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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 aerosol **particles** on shallow precipitation events.  
20 Using eight years of **data from dual-frequency precipitation radar measurements,**  
21 aerosol **reanalysis,** and **atmospheric** reanalysis, this study investigates the potential  
22 influence of coarse and fine aerosol **particles** on the structure of the precipitation and  
23 the microphysical processes of shallow precipitation in South China. Statistical results  
24 indicate that during coarse aerosol-polluted conditions, shallow precipitation clouds  
25 have a lower mean height of the storm top (STH,  $\sim 3.2$  km), but a higher mean near-  
26 surface rainfall (RR,  $\sim 1.78$  mm h<sup>-1</sup>), characterized by high concentrations of large  
27 raindrops, driven mainly by significant collision-coalescence processes (accounting for  
28 74.1%). In contrast, during fine aerosol-polluted conditions, shallow precipitation  
29 clouds develop a deeper median STH  $\sim 3.7$  km with lower surface RR characterized by  
30 a low concentration of small **raindrops,** resulting from increased breakup processes  
31 (33.1%) and reduced collision-coalescence processes (69.6%). The coarse (fine)  
32 aerosol **particles** act as promoters (inhibitors) of radar reflectivity in the profile of  
33 shallow precipitation, regardless of dynamic and humid conditions. The effect of coarse  
34 aerosol **particles** in promoting precipitation and the inhibiting effect of fine **aerosol**  
35 **particles** are the most significant under low humidity conditions, mainly attributed to  
36 significantly enhanced collision-coalescence processes, exceeding 22.2%. Furthermore,  
37 the increase in RR above 3 km during coarse aerosol-polluted environments is mainly  
38 driven by the high concentration of hydrometeors in low instability conditions, whereas  
39 by large hydrometeors in high instability environments.  
40

41 **Short Summary:** The microphysical mechanisms of precipitation responsible for the  
42 varied impacts of aerosol [particles](#) on shallow precipitation remain unclear. This study  
43 reveals that coarse [aerosol particles](#) invigorate shallow rainfall through enhanced  
44 coalescence processes, whereas fine [aerosol particles](#) suppress shallow rainfall through  
45 intensified microphysical breaks. These impacts are independent of thermodynamic  
46 environments, but are more significant in low-humidity conditions.  
47

## 48 *1 Introduction*

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

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

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

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

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



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

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## 113 ***2 Data and Methods***

### 114 ***2.1 Data***

115 In this study, four different data set are used to illustrate the potential impact of  
116 [aerosol particles](#) on microphysical precipitation structures and shallow precipitation  
117 processes over southern China during the summers between 2014 and 2021.

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

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

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

150 The GPM DPR consists of two precipitation radars operating in the Ka and Ku  
151 bands, providing a unique opportunity to obtain information on three-dimensional  
152 precipitation and particle drop size distributions (DSDs) at the same time. In the present  
153 study, the official [DPR](#) (version 7) dataset covering the summers (June to August) of  
154 2014 and 2021 is also used, which provides information on the observation time, near-  
155 surface rain rate (RR), liquid water path (LWP), the three-dimensional profiles of  
156 attenuation-corrected reflectivity ( $Z_e$ ), rainfall, the mass-weighted mean diameter  $D_m$   
157 (in mm) and the generalized intercept  $N_w$  (i.e., [number concentration of droplets](#) in  $\text{mm}^{-3}$

158  $^1 \text{ m}^{-3}$ ) of the normalized gamma distributions with a vertical resolution of 125 m in each  
159 scanning pixel (Iguchi et al., 2017). The reliability of DSDs and precipitation has been  
160 validated by many previous studies (Huang et al., 2021; Radhakrishna et al., 2016). Due  
161 to the high spatial resolution (125 m in **vertical** and 4.5 km in horizontal resolution),  
162 the official 2ADPR (version 7) dataset has been widely used in the field of climatology  
163 (Chen et al., 2024; Zhang et al., 2020a; Chen et al., 2020). Shallow precipitation clouds  
164 are defined by their near-surface RR exceeding  $0.1 \text{ mm h}^{-1}$  and **storm top height (STH)**  
165 below 5 km in altitude. The **STH** is defined as the maximum height where the  $Z_e$   
166 exceeds 20dBZ (Liu and Zipser, 2013).

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

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## 175 **2.2 Methods**

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

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

192

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

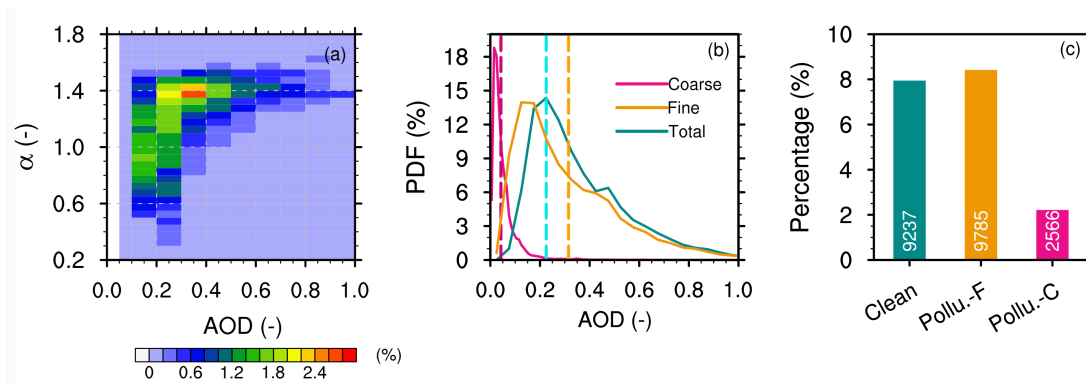
203 There are three types of aerosol conditions discussed in the present study: clean  
204 environment, fine aerosol-polluted environment, and coarse aerosol-polluted  
205 environment. To classify clean and aerosol-polluted conditions over South China, PDFs  
206 of AOD for fine, coarse, and total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

