The evolution of cloud and aerosol microphysics at the summit of Mt. Tai, China

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Abstract. The influence of aerosols, both natural and anthropogenic, remains a major area of uncertainty when predicting the properties and the behaviours of clouds and their influence on climate. In an attempt to better understand the microphysical properties of cloud droplets, the simultaneous variations in aerosol microphysics and their potential interactions during cloud life cycles in the North China Plain, an intensive observation took place from 17 June to 30 July 2018 at the summit of Mt. Tai. Cloud microphysical parameters were monitored simultaneously with number concentrations of cloud condensation nuclei (NCCN) at different supersaturations, PM2.5 mass concentrations, particle size distributions and meteorological parameters. Number concentrations of cloud droplets (NC), liquid water content (LWC) and effective radius of cloud droplets (reff) show large variations among 40 cloud events observed during the campaign. The low values of reff and LWC observed at Mt. Tai are comparable with urban fog. Clouds on clean days are more susceptibility to the change in concentrations of particle number (Np), while clouds formed on polluted days might be more sensitive to meteorological parameters, such as updraft velocity and cloud base height. Through studying the size distributions of aerosol particles and cloud droplets, we find that particles larger than 150 nm play important roles in forming cloud droplets with the size of 5–10 µm. In general, LWC consistently varies with reff. As NC increases, reff changes from a trimodal distribution to a unimodal distribution and shifts to smaller size mode. By assuming a constant cloud thickness and ignoring any lifetime effects, increase in NC and decrease in reff would increase cloud albedo, which may induce a cooling effect on the local climate system. Our results contribute valuable information to enhance the understanding of cloud and aerosol properties, along with their potential interactions on the North China plain.
1 Introduction

Clouds are key factors in the atmospheric hydrological cycle, which play an important role in the atmospheric energy budget and significantly influence the global and regional climate (Chang et al., 2019b; Y. Zhang et al., 2004). Clouds can be physically described by their liquid water content (LWC), number concentrations of droplets (NC) and effective radius of droplets (reff). These parameters may show small interannual variations for the same monitoring station (Möller et al., 1996), but they vary over a large range for different cloud types (Quante, 2004), for different cloud altitudes (Padmakumari et al., 2017; Zhao et al., 2018) and in different parts of a cloud (Deng et al., 2009).

The interactions between the clouds and the aerosols are complex. Clouds can efficiently remove aerosols by activating cloud condensation nuclei (CCN) to form cloud droplets (Croft et al., 2010; L. M. Zhang et al., 2004). The cloud processes can incorporate large amount of fine particulate mass (Heintzenberg et al., 1989), change their size distributions (Drewnick et al., 2007; Schroder et al., 2015), and alter the CCN compositions through homogeneous and heterogeneous reactions (Roth et al., 2016). In addition, the variation of aerosol number concentrations and aerosol size distributions could alter the cloud microphysics. Through studying microphysical characteristics of cloud droplet residuals at Mt. Åreskutan, Noone et al. (1990) found that larger cloud microphysical properties and their potential links to aerosol particles makes it difficult to evaluate the cloud microphysics in small-scale regions (Fan et al., 2016; Khain et al., 2015; Sant et al., 2013). Discrepancy still exists between the widths of observed and simulated size distributions of cloud droplets (Grabowski and Wang, 2013). In addition to this, incomplete knowledge of the impact of cloud–aerosol interactions (Rosenfeld et al., 2014b), unresolved processes of cloud formation (Stevens and Bony, 2013) and the lack of research into the variation of cloud microphysical parameters at different cloud stages still hinder modelling studies.

In this study, in situ observations at the summit of Mt. Tai were presented to investigate the evolution of cloud microphysics coupled to simultaneous monitoring of aerosol size distributions, PM2.5 mass and CCN concentrations within non-precipitating clouds. The summit of Mt. Tai is the highest point in the centre of the North China Plain (NCP). Sufficient moisture in summer and dramatic temperature differences between day and night make it ideal for in situ orographic cloud monitoring (J. Li et al., 2017). The summit of Mt. Tai is far away from anthropogenic emission sources on the ground. However, high concentrations of inorganic ions in PM2.5 (Zhou et al., 2009), abundant bacterial communities (Zhu et al., 2018), and NH3 and NOx emissions from biomass burning (Chang et al., 2019a) have been observed at the summit, and thus a strong anthropogenic influence must exist. Previous studies of cloud samples collected at the same position showed high inorganic ion concentrations (J. Li et al., 2017; Wang et al., 2011), which can be attributed to the increase in anthropogenic aerosol. In this study, two typical cloud processes are discussed in detail to elucidate the relationship of NC, reff, and LWC under clean or polluted conditions (indicated by NP and NCCN) and during the cloud life cycle. This paper provides comprehensive information about cloud microphysical properties and their potential links to aerosol concentrations and size distribution. Implications of cloud and aerosol microphysics for cloud albedo and climate are discussed.
2 Experiments

2.1 Observation duration and site

From 17 June to 30 July 2018, 40 cloud events in total were monitored at the Shandong Taishan Meteorological Station at summit of Mt. Tai (Tai’an, China; 36°18’ N, 117°13’ E; 1545 m a.s.l.; Fig. S1). Mt. Tai is the highest point in the central of North China Plain (NCP) and located within the transportation channel between the NCP and the Yangtze River Delta (Shen et al., 2019). The altitude of Mt. Tai is close to 1.6 km. This height is close to the top of the planetary boundary layer in central eastern China and is suitable for studying the characteristics of particles inputting to clouds (Hudson, 2007). Orographic clouds, which are mainly formed in the boundary layer via air approaching the ridge, being forced to rise up and then being cooled by adiabatic expansion (Choularton et al., 1997), frequently occur at the summit of Mt. Tai, especially in summer. Previous studies concentrated on cloud chemistry found that Mt. Tai is significantly influenced by anthropogenic emissions (J. Li et al., 2017; Wang et al., 2011). In addition, fixed observation location is mainly applied to study the evolution of aerosol properties and cloud processing (Mertes et al., 2005; Roth et al., 2016). Thus, Mt. Tai is a good site for monitoring orographic clouds and simultaneously investigating aerosol and cloud microphysics. The arrangement of instruments is presented in Fig. S1c. As shown in Fig. S2, the prevailing wind direction during this summer campaign was easterly wind (23.3%), southwesterly wind (22.8%) and southerly wind (21.9%), respectively. About 85.6% of wind speed was less than 8 m s\(^{-1}\). The monitored cloud events in this study were mainly influenced by southerly wind (34.7%) and southwesterly wind (22%).

2.2 Cloud microphysical parameters

A Fog Monitor (Model FM-120, Droplet Measurement Technologies Inc., USA), a forward-scattering optical spectrometer with a sampling flow of 1 m\(^3\) min\(^{-1}\), was applied in situ for real-time size distributions of cloud droplets and computing \(N_C\), LWC, median volume diameter (MVD), and effective diameter (ED) in the size range of 2 to 50 \(\mu\)m (Spiegel et al., 2012). The corresponding equations are

\[
N_C = \Sigma N_i,
\]

\[
\text{LWC} = \frac{4\pi}{3} \Sigma N_i r_i^3 \rho_w,
\]

\[
\text{MVD} = 2 \times \left( \frac{\Sigma N_i r_i^3}{\Sigma N_i} \right)^{\frac{1}{3}},
\]

\[
\text{ED} = 2 \times r_{\text{eff}} = 2 \times \Sigma n_i r_i^3 / \Sigma n_i r_i^2,
\]

where \(N_i\) is the cloud number concentration at the \(i\)th bin, \(r_i\) represents the radius at the \(i\)th bin and \(\rho_w = 1 \text{ g cm}^{-3}\) stands for the density of liquid water. Droplets are categorised into the manufacturer’s 30 predefined size bins with a sampling resolution of 1 s. The size bin widths using this configuration were 1 \(\mu\)m for droplets < 15 and 2 \(\mu\)m for droplets >15 \(\mu\)m. The true air speed calibration and size distribution calibration of FM-120 were carried out by the manufacturer using borosilicate glass microspheres of various sizes (5.0, 8.0, 15.0, 30.0, 40.0 and 50.0 \(\mu\)m, Duke Scientific Corporation, USA). The difference in optical properties between the glass beads and water was taken into account during the calibration process. In this study, the sampling inlet nozzle faced the main wind direction and was horizontally set. Cloud events are defined by the universally accepted threshold values in \(N_C\) and LWC, i.e. \(N_C > 10 \text{ cm}^{-3}\) and LWC > 0.001 g m\(^{-3}\) (Demoz et al., 1996). Too short cloud events with a duration < 15 min were excluded.

