) (Koren et al., 2014; Fan et al., 2020; Christensen and  
59 Stephens, 2012).

60 The impact of aerosols on precipitation has been widely investigated in many  
61 previous studies (Sun and Zhao, 2021; Miltenberger et al., 2018; Liu et al., 2022; Fan  
62 et al., 2018). Regional differences show that aerosols can delay the start time of  
63 precipitation by 2 hours in the Pearl River Delta but advance by 3 hours in the North  
64 China Plain (Sun and Zhao, 2021). Furthermore, precipitation is suppressed for  
65 stratocumulus and small cumulus clouds in [highly polluted environments](#), but enhanced  
66 for heavy precipitation events and deep convective clouds (Yuan et al., 2011; Rosenfeld  
67 et al., 2008; Xiao et al., 2022; Miltenberger et al., 2018). However, convective rainfall  
68 invigoration depends on aerosol concentrations, which turns into suppression at the  
69 turning zone of aerosol optical depth in 0.25-0.30 (Guo et al., 2019), potentially linked

70 to a change from aerosol microphysical effects to aerosol radiative effects (Jiang et al.,  
71 2016). Liu et al. (2022) examined various aerosol types and discovered that marine  
72 warm clouds experienced a fourfold increase in rainfall flux in the presence of high  
73 levels of coarse spray aerosols, while there was a reduction by 75% in conditions with  
74 high concentrations of fine aerosols. Additionally, these contrast effects are independent  
75 of meteorological conditions. Another study suggests that the improvement of rainfall  
76 in orographic regions with high mineral dust concentrations is more significant in  
77 humid environments (Zhang et al., 2020b). Overall, the effects of aerosols on  
78 precipitation depend on numerous elements such as weather conditions, types of  
79 aerosols, their concentration, types of clouds, among others, and thus need to be  
80 carefully analyzed.

81 Most of these studies on the interactions between aerosols and precipitation have  
82 focused on the intensity, frequency of precipitation, and start and peak times of  
83 precipitation, but few studies have reported on how aerosols impact rainfall through  
84 modulating microphysical structures and processes of precipitation. Using three-  
85 dimensional observations of precipitation and microphysics from dual frequency  
86 precipitation radar (DPR) onboard the Global Precipitation Mission (GPM), recent  
87 studies have revealed that aerosol mainly reduces mean droplet concentration and  
88 increases the effective radius of precipitation in most regions of eastern China (except  
89 Northeast China) (Sun et al., 2022); Xiao et al. (2022) found that the aerosol  
90 invigoration effect on convective rainfall is characterized by higher droplet  
91 concentration with smaller size under polluted conditions in Northeast China. However,  
92 the impact of different aerosol species on precipitation microphysical structures and  
93 microphysical processes (i.e., coalescence efficiency of rain droplets) has been scarcely  
94 examined, which is essential for comprehending the full picture of the connections  
95 between aerosols, precipitation microphysics, and precipitation.

96 South China (18~29°N, 110~123°E) is a region where shallow precipitation occurs  
97 frequently (occurrence frequency up to 20%), and different types of aerosols prevail  
98 during summer (Yang et al., 2021), making it an ideal region for the study of the aerosol  
99 effect on shallow precipitation. Using the combined data set of GPM DPR and

100 MERRA-2 (Modern-Era retrospective analysis for Research and Applications,  
101 Versions2), this study aims to answer the following questions: 1) Do coarse and fine  
102 aerosols enhance or diminish the surface precipitation associated with shallow  
103 precipitation? 2) In what manner do aerosols influence the microphysical structures or  
104 processes of precipitation (such as break-up and collision-coalescence)? 3) To what  
105 extent are the relationships between aerosols and rainfall, microphysical structures, and  
106 processes sensitive to the dynamical and vapor components? The data and methods are  
107 introduced in Section 2. Section 3 discusses the impacts of fine and coarse aerosols on  
108 the microphysical properties and processes for shallow precipitation. A summary and  
109 conclusions are presented in Section 4.

110

## 111 ***2 Data and Methods***

### 112 ***2.1 Data***

113 In this study, four different data set are used to illustrate the potential impact of  
114 aerosols on microphysical precipitation structures and shallow precipitation processes  
115 over southern China during the summers between 2014 and 2021.

116 In the present study, the hourly MERRA-2 aerosol dataset  
117 (MERRA2\_400.tavg1\_2d\_aer\_Nx) at  $0.5 \times 0.625$  spatial resolution is used, which has  
118 been widely utilized with the advantage of high temporal and spatial resolution.  
119 MERRA-2 is produced using the Goddard Earth Observing System, Version 5 (GEOS-  
120 5) atmospheric model and the Gridpoint Statistical Interpolation (GSI) assimilation  
121 system (Molod et al., 2015). GEOS-5 integrates a radiatively coupled version of the  
122 Goddard Chemical Aerosol Radiation and Transport (GOCART) model to simulate  
123 aerosol components (Chin et al., 2002). In the estimation of aerosol properties,  
124 MERRA-2 assimilates aerosol data from ground-based observations from Aerosol  
125 Robotic NETWORK (AERONET) and spaceborne aerosol products from Advanced Very  
126 High Resolution Radiometer (AVHRR), Multiangle Imaging Spectro Radiometer

127 (MISR) (Randles et al., 2017; Buchard et al., 2017). Previous studies have shown a  
128 relatively good consistency of AOD from MERRA-2 and ground-based observations,  
129 i.e., AERONET, Sun sky radiometer Observation NETwork (SONET) (Ou et al., 2022;  
130 Buchard et al., 2015; Sun et al., 2019a). The correlation coefficient between MERRA-  
131 2 AOD and AERONET could reach 0.92 in summer China (Sun et al., 2019a). However,  
132 there is a slight underestimation of MERRA-2 AOD when compared to situ  
133 observations. Ou et al. (2022) revealed that the MERRA-2 AOD is underestimated by  
134 approximately 0.1 compared to a SONET station over South China. This is mainly  
135 because MERRA-2 lacks nitrate aerosols, leading to underestimations in the estimation  
136 of total AOD and fine aerosols (Sun et al., 2019b; Ou et al., 2022). The fine and coarse  
137 aerosol environment is defined by not only the AOD thresholds but also the AOD  
138 fractions to the total AOD, which may reduce uncertainties caused by underestimating  
139 AOD to some extent.

140 Aerosol species, including black carbon, organic carbon, sulfate, sea salt, and dust,  
141 are assumed to be external mixtures that do not interact with each other. In this present  
142 study, we consider the aerosol optical thickness and the extinction at 550 nm for five  
143 species, i.e. black carbon, organic carbon, sulfate, sea salt, and dust, as well as the  
144 Angstrom exponent ( $\alpha$ ) between 470 and 870 nm.  $\alpha$  is a significant parameter in aerosol  
145 science, which elucidates the AOD dependency on wavelength. A higher  $\alpha$  is related to  
146 a higher concentration of fine particles, whereas a lower  $\alpha$  suggests a higher  
147 concentration of coarse particles (Lolli et al., 2023).

148 The GPM DPR consists of two precipitation radars operating in the Ka and Ku  
149 bands, providing a unique opportunity to obtain information on three-dimensional  
150 precipitation and particle drop size distributions (DSDs) at the same time. In the present  
151 study, the official 2ADPR (version 7) dataset covering the summers (June to August)  
152 of 2014 and 2021 is also used, which provides information on the observation time,  
153 near-surface rain rate (RR), liquid water path (LWP), the three-dimensional profiles of  
154 attenuation-corrected reflectivity ( $Z_e$ ), rainfall, the mass-weighted mean diameter  $D_m$   
155 (in mm) and the generalized intercept  $N_w$  (in  $\text{mm}^{-1} \text{m}^{-3}$ ) of the normalized gamma  
156 distributions with a vertical resolution of 125 m in each scanning pixel (Iguchi et al.,

157 2017). The reliability of DSDs and precipitation has been validated by many previous  
158 studies (Huang et al., 2021; Radhakrishna et al., 2016). Due to the high spatial  
159 resolution (125m in vertically and 4.5 km in horizontal resolution), the official 2ADPR  
160 (version 7) dataset has been widely used in the field of climatology (Chen et al., 2024;  
161 Zhang et al., 2020a; Chen et al., 2020). Shallow precipitation clouds are defined by  
162 their near-surface RR exceeding  $0.1 \text{ mm h}^{-1}$  and STH below 5 km in altitude. The storm  
163 top height (STH) is defined as the maximum height where the  $Z_e$  exceeds 20dBZ (Liu  
164 and Zipser, 2013).

165 In this study, convective available potential energy (CAPE) and relative humidity  
166 (RH) at 850 hPa from the [fifth-generation](#) global reanalysis of the European Center for  
167 Medium-Range Weather Forecasts (ERA5) covering the period from 2014 to 2021 are  
168 also used to investigate the meteorological dependence on the relationship between  
169 aerosols and precipitation. Additionally, the global 1km grid quality-controlled global  
170 digital elevation model (DEM) (<https://ngdc.noaa.gov/mgg/topo/globe.html>) is also  
171 used to exclude the influence of topography in the present study.

