

**Spatial distribution of aerosol microphysical and optical properties and direct radiative effect from the China Aerosol Remote Sensing Network**

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37 **Abstract**

38 Multi-year observations of aerosol microphysical and optical properties  
39 obtained through ground-based remote sensing at 50 China Aerosol Remote  
40 Sensing Network (CARSNET) sites were used to characterize the aerosol  
41 climatology for representative remote, rural, and urban areas over China to  
42 assess effects on climate. The annual mean effective radii for total particles  
43 ( $R_{\text{eff}}$ ) decreased from north to south and from rural to urban sites, and high  
44 total particle volumes were found at the urban sites. The aerosol optical depth  
45 at 440 nm ( $AOD_{440\text{nm}}$ ) increased from remote/rural sites (0.12) to urban sites  
46 (0.79), and the extinction Ångström exponent ( $EAE_{440-870\text{nm}}$ ) increased from  
47 0.71 at the arid/semi-arid sites to 1.15 at the urban sites, presumably due to  
48 anthropogenic emissions. Single scattering albedos ( $SSA_{440\text{ nm}}$ ) ranged from  
49 0.88 to 0.92 indicating slightly to strongly absorbing aerosols. Absorption  
50  $AOD_{440\text{nm}}$ 's were 0.01 at the remote sites versus 0.07 at the urban sites. The  
51 average direct aerosol radiative effect (DARE) at the bottom of atmosphere  
52 increased from the sites in the remote ( $-24.40\text{ W/m}^2$ ) to the urban area  
53 ( $-103.28\text{ W/m}^2$ ) indicating increased cooling at the latter. The DARE for the top  
54 of the atmosphere increased from  $-4.79\text{ W/m}^2$  at the remote sites to  $-30.05$   
55  $\text{W/m}^2$  at the urban sites, indicating overall cooling effects for the  
56 earth-atmosphere system. A classification method based on  $SSA_{440\text{ nm}}$ , fine  
57 mode fraction (FMF), and  $EAE_{440-870\text{ nm}}$  showed that coarse mode particles  
58 (mainly dust) were dominant at the rural sites near the northwestern deserts,  
59 while light-absorbing, fine-mode particles were important at most urban sites.  
60 This study will be important for understanding aerosol climate effects and  
61 regional environmental pollution, and the results will provide useful information  
62 for satellite validation and the improvement of climate modelings.

63 **Keywords:** aerosol optical properties; direct aerosol radiative effect; aerosol  
64 type; climatology; China Aerosol Remote Sensing Network

65

## 66 **1. Introduction**

67 Atmospheric aerosols have important direct effects on climate because  
68 they can scatter and absorb radiant energy and in so doing affect the Earth's  
69 energy balance (Charlson et al., 1992; Yang et al., 2016). Meanwhile, the  
70 aerosols can be served as cloud condensation nuclei or ice nuclei to affect the  
71 climate indirectly through aerosol–cloud interactions (Twomey et al., 1984;  
72 Garrett et al., 2006; Zhao et al., 2015; Xie et al., 2013). The optical properties  
73 of the aerosol determine the particles' direct effects on the Earth's radiative  
74 balance and weather-climate change (Ramanathan et al., 2001; Eck et al.,  
75 2005; Myhre, 2009; Zhao et al., 2018). Aerosol optical depth (AOD) is one of  
76 the key measures of the total aerosol extinction effects on climate (Breon et al.,  
77 2002), and the extinction Ångström exponent (EAE) with spectral dependence  
78 can be used to obtain the information about aerosol size distributions (Gobbi et  
79 al., 2007; Eck et al., 1999; Zheng et al., 2017). The aerosols' absorptivity  
80 depends on particle composition is a key determinant to calculate the direct  
81 aerosol radiative effect (Haywood and Shine, 1995; Li et al., 2016; Zheng et al.,  
82 2018), and the single scattering albedo (SSA) is the parameter has spectral  
83 dependence to distinguish major aerosol particle types (Jacobson et al., 2000;  
84 Dubovik et al., 2002; Gelencser et al., 2004; Russell et al., 2010; Giles et al.,  
85 2012).

86 With the recognition of the importance for climate, the aerosol optical  
87 properties have been obtained from ground-based monitoring networks  
88 worldwide; some of the major networks include AERONET-Aerosol Robotic  
89 Network) (Holben et al., 1998) and its sub-networks of  
90 PHOTONS-PHOTométrie pour le Traitement Opérationnel de Normalisation  
91 Satellitaire, AEROCAN-Canadian Sun-Photometer Network, and RIMA-Iberian  
92 Network for aerosol measurements ((Goloub et al., 2007; Bokoye et al., 2001;  
93 Prats et al., 2011), SKYNET-SKYrad Network (Takamura and Nakajima, 2004;  
94 Che et al., 2008), EARLINET-European aerosol Lidar network (Pappalardo et  
95 al., 2014), the GAW-PFR Network-Global Atmosphere Watch

96 Programmer-Precision Filter Radiometers (Wehrli, 2002; Estellés et al., 2012),  
97 The CARSNET-China Aerosol Remote Sensing NETwork, the  
98 CSHNET-Chinese Sun Hazemeter Network and the SONET-Sun–Sky  
99 Radiometer Observation Network have been established to measure aerosol  
100 optical properties in China (Che et al., 2009a, 2015; Xin et al., 2007; Li et al.,  
101 2018). Furthermore, the aerosol optical properties have also been used in  
102 comprehensive studies of aerosol physical characteristics and chemical  
103 composition in many regions of China (Che et al., 2009c, 2018; Zhao et al.,  
104 2018).