2.3 Aerosol size distribution

A Scanning Mobility Particle Sizer (SMPS, Model 3938, TSI Inc., USA) consisting of a Differential Mobility Analyser (DMA, Model 3082, TSI Inc., USA) and a Condensation Particle Counter (CPC, Model 3775, TSI Inc., USA) was applied to monitor the size distributions of dehumidified aerosols through a PM\(_{10}\) inlet. The neutralised aerosols were classified by DMA to generate a monodisperse stream of known size according to their electrical mobility. The CPC placed downstream counts the particles and gives the number of particles with different sizes. In this study, each scan was fixed at 5 min for every loop with a flow rate of 1.5 L min\(^{-1}\) sizing particles in the range of 13.6–763.5 nm in 110 size bins.

2.4 CCN number concentration

The \(N_{\text{CCN}}\) at certain supersaturations (SSs) were quantified by a Cloud Condensation Nuclei Counter (Model CCN-100, DMT Inc., USA). The CCN counter was set at five SS values sequentially for 10 min each at 0.2 %, 0.4 %, 0.6 %, 0.8 % and 1.0 % with a full scan time resolution of 50 min. Data collected during the first 5 min of each SS were excluded since the CCN counter needs time for temperature stabilisation after the change in SS. The ratio of sample flow to sheath flow was set at 1 : 10 with a total airflow of 500 ccm. The SSs of the CCN counter were calibrated before the campaign and checked at the end of the campaign with monodisperse ammonium sulfate particles of different sizes (Rose et al., 2008).

2.5 PM\(_{2.5}\) concentrations and meteorological parameters

The PM\(_{2.5}\) mass concentration was measured using a beta attenuation and optical analyser (SHARP monitor, model 5030i, Thermo Scientific Inc., USA). Meteorological parameters, including the ambient temperature (\(T_a\ °\)), relative humidity (RH), wind speed (WS, m s\(^{-1}\)) and wind direction
thickness, droplet size distribution (Stephens, 1978), the cloud optical and the cloud is vertically uniform with respect to below (Seinfeld and Pandis, 2006). Assuming the cloud Cloud albedos can be calculated using the equations shown below (J. Li et al., 2017), the large LWC of clouds in CE-20. In the following sections, the evolution of cloud and aerosol microphysical properties are presented. The influence of meteorological parameters

\[
\text{Albedo} = \frac{\sqrt{3}(1 - g)\tau_c}{2 + \sqrt{3}(1 - g)\tau_c},
\]

where \( g \) is the asymmetry factor. The radius of cloud droplets was much greater than the wavelength of visible light, and hence \( g = 0.85 \). The equation before becomes

\[
\text{Albedo} = \frac{\tau_c}{\tau_c + 7.7}.
\]

3 Results and discussion

3.1 Overview of the cloud microphysics

From 17 June to 30 July 2018, 40 cloud events were captured at the summit of Mt. Tai. Large ranges of cloud microphysics were observed during the campaign. The averaged \( N_C \), LWC and \( r_{	ext{eff}} \) of the 40 cloud events at the summit of Mt. Tai varied over the ranges of 59–1519 cm\(^{-3}\), 0.01–0.59 g m\(^{-3}\) and 2.6–7.4 \( \mu \)m, respectively (Table S1). The monitored number concentration of cloud droplets at Mt. Tai both in this study and in 2014 can reach 2000–3000 cm\(^{-3}\) (J. Li et al., 2017), which is much higher than those values (with a range of 10–700 cm\(^{-3}\)) for city fog and convective and orographic clouds (Allan et al., 2008; Li et al., 2011; Padmakumari et al., 2017) (Table 1). This suggests that clouds at Mt. Tai were characterised with high \( N_C \).

The microphysics of different clouds and fog could generally be distinguished in a plot of \( r_{	ext{eff}} \) (or MVD) against LWC. As illustrated in Fig. 1, the LWC generally increased on the order of city fog, orographic clouds, and convective clouds, and Mt. Tai generally follows this rule. It was consistent with the study by Penner et al. (2004) that LWC within clouds increases linearly with altitude. For LWC values of clouds at Mt. Tai, both the high values, which were comparable with convective clouds, and the low values, which were similar to city fog (Fig. 1), were monitored. It indicated that clouds at Mt. Tai appeared to show a larger range of LWC values. The increase in LWC at Mt. Tai was determined by the increase in \( r_{	ext{eff}} \), and/or \( N_C \). However, sometimes only one factor played the dominant role. As illustrated in Table S1, \( N_C \), \( r_{	ext{eff}} \) and LWC in cloud event 20 (CE-20) were 1519 cm\(^{-3}\), 5.2 \( \mu \)m and 0.54 g m\(^{-3}\), respectively, while the corresponding values in cloud event 16 (CE-16) were 59 cm\(^{-3}\), 9.8 \( \mu \)m and 0.14 g m\(^{-3}\), respectively. Even though \( r_{	ext{eff}} \) of CE-20 was smaller when compared to CE-16, the higher \( N_C \) determined the larger LWC of clouds in CE-20. In the following sections, the evolution of cloud and aerosol microphysical properties are presented. The influence of meteorological parameters

\[
\tau_c = h \left( \frac{g r_{	ext{eff}} LWC^2 N_C}{2 \rho_w} \right)^{\frac{1}{3}},
\]

where \( h \) is the thickness of the cloud and \( \rho_w \) is the density of cloud water.

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Table 1. Comparison of clouds monitored at Mt. Tai with city fog, convective clouds monitored by research aircrafts and other orographic clouds. Including sampling information (site, period and altitude); the range of PM$_{2.5}$ mass concentrations; the range of microphysical parameters, i.e. number concentrations of cloud droplets, $N_C$; liquid water content, LWC; median volume diameter, MVD; and effective radius, $r_{\text{eff}}$; and the number of monitored clouds, cloud events, and fog events.