172

## 173 **2.2 Methods**

174 Due to the different spatial and temporal resolutions of DPR, MERRA-2, and  
175 ERA5, [Prior to](#) examining the potential influence of various aerosol types on shallow  
176 precipitation, it is necessary to [harmonize](#) these three datasets. Since the DPR detects  
177 the rainy pixels at approximately 4.5 km spatial resolution, both MERRA-2 at  $0.5 \times$   
178  $0.625^\circ$  resolution and ERA5 at  $0.25^\circ$  resolution are first linearly interpolated to  $0.05^\circ$   
179 resolution. [To accurately depict the aerosol conditions preceding shallow precipitation,](#)  
180 [observations of AOD from MERRA-2, corresponding closely to the timing of DPR](#)  
181 [observations and with a spatial resolution of  \$0.05^\circ\$ , are utilized. Concurrently,](#)  
182 [atmospheric data derived from ERA5 at a  \$0.05^\circ\$  resolution, which is in closest proximity](#)  
183 [to the center and observation time of the DPR pixel, are also used.](#) The aerosol fine  
184 mode AOD is defined as the total AOD sum of partial AOD of black carbon, organic

185 carbon, and sulfate, while the AOD of coarse aerosols is the total value of the sum of  
186 AOD values of sea salt and dust particles (Gelaro et al., 2017). Additionally, to  
187 eliminate the potential impact of topography on precipitation and aerosol analysis, the  
188 study includes only shallow precipitation pixels that occur over regions with a  
189 topographic elevation of less than 100 meters.

190

191 Figure 1a illustrates the probability density of the joint distribution of AOD and  $\alpha$   
192 prior to the occurrence of the shallow precipitation event. Shallow precipitation is most  
193 probable when the AOD is approximately 0.4 and  $\alpha$  is approximately 1.4, which  
194 suggests a predominance of the fine aerosol mode. This can be primarily attributed to  
195 the increased presence of fine aerosols in South China during summer season, as  
196 represented in Figure 1b, where the probability density distributions (PDF) of AOD for  
197 fine aerosols and total aerosols reveal comparable values. Nonetheless, shallow  
198 precipitation is also evident in settings characterized by coarse aerosols, exhibiting a  
199 significant frequency when  $\alpha$  is less than 1 and AOD is less than 0.3, as shown in Figure  
200 1a.

201 There are three types of aerosol conditions discussed in the present study: clean  
202 environment, fine aerosol-polluted environment, and coarse aerosol-polluted  
203 environment. To classify clean and aerosol-polluted conditions over South China, PDFs  
204 of AOD for fine, coarse, and total aerosols are calculated before shallow precipitation,  
205 as shown in Figure 1b. It can be observed that the coarse mode AOD is relatively small,  
206 primarily distributed between 0 and 0.2, while fine mode AOD and total AOD are  
207 almost equal, mainly concentrated between 0 and 1.0. Specifically, the peak frequency  
208 occurs at an AOD of approximately 0.1 for coarse aerosols, 0.15 for fine aerosols, and  
209 0.2 for total aerosols. We define a clean environment as one in which the AOD of the  
210 total aerosols falls below the 30th percentile in all the data sampled, specifically the  
211 AOD of the total aerosols  $< 0.225$  (see Table 1 for reference). A fine (or coarse) aerosol-  
212 polluted environment must not only exceed 60% quantiles across all sampled data but  
213 also have the AOD of fine (or coarse) particles exceeding 50% of the total aerosol AOD.  
214 This approach ensures that in fine (or coarse) aerosol-polluted environments, fine (or

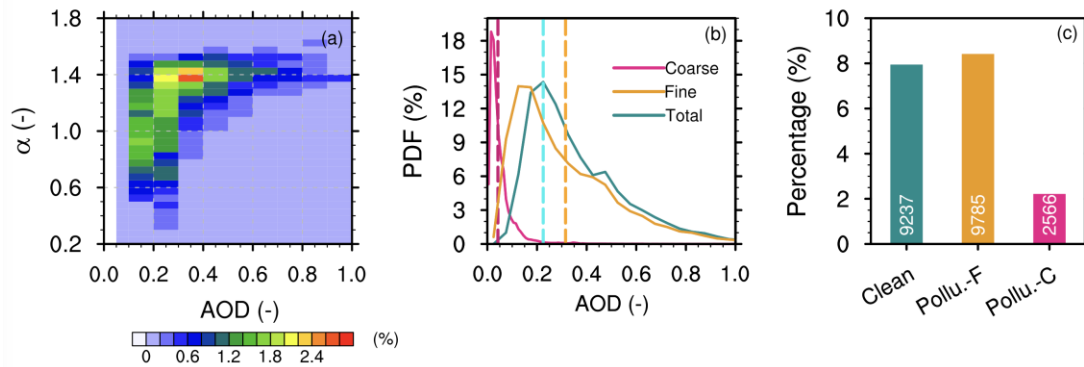


215 coarse) particles are the primary influencing factor. Based on these standards, a coarse  
216 aerosol-polluted environment is classified as having a coarse AOD > 0.0425, as well as  
217 the proportion of coarse AOD to total aerosols exceeds 50%. Similarly, a fine aerosol-  
218 polluted environment is defined by a fine AOD > 0.315, with the proportion of fine  
219 AOD to total aerosols exceeding 50% (see Table 1 for reference). A sensitivity test was  
220 conducted with different thresholds to ensure the robustness of the present study. The  
221 results indicate that varying the thresholds does not significantly affect the conclusions  
222 of the work. During the study period, there are 9237, 9785, and 2566 shallow  
223 precipitation samples under clean, fine aerosol, and coarse aerosol-polluted conditions,  
224 respectively (Figure 1c). The mean AODs of five aerosol species under various  
225 environmental conditions are calculated to understand the contributions of different  
226 aerosol types (not shown). In South China, the primary contributors to aerosol species  
227 are sulfate aerosol, sulfate aerosol, and sea salt aerosols in clean, fine, and coarse  
228 aerosol-polluted environments, respectively. The shallow precipitation accounts for a  
229 higher proportion with respect to the total precipitation samples, reaching ~8% in clean  
230 and fine aerosol-polluted conditions (Figure 1c). However, under coarse aerosol-  
231 polluted conditions, the proportion of shallow precipitation samples is much lower, at  
232 around ~2%. Due to the lower AOD of coarse aerosol mode, occurrences, where the  
233 AOD of coarse aerosols accounts for more than 50% of the total AOD are less frequent,  
234 which explains the lower shallow precipitation samples in coarse aerosol-polluted  
235 conditions. However, the approximately 2500 samples ensure the reliability of our  
236 research results to some extent.

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**Figure 1** The observed frequency of AOD and  $\alpha$  prior to the occurrence of shallow precipitation is illustrated in (a). The probability distribution functions of AOD for fine, coarse, and total aerosols before the shallow precipitation event are depicted in (b). The proportion of shallow precipitation samples relative to total precipitation samples, categorized by different aerosol conditions, is shown in (c), as recorded by DPR in southern China during the summers from 2014 to 2021. The pink vertical line (orange) in (b) represents the upper 60% threshold for fine (coarse) aerosols, respectively. The cyan vertical line in (b) denotes the lower 30% threshold for the total AOD. The shallow precipitation samples are represented by white text in (c).

**Table 1** Definitions of polluted and clean conditions of coarse and fine aerosol modes in southern China during the summers from 2014 to 2021.

Environment	Definition
Clean	Total AOD < 0.225
Polluted_Fine	Fine AOD > 0.315 & Fine AOD ratio > 50%
Polluted_Coarse	Coarse AOD > 0.0425 & Coarse AOD ratio > 50%

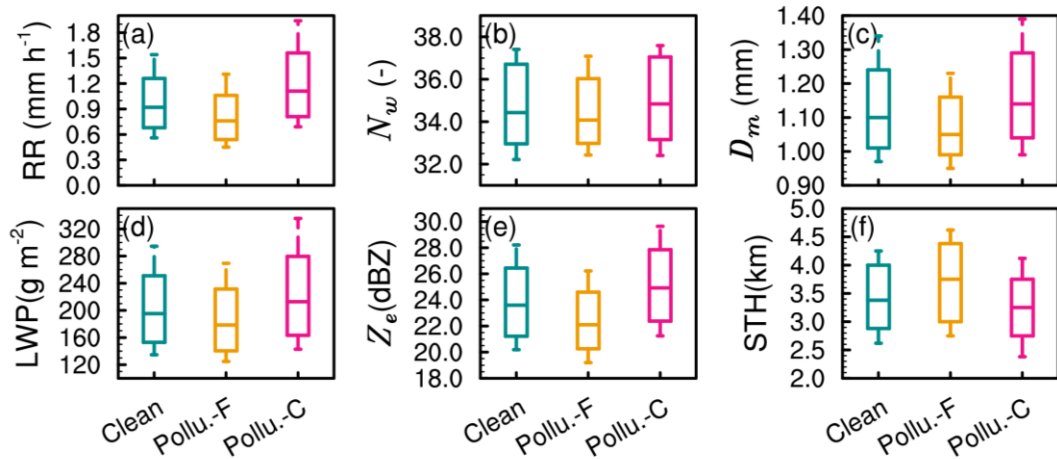
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### 254 3 Results