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

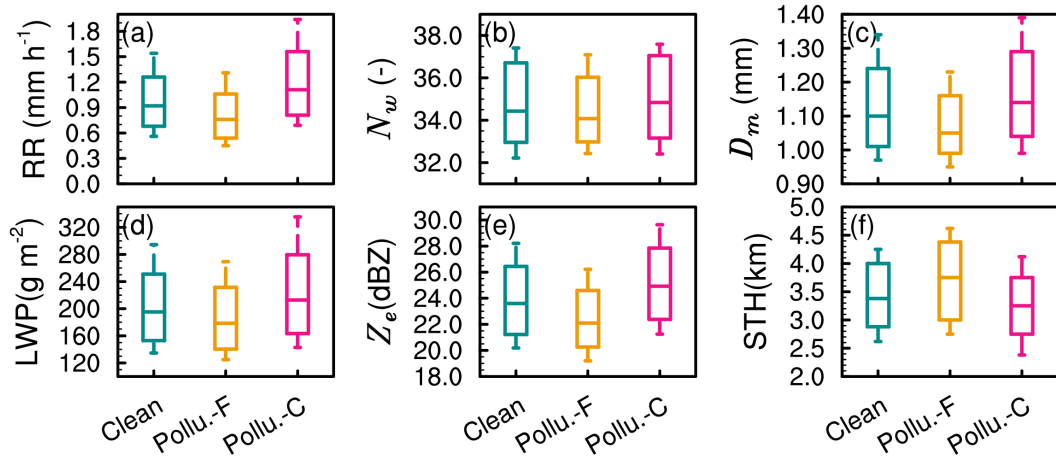
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### 257 3 Results

#### 258 3.1 Influence of aerosol particles on rainfall and microphysical characteristics

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

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



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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.

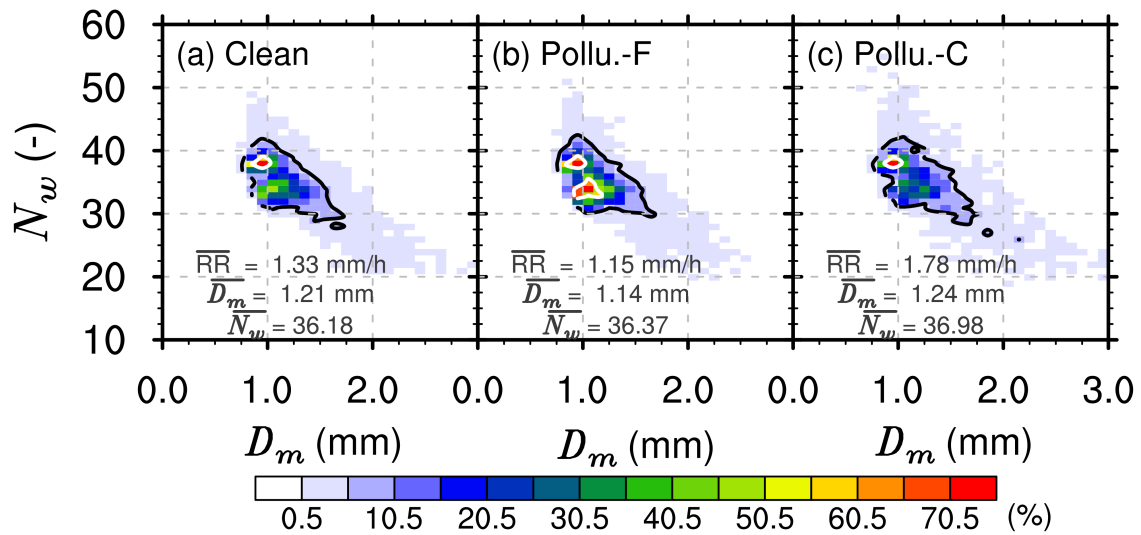
DSDs directly impact RR. Therefore, the DSDs at 2.5 km altitude for shallow precipitation clouds over southern China under three aerosol **particle** conditions are illustrated in Figure 3. Irrespective of the aerosol **particle** background, the DSDs are characterized by a high concentration of small **droplets** and a low concentration of large **droplets**, 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 mm h<sup>-1</sup>) 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



310 3c). Furthermore, the DSD reveals more samples with  $D_m$  exceeding 2 mm or  $N_w$   
 311 exceeding 40, further indicating the enhancement of RR for shallow precipitation in  
 312 coarse aerosol-polluted environments.