| Sampling site                  | Period                        | Altitude (m a.s.l) | PM$_{2.5}$ (µg m$^{-3}$) | $N_C$ (cm$^{-3}$) | LWC (g m$^{-3}$) | MVD (µm) | $r_{\text{eff}}$ (µm) | Number of clouds, cloud events, and fog events | Reference         |
|--------------------------------|-------------------------------|--------------------|--------------------------|------------------|------------------|----------|-----------------------|------------------------------------------------|------------------|
| City fog                       |                               |                    |                          |                  |                  |          |                       |                                                 |                  |
| Shanghai, China                | November 2009                 | 7                  | –                        | 11–565           | 0.01–0.14        | 5.0–20.0 | –                     | 1 Li et al. (2011)                               |                  |
| Nanjing, China                 | December 2006–December 2007   | 22                 | 0.03$^a$–0.60$^a$        | –                | 2.69 × 10$^{-3}$–0.16 | –       | 1.6$^b$–2.7$^b$       | 7 Lu et al. (2010)                               |                  |
| Convective clouds              |                               |                    |                          |                  |                  |          |                       |                                                 |                  |
| Amazon Basin and Cerrado regions, Brazil | August–September 1995         | 90–4000            | –                        | –                | 0.1d–2.10d       | –        | 2.8d–9.2d             | > 1000 Reid et al. (1999)                        |                  |
| Hyderabad, Bay of Bengal, India | 29 October 2010               | 1300–6300          | 10$^d$–380              | 0$^d$–1.80       | 3.8d–17.0        |          |                       | 1 Padmakumari et al. (2017)                     |                  |
| Orographic clouds              |                               |                    |                          |                  |                  |          |                       |                                                 |                  |
| Mt. Schmücke, Germany          | September–October 2010        | 937                | –                        | 193–519          | 0.14–0.37        | 5.7–8.7  |                       | 8 van Pinxteren et al. (2016)                   |                  |
| East Peak, Puerto Rico         | December 2004                 | 1040               | –                        | 1040             | 0.24–0.31        | 14.0–20.0 | –                     | 2 Allan et al. (2008)                            |                  |
| Mt. Tai, China                 | July–August 2014              | 1545               | 11.1–173.3              | 4–2186           | 0.01–1.52        | 1.6–43.0 | 0.8–18.9              | 24 Unpublished data from J. Li et al. (2017)    |                  |
| Mt. Tai, China                 | June–July 2018                | 1545               | 1.2–127.1               | 10–3163          | 1.01 × 10$^{-3}$–1.47 | 4.4–25.0 | 2.4–13.4              | 40 This study                                  |                  |
| Mt. Tai, China (CP-1$^c$)      | 10–13 July 2018               | 1545               | 1.3–40.7                | 11–2470          | 1.12 × 10$^{-3}$–1.47 | 4.6–17.4 | 2.5–10.7              | 12 This study                                  |                  |
| Mt. Tai, China (CP-2$^c$)      | 13–20 July 2018               | 1545               | 1.2–66.2                | 10–3163          | 1.03 × 10$^{-3}$–1.10 | 4.6–13.5 | 2.4–7.9               | 12 This study                                  |                  |

$^a$ Represents the mass concentrations of PM$_{10}$. $^b$ Represents the range of averaged radius. $^c$ Two cloud processes which are detailedly discussed in this study. $^d$ Values were read from the graphs.
3.2 Analysis of typical cloud processes

By assuming a density of particles $\rho = 1.58$ g cm$^{-3}$ (Cross et al., 2007), the mass concentrations of particles, which were calculated from the aerosol number size distribution measured by SMPS and named PM$_{0.8}$, were highly consistent with PM$_{2.5}$, especially when PM$_{2.5}$ was less than 20 µg m$^{-3}$ (Fig. 2c). Based on the mass concentration (PM$_{2.5}$) and the number concentration (N$_P$, which represented the total number concentration of aerosol particles measured by SMPS) of aerosols, two typical cloud processes, “cloud process-1” (CP-1) and “cloud process-2” (CP-2), were selected and analysed with respect to their special characteristics. The variations in updraft velocity $v_{up}$ on cloud microphysics during CP-1 and CP-2 were ignored for simplicity (Table S2, Figs. S3 and S4). The sampling angle ($\theta_s$) and $v_{up}$ for CP-1 and CP-2 were 11.9° and 0.82 ± 0.29 m s$^{-1}$ and 10.6° and 0.92 ± 0.36 m s$^{-1}$, respectively (Table S2). According to the calculations provided by Spiegel et al. (2012), the aspiration efficiency and transmission efficiency of the fog monitor were all close to 1. In CP-1 (which only contained cloud event 19, CE-19), cloud droplets formed under a relatively stable (wind speed < 4 m s$^{-1}$) and clean (PM$_{2.5}$ ≈ 10.9 µg m$^{-3}$ and N$_P$ ≈ 1425 cm$^{-3}$) conditions accompanied by a slow increase in $T_a$ (Figs. 2 and 3). During daytime, especially in the afternoon, the PM$_{2.5}$ mass concentration dramatically increased with few changes in wind speed and wind direction, meanwhile, N$_P$ reached to about 5000 cm$^{-3}$ (Fig. 3). CP-1 persisted for 74 h, making it the longest cloud event during the presented campaign. Quite different from CP-1, CP-2 contained eight cloud events (CE-20 to cloud event 26, CE-26; see Fig. 3) and occurred periodically under high PM$_{2.5}$ (Fig. 2, 50.7 µg m$^{-3}$ on average) and high N$_P$ (Fig. 3, 1694 cm$^{-3}$ on average) conditions. Cloud events in CP-2 formed after sunset, with a sharp decrease in PM$_{2.5}$ and N$_P$, and transitorily dissipated at noon accompanied with the increase in PM$_{2.5}$, N$_P$, $T_a$ and cloud base height (CBH).

For cloud water samples collected during CP-1 and CP-2, the percentage of chemical compositions did not change a lot (Fig. S5). The total measured dominant ions (sulfate, nitrate and ammonia) were 93.39% in CP-1 and 90.37% in CP-2. The high concentration of secondary ions in the cloud water samples indicated that clouds at Mt. Tai were dramatically influenced by anthropogenic emissions.

CP-1 was separated into four stages, SL1 (stage: low 1), SH1 (stage: high 1), SL2 (stage: low 2) and SH2 (stage: high 2), based on the aerosol concentrations (Fig. 3). The characteristics of SL1 and SL2 were low N$_C$ (383 and 347 cm$^{-3}$, respectively), large $r_{eff}$ (7.26 and 6.36 µm, respectively) and high LWC / N$_C$ (which represents the average water each cloud droplet contained: 1.01 and 0.75 ng, respectively). During SH1 and SH2, a dramatic increase in N$_C$ (to 949 and 847 cm$^{-3}$, respectively), decrease in $r_{eff}$ (to 4.90 and 4.88 µm, respectively) and decrease in LWC / N$_C$ (to 0.35 and 0.36 ng, respectively) were found with the increase in N$_P$ (to 4196 and 4665 cm$^{-3}$, respectively).

Each cloud event of CP-2 was separated into an activation stage (S1), collision–coalescence stage (S2), stable stage (S3) and dissipation stage (S4) according to the regular changes in N$_C$ and LWC / N$_C$ (Fig. 3a). In S1, N$_C$ dramatically increased to its maximum value among the cloud events. In S2, N$_C$ declined sharply to a stable value, meanwhile LWC / N$_C$ reached the maximum value. In S3, N$_C$ was stable or slightly varied and LWC / N$_C$ started to decrease. In
S4, both $N_C$ and $LWC / N_C$ decreased sharply and finally arrived at zero. Even though the two stages (S2 and S3) in cloud event 25 (CE-25) did not totally follow the division rules, the other six cloud events followed them closely. This indicated that the division was helpful for the study of the variations of cloud microphysical properties during CP-2. The newly formed cloud droplets during S1 were characterised by small size, high $N_C$ and low $LWC / N_C$ values (Figs. 2f and 3b). For example, about $2310 \text{ cm}^{-3}$ of cloud droplets can quickly form in the first 2 h of CE-20. The $r_{\text{eff}}$ of these droplets was smaller than 4.1 $\mu$m, and $LWC / N_C$ was about 0.2 ng. In going from S2 to S3, the strong collision–coalescence between cloud droplets caused the increase in both $r_{\text{eff}}$ and $LWC / N_C$. In S4, the increase in PM$_{2.5}$, through evaporation of cloud droplets or lifting of CBH (Fig. 2), was observed to coincide with the vanishing of cloud events (Mazoyer et al., 2019; J. Li et al., 2017).