#### 255 3.1 Influence of aerosol on rainfall and microphysical characteristics

256 Figure 2 exhibits boxplots illustrating the near-surface RR,  $N_w$ ,  $D_m$ , and  $Z_e$  at an

257 altitude of 2.5 km, alongside LWP and STH, for shallow precipitation under varying  
258 aerosol conditions in South China. Compared to clean environment, the RR decreases  
259 slightly during fine mode aerosol pollution conditions, with a median value of only 0.7  
260 mm h<sup>-1</sup>, while in presence of coarse mode aerosol-polluted environment, the median  
261 value of RR increases, reaching 1.0 mm h<sup>-1</sup>. This is consistent with a higher median Z<sub>e</sub>  
262 at 2.5 km in altitude (25 dBZ) under coarse aerosol-polluted conditions and a lower one  
263 (22 dBZ) under fine aerosol-polluted conditions, suggesting the inhibition effect of fine  
264 particles and the invigoration effect of coarse particles on the near-surface RR for  
265 shallow precipitation. Nevertheless, the presence of coarse aerosol-polluted conditions  
266 appears to inhibit the vertical development of shallow precipitation clouds (Figure 2f),  
267 with a significantly lower median STH (~3.2 km) than that (~3.7 km) for fine aerosol-  
268 polluted environments. Examining the situation from a microphysical standpoint, it is  
269 observed that in comparison to a clean environment, there is a reduction in the median  
270 values of LWP at approximately 170 g m<sup>-2</sup>, number concentration of droplets (N<sub>w</sub>) at  
271 34, and mass-weighted mean diameter (D<sub>m</sub>) at 1.05 mm at an altitude of 2.5 km in fine  
272 mode aerosol environments. On the contrary, under coarse aerosol-polluted conditions,  
273 the median values of LWP, N<sub>w</sub>, and D<sub>m</sub> at 2.5 km altitude increase, reaching 210 g m<sup>-2</sup>,  
274 35, and 1.15 mm, respectively. This indicates that the enhancement of near-surface RR  
275 under coarse aerosol-polluted conditions is contributed by higher concentrations of  
276 large rain droplets, while the weakening under fine aerosol-polluted conditions is  
277 influenced by lower concentrations of small rain droplets. In South China, sea salt  
278 aerosols are the primary components of coarse particles, and a recent study by Liu et al.  
279 (2022) has shown that sea salt aerosols are more likely to form large cloud droplets  
280 through hygroscopic growth, which are more likely to form rain droplets through  
281 condensation and other microphysical processes, resulting in higher cloud water  
282 content within shallow precipitation clouds. On the contrary, fine aerosols tend to  
283 reduce the effective radius of cloud droplets, with small cloud droplets being prone to  
284 evaporation and subsequent loss of cloud water. Our results fill the gap between cloud  
285 microphysics, precipitation microphysics, and precipitation.



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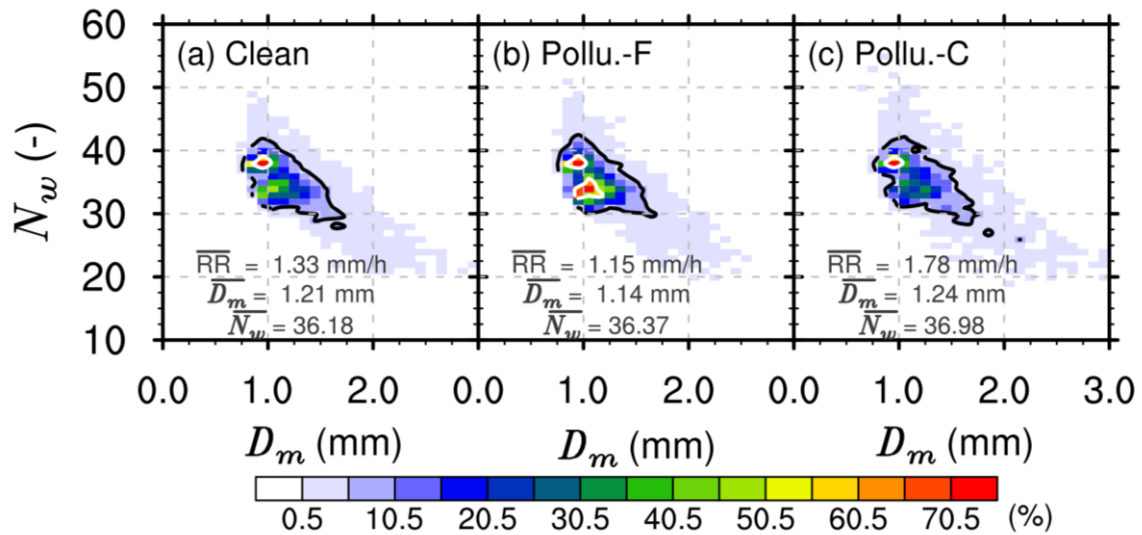
**Figure 2** The box plot presents the near-surface rain rate (a),  $N_w$  (b),  $D_m$  (c), LWP (d),  $Z_e$  (e), and STH (f) for shallow precipitation across varying aerosol conditions in southern China during the summer seasons from 2014 to 2021. The top and bottom edges of the boxes indicate the upper and lower tritile, respectively. The line inside the box denotes the median. The whiskers extending from the box illustrate the upper and lower quartiles.

DSDs directly impact RR. Therefore, the DSDs at 2.5 km altitude for shallow precipitation clouds over southern China under three aerosol conditions are illustrated in Figure 3. Irrespective of the aerosol background, the DSDs are characterized by a high concentration of small particles and a low concentration of large particles, aligning with prior research findings (Wang et al., 2016; Chen et al., 2022). In a clean environment (Figure 3a), the DSD of shallow precipitation exhibits a high-frequency center around  $N_w$  of approximately 40, with  $D_m$  around 1.0 mm, reaching a frequency exceeding 70%. A secondary peak (40%) slightly shifts towards the lower right, located at  $D_m$  around 1.2 mm and  $N_w$  around 32. In the case of fine aerosol-polluted environments (Figure 3b), the average RR ( $1.15 \text{ mm h}^{-1}$ ) and  $D_m$  (1.14 mm) are slightly reduced compared to the clean environment, while the mean  $N_w$  increases slightly to 36.37. Furthermore, the secondary peak observed in a clean environment becomes more pronounced under fine aerosol-polluted conditions, with a frequency exceeding 50%. In contrast to clean and fine aerosol-polluted environments, both the mean values of RR and  $N_w$  increase under coarse aerosol-polluted conditions (Figure 3c). Furthermore,

309 the DSD reveals more samples with  $D_m$  exceeding 2 mm or  $N_w$  exceeding 40, further  
 310 indicating the enhancement of RR for shallow precipitation in coarse aerosol-polluted  
 311 environments.

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315 **Figure 3** DSDs at 2.5 km altitude for shallow precipitation in clean (a), fine (b) aerosol-  
 316 polluted and coarse (c) aerosol-polluted environments over southern China during the  
 317 summers from 2014 to 2021. The mean values of  $D_m$  and  $N_w$  under different aerosol  
 318 conditions are presented in each panel. The 5% and 50% contours are indicated by black  
 319 and white solid lines, respectively.