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

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### 322 3.2 Influence of aerosol particles on microphysical structures and processes

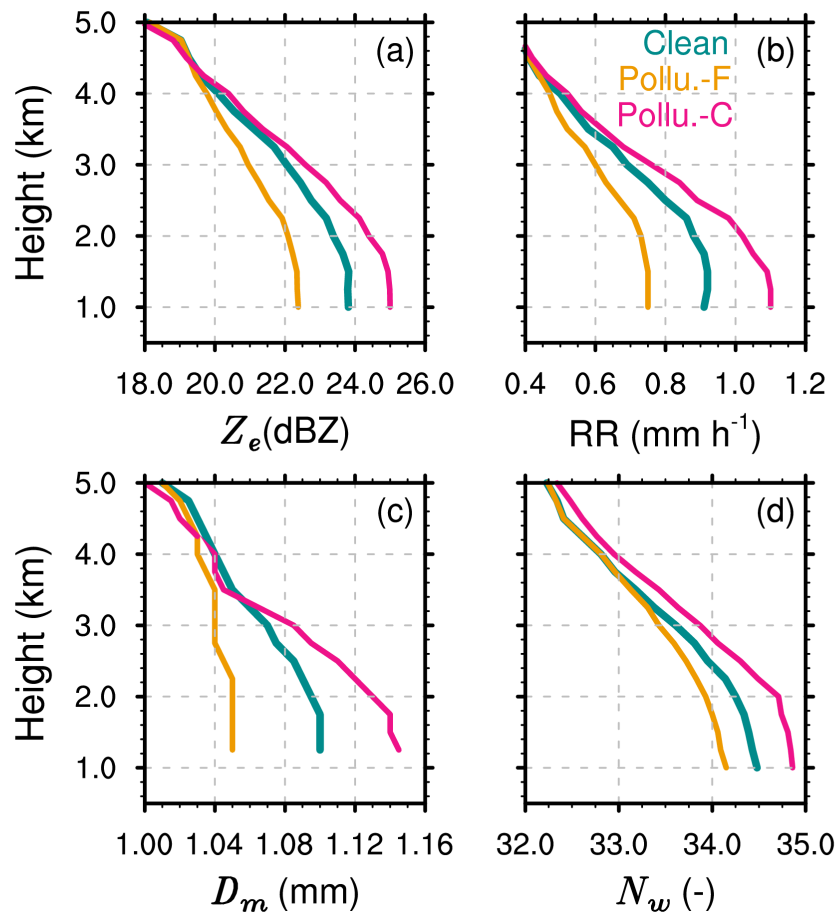
323 The above analysis has shown significant differences in near-surface RR and DSD  
 324 for shallow precipitation under different aerosol particle environments. The vertical  
 325 structure of precipitating clouds is closely related to near-surface RR and DSD,  
 326 reflecting the thermal and dynamic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

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

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

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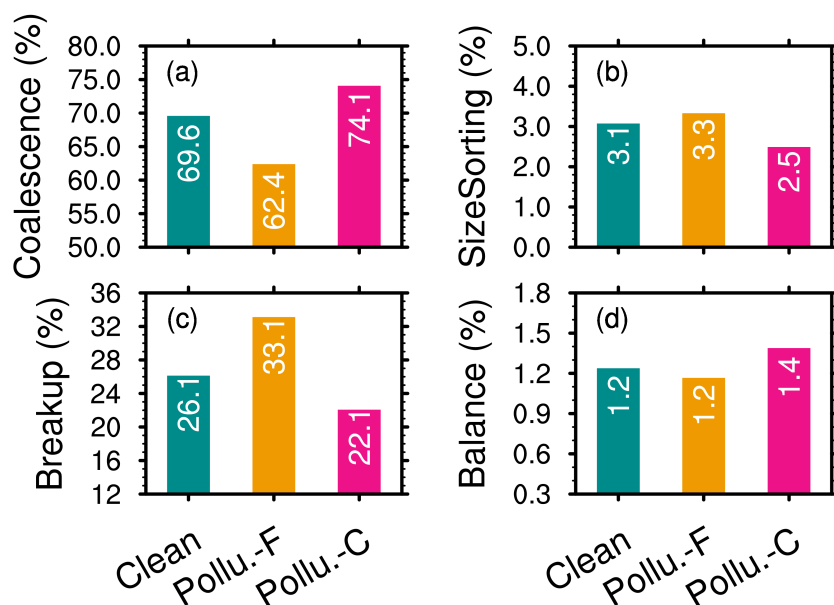
363 **Figure 4** The profiles of the median  $Z_e$  (a), rain rate (b),  $D_m$  (c), and  $N_w$  (d) for  
364 shallow precipitation in different aerosol [particle](#) conditions over southern China  
365 during the summers from 2014 to 2021.

366

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

375 signature of a "balance" between collision-coalescence and breakup is shown by a  
376 minor reduction in  $D_m$  and a rise in  $Z_e$ .

377 Figure 5 shows the proportions of collision-coalescence, size sorting, breakup, and  
378 balance processes of **raindrops** in shallow precipitation clouds under three different  
379 aerosol backgrounds. In general, the microphysical process of collision-coalescence of  
380 **raindrops** dominates shallow precipitation, accounting for more than 60%. This is  
381 followed by the **raindrop** breakup process, which accounts for more than 20%, while  
382 size sorting and balance processes account for the smallest proportions, only about 3%  
383 and 1%, respectively. In presence of fine aerosol-mode, the proportion of the collision-  
384 coalescence process is only 62.4%, while this proportion reaches 74.1% in coarse  
385 aerosol-polluted environments, with an increase of about 11.7%. Similarly, the  
386 proportion of the **raindrop** breakup process is 33.1% , with a decrease of 10% compared  
387 to 22.1% in coarse aerosol-polluted environments. This indicates the increase in the  
388 proportion of raindrop breakup processes and the weakening of the collision-  
389 coalescence process in fine aerosol-polluted environments, which may be the reason  
390 for the weakened near-surface RR. Conversely, in coarse aerosol-polluted mode  
391 environments, **raindrops** undergo more collision-coalescence growth processes and  
392 fewer breakup and evaporation processes, which contributes to the enhancement of  
393 surface RR.