3.3 Relationships among $N_P$, $N_{CCN}$ and $N_C$

In this study, both consistent variation and inverse variation between $N_P$ and $N_C$ were observed. $N_P$ and $N_C$ showed consistent variation in CP-1. However, in CP-2, an obviously inverse variation was found between $N_P$ and $N_C$ in S1 and S4, while a consistent variation was found between $N_P$ and $N_C$ in S2 and S3 (Figs. 3a, 4b, and c). Some in situ observations (Lu et al., 2007; Mazoyer et al., 2019) and modelling studies (Heikenfeld et al., 2019; Zhang et al., 2014) supported the viewpoint that the increase in $N_P$ brings more CCN and further increases $N_C$, which could cause the consistent variation between $N_P$ and $N_C$. In contrast, some recent studies of fog also suggested that the increase in $N_P$ could decrease the ambient supersaturation and then decrease $N_C$ (Boutle et al., 2018; Mazoyer et al., 2019). Besides, Modini et al. (2015) found inverse variation between $N_C$ and the number of particles with diameters larger than 100 nm due to the reduction of supersaturation by coarse primary marine aerosol parti-
13742 J. Li et al.: The evolution of cloud and aerosol microphysics

Figure 3. Variation of (a) $N_C$, $N_p$, and $N_{CCN,0.2}$ and (b) $N_{CCN,0.2}/N_p$ and LWC$/N_C$ during CP-1 and CP-2.

In general, the covariation between $N_p$ and $N_C$ could be affected by many factors, including competition of water vapour between aerosol particles and/or cloud droplets, the scavenging of particles by cloud droplets, and new particle formation through cloud processes. In this study, consistent variation between $N_p$ and $N_C$ was characterised with higher LWC$/N_C$, while inverse variation between $N_p$ and $N_C$ appeared with lower LWC$/N_C$. The average LWC$/N_C$ was 0.61 ng in CP-1 and was 0.15, 0.42, 0.39 and 0.16 ng in S1, S2, S3 and S4, respectively, in CP-2. The relatively higher LWC$/N_C$ values could indicate conditions where the amount of water vapour was not limiting cloud droplet formation. Once $N_p$ increased, part of the cloud water was taken away by the CCN in the particles to form new droplets, and the remaining amount of water was still sufficient to maintain the previous droplets in liquid state. Thus, both $N_p$ and $N_C$ simultaneously increased. On the other hand, relatively low LWC$/N_C$ values could, to some extent, limit the formation of new cloud droplets. The activated particles that grew at the beginning of the cloud cycle would lower the surrounding supersaturation and to some extent limit further aerosol activation (Ekman et al., 2011). The part of the water taken by the CCN in the particles was not enough to change all of them into new droplets, and the remaining amount of water was also insufficient to maintain all of the previous droplets in a liquid state. Following this, the $N_C$ would decrease and the higher the $N_p$ becomes, the sharper decrease the $N_C$ will be. Thus, the inverse variation between $N_p$ and $N_C$ was observed.

The ratio between $N_{CCN}$ and $N_p$ reflects the activation ratio of aerosol particles. As shown in Fig. S6, $N_{CCN}$ increased with the increase in SS. In addition, $N_{CCN}$ of CP-2 was higher than that of CP-1 at the same SS. In order to compare with previous studies as discussed below, SS = 0.2 % was chosen to calculate $N_{CCN}/N_p$, which represented the activation ratio of aerosol particles. As shown in Fig. 3b, $N_{CCN,0.2}/N_p$ (activation ratio at a certain SS = 0.2 %) ranged from 0.06 to 0.69 in CP-1, yet it ranged from 0.22 to 0.66 in CP-2. The average value of 0.30 in CP-1 was smaller than that of 0.38 in CP-2, and values lower than 0.22 did not appear during CP-2. This indicated that the activation of aerosol particles in CP-2 was relatively easy. Both the size distribution and the chemical composition could impact the cloud-nucleating ability of aerosol particles (Dusek et al., 2006; Mazoyer et al., 2019). Figure S7 shows the relation between $N_{CCN,0.2}/N_p$ and GMrp during CP-1 and CP-2. As can be seen in Fig. S7, higher correlation of $N_{CCN,0.2}/N_p$ with GMrp was found during CP-1 than during CP-2, which suggested that the physical properties might have more influence on the activation of aerosols during CP-1. Aside from this, Asmi et al. (2012) found that...
higher $N_{CCN} / N_p$ and a more concentrated plot of $N_{CCN}$ versus $N_p$ usually occurred during winter when a higher fraction of aged organics was observed during the observation programme at Puy-de-Dôme, France. In this study, the plot of $N_{CCN,0.2}$ versus $N_p$ was found to be more scattered in CP-1 than in CP-2 (Fig. S8). Even though the chosen SS in this study (SS = 0.2%) is different from that at Puy-de-Dôme (SS = 0.24%), most of the data points of CP-1 and CP-2 were distributed between the two recommended dashed lines (the visually defined boundaries within most of the data are centred, Fig. S8) by Asmi et al. (2012). This suggested that the difference in aerosol organic chemical compositions during CP-1 and CP-2 might also explain the different activation ratio of aerosol particles during these two cloud processes.

3.4 The influence of $N_p$, CBH and $v_{up}$ on cloud microphysics

No negative AIE$_r$ or AIE$_N$ were found in this study (Fig. 4). The positive AIE$_r$ and AIE$_N$ at Mt. Tai mean that the increase in $N_p$ is accompanied by decreased $r_{eff}$ and increased $N_C$. However, in the studies of Yuan et al. (2008) and Tang et al. (2014), AOD was applied to represent aerosol loading and negative AIE$_r$ (indicating $r_{eff}$ increased with the increasing of AOD) near the coastlines of the Gulf of Mexico, the South China Sea and over eastern China. The reason proposed by Yuan et al. (2008) was the increasing slightly soluble organic (SSO) particles, which would increase the critical supersaturation and hinder the activation of the particles. Meanwhile, Tang et al. (2014) showed that the meteorological conditions, which favoured the transportation of pollutants and water vapour, led to simultaneous increases in both AOD and $r_{eff}$. Different from the coastal area, the summit of Mt. Tai is relatively far from the sea (around 230 km from the Bohai Sea and Yellow Sea) (Guo et al., 2012), leading to less moisture in the air. This might hinder the growth of cloud droplets and cause the positive AIE$_r$. In addition, the increase in LWC was found to covary with the decrease in AIE, especially at coastal sites (McComiskey et al., 2009; Zhao et al., 2012). However, non-obvious variation was found between AIE$_r$ and LWC at Mt. Tai (Fig. 4a). It might due to the high aerosol loading during cloud processes (Zhao et al., 2012).