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### 321 *3.2 Influence of aerosol on microphysical structures and processes*

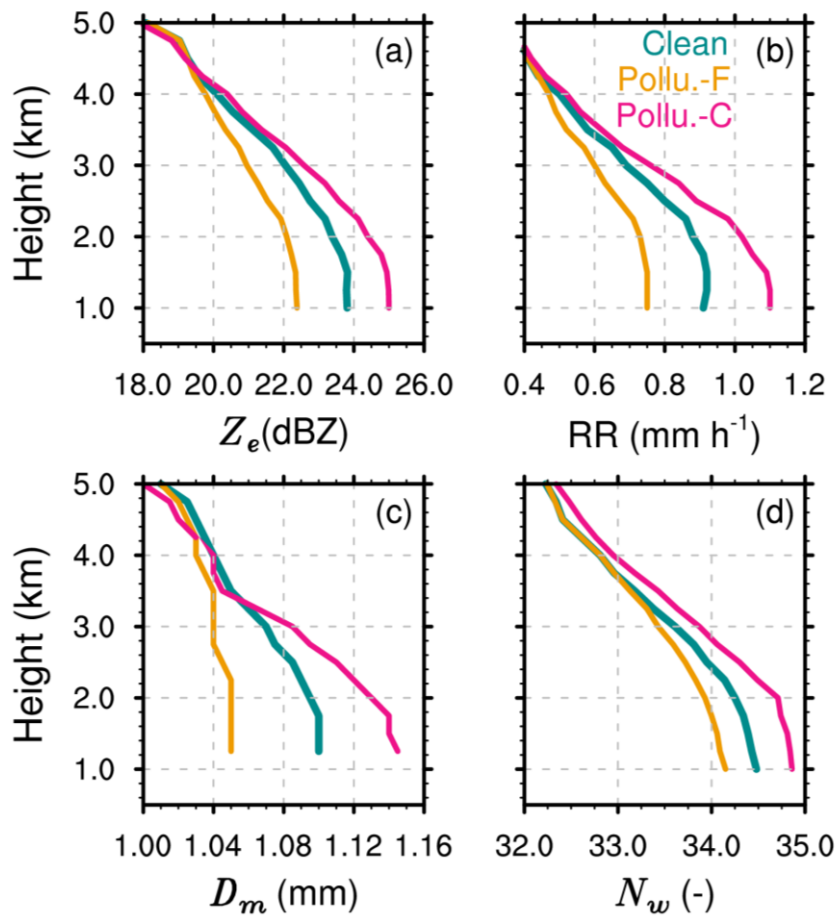
322 The above analysis has shown significant differences in near-surface RR and DSD  
 323 for shallow precipitation under different aerosol environments. The vertical structure of  
 324 precipitating clouds is closely related to near-surface RR and DSD, reflecting the  
 325 thermal and dynamic structure within the clouds. Investigating the precipitation and  
 326 microphysical structures under different aerosol backgrounds can further deepen our  
 327 understanding of the thermodynamic and microphysical mechanisms by which aerosols  
 328 affect shallow precipitation near the surface.

329 Figure 4 presents the profiles of the median values of  $Z_e$ , RR,  $D_m$ , and  $N_w$  for  
330 shallow precipitation over southern China in summer in three different aerosol  
331 environments. In general, shallow precipitation exhibits an increase in  $Z_e$ , RR,  $D_m$ , and  
332  $N_w$  with a decrease in altitude across various aerosol environments, suggesting that the  
333 growth process of shallow precipitation is predominantly governed by warm rain  
334 collision-coalescence mechanisms. This is similar to the precipitation structures for  
335 shallow precipitation in the Yangtze-Huaihe River Basin (Chen et al., 2024). However,  
336 the median values of  $Z_e$ , RR,  $D_m$ , and  $N_w$  at each altitude differ under different aerosol  
337 environments. The promotion effect of coarse aerosols and the inhibition effect of fine  
338 aerosols are present throughout the profile. For example, the median values of  $Z_e$ , RR,  
339  $D_m$ , and  $N_w$  at any given altitude are the largest in a coarse aerosol-polluted environment  
340 and the smallest in a fine aerosol-polluted pollution. Furthermore, the most significant  
341 differences in precipitation microphysical structures under different aerosol  
342 backgrounds occur near the surface (below 2 km). For example, at 1 km altitude, the  
343 differences in  $Z_e$ , RR,  $D_m$ , and  $N_w$  are approximately 3 dBZ, 0.4 mm h<sup>-1</sup>, 0.12 mm and  
344 1, respectively.

345 Considering the increasing amplitude of the median values of  $Z_e$ , RR,  $D_m$ , and  $N_w$   
346 with decreasing altitude, there are significant differences under different aerosol  
347 backgrounds, reflecting different microphysical precipitation processes within shallow  
348 precipitation systems. Specifically, in coarse aerosol-polluted environments, the  
349 increases in  $Z_e$ , RR,  $D_m$ , and  $N_w$  within the same altitude layer are the largest, while the  
350 increases in these variables are the smallest in fine aerosol-polluted environments. This  
351 explains why a concentration increase of coarse particles results in an enhancement of  
352 RR compared to a clean environment, whereas an increase in fine aerosols leads to a  
353 precipitation suppression. For instance, the median  $D_m$  in pristine environments shows  
354 an increment from 1.07 mm at 3 km altitude to 1.1 mm at 1 km. In environments  
355 polluted by coarse aerosols,  $D_m$  exhibits a more pronounced increasing trend, with the  
356 median  $D_m$  rising from 1.08 mm at 3 km to 1.14 mm at 1 km. Conversely, with fine  
357 mode aerosols, the change in the median  $D_m$  from 3 km to 1 km is negligible, almost  
358 remaining constant at approximately 1.04 mm.

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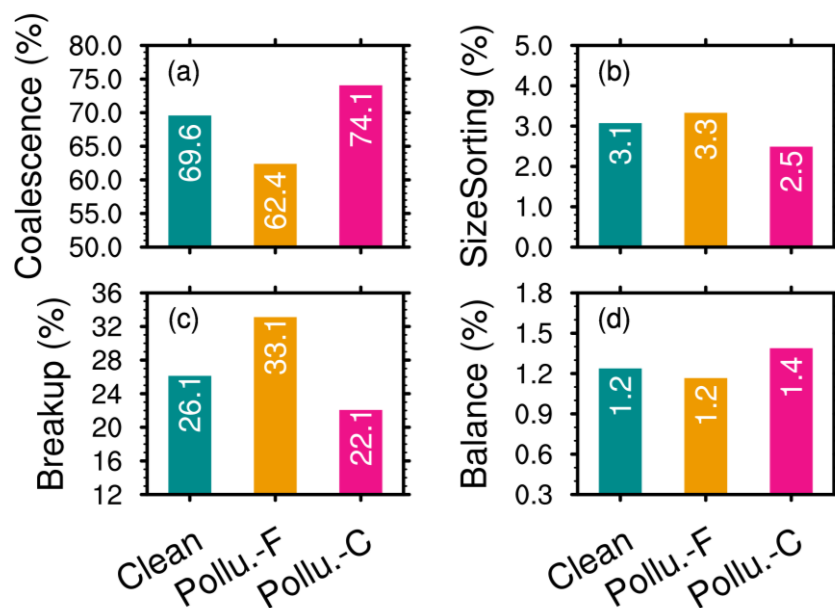
373

**Figure 4** The profiles of the median  $Z_e$  (a), rain rate (b),  $D_m$  (c), and  $N_w$  (d) for shallow precipitation in different aerosol conditions over southern China during the summers from 2014 to 2021.

To more intuitively reflect the potential impact of different aerosol types on the near-surface microphysical processes of shallow precipitation, the methods of Kumjian et al. (2014) are adopted to quantify the near-surface microphysical processes using changes in  $Z_e$  ( $\Delta Z_e = Z_e^{1\text{km}} - Z_e^{3\text{km}}$ ) and  $D_m$  ( $\Delta D_m = D_m^{1\text{km}} - D_m^{3\text{km}}$ ) at 3 km and 1 km. For example, collision-coalescence typically causes increases in  $Z_e$  and  $D_m$ , while breakup causes decreases. Similarly, an upward trend in  $D_m$  combined with a downward trend in  $Z_e$  as they approach the ground (positive  $\Delta D_m$  and negative  $\Delta Z_e$ ) indicates evaporation or size sorting is the dominant process. The signature of a "balance"

374 between collision-coalescence and breakup is shown by a minor reduction in  $D_m$  and a  
375 rise in  $Z_e$ .

376 Figure 5 shows the proportions of collision-coalescence, size sorting, breakup, and  
377 balance processes of raindrop particles in shallow precipitation clouds under three  
378 different aerosol backgrounds. In general, the microphysical process of collision-  
379 coalescence of hydrometeors dominates shallow precipitation, accounting for more  
380 than 60%. This is followed by the hydrometeor breakup process, which accounts for  
381 more than 20%, while size sorting and balance processes account for the smallest  
382 proportions, only about 3% and 1%, respectively. In presence of fine aerosol-mode, the  
383 proportion of the collision-coalescence process is only 62.4%, while this proportion  
384 reaches 74.1% in coarse aerosol-polluted environments, with an increase of about  
385 11.7%. Similarly, the proportion of the hydrometeor particle breakup process is 33.1%  
386 (a decrease of 10%). This indicates the increase in the proportion of raindrop breakup  
387 processes and the weakening of the collision-coalescence process in fine aerosol-  
388 polluted environments, which may be the reason for the weakened near-surface RR.  
389 Conversely, in coarse aerosol-polluted mode environments, raindrop hydrometeors  
390 undergo more collision-coalescence growth processes and fewer breakup and  
391 evaporation processes, which contributes to the enhancement of surface RR.