394

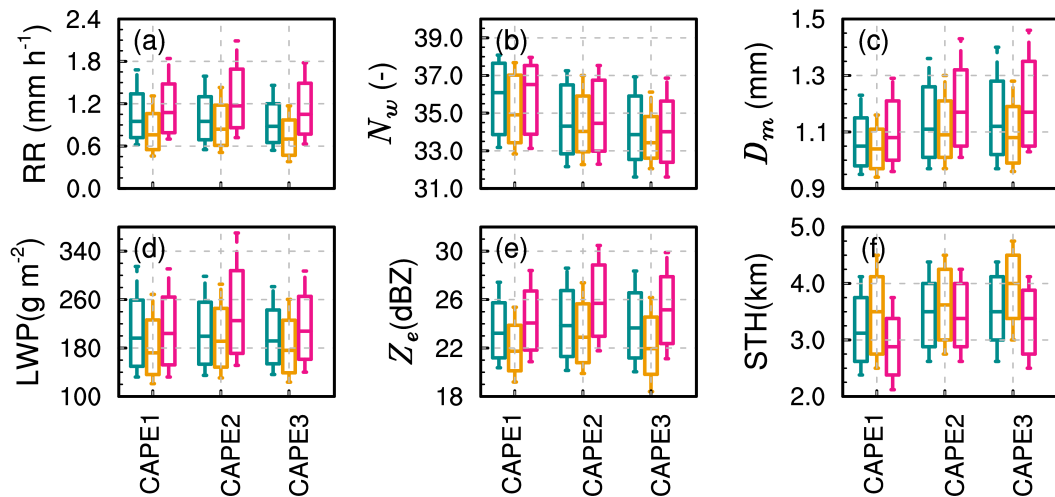
395 **Figure 5** The percentages of coalescence (a), size sorting (b), break up(c), and  
 396 balance (d) for shallow precipitation shallow precipitation **raindrops** under different  
 397 aerosol **particle** conditions in southern China during the summers from 2014 to 2021.

398

### 399 **3.3 Sensitivities of aerosol impacts on precipitation to meteorological factors**

400 The findings from the prior section demonstrate that shallow precipitation shows  
 401 notable variations in surface RR, precipitation structures, and microphysical processes  
 402 depending on different aerosol **particle** conditions. However, precipitation itself is a  
 403 complex process influenced by multiple thermal and dynamic environmental factors,  
 404 such as instability, humidity, temperature, and wind vectors. Among these, dynamic  
 405 conditions and moisture levels are particularly important indicators. Therefore, CAPE  
 406 and RH at 850 hPa, which, respectively, reflect atmospheric instability and moisture,  
 407 are used to isolate and assess the impact of aerosol particles. CAPE is divided into three  
 408 intervals based on the terciles of CAPE values during precipitation events in southern  
 409 China: CAPE  $333 \text{ J kg}^{-1}$  (CAPE1),  $333 < \text{CAPE} < 1031 \text{ J kg}^{-1}$  (CAPE2), and CAPE  
 410  $1031 \text{ J kg}^{-1}$  (CAPE3). Similarly, RH at 850 hPa is divided into three intervals, that is,  
 411 RH 83% (RH1),  $83\% < \text{RH} < 91\%$  (RH2), and RH 91% (RH3).

412 The box plots of RR, LWP, and STH, as well as  $N_w$ ,  $D_m$ , and  $Z_e$  at 2.5 km altitude  
 413 for shallow precipitation in southern China under different aerosol [particle](#) backgrounds  
 414 and CAPEs are presented in Figure 6. Consistent with the conclusions of Figure 2, it  
 415 becomes apparent that under varying CAPE conditions, the median STH of shallow  
 416 precipitation clouds attains its lowest values in coarse aerosol-polluted environments,  
 417 whereas the median RR and  $Z_e$  at an altitude of 2.5 km reach their highest levels. On  
 418 the contrary, the median STH is the highest, but the median RR and  $Z_e$  at 2.5 km are the  
 419 lowest in a fine aerosol-polluted environment. This indicates that the suppression of RR  
 420 in fine aerosol-polluted environments and the invigoration of RR in coarse aerosol-  
 421 polluted environments are independent of the dynamic conditions (CAPE in this case).  
 422 Furthermore, when seen from microphysics, under different CAPE conditions, shallow  
 423 precipitation clouds in coarse aerosol-polluted environments exhibit the highest median  
 424 values of values of LWP,  $N_w$ , and  $D_m$  at 2.5 km, while these variables are the lowest in  
 425 fine aerosol-polluted environments. This helps explain why shallow precipitation has  
 426 the highest near-surface RR in coarse aerosol-polluted environments and the lowest  
 427 surface RR in fine aerosol-polluted environments from the microphysical perspective.  
 428



429

430 **Figure 6** Box plot of the near-surface rain rate (a),  $N_w$  (b),  $D_m$  (c), LWP (d),  $Z_e$  (e),  
 431 and STH (f) under different aerosol [particle](#) and CAPE conditions for shallow  
 432 precipitation over southern China during the summers of 2014-2021. [The blue, orange](#)

433 and pink colors indicate the clean, fine-aerosol polluted and coarse-aerosol polluted  
434 environments, respectively. The boxes' top and bottom edges indicate the upper and  
435 lower tritile, respectively. The median is depicted by the line inside the box. The  
436 whiskers extending from the box illustrate the lower and upper quartiles.