Although all positive AIE$_r$ and AIE$_N$ were found in CP-1 and CP-2, the specific values were different. According to the studies of AIE$_r$ and AIE$_N$ of CP-1 and CP-2, our results suggested that cloud droplet numbers were more sensitive to $N_p$ under conditions with lower aerosol concentrations. The calculation of AIE$_r$ was shown in Fig. S9 and summarised in Fig. 4. As shown in Fig. 4a, except for the out-of-bounds AIE$_r$ values calculated with insufficient data points when LWC was larger than 0.7 g m$^{-3}$, AIE$_r$ of 0.181–0.269 for CP-1 was always higher than that of 0.025–0.123 for CP-2 in corresponding narrow LWC ranges. When calculating AIE$_N$, the number of cloud droplets may be underestimated during the activation and dissipation stages (Mazoyer et al., 2019), due to the limitations of the fog monitor, which could be a cause of the low $R^2$ of AIE$_N$ in CP-1. In CP-2, only the data of S2 and S3 were employed to calculate AIE$_N$ for excluding the points in S1 and S4, which may be an underestimation. As shown in Fig. 4b and c, both the slope (0.144) and $R^2$ (0.050) of CP-2 are lower than those (0.544 and 0.282, respectively) of CP-1. It was verified that cloud droplets in CP-2 were influenced little by aerosols. In the previous studies, both observation and modelling studies also found that AIE$_r$ was higher under smaller aerosol amount conditions. Twyoh et al. (2005) measured the equivalent AIE$_r$ of 0.27 on the California coast, while Zhao et al. (2018) used satellite observations to attribute lower values of 0.10–0.19 for convective clouds over Hebei, a polluted region in China. Using an adiabatic cloud parcel model, Feingold (2003) found that AIE$_r$ increased from 0.199 to 0.301 when $N_p$ decreased to less than 1000 cm$^{-3}$. By using the Community Atmospheric Model version 5 (CAM5), Zhao et al. (2012) also found high AIE$_r$ values in the tropical western Pacific at Darwin (TWP) due to the low $N_p$ in December, January and February. Through studying the impact of ship-produced aerosols on the microstructure and albedo of warm marine stratocumulus clouds, Durkee et al. (2000) found that the clean and shallow boundary layers would be more readily perturbed by the addition of ship particle effluent. In this study, the higher values of AIE$_r$ and AIE$_N$ of CP-1 indicated that if the same amount of aerosol particles entered into the cloud, the microphysical parameters would be influenced more in CP-1 than in CP-2.

In addition, the meteorological conditions and the topography during the monitoring period would also affect the microphysical properties of clouds. During the observation period, CBH ranged from 460.3 to 3639.1 m with an average value of 1382.5 m. The observation station would be totally enveloped in clouds when cloud events occurred, and the corresponding distance between the observation point and CBH is represented in Fig. 2b. The sensitivity analysis of $N_C$ to CBH and $v_{up}$ was estimated by applying the equation as $S(X_i) = \partial \ln N_C / \partial \ln X_i$, where $X_i$ represented CBH and $v_{up}$. As shown in Table S2, CP-2 was more sensitive to the variation of meteorological parameters if compared with CP-1. This is consistent with the study done by McFiggans et al. (2006). They found that the sensitivity of $N_C$ to $v_{up}$ increased, while the sensitivity of $N_C$ to $N_p$ decreased when $N_p > 1000$ cm$^{-3}$. The higher values of $S(CBH)$ and $S(v_{up})$ of CP-2 indicated that CP-2 was more sensitive to the change in CBH and $v_{up}$. This might cause the periodical variations of cloud microphysical properties during CP-2.

3.5 Size distribution of cloud droplets and particles

To illustrate the evolution of the aerosol particles and cloud droplets during the cloud processes, the size distributions of $N_p$ and $N_C$ during different cloud stages are plotted in Fig. 5. For each of the four size bins, ranging from 2 to 13 µm,
Figure 5. Size distribution of particles and cloud droplets during CP-1 and CP-2. (a) Time series plot of $N_C$ in five size ranges, [2, 5) µm, [5, 7) µm, [7, 10) µm, [10, 13) µm, and [13, 50) µm, and $N_P$ in five size ranges, (15, 50) nm, [50, 100) nm, [100, 150) nm, [150, 200) nm, and [200, 765) nm. (b) Five size ranges of $N_C$ and five size ranges of $N_P$ in SL1, SH1, SL2, SH2, and CP-2; (c) five size ranges of $N_C$ and five size ranges of $N_P$ in S1, S2, S3, S4, and NC (“NC” in c represents particle size distributions during the cloudless period).
cloud number concentrations of SL1 and SL2 were lower than those of SH1 and SH2. However, in the size bin of 13–50 µm, Nc of SL1 and SL2 were the largest (Fig. 5b). The size distributions of cloud droplets in SL1 and SL2 resulted in the larger r_eff during the two stages, which was consistent with the result shown in Fig. 3b. During SH1 and SH2 in CP-1, the numbers of aerosol particles in all size bins increased. However, the increase in aerosol particles larger than 150 nm was the smallest, indicating that aerosols larger than 150 nm were more easily activated into cloud droplets. The activation of aerosol particles with sizes larger than 150 nm in this study dramatically increased Nc of 5–10 µm and made Nc of SH1 and SH2 in different size bins all comparable with those of CP-2 (Fig. 5b).

As shown in Fig. 5c, cloud droplets with DC ranging from 5 to 10 µm had high Nc in each stage in CP-2, and cloud droplets with DC ranging from 13 to 50 µm had low Nc in each stage if compared to CP-1. This caused the lower r_eff in CP-2 than in CP-1. During CP-2, aerosol particles with diameters larger than 150 nm quickly decreased by activation when cloud events occurred, while the number of aerosol particles in the size of 50–150 nm were slightly influenced by cloud events (see the left-most graph of Fig. 5a). This was consistent with the study of Targino et al. (2007), who found aerosol size distributions of cloud residuals, which represented aerosol particles activated to cloud droplets, peaked at about 0.15 µm at Mt. Åreskutan. Mertes et al. (2005) also found that particles centred at dp = 200 nm could be efficiently activated to droplets, while most Aitken mode particles remained in the interstitial phase. Compared with other stages, S1 had the highest Nc in two size bins of (2, 5) and [5, 7) µm. This indicated that large numbers of cloud droplets with small sizes were formed in the beginning of cloud events in CP-2.

3.6 Relations among LWC, r_eff and Nc

The 5 min averaged LWC for CP-1 and CP-2 is plotted against corresponding r_eff in Fig. 6a. Large cloud droplets (r_eff > 8 µm) were observed in CP-1, while the r_eff for CP-2 varied narrowly in the range of 2.5–8 µm.

Cloud droplets with r_eff > 8 µm only occurred in the two relatively clean stages, SL1 and SL2, during CP-1. It was due to the weaker competition among droplets at lower NCCN conditions. This was also observed in the US mid-Atlantic region where cloud droplets with larger sizes are more easily formed with lower NCCN (S. Li et al., 2017). At the same LWC level, the growth of cloud droplets during SH1 and SH2 was obviously limited if compared with SL1 and SL2, which is referred to as the “Twomey effect” (Twomey, 1977). This is consistent with the illustration in Fig. 3 that cloud droplets in SH1 and SH2 were smaller.

The variation r_eff and/or Nc can influence LWC, while the key factor may be different in different stages of the cloud. As shown in the lower row of Fig. 6a, CE-20 was taken as an example to discuss the relation among LWC, r_eff and Nc in different cloud stages. During S1, the existing numerous CCN (Fig. 3a) were quickly activated to form cloud droplets. The newly formed droplets were characterised with small sizes but large numbers. They will suppress the beginning of collision-coalescence processes (Rosenfeld et al., 2014a) and may further significantly delay raindrop formation (Qian et al., 2009). In S1, Nc and r_eff consistently varied. Both the increase in Nc (from 1188 to 2940 cm^{-3}) and the growth of r_eff (from ~3.5 µm to ~4.5 µm) boosted the LWC in this stage. This is different from the results of Mazoyer et al. (2019), who found a clearly inverse relationship between the number and the size of droplets at the beginning of the first hour of fog events during an observation in suburban Paris. When compared with fog, cloud is usually formed under conditions with more condensible water vapour (Fig. 1). The limited growth of droplets in fog will not occur in cloud. This causes the positive relationship with cloud droplet number and droplet size. At the beginning of S2, Nc reached the maximum. The high Nc yielded a great coalescence rate between cloud droplets. Meanwhile, the coalescence process was self-accelerating (Freud and Rosenfeld, 2012) and thus caused the quick decrease in Nc (Fig. 3a). This made cloud droplets in S2 that were characterised by larger sizes and lower number concentrations, whilst LWC simply varied in a relatively narrow range (Fig. 6a). During S3, Nc was almost constant due to the formation, coagulation and evaporation of the cloud droplets reaching a balance. As shown in Fig. 6a, the relationship between r_eff and LWC in this stage could be fitting as r_eff = a × LWC^{0.34±0.02}, which meant under the increase in LWC, the Nc was almost unchanged. The variation of LWC values was mainly due to the changes in droplet sizes. At the dissipation stage of S4, the increase in CBH brought air with low RH and high Np to the summit of Mt. Tai and caused the dissipation of cloud events (Figs. 2c and 3a). The previously activated CCN returned to the interstitial aerosol phase due to the evaporation of the droplets (Verheggen et al., 2007). Both Nc and r_eff declined. It was also illustrated in Fig. 5c that all of the Nc in the five size bins of cloud droplets decreased in S4.