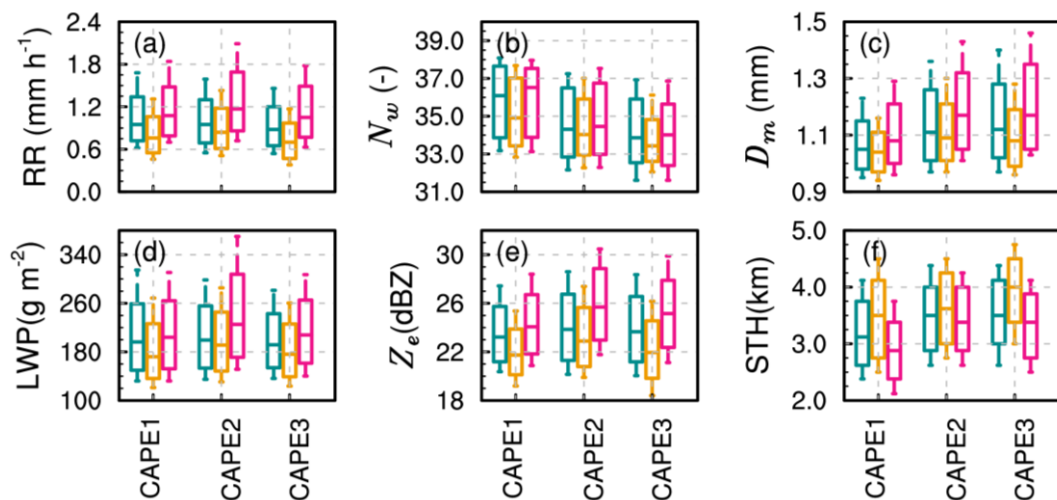
393 **Figure 5** The percentages of coalescence (a), size sorting (b), break up(c), and  
394 balance (d) for shallow precipitation shallow precipitation rain hydrometeors under  
395 different aerosol conditions in southern China during the summers from 2014 to 2021.  
396

### 397 *3.3 Sensitivities of aerosol impacts on precipitation to meteorological factors*

398 The findings from the prior section demonstrate that shallow precipitation shows  
399 notable variations in surface RR, precipitation structures, and microphysical processes  
400 depending on different aerosol conditions. However, precipitation itself is a complex  
401 process influenced by multiple thermal and dynamic environmental factors, such as  
402 instability, humidity, temperature, and wind vectors. Among these, dynamic conditions  
403 and moisture levels are particularly important indicators. Therefore, CAPE and RH at  
404 850 hPa, which, respectively, reflect atmospheric instability and moisture, are used to  
405 isolate and assess the impact of aerosols. CAPE is divided into three intervals based on  
406 the terciles of CAPE values during precipitation events in southern China: CAPE 333  
407  $\text{J kg}^{-1}$  (CAPE1),  $333 < \text{CAPE} < 1031 \text{ J kg}^{-1}$  (CAPE2), and CAPE 1031  $\text{J kg}^{-1}$  (CAPE3).  
408 Similarly, RH at 850 hPa is divided into three intervals, that is, RH 83% (RH1),  $83\% <$   
409  $\text{RH} < 91\%$  (RH2), and RH 91% (RH3).

410 The box plots of RR, LWP, and STH, as well as  $N_w$ ,  $D_m$ , and  $Z_e$  at 2.5 km altitude  
411 for shallow precipitation in southern China under different aerosol backgrounds and  
412 CAPEs are presented in Figure 6. Consistent with the conclusions of Figure 2, it  
413 becomes apparent that under varying CAPE conditions, the median STH of shallow  
414 precipitation clouds attains its lowest values in coarse aerosol-polluted environments,  
415 whereas the median RR and  $Z_e$  at an altitude of 2.5 km reach their highest levels. On  
416 the contrary, the median STH is the highest, but the median RR and  $Z_e$  at 2.5 km are the  
417 lowest in a fine aerosol-polluted environment. This indicates that the suppression of RR  
418 in fine aerosol-polluted environments and the invigoration of RR in coarse aerosol-  
419 polluted environments are independent of the dynamic conditions (CAPE in this case).  
420 Furthermore, when seen from microphysics, under different CAPE conditions, shallow

421 precipitation clouds in coarse aerosol-polluted environments exhibit the highest median  
 422 values of values of LWP,  $N_w$ , and  $D_m$  at 2.5 km, while these variables are the lowest in  
 423 fine aerosol-polluted environments. This helps explain why shallow precipitation has  
 424 the highest near-surface RR in coarse aerosol-polluted environments and the lowest  
 425 surface RR in fine aerosol-polluted environments from the microphysical perspective.  
 426



427

428 **Figure 6** Box plot of the near-surface rain rate (a),  $N_w$  (b),  $D_m$  (c), LWP (d),  $Z_e$  (e),  
 429 and STH (f) under different aerosol and CAPE conditions for shallow precipitation over  
 430 southern China during the summers of 2014-2021. The boxes' top and bottom edges  
 431 indicate the upper and lower tritile, respectively. The median is depicted by the line  
 432 inside the box. The whiskers extending from the box illustrate the lower and upper  
 433 quartiles.

434

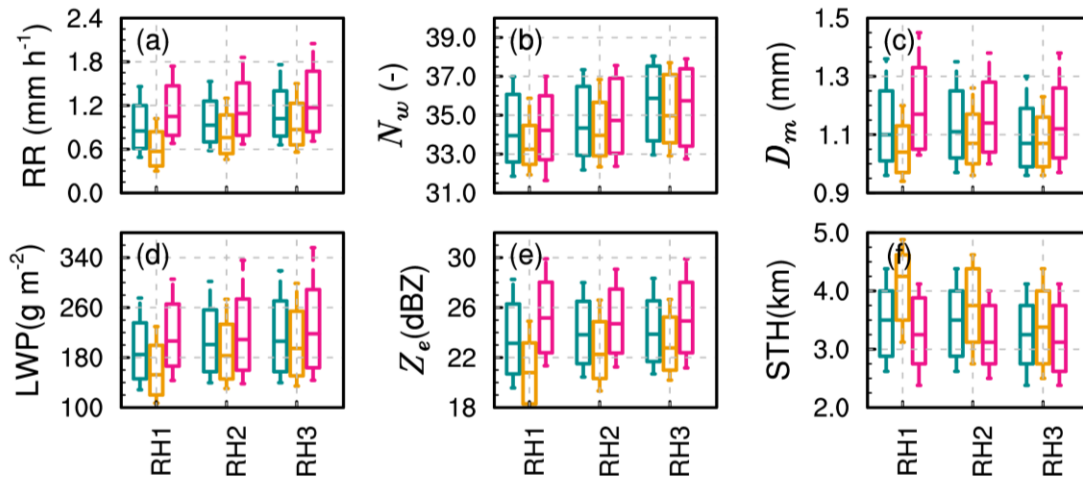
435 Similarly, the sensitivity of humidity to the impact of aerosol on shallow  
 436 precipitation is examined by presenting the box plots of precipitation parameters, as  
 437 illustrated in Figure 7. Regardless of 850hPa-RH, the vertical development of shallow  
 438 precipitation clouds is hindered in coarse aerosol-polluted environments, with the  
 439 median STH being the smallest. However, the near-surface RR is the highest,  
 440 corresponding to the highest median  $Z_e$  at 2.5 km. On the contrary, in fine particle  
 441 pollution environments, the vertical development of shallow precipitation clouds is

442 enhanced (with the highest median STH), but the near-surface RR and  $Z_e$  are the  
443 weakest. This further confirms that the impact of coarse and fine aerosols on near-  
444 surface RR and LWP is independent of moisture and dynamic conditions.

445 It is important to note that the degree of enhancement or suppression of RR by  
446 coarse and fine aerosols varies under different humidity conditions. Compared to high-  
447 humidity environments, coarse aerosols have the most significant enhancement effect  
448 on RR, while fine aerosols have the most significant suppression effect in relatively  
449 low-humidity environments (RH1). In fine aerosol-polluted environments, the box plot  
450 of RR shows a significant decrease compared to that in clean environments, while [that](#)  
451 [in coarse aerosol-polluted environments](#) shows a significant increase. Specifically, the  
452 median RR in the coarse aerosol-polluted environment is around  $1.1 \text{ mm h}^{-1}$ , [while](#) it is  
453 around  $0.7 \text{ mm h}^{-1}$  in the fine aerosol-polluted environment.

454 Regarding STH, under low relative humidity and fine aerosol pollution conditions,  
455 shallow precipitation clouds develop more deeply, with the 25th percentile of STH  
456 reaching 5 km, significantly higher than in clean and coarse aerosol-polluted  
457 environments. This may be because there is a reduction in the effective radius of cloud  
458 droplets in fine aerosol-polluted and low-humidity conditions. Smaller cloud droplets  
459 are more prone to evaporation, resulting in a lower LWP, which does not favor an  
460 increase in near-surface RR. This is also reflected in the near-surface DSD, which is  
461 characterized by lower  $N_w$  and smaller  $D_m$ . However, although the humidity is relatively  
462 low, the coarse particles, being more hygroscopic, can form larger cloud droplets,  
463 reducing the loss of cloud water due to evaporation (resulting in a higher LWP), and  
464 thereby enhancing surface RR. This is also reflected in the near-surface DSD, which is  
465 characterized by a higher  $N_w$  and larger  $D_m$ . In high humidity environments, a high  
466 concentration of fine particles can promote the formation of more cloud condensation  
467 nuclei, which to some extent reduces the loss of cloud water due to the evaporation of  
468 small particles. Therefore, the LWP in fine-particle pollution environments does not  
469 differ much from that in coarse aerosol-polluted environments. This may [also](#) lead to  
470 smaller differences in RR,  $Z_e$ , and other variables between coarse and fine aerosol-  
471 polluted environments under relatively high humidity conditions.