437

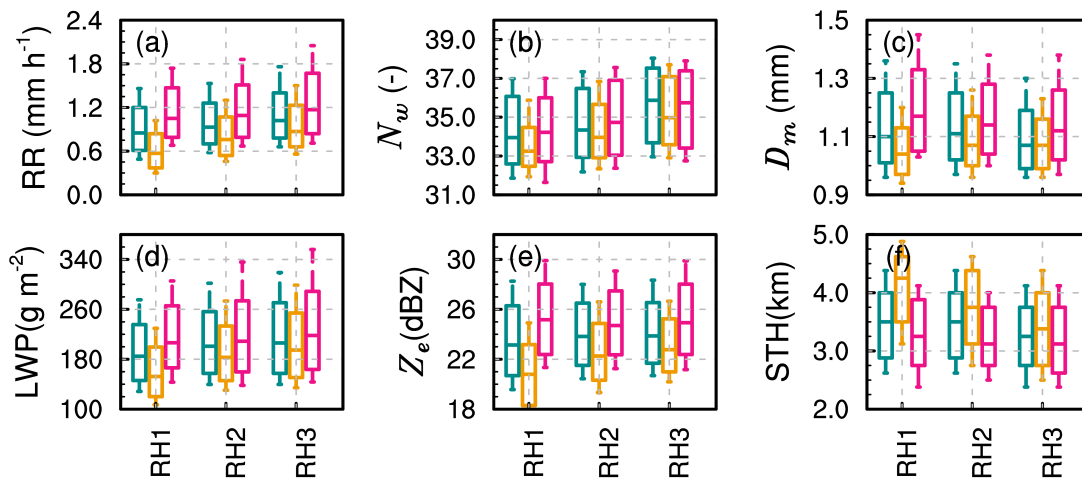
438 Similarly, the sensitivity of humidity to the impact of aerosol particles on shallow  
439 precipitation is examined by presenting the box plots of precipitation parameters, as  
440 illustrated in Figure 7. Regardless of 850hPa-RH, the vertical development of shallow  
441 precipitation clouds is hindered in coarse aerosol-polluted environments, with the  
442 median STH being the smallest. However, the near-surface RR is the highest,  
443 corresponding to the highest median  $Z_c$  at 2.5 km. On the contrary, in fine particle  
444 pollution environments, the vertical development of shallow precipitation clouds is  
445 enhanced (with the highest median STH), but the near-surface RR and  $Z_c$  are the  
446 weakest. This further confirms that the impact of coarse and fine aerosol particles on  
447 near-surface RR and LWP is independent of moisture and dynamic conditions.

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

458 Regarding STH, under low relative humidity and fine aerosol-polluted conditions,  
459 shallow precipitation clouds develop more deeply, with the 25th percentile of STH  
460 reaching 5 km, significantly higher than in clean and coarse aerosol-polluted  
461 environments. This may be because there is a reduction in the effective radius of cloud  
462 droplets in fine aerosol-polluted and low-humidity conditions. Smaller cloud droplets

463 are more prone to evaporation, resulting in a lower LWP, which does not favor an  
 464 increase in near-surface RR. This is also reflected in the near-surface DSD, which is  
 465 characterized by lower  $N_w$  and smaller  $D_m$ . However, although the humidity is relatively  
 466 low, the coarse particles, being more hygroscopic, can form larger cloud droplets,  
 467 reducing the loss of cloud water due to evaporation (resulting in a higher LWP), and  
 468 thereby enhancing surface RR. This is also reflected in the near-surface DSD, which is  
 469 characterized by a higher  $N_w$  and larger  $D_m$ . In high humidity environments, a high  
 470 concentration of fine particles can promote the formation of more cloud condensation  
 471 nuclei, which to some extent reduces the loss of cloud water due to the evaporation of  
 472 small particles. Therefore, the LWP in fine-particle pollution environments does not  
 473 differ much from that in coarse aerosol-polluted environments. This may also lead to  
 474 smaller differences in RR,  $Z_e$ , and other variables between coarse and fine aerosol-  
 475 polluted environments under relatively high humidity conditions.

476



477

478 **Figure 7** Same as Figure 6, but for RH at 850hPa. The blue, orange and pink colors  
 479 indicate the clean, fine-aerosol polluted and coarse-aerosol polluted environments,  
 480 respectively.