In order to investigate the variation of r_eff upon Nc, the distribution of r_eff was classified with different Nc ranges in Fig. 6b. For Nc < 1000 cm^{-3}, r_eff displayed a trimodal distribution and concentrated on 3.25 µm (Peak-1), 4.86 µm (Peak-2) and 7.52 µm (Peak-3). Peak-1 corresponded to cloud droplets with low Nc, LWC and r_eff values, while the NcCCN0.2 was very high (Fig. 6c). These points represented cloud droplets in the incipient stage or the dissipation stage of cloud events where large numbers of CCN exist in the atmosphere. Peak-2 and Peak-3 represented the mature stages for cloud events with different environmental conditions. Peak-3 represented cloud droplets formed under a relatively clean atmosphere. In this circumstance, CCN were efficiently activated and had a lower concentration remaining in the atmosphere (Fig. 6c). The sufficient ambient water vapour ac-
Figure 6. The plot of LWC versus $r_{\text{eff}}$ (a) in different cloud stages of CP-1 and CP-2, (b) under different $N_C$ ranges, and (c) under different $N_{\text{CCN}}$. The time resolution of the corresponding data was 5 min in (a) and (b) and 50 min in (c). (d) The plot of albedo versus the variation of cloud thickness during CP-1 and CP-2. The averaged values of LWC and $N_C$ of CP-1 and CP-2 were applied to calculate albedo according to the equations in Sect. 2.8.

Accelerated the growth of the formed droplets, which were characterized with low $N_C$ and LWC but large $r_{\text{eff}}$. Peak-2 represented cloud droplets formed under relatively polluted conditions and was the only peak found for $N_C$ larger than 1000 cm$^{-3}$. With the increase in $N_C$, the distribution of this peak narrowed and slightly moved to lower $r_{\text{eff}}$ mode.

The thickness of orographic cloud can be easily influenced by the specific topography and environmental conditions (Barros and Lettenmaier, 1994; Welch et al., 2008). When assuming the cloud thickness during CP-1 and CP-2 to be equal, albedo would depend on the values of LWC and $N_C$, as described in Sect. 2.8. Cloud albedo during CP-2 was always higher than that during CP-1, especially when the cloud thickness was lower than about 2500 m (Fig. 6d).

Note that the increase in $N_C$ could enhance the evaporation and further reduce the lifetime of cloud, which was not taken into account when calculating the induced albedo. Through studying marine stratocumulus clouds in the north-eastern Pacific Ocean, Twohy et al. (2005) also found that the increase in $N_C$ by a factor of 2.8 would lead to 40% increase in albedo, going from 0.325 to 0.458. This indicated that the higher $N_C$ would increase the cloud albedo when assuming no change to cloud thickness and when disregarding any cloud lifetime effects.
4 Conclusions

From 17 June to 30 July 2018, in situ observations of number concentrations and size distributions of aerosol particles and cloud droplets were used to study aerosol–cloud interactions at the summit of Mt. Tai. Large variations of the characteristics in terms of \( N_C \), LWC and \( r_{\text{eff}} \) were found during the observation period. Cloud events with small \( r_{\text{eff}} \) and low LWC similar to urban fog were also observed at Mt. Tai.

Two typical cloud processes, CP-1 and CP-2, were applied to study the cloud–aerosol interactions based on the aerosol characteristics (especially \( N_P \) and \( N_{\text{CCN}} \)) before cloud onsets. For the CP-1, which corresponded to relatively clean conditions, water content was sufficient, while \( N_{\text{CCN}} \) was considered the limitation of cloud droplet formation according to the observation results. The newly formed cloud droplets were characterised with low \( N_C \), high LWC / \( N_C \) and large \( r_{\text{eff}} \). With the increase in aerosol concentration, \( N_C \) was found to dramatically increase. Large numbers of \( N_{\text{CCN}} \) competed for the water content with the formed cloud droplets and, as a result, further dramatically decrease the LWC / \( N_C \) and \( r_{\text{eff}} \) values of cloud droplets. In CP-2, \( N_P \) before the cloud onset was high and \( N_{\text{CCN}} \) was sufficient. Water vapour was considered the limitation for cloud formation. Large numbers of small cloud droplets with low LWC / \( N_C \) were observed in the incipient stage of cloud formation. In addition, periodical changes in cloud microphysical properties were found. Both consistent variation and inverse variation between \( N_P \) and \( N_C \) were observed in this study, which were characterised with relatively high and low LWC / \( N_C \) values, respectively.

Both positive AIE\(_P\) and AIE\(_N\) values observed at Mt. Tai indicate that the increase in \( N_P \) will decrease \( r_{\text{eff}} \) and increase \( N_C \) of cloud droplets. The lower values of AIE\(_P\) and AIE\(_N\) with higher \( N_P \) and \( N_{\text{CCN}} \) suggest that the increase in \( N_P \) will more strongly decrease the size and increase the number of cloud droplets under the conditions with smaller aerosol amounts. Through studying the size distributions of aerosol particles and cloud droplets, higher \( N_C \) in the size bin of 13–50 µm resulted in the larger \( r_{\text{eff}} \) during the two clean stages in CP-1. Particles larger than 150 nm were able to be efficiently activated to cloud droplets and made important contributions to the increase in cloud droplets in the size range of 5–10 µm. The LWC of cloud depended on the change in \( r_{\text{eff}} \) and \( N_C \). However, the decisive factor may differ at different stages of the cloud. In general, the \( r_{\text{eff}} \) of cloud droplets consistently varied with LWC. However, in different \( N_C \) ranges, the \( r_{\text{eff}} \) of cloud droplets were observed with different distribution shapes. For \( N_C < 1000 \text{ cm}^{-3} \), \( r_{\text{eff}} \) displayed a trimodal distribution. Three peaks at 3.25, 4.86 and 7.52 µm were observed. With the increase in \( N_C \), a narrowed unimodal distribution of \( r_{\text{eff}} \) appeared, and the peak value slightly moved towards lower \( r_{\text{eff}} \) mode. For a constant cloud thickness and disregarding any cloud lifetime effects, the increased \( N_C \) and decreased \( r_{\text{eff}} \) dramatically increase the cloud albedo, which may further influence the regional climate in the North China Plain.

The local topography of the surrounding areas at Mt. Tai supplies potential access for aerosol transportation and can affect the measured cloud droplet distributions by increasing turbulence or causing orographic flows. Even though the summit of Mt. Tai is far away from the polluted sources, the transported CCN could change the cloud microphysical properties (e.g. during CP-1). The cloud microphysical parameters derived in our study characterised the cloud features in the North China Plain and provided valuable data for modelling studies of cloud microphysics in the future.

Data availability. All data used to support the conclusion are presented in this paper. Additional data are available upon request. Please contact the corresponding authors Jianmin Chen (jm-chen@fudan.edu.cn) and Hui Chen (hui_chen@fudan.edu.cn).