472



473

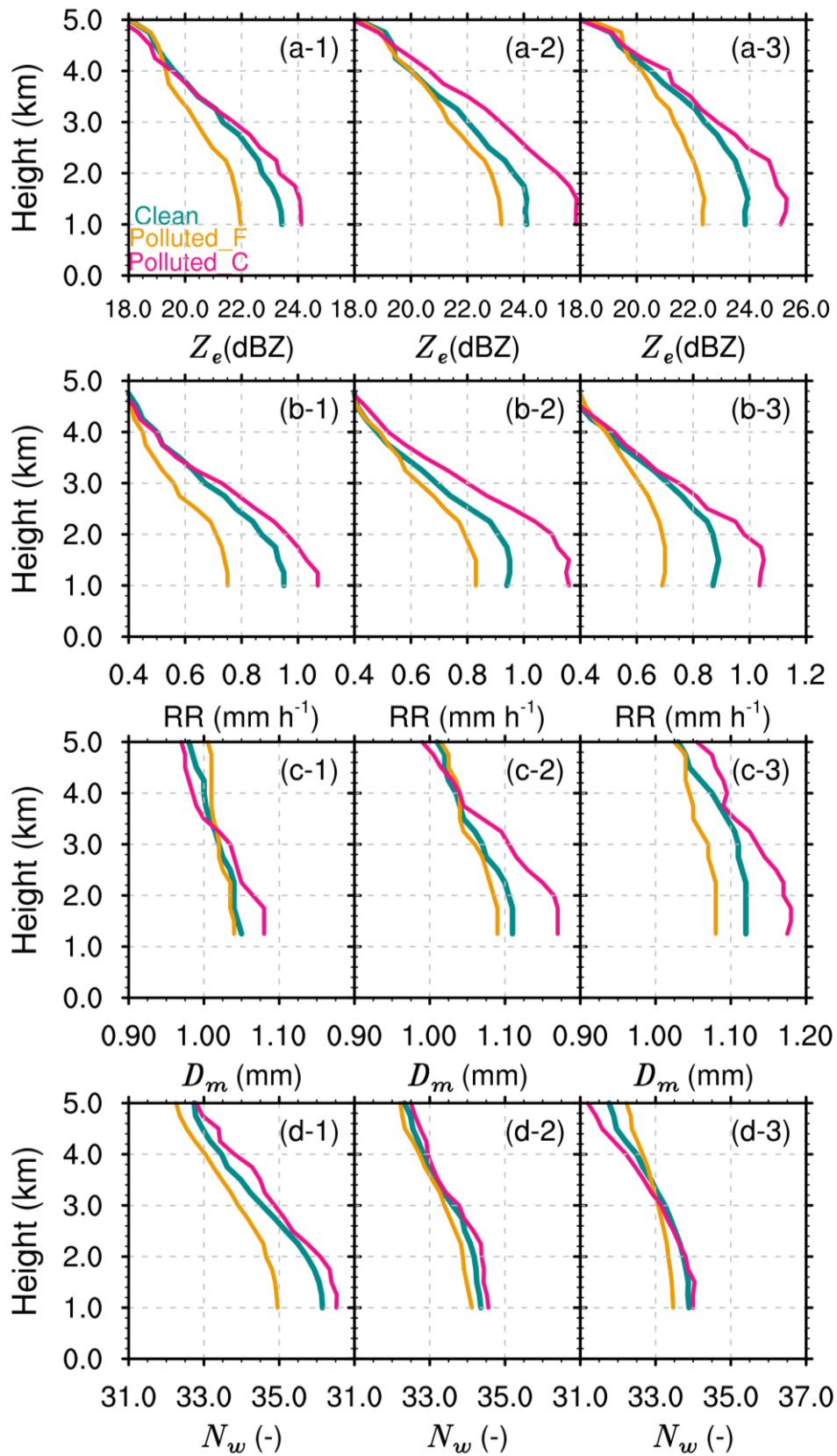
474 **Figure 7** Same as Figure 6, but for RH at 850hPa.

475

476 **3.4 Sensitivities of aerosol impacts on microphysical structures and processes to**  
477 **meteorological factors**

478 This part continues examining how coarse and fine aerosol modes affect  
479 precipitation structure and the microphysical processes in different environmental  
480 settings. As shown in Figure 8, under different CAPE and aerosol backgrounds, shallow  
481 precipitation profiles consistently exhibit increasing trends in Z<sub>e</sub>, RR, N<sub>w</sub>, and D<sub>m</sub> with  
482 decreasing altitude. Furthermore, irrespective of CAPE values, at a specified altitude,  
483 the parameters Z<sub>e</sub> and RR are observed to be at their maximum in aerosol coarse mode  
484 environments polluted with coarse aerosols, followed by those in a clean environment,  
485 and at their minimum in environments polluted with fine aerosols. This is consistent  
486 with the results in Figure 4. When compared between different CAPE conditions, the  
487 Z<sub>e</sub>, RR, and D<sub>m</sub> of shallow precipitation in CAPE2 are the highest at different altitudes,  
488 while as the CAPE increases further (CAPE3), these values even decrease. Apart from  
489 instability, precipitation can be influenced by moisture, topography, and other factors;  
490 therefore, it is possible for an even lower RR in high CAPE conditions.

491           When seen from  $D_m$  and  $N_w$  (Figures 8c1-c3, d1-d3), the promotion effect of  
492 coarse aerosols and the suppression effect of fine aerosols can vary under different  
493 dynamic environmental conditions. Under moderate CAPE conditions (CAPE2),  $D_m$   
494 and  $N_w$  in coarse aerosol-polluted environments are the largest at different altitudes,  
495 while  $D_m$  and  $N_w$  in a fine aerosol-polluted environment are the smallest. This indicates  
496 that under moderate CAPE conditions, the enhancement of RR in coarse aerosol-  
497 polluted environments is contributed by large particles and high concentrations. For  
498 low CAPE conditions (CAPE1), the median  $D_m$  above 3 km is even the smallest in  
499 coarse aerosol-polluted environments, compared to clean and fine aerosol-polluted  
500 environments. Therefore, the maximum values of RR and  $Z_e$  at this layer are mainly  
501 contributed by high concentrations of raindrop particles (with large median  $N_w$ , as  
502 shown in Figure 8d-1). For high CAPE conditions (CAPE3), the median  $N_w$  above the  
503 3 km altitude layer in coarse aerosol-polluted environments is even the smallest.  
504 Therefore, the maximum values of RR and  $Z_e$  at this altitude are mainly contributed by  
505 high concentrations of raindrop particles (with large median  $D_m$ , as shown in Figure 8c-  
506 3).



507

508 **Figure 8** The  $Z_e$  (a), rain rate (b),  $D_m$  (c), and  $N_w$  (d) profiles for shallow precipitation  
 509 in different aerosol and CAPE conditions over southern China during the summers from

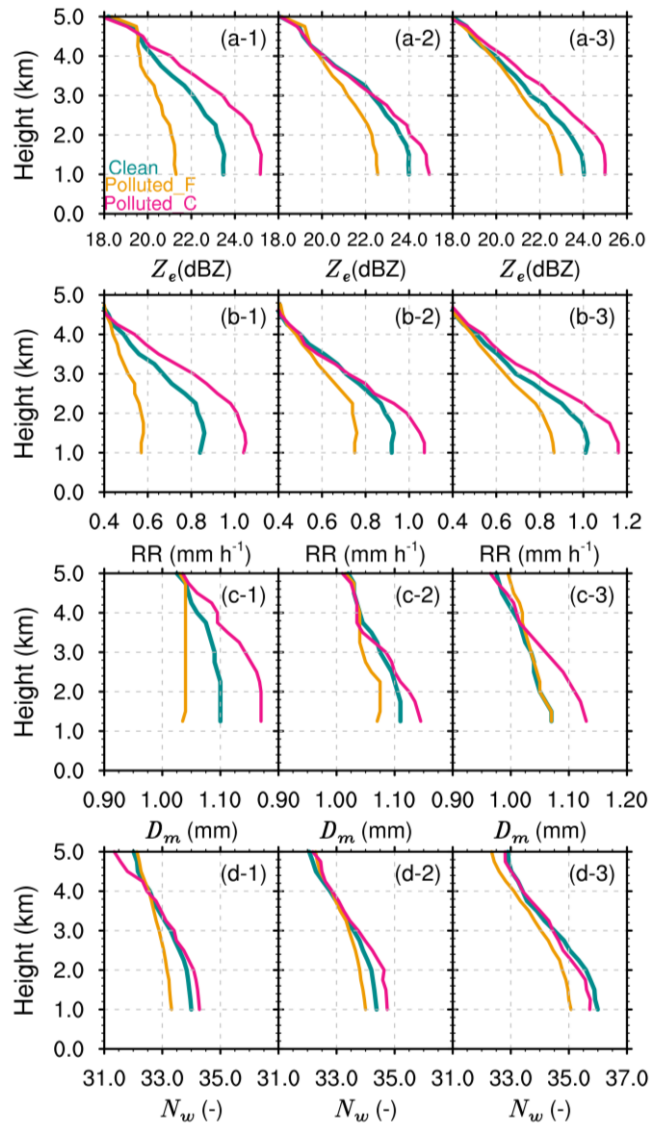
510 2014 to 2021. CAPE1, CAPE2, and CAPE3 are shown in the left, middle, and right  
511 panels, respectively.