481

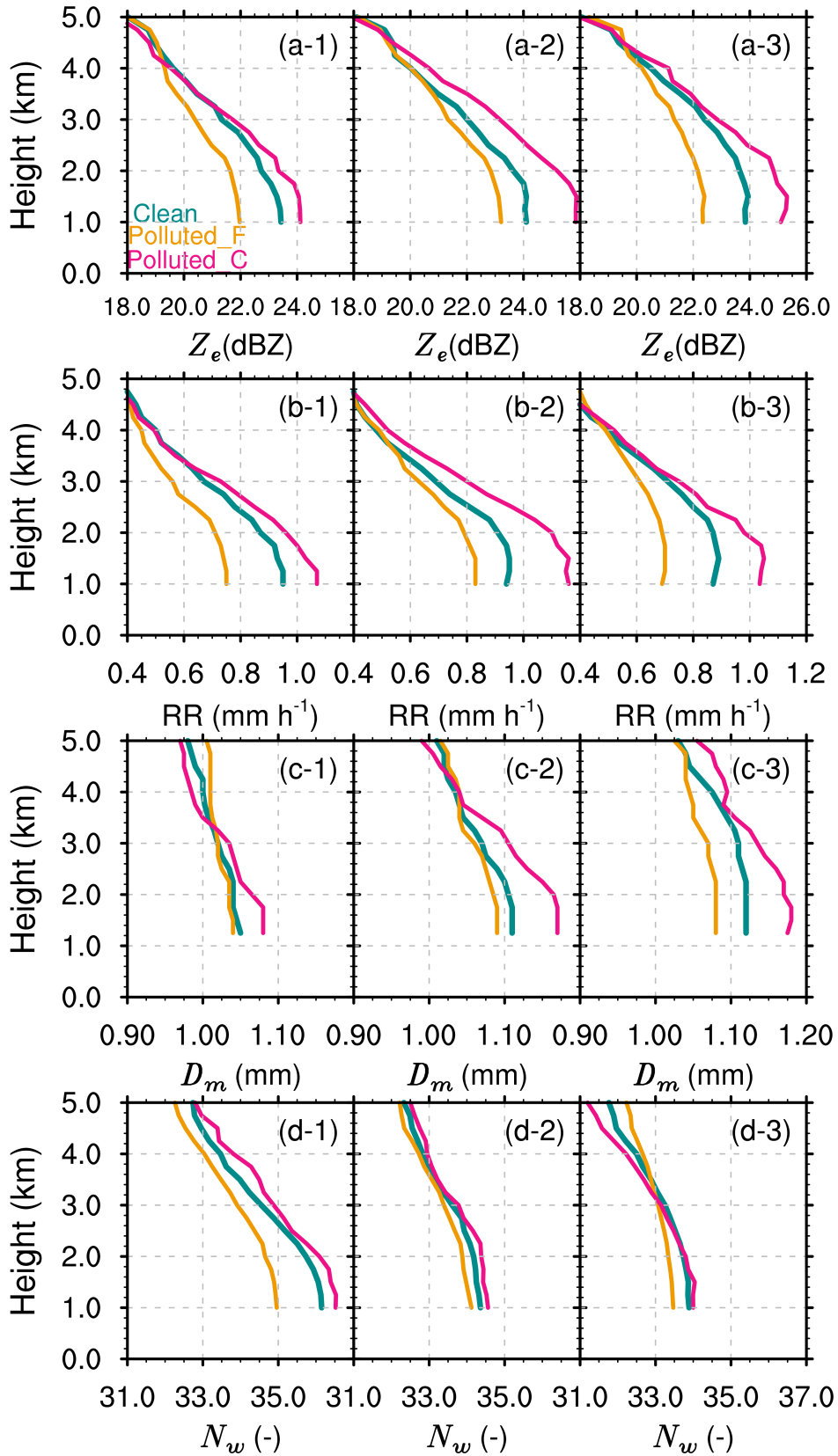


482 ***3.4 Sensitivities of aerosol impacts on microphysical structures and processes to***  
483 ***meteorological factors***

484 This part continues examining how coarse and fine aerosol modes affect  
485 precipitation structure and the microphysical processes in different environmental  
486 settings. As shown in Figure 8, under different CAPE and aerosol particle backgrounds,  
487 shallow precipitation profiles consistently exhibit increasing trends in  $Z_c$ , RR,  $N_w$ , and  
488  $D_m$  with decreasing altitude. Furthermore, irrespective of CAPE values, at a specified  
489 altitude, the parameters  $Z_c$  and RR are observed to be at their **maximum in environments**  
490 **polluted** with coarse aerosol particles, followed by those in a clean environment, and at  
491 their minimum in environments polluted with fine aerosol particles. This is consistent  
492 with the results in Figure 4. When compared between different CAPE conditions, the  
493  $Z_c$ , RR, and  $D_m$  of shallow precipitation in CAPE2 are the highest at different altitudes,  
494 while as the CAPE increases further (CAPE3), these values even decrease. Apart from  
495 instability, precipitation can be influenced by moisture, topography, and other factors;  
496 therefore, it is possible for an even lower RR in high CAPE conditions.

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

511 concentrations of **raindrops** (with large median  $D_m$ , as shown in Figure 8c-3).