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Competing interests. The authors declare that they have no conflict of interest.

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References

Albrecht, B. A.: Aerosols, cloud microphysics, and fractional cloudiness, Science, 245, 1227–1230, https://doi.org/10.1126/science.245.4923.1227, 1989.

Allan, J. D., Baumgardner, D., Raga, G. B., Mayol-Bracero, O. L., Morales-Garcia, F., García-Garcia, F., Montero-Martínez, G., Bormann, S., Schneider, J., Mertes, S., Walter, S., Gysel, M., Dusek, U., Frank, G. P., and Krämer, M.: Clouds and aerosols in Puerto Rico – a new evaluation, Atmos. Chem. Phys., 8, 1293–1309, https://doi.org/10.5194/acp-8-1293-2008, 2008.

Andreae, M. O., Rosenfeld, D., Artaxo, P., Costa, A. A., Frank, G. P., Longo, K. M., and Silva-Dias, M. A. F.: Smok- ing rain clouds over the Amazon, Science, 303, 1337–1342, https://doi.org/10.1126/science.1092779, 2004.

Barros, A. P. and Lettenmaier, D. P.: Dynamic modeling of orographically induced precipitation, Rev. Geophys., 32, 265–284, https://doi.org/10.1029/94rg00623, 1994.

Boute, I., Price, J., Kudzotsa, I., Kokkola, H., and Ro- makkainen, S.: Aerosol–fog interaction and the transition to well-mixed fog clouds, Atmos. Chem. Phys., 18, 7827–7840, https://doi.org/10.5194/acp-18-7827-2018, 2018.

Chang, Y., Zhang, Y.-L., Li, J., Tian, C., Song, L., Zhai, X., Zhang, W., Huang, T., Lin, Y.-C., Zhu, C., Fang, Y., Lehmann, M. F., and Chen, J.: Isotopic constraints on the atmospheric sources and formation of nitrogenous species in clouds influenced by biomass burning, Atmos. Chem. Phys., 19, 12221–12234, https://doi.org/10.5194/acp-19-12221-2019, 2019a.

Chang, Y., Guo, X., Tang, J., and Lu, G.: Aircraft measurement campaign on summer cloud microphysical properties over the Tibetan Plateau, Sci. Rep., 9, 49122, https://doi.org/10.1038/s41598-019-41544-5, 2019b.

Choularton, T. W., Colville, R. N., Bower, K. N., Gallagher, M. W., Wells, M., Beswick, K. M., Arends, B. G., Mols, J. J., Kos, G. P. A., Fuzzi, S., Lind, J. A., Orsi, G., Facchini, M. C., Laj, P., Gieray, R., Wieser, P., Engelhardt, T., Berner, A., Kruijss, C., Moller, D., Acker, K., Wieprecht, W., Lutke, J., Lesven, K., Bizjak, M., Hansson, H. C., Cederfelt, S. I., Frank, G., Mentes, B., Martinsson, B., Orsini, S., Svenningsson, B., Siewicki, E., Wiedensohler, A., Noone, K. J., Pahl, S., Winkler, P., Seyffer, E., Helas, G., Jaeschke, W., Georgii, H. W., Wobrock, W., Preiss, M., Maser, R., Schell, D., Dollard, G., Jones, B., Davies, T., Sedlak, D. L., David, M. M., Wendisch, M., Cape, J. N., Hargreaves, K. J., Sutton, M. A., Storeton-West, R. L., Fowler, D., Hallberg, A., Harrison, R. M., and Peak, J. D.: The Great Dun Fell Cloud Experiment 1993: An overview, Atmos. Environ., 31, 2393–2405, https://doi.org/10.1016/s1352-2310(96)00316-0, 1997.

Croft, B., Lohmann, U., Martin, R. V., Siter, P., Wurzler, S., Feichter, J., Hoose, C., Heikkilä, U., van Donkelaar, A., and Ferrachat, S.: Influences of in-cloud aerosol scavenging parameterizations on aerosol concentrations and wet deposition in ECHAM5-HAM, Atmos. Chem. Phys., 10, 1511–1543, https://doi.org/10.5194/acp-10-1511-2010, 2010.

Cross, E. S., Slowik, J. G., Davidovits, P., Allan, J. D., Worsnop, D. R., Jayne, J. T., Lewis, T. D., K., Canagaratna, M., and Onasch, T. B.: Laboratory and Ambient Particle Density Determinations using Light Scattering in Conjunction with Aerosol Mass Spectrometry, Aerosol Sci. Technol., 41, 343–359, https://doi.org/10.1080/02786820701199736, 2007.

J. Li et al.: The evolution of cloud and aerosol microphysics

Demoz, B. B., Collett, J. L., and Daube, B. C.: On the Caltech Active Strand Cloudwater Collectors, Atmos. Res., 41, 47–62, https://doi.org/10.1016/0169-8095(95)00044-5, 1996.

Deng, Z., Zhao, C., Zhang, Q., Huang, M., and Ma, X.: Statistical analysis of microphysical properties and the parameterization of effective radius of warm clouds in Beijing area, Atmos. Res., 93, 888–896, 2009.

Drewnick, F., Schneider, J., Hings, S. S., Hock, N., Noone, K., Targino, A., Weimer, S., and Bormann, S.: Measurement of ambient, interstitial, and residual aerosol particles on a mountaintop site in central Sweden using an aerosol mass spectrometer and a CVI, J. Atmos. Chem., 56, 1–20, https://doi.org/10.1007/s10587-006-9036-8, 2007.

Durkee, P. A., Noone, K. J., Ferek, R. J., Johnson, D. W., Taylor, J. P., Garrett, T. J., Hobbs, P. V., Hudson, J. G., Bretherton, C. S., Innis, G., Frick, G. M., Hoppel, W. A., O’Dowd, C. D., Russell, L. M., Gasparovic, R., Nielsen, K. E., Tessler, S. A., Ostrom, E., Osborne, S. R., Flagan, R. C., Seinfeld, J. H., and Rand, H.: The impact of ship-produced aerosols on the microstructure and albedo of warm marine stratuscumulus clouds: A test of MAST hypotheses li and lii, J. Atmos. Sci., 57, 2554–2569, https://doi.org/10.1175/1520-0469(2000)057<2554:Tiasea>2.0.Co;2, 2000.

Dusek, U., Frank, G. P., Hildebrandt, L., Curtius, J., Schneider, J., Walter, S., Chand, D., Drewnick, F., Hings, S., Jung, D., Borm- mann, S., and Andreae, M. O.: Size matters more than chemistry for cloud-nucleating ability of aerosol particles, Science, 312, 1375–1378, https://doi.org/10.1126/science.1125261, 2006.

Ekman, A. M. L., Engstrom, A., and Soderberg, A.: Impact of Two-Way Aerosol-Cloud Interaction and Changes in Aerosol Size Distribution on Simulated Aerosol-Induced Deep Convective Cloud Sensitivity, J. Atmos. Sci., 68, 685–698, https://doi.org/10.1175/2010jas3651.1, 2011.

Fan, J., Leung, L. R., Rosenfeld, D., Chen, Q., Li, Z., Zhang, J., and Yan, H.: Microphysical effects determine macrophysical response for aerosol impacts on deep convective clouds, P. Natl. Acad. Sci. USA, 110, 4581–4590, https://doi.org/10.1073/pnas.1316830110, 2013.

Fan, J., Wang, Y., Rosenfeld, D., and Liu, X.: Review of Aerosol-Cloud Interactions: Mechanisms, Significance, and Challenges, J. Atmos. Sci., 73, 4221–4252, https://doi.org/10.1175/jas-d-16-0037.1, 2016.