512

513 Similarly, the profiles of  $Z_e$ , RR,  $D_m$ , and  $N_w$  in different 850hPa-RH and aerosol  
514 backgrounds are illustrated in Figure 9. Consistent with previous research results, the  
515 median values of  $Z_e$ , RR,  $D_m$ , and  $N_w$  of shallow precipitation exhibit a gradual increase  
516 with decreasing altitude, reflecting the warm rain collision-coalescence growth process.  
517 However, the microphysical structures of shallow precipitation vary under different RH  
518 conditions with similar aerosol backgrounds. As RH at 850hPa increases, the median  
519 values of  $Z_e$ , RR,  $D_m$ , and  $N_w$  of shallow precipitation increase more significantly with  
520 decreasing altitude. For example, under low humidity conditions (RH1), the median  $D_m$   
521 increases slightly when hydrometeors fall from 3 km to 1 km (Figure 9c-1), and even  
522 decreases under fine aerosol-polluted conditions, indicating more breakup processes.  
523 Subsequently, with increasing humidity, the increase in  $D_m$  becomes more apparent  
524 (Figure 9c-3). For example, the median  $D_m$  increases from 1.05 mm to 1.15 mm in  
525 coarse aerosol-polluted environments.

526 The median values of  $Z_e$  and RR across various aerosol backgrounds are markedly  
527 elevated in environments contaminated with coarse aerosols across all altitude layers,  
528 demonstrating a notable increase with decreasing altitude. Conversely, in conditions  
529 contaminated by fine aerosols, the median values of  $Z_e$  and RR are at their lowest at  
530 each altitude layer, exhibiting minimal increases as altitude decreases. This is consistent  
531 with previous conclusions (Figures 4 and 8), further indicating that the impact of coarse  
532 and fine aerosols on the near-surface RR and the precipitation structure is not sensitive  
533 to dynamic and moisture conditions. However, from a microphysical structure  
534 perspective, there are still some differences in aerosol backgrounds. Under low and  
535 moderate humidity conditions (RH1 and RH2), at a given altitude,  $D_m$  and  $N_w$  are the  
536 largest in coarse aerosol-polluted environments and the smallest in fine aerosol-polluted  
537 environments. In RH3 conditions at the same altitude, a clean environment has the  
538 highest  $N_w$  and a relatively small  $D_m$ ; for coarse mode,  $N_w$  is moderate with the largest  
539  $D_m$ ; and for fine mode,  $N_w$  is the lowest with a relatively small  $D_m$ . This indicates that  
540 in high RH environments, fine aerosols mainly reduce RR by suppressing the  
541 concentration of raindrops, while coarse aerosols increase RR by increasing the size of  
542 hydrometeors. Furthermore, the differences in precipitation structures in aerosol-  
543 polluted coarse and fine environments depend on humidity conditions, consistent with

544 the conclusions in Figure 7. The differences are the greatest under RH1 conditions, with  
 545 the differences in RR,  $Z_e$ ,  $D_m$ , and  $N_w$  at 1 km altitude being  $0.42 \text{ mm h}^{-1}$ , 4.5 dBZ, 0.19  
 546 mm, and about 1.3, respectively. Under RH3 conditions, the differences are smallest,  
 547 with the differences in the aforementioned variables being  $0.35 \text{ mm h}^{-1}$ , 2 dBZ, 0.05  
 548 mm, and approximately 0.8, respectively.



549

550 **Figure 9** The  $Z_e$  (a), rain rate (b),  $D_m$  (c), and  $N_w$  (d) profiles for shallow precipitation  
 551 in different aerosol conditions and 850 hPa-RH over southern China during the  
 552 summers from 2014 to 2021. RH1, RH2, and RH3 are shown in left, middle, and right  
 553 panels, respectively.

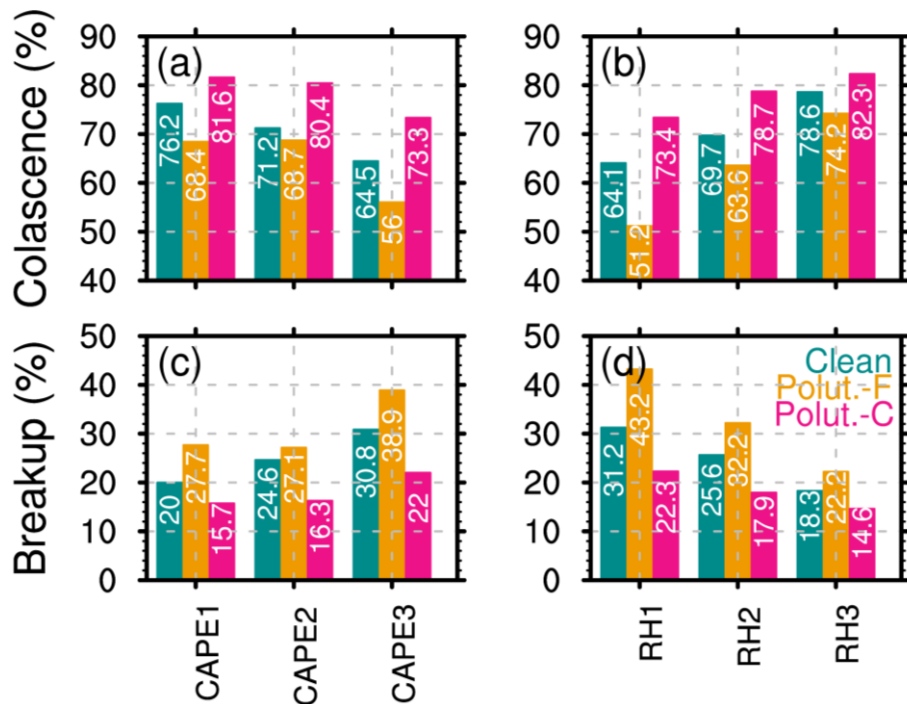
554



555 To quantitatively analyze the dependence of microphysical processes on dynamics  
556 and moisture under different aerosol backgrounds, we examined the differences in the  
557 two primary microphysical processes, i.e., collision-coalescence and breakup. As a  
558 result of the low proportions of size sorting and balance, further analysis of these  
559 microphysical processes is not included. The microphysical processes of precipitation  
560 depend on the dynamic and moisture conditions. For instance, with decreasing CAPE  
561 and increasing RH, the proportion of collision-coalescence increases, while the  
562 proportion of breakup decreases in clean, coarse, and fine aerosol-polluted  
563 environments. **Environments with high RH and low CAPE encourage aerosol particles**  
564 **in the boundary layer to gather moisture, leading to the formation of additional cloud**  
565 **droplets. These droplets then condense further to create more raindrops, thus enhancing**  
566 **the collision-coalescence process.**

567 After comparing various aerosol backgrounds, it is possible to determine certain  
568 overarching patterns that remain consistent regardless of thermodynamic conditions.  
569 **Initially, irrespective of CAPE, RH, or aerosol background, shallow precipitation**  
570 **systems predominantly exhibit the warm rain collision-coalescence process, with its**  
571 **occurrence proportion spanning from a minimum of 51.2% to a maximum of**  
572 **82.3%. There is also a certain proportion of break-up processes, ranging from 14.6% to**  
573 **43.2%. Second, regardless of the value of CAPE and RH, the proportion of the**  
574 **collision-coalescence process is always the highest in coarse aerosol-polluted**  
575 **environments, while the proportion of the breakup process is always the highest in fine**  
576 **aerosol-polluted environments. These conclusions are consistent with the results in**  
577 **Figure 5.** However, the increase in the proportion of collision coalescence in coarse  
578 aerosol-polluted environments and the increase in the proportion of breakup in fine  
579 aerosol-polluted environments depend on dynamic and moisture conditions. For  
580 example, under low relative humidity (RH1) conditions, the proportion of the collision-  
581 coalescence process in coarse aerosol-polluted environments (73.4%) is significantly  
582 higher than that in fine aerosol-polluted environments (51.2%), **with an enhancement**  
583 **of 22.2%.** On the contrary, the proportion of the breakup process in fine aerosol-  
584 polluted environments (43.2%) is significantly higher than in coarse aerosol-polluted  
585 environments (22.3%). This is consistent with previous findings that under RH1  
586 conditions,  $D_m$  in fine aerosol-polluted environments rapidly decreases with decreasing  
587 altitude.