512

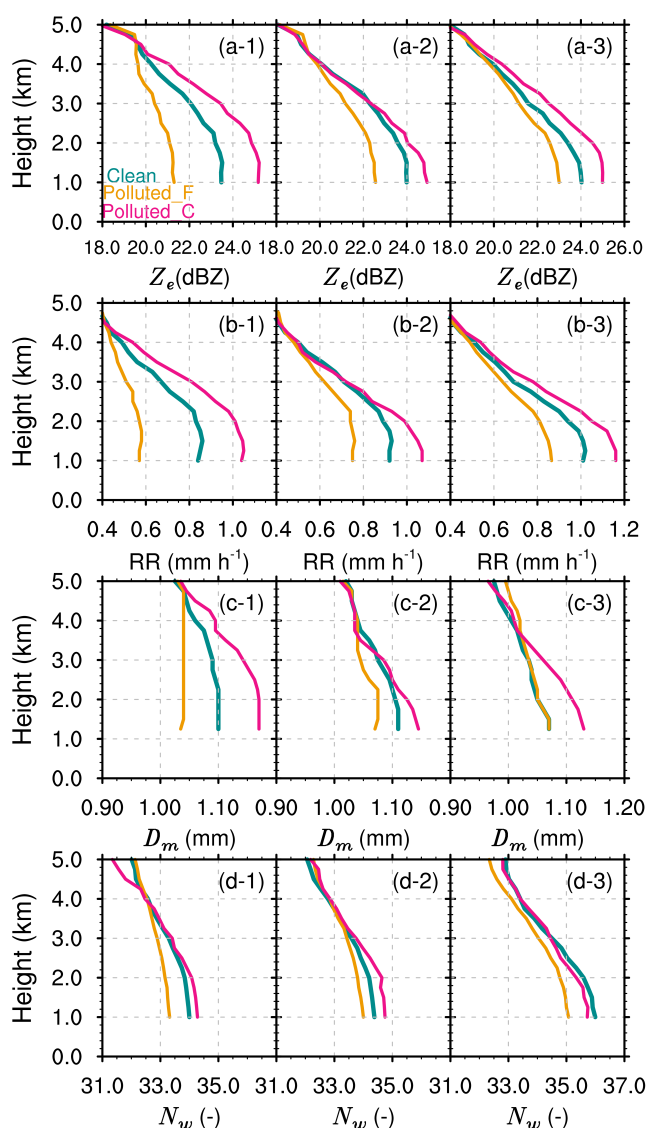
513 **Figure 8** The  $Z_e$  (a), rain rate (b),  $D_m$  (c), and  $N_w$  (d) profiles for shallow precipitation  
514 in different aerosol particle and CAPE conditions over southern China during the  
515 summers from 2014 to 2021. CAPE1, CAPE2, and CAPE3 are shown in the left, middle,  
516 and right panels, respectively.

517

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

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

546 fine aerosol particles mainly reduce RR by suppressing the concentration of raindrops,  
 547 while coarse aerosol particles increase RR by increasing the size of raindrops.  
 548 Furthermore, the differences in precipitation structures in aerosol-polluted coarse and  
 549 fine environments depend on humidity conditions, consistent with the conclusions in  
 550 Figure 7. The differences are the greatest under RH1 conditions, with the differences in  
 551 RR,  $Z_e$ ,  $D_m$ , and  $N_w$  at 1 km altitude being  $0.42 \text{ mm h}^{-1}$ , 4.5 dBZ, 0.19 mm, and about  
 552 1.3, respectively. Under RH3 conditions, the differences are smallest, with the  
 553 differences in the aforementioned variables being  $0.35 \text{ mm h}^{-1}$ , 2 dBZ, 0.05 mm, and  
 554 approximately 0.8, respectively.



555

556 **Figure 9** The  $Z_e$  (a), rain rate (b),  $D_m$  (c), and  $N_w$  (d) profiles for shallow precipitation  
 557 in different aerosol particle conditions and 850 hPa-RH over southern China during the

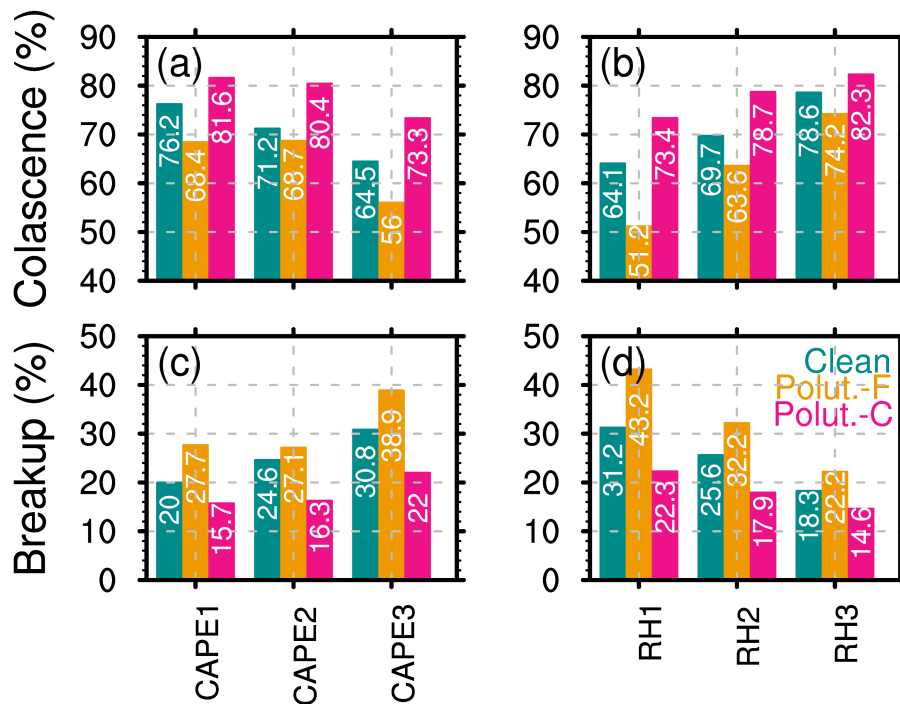
558 summers from 2014 to 2021. RH1, RH2, and RH3 are shown in left, middle, and right  
559 panels, respectively.

560

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

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

592 conditions,  $D_m$  in fine aerosol-polluted environments rapidly decreases with decreasing  
 593 altitude.  
 594



595  
 596 **Figure 10** The percentages of coalescence (a, b) and break-up (c,d) for shallow  
 597 precipitation raindrops under different aerosol particle, CAPE and 850 hPa-RH  
 598 conditions in southern China during the summers from 2014 to 2021.