Feingold, G., Remer, L. A., Ramaprasad, J., and Kaufman, Y. J.: Analysis of smoke impact on clouds in Brazilian biomass burning regions: An extension of Twomey’s approach, J. Geophys. Res.-Atmos., 106, 22907–22922, https://doi.org/10.1029/2001jd000732, 2001.

Feingold, G.: Modeling of the first indirect effect: Analysis of measurement requirements, Geophys. Res. Lett., 30, 1997, https://doi.org/10.1029/2003gl017967, 2003.

Freud, E. and Rosenfeld, D.: Linear relation between convective cloud drop number concentration and depth for rain initiation, J. Geophys. Res.-Atmos., 117, 13, https://doi.org/10.1029/2011jd016457, 2012.

Frey, L., Bender, F. A.-M., and Svensson, G.: Cloud albedo changes in response to anthropogenic sulfate and non-sulfate aerosol forcings in CMIP5 models, Atmos. Chem. Phys., 17, 9145–9162, https://doi.org/10.5194/acp-17-9145-2017, 2017.
Lu, M.-L., Conant, W. C., Jonsson, H. H., Varutbangkul, Li, S., Joseph, E., Min, Q., and Yin, B.: Multi-year ground-based
observations of aerosol-cloud interactions in the Mid-Atlantic of the United States, J. Quant. Spectrosc. Radiat. Transfer, 188, 192–199, https://doi.org/10.1016/j.jqsrt.2016.02.004, 2017.

Lohmann, U. and Feichter, J.: Global indirect aerosol effects: a review, Atmos. Chem. Phys., 5, 715–737, https://doi.org/10.5194/acp-5-715-2005, 2005.

Lu, C., Niu, S., Tang, L., Lv, J., Zhao, L., and Zhu, B.: Chemical composition of fog water in Nanjing area of China and its related fog microphysics, Atmos. Res., 97, 47–69, 2010.

Lu, M.-L., Conant, W. C., Jonsson, H. H., Varutbangkul, V., Flagan, R. C., and Seinfeld, J. H.: The Marine Stratus/Stratocumulus Experiment (MASE): Aerosol-cloud relationships in marine stratocumulus, J. Geophys. Res.-Atmos., 112, https://doi.org/10.1029/2006JD007985, 2007.

Mazoyer, M., Burnet, F., Denjean, C., Roberts, G. C., Haeffelin, M., Dupont, J.-C., and Elias, T.: Experimental study of the aerosol impact on fog microphysics, Atmos. Chem. Phys., 19, 4323–4344, https://doi.org/10.5194/acp-19-4323-2019, 2019.

McComiskey, A., Feingold, G., Frisch, A. S., Turner, D. D., Miller, M. A., Chiu, J. C., Min, Q., and Ogren, J. A.: An assessment of aerosol-cloud interactions in marine stratus clouds based on surface remote sensing, J. Geophys. Res.-Atmos., 114, D09203, https://doi.org/10.1029/2008JD011006, 2009.

McGfigns, G., Artaxo, P., Baltensperger, U., Coe, H., Facchini, M. C., Feingold, G., Fuzzi, S., Gysel, M., Laaksonen, A., Lohmann, U., Mentel, T. F., Murphy, D. M., O’Dowd, C. D., Snider, J. R., and Weingartner, E.: The effect of physical and chemical aerosol properties on warm cloud droplet activation, Atmos. Chem. Phys., 6, 2593–2649, https://doi.org/10.5194/acp-6-2593-2006, 2006.

Mertes, S., Galgon, D., Schwirm, K., Nowak, A., Lehmann, K., Massling, A., Wiedensohler, A., and Wieprecht, W.: Evolution of particle concentration and size distribution observed upwind, inside and downwind hill cap clouds at connected flow conditions during FEBUKO, Atmos. Environ., 39, 4233–4245, https://doi.org/10.1016/j.atmosenv.2005.02.009, 2005.

Modini, R. L., Frossard, A. A., Ahln, L., Russell, L. M., Corrigan, C. E., Roberts, G. C., Hawkins, L. N., Schroder, J. C., Bertram, A. K., Zhao, R., Lee, A. K. Y., Abbatt, J. P. D., Lin, J., Nenes, A., Wang, Z., Wonaschuetz, A., Sorooshian, A., Noone, K. J., Jonsson, H., Seinfeld, J. H., Toom-Sauntry, D., Macdonald, A. M., and Leaitch, W. R.: Primary marine aerosol-cloud interactions off the coast of California, J. Geophys. Res.-Atmos., 120, 4282–4303, https://doi.org/10.1002/2014jd022963, 2015.

Möller, D., Acker, K., and Wieprecht, W.: A relationship between liquid water content and chemical composition in clouds, Atmos. Res., 41, 321–335, 1996.

Noone, K. J., Ogren, J. A., and Heinzenberger, J.: An Examination of Clouds at a Mountain-Top Site in Central Sweden: The Distribution of Solute within Cloud Droplets, Atmos. Res., 25, 3–15, https://doi.org/10.1016/0169-8095(90)90002-T, 1990.

Padmakumari, B., Maheshrumar, R. S., Anand, V., and Axtia, D.: Microphysical characteristics of convective clouds over ocean and land from aircraft observations, Atmos. Res., 195, 62–71, https://doi.org/10.1016/j.atmosres.2017.05.011, 2017.

Penner, J. E., Dong, X. Q., and Chen, Y.: Observational evidence of a change in radiative forcing due to the indirect aerosol effect, Nature, 427, 231–234, https://doi.org/10.1038/nature02234, 2004.

Qian, Y., Gong, D. Y., Fan, J. W., Leung, L. R., Bennartz, R., Chen, D. L., and Wang, W. G.: Heavy pollution suppresses light rain in China: Observations and modeling, J. Geophys. Res.-Atmos., 114, 16, https://doi.org/10.1029/2008JD011575, 2009.

Quante, M.: The role of clouds in the climate system, J. Phys. IV, 6153, 1999.

https://doi.org/10.5194/acp-20-13735-2020

Atmos. Chem. Phys., 20, 13735–13751, 2020
Zhang, Y., Rossow, W. B., Lacis, A. A., Oinas, V., and Mishchenko, M. I.: Calculation of radiative fluxes from the surface to top of atmosphere based on ISCCP and other global data sets: Refinements of the radiative transfer model and the input data, J. Geophys. Res.-Atmos., 109, https://doi.org/10.1029/2003JD004457, 2004.

Zhao, C., Klein, S. A., Xie, S., Liu, X., Boyle, J. S., and Zhang, Y.: Aerosol First Indirect Effects on Non-Precipitating Low-Level Liquid Cloud Properties as Simulated by CAM5 at ARM Sites, Geophys. Res. Lett., 39, L08806, https://doi.org/10.1029/2012GL051213, 2012.

Zhao, C., Qiu, Y., Dong, X., Wang, Z., Peng, Y., Li, B., Wu, Z., and Wang, Y.: Negative Aerosol-Cloud Relationship from Aircraft Observations over Hebei, China, Earth Space Sci., 5, 19–29, https://doi.org/10.1002/2017EA000346, 2018.

Zhou, Y., Wang, T., Gao, X., Xue, L., Wang, X., Wang, Z., Gao, J., Zhang, Q., and Wang, W.: Continuous observations of watersoluble ions in PM$_{2.5}$ at Mount Tai (1534 m a.s.l.) in central-eastern China, J. Atmos. Chem., 64, 107–127, 2009.

Zhu, C., Chen, J., Wang, X., Li, J., Wei, M., Xu, C., Xu, X., Ding, A., and Collett, J. L.: Chemical Composition and Bacterial Community in Size-Resolved Cloud Water at the Summit of Mt. Tai, China, Aerosol Air Qual. Res., 18, 1–14, https://doi.org/10.4209/aaqr.2016.11.0493, 2018.