588



589

590 **Figure 10** The percentages of coalescence (a), size sorting (b), break-up(c), and balance  
 591 (d) for shallow precipitation rain hydrometeors under different aerosol conditions in  
 592 southern China during the summers from 2014 to 2021.

#### 593 **4 Conclusion and Discussion**

594 Using the combined data of DPR, MERRA-2 aerosol datasets, and ERA5 during  
 595 the summers of 2014-2021, this study investigates the potential impacts of coarse and  
 596 fine aerosol modes on the Rain Rate (RR), microphysical structure, and processes for  
 597 shallow precipitation in South China. Clean, coarse, and fine aerosol-polluted modes  
 598 are classified according to the AOD for total aerosols, coarse aerosols, and fine aerosols  
 599 derived from MERRA-2 products. ERA5 reanalysis data are used to explore the  
 600 sensitivity of aerosol impacts on shallow precipitation to dynamic and moisture  
 601 conditions in South China. The main findings are summarized as follows.

602 In comparison to clean environments, coarse aerosol-polluted environments  
 603 enhance near-surface rainfall rates of shallow precipitation, characterized by stronger  
 604 near-surface RR (average precipitation intensity of  $1.78 \text{ mm h}^{-1}$ ), higher concentrations

605 (average  $N_w = 36.98$ ) and larger raindrop sizes (average  $D_m = 1.24$  mm) of hydrometeor  
606 particles. This can be ascribed to the high presence of sea salt aerosols in South China,  
607 which tend to form larger cloud droplets through hygroscopic growth, leading to larger  
608 raindrop particles through microphysical processes such as condensation. On the  
609 contrary, fine aerosol mode suppress near-surface RR, with an average near-surface RR  
610 of only  $1.33$  mm h<sup>-1</sup> and lower concentrations and smaller sizes of hydrometeors  
611 (average  $N_w = 36.37$ , average  $D_m = 1.14$  mm). Liu et al. (2022) noted similar opposing  
612 effects of fine aerosols and coarse sea spray on warm marine clouds. Deep clouds show  
613 increased rainfall with high liquid water content but reduced rainfall if water content is  
614 low (Li et al., 2011). This underscores the distinct behavior of shallow precipitation and  
615 the varied impacts of aerosol types on it. However, fine aerosol-polluted environments  
616 promote vertical development of shallow precipitation clouds (median STH of 3.7 km),  
617 approximately 0.5 km higher than in coarse aerosol-polluted conditions. The inhibition  
618 of the vertical development of precipitation clouds by coarse aerosol particles explains  
619 their suppressive effect on lightning activity to some extent (Pan et al., 2022).

620 From the perspective of precipitation vertical structure and microphysical  
621 processes, shallow precipitation is dominated by warm-rain collision-coalescence  
622 processes under different aerosol backgrounds, with the collision-coalescence process  
623 accounting for over 62%. However, there are significant differences in the efficiency  
624 of hydrometeor collision-coalescence growth under different aerosol conditions. In  
625 contrast to clean conditions, the median values of  $Z_e$ , RR,  $D_m$ , and  $N_w$  are highest in  
626 presence of aerosol coarse mode and lowest in conditions for fine aerosol mode at all  
627 altitude levels. Looking at it from a microphysical standpoint, the increase in  $D_m$  with  
628 decreasing altitude is most pronounced under coarse aerosol-polluted conditions,  
629 reflecting more significant collision-coalescence growth processes, accounting for  
630 74.1%. In contrast, the increase in  $D_m$  with decreasing altitude is weakest under fine  
631 aerosol-polluted conditions, due to the higher proportion of breakup processes  
632 (accounting for 33.1%) and a decrease of approximately 12% in the collision-  
633 coalescence process (accounting for 62.4%). Overall, the promotion of RR is associated  
634 with more significant collision-coalescence processes by coarse aerosols, while the

635 suppression of RR is characterized by more significant breakup processes with fine  
636 aerosols.

637 The effects of fine and coarse aerosols on the suppression and enhancement of RR  
638 are independent of CAPE and humidity, consistent with the findings by Liu et al. (2022).  
639 However, our results show that the extent of suppression or enhancement varies with  
640 CAPE and humidity. Additionally, the analysis of aerosol-precipitation interactions  
641 under different surface air temperatures yields results similar to those observed for  
642 CAPE and RH at 850 hPa (figures not shown). The promotion and suppression effects  
643 are the most pronounced under low relative humidity conditions (RH1). This is mainly  
644 contributed by the stronger suppression of fine aerosols in low-humidity environments.  
645 For instance, the median RR is around 1.12 mm h<sup>-1</sup> under coarse aerosol-polluted  
646 conditions, while it is around 0.7 mm h<sup>-1</sup> under fine aerosol-polluted conditions, with a  
647 difference of approximately 0.42 mm h<sup>-1</sup>. The collision-coalescence and breakup  
648 microphysical processes play an important role in these differences, with the collision-  
649 coalescence accounting for 73.4% under coarse aerosol-polluted conditions, which is  
650 22.2% higher than the 51.2% observed under fine aerosol-polluted conditions.  
651 Correspondingly, the breakup microphysical processes account for 43.2% under fine  
652 aerosol-polluted conditions, significantly higher than the 22.3% in coarse aerosol-  
653 polluted conditions. Under high relative humidity conditions, fine aerosol-polluted  
654 environments primarily reduce RR by inhibiting hydrometeor concentration (possibly  
655 as a result of the evaporation effects of small cloud droplets), while coarse aerosols  
656 invigorate RR by increasing the size of hydrometeor particles. Additionally, the  
657 increase in RR above 3 km in coarse aerosol-polluted environments is mainly driven  
658 by the high concentration of hydrometeors in low instability conditions, while by large  
659 hydrometeors in high instability environments. It is important to note that precipitation  
660 is a complex process influenced by multiple meteorological factors, including  
661 instability, moisture, and temperature. Additionally, other factors such as wind vectors  
662 and pressure may also affect the impact of aerosols on precipitation, which is worthy  
663 of further study.

664 This study primarily elucidates the microphysical processes within shallow

665 precipitation systems under varying aerosol conditions. However, the methods and data  
666 utilized have broad application potential. Future research could extend these  
667 approaches to explore the relationship between deep convection or mixed-phase clouds  
668 and aerosols. Such investigations could reveal the complex effects of aerosols on the  
669 precipitation process and further enhance our scientific understanding of the physical  
670 connections between aerosols and precipitation microphysics. However, it is important  
671 to note that the spatial resolution of MERRA-2 and ERA5 is much coarser than that of  
672 DPR. The interpolation methods employed in the present study may introduce errors  
673 and may not fully capture the true conditions, making it challenging to accurately assess  
674 fine-scale processes in aerosol-cloud interactions. Furthermore, MERRA-2 shows a  
675 slight underestimation of approximately 0.1 compared to in-situ observations in South  
676 China (Ou et al., 2022), probably due to the absence of nitrate aerosols in the MERRA-  
677 2 dataset. Consequently, the fine aerosol-polluted environments examined in this study  
678 may not fully capture conditions with high nitrate loading. There is an urgent need for  
679 long-term observational data on aerosol concentrations with high spatiotemporal  
680 resolution and accuracy to fully capture the samples of high aerosol loading and more  
681 effectively capture fine-scale processes in aerosol-cloud interactions.

682

### 683 **Data availability**

684 The GPM DPR data provided by NASA Goddard Space Flight Center's Mesoscale  
685 Atmospheric Processes Laboratory and Precipitation Processing System (PPS) can be  
686 downloaded from <https://pmm.nasa.gov/dataaccess/downloads/gpm>. MERRA-2 data  
687 can be downloaded from [https://gmao.gsfc.nasa.gov/reanalysis/MERRA-  
688 2/data\\_access/](https://gmao.gsfc.nasa.gov/reanalysis/MERRA-2/data_access/). The ERA5 data can be downloaded from  
689 <https://www.ecmef.int/en/forecasts/dataset/ecmwf-reanalysis-v5>. The ancillary digital  
690 terrain data is from the National Geophysical Data Center (NGDC) (available online at  
691 <http://www.ngdc.noaa.gov>, accessed in May 2023).

692

693 **Author contributions**

694 YY designed the manuscript and led the data analysis; FC performed the analysis  
695 and wrote the manuscript draft; YL and LY collected the data; GL, LY, and SL  
696 reviewed and edited the manuscript; SL helped with the data analysis.

697 **Declaration of competing interest**

698 The authors declare no competing interests.

699

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709 quantum technology in meteorological detection" Youth Innovation Team Project  
710 \(No.CMA2024QN11\)](#), and the Open Project of State Key Laboratory of Severe  
711 [Weather \(2024LASW-B11\)](#).

712

713

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