599 **4 Conclusion and Discussion**

600 Using the combined data of DPR, MERRA-2 aerosol datasets, and ERA5 during  
 601 the summers of 2014-2021, this study investigates the potential impacts of coarse and  
 602 fine aerosol particles on the Rain Rate (RR), microphysical structure, and processes for  
 603 shallow precipitation in South China. Clean, coarse, and fine aerosol-polluted modes  
 604 are classified according to the AOD for total aerosol particles, coarse aerosol particles,  
 605 and fine aerosol particles derived from MERRA-2 products. ERA5 reanalysis data are  
 606 used to explore the sensitivity of aerosol impacts on shallow precipitation to dynamic  
 607 and moisture conditions in South China. The main findings are summarized as follows.  
 608 In comparison to clean environments, coarse aerosol-polluted environments enhance

609 near-surface rainfall rates of shallow precipitation, characterized by stronger near-  
610 surface RR (average precipitation intensity of  $1.78 \text{ mm h}^{-1}$ ), higher concentrations  
611 (average  $N_w = 36.98$ ) of raindrops at larger sizes (average  $D_m = 1.24 \text{ mm}$ ). This can be  
612 ascribed to the high presence of sea salt aerosol particles in South China, which tend to  
613 form larger cloud droplets through hygroscopic growth. This, in turn, enhances the  
614 condensation process, leading to the formation of additional rain droplets. As a result,  
615 it facilitates the coalescence and growth of raindrops, ultimately contributing to the  
616 formation of larger raindrops. On the contrary, fine aerosol particles suppress near-  
617 surface RR, with an average near-surface RR of only  $1.33 \text{ mm h}^{-1}$  and lower  
618 concentrations and smaller sizes of raindrops (average  $N_w = 36.37$ , average  $D_m = 1.14$   
619 mm). Liu et al. (2022) noted similar opposing effects of fine aerosol particles and  
620 coarse sea spray on warm marine clouds. Deep clouds show increased rainfall with high  
621 liquid water content but reduced rainfall if water content is low (Li et al., 2011). This  
622 underscores the distinct behavior of shallow precipitation and the varied impacts of  
623 aerosol types on it. However, fine aerosol-polluted environments promote vertical  
624 development of shallow precipitation clouds (median STH of 3.7 km), approximately  
625 0.5 km higher than in coarse aerosol-polluted conditions. The inhibition of the vertical  
626 development of precipitation clouds by coarse aerosol particles explains their  
627 suppressive effect on lightning activity to some extent (Pan et al., 2022).

628 From the perspective of precipitation vertical structure and microphysical  
629 processes, shallow precipitation is dominated by warm-rain collision-coalescence  
630 processes under different aerosol particle backgrounds, with the collision-coalescence  
631 process accounting for over 62%. However, there are significant differences in the  
632 efficiency of raindrop collision-coalescence growth under different aerosol particle  
633 conditions. In contrast to clean conditions, the median values of  $Z_c$ , RR,  $D_m$ , and  $N_w$  are  
634 highest in presence of aerosol coarse mode and lowest in conditions for fine aerosol  
635 mode at all altitude levels. Looking at it from a microphysical standpoint, the increase  
636 in  $D_m$  with decreasing altitude is most pronounced under coarse aerosol-polluted  
637 conditions, reflecting more significant collision-coalescence growth processes,  
638 accounting for 74.1%. In contrast, the increase in  $D_m$  with decreasing altitude is weakest

639 under fine aerosol-polluted conditions, due to the higher proportion of breakup  
640 processes (accounting for 33.1%) and a decrease of approximately 12% in the collision-  
641 coalescence process (accounting for 62.4%). Overall, the promotion of RR is associated  
642 with more significant collision-coalescence processes by coarse [aerosol particles](#), while  
643 the suppression of RR is characterized by more significant breakup processes with fine  
644 [aerosol particles](#).

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



669 including instability, moisture, and temperature. Additionally, other factors such as  
670 wind vectors and pressure may also affect the impact of [aerosol particles](#) on  
671 precipitation, which is worthy of further study.

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

691

## 692 **Data availability**

693 The GPM DPR data provided by NASA Goddard Space Flight Center's Mesoscale  
694 Atmospheric Processes Laboratory and Precipitation Processing System (PPS) can be  
695 downloaded from <https://pmm.nasa.gov/dataaccess/downloads/gpm>. MERRA-2 data  
696 can be downloaded from <https://gmao.gsfc.nasa.gov/reanalysis/MERRA->

697 [2/data\\_access/](#). The ERA5 data can be downloaded from  
698 <https://www.ecmef.int/en/forecasts/dataset/ecmwf-reanalysis-v5>. The ancillary digital  
699 terrain data is from the National Geophysical Data Center (NGDC) (available online at  
700 <http://www.ngdc.noaa.gov>, accessed in May 2023).

701

## 702 **Author contributions**

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

## 706 **Declaration of competing interest**

707 The authors declare no competing interests.

708

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711 Processes Laboratory and PPS, NGDC, and ECMWF for providing the analysis data.

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717 FEMD (KLME202303), China Meteorological Administration "Application of  
718 quantum technology in meteorological detection" Youth Innovation Team Project  
719 (No.CMA2024QN11), and the Open Project of State Key Laboratory of Severe  
720 Weather (2024LASW-B11).

721

722

## Reference

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