105 China has become one of the largest aerosol sources in the world  
106 associated with its rapid economic development, and this has caused  
107 significant effects on local environments and regional climate (Che et al., 2005;  
108 Xia, 2010; Li et al., 2016; Yang et al., 2018, 2019; Zhao et al., 2019). There  
109 have been numerous studies that have focused on aerosol optical properties  
110 obtained through ground-based remote sensing methods in China (Luo et al.,  
111 2002; Li et al., 2003; Duan and Mao, 2007). A few researches have paid more  
112 attention to the aerosol optical properties and its radiative effects over the  
113 urban-industrial areas as well as at coastal sites in northeastern and eastern  
114 China (Wang et al., 2010; Xin et al., 2011; Xia et al., 2007; Zhao et al., 2016;  
115 Wu et al. 2012; Shen et al., 2019). Many studies of aerosol optical properties  
116 were conducted in northern China with high aerosol loadings, such as the  
117 Beijing-Tianjin-Hebei region (Che et al. 2014; Xia et al., 2013; Fan et al., 2006;  
118 Xie et al., 2008; Zhang et al., 2019; Yang et al., 2019; Zhao et al., 2018).  
119 Aerosol optical properties also have been investigated at Hefei, Shouxian,  
120 Nanjing, Taihu, Shanghai and other sites in eastern China (Lee et al., 2010; He  
121 et al., 2012; Zhuang et al., 2014; Wang Z et al., 2015; Che et al., 2018). Some  
122 studies of aerosol optical properties have been made in southern China (Wang  
123 et al., 2015; Tao et al., 2014), and those at remote and rural sites in China  
124 provide information on regional background conditions (Che et al., 2009b;  
125 Wang et al., 2010; Xue et al., 2011; Zhu et al., 2014; Yuan et al., 2014).

126 China's vast size, varied terrain, and heterogeneity of aerosol sources,  
127 has led to strong temporal and spatial variability in aerosol optical and physical  
128 properties. The mixtures of aerosol types at most sites are complex, and  
129 aerosol populations' size and composition are affected by their sources,  
130 transformations that occurring during transportation and removing processes  
131 (Cao et al., 2007; Wang et al., 2007; Zhang et al., 2013; Wan et al., 2015).  
132 National scale, ground-based measurements of aerosol microphysical and its  
133 optical properties obtained from the sunphotometer provide for a better  
134 understanding of the aerosols' climate effects over the different regions of  
135 China. The measurements of greatest interest include aerosol size  
136 distributions (volume and aerosol effective radii), optical properties (AOD, AE,  
137 SSA, absorption AOD) because those data can at least be used to evaluate  
138 direct radiative effect.

139 The aim of this study was focused on the investigation of the  
140 climatological spatial distribution of aerosol microphysical and optical  
141 properties over regional-scales using spatial distribution data from the national  
142 CARSNET network. The data were collected at CARSNET sites, which include  
143 sites in the remote, rural and urban area, with the same calibration procedures  
144 and calculation algorithms were used at all sites. As a result, the data are  
145 directly comparable among sites (Che et al., 2009a), and the results can be  
146 provided to characterize the regional distribution and temporal variation of  
147 aerosol optical properties. This research focused on aerosol climate effects  
148 and regional environmental pollution, and the results should be useful for  
149 satellite validations and for the improvement of models in the future. The  
150 remainder of this paper is organized as following: Section 2 firstly describes  
151 the sites in detail, and then introduced the methods in data processing of the  
152 aerosol optical properties as well as the direct aerosol radiative effect  
153 calculation through the retrieved aerosol optical parameters. Section 3  
154 illustrates the aerosol microphysical and optical properties, as well as its direct  
155 aerosol radiative effect. An aerosol type classification method is proposed

156 according to the aerosol optical parameters. Section 4 presents the  
157 conclusions of the study.

## 158 **2 Site description, instruments, and data**

### 159 **2.1 Site description**

160 Sunphotometers (CE-318, Cimel Electronique, Paris, France, see  
161 Appendix A,) were installed at 50 CARSNET sites (Fig. 1) from 2010 to 2017.  
162 The stations were classified as remote, rural, or urban sites based on  
163 administrative division (Appendix Table 1). Three of the remote stations were  
164 about more than 3000 m above the sea level on the Tibetan Plateau far from  
165 the anthropogenic influences, and one of them was a northwestern regional  
166 background site in China. The 23 rural sites represent (a) five sites of desert  
167 regions affected by most of dust aerosols rather than anthropogenic particles,  
168 (b) two sites affected by both dust and anthropogenic activities on the Loess  
169 Plateau, and (c) 16 sites located near or surrounding the large cities relatively  
170 strong to the impacts of anthropogenic activities in the central and eastern  
171 China. The last category is 24 urban sites located in provincial capitals or  
172 heavily populated cities.

### 173 **2.2 Instruments and calibration**

174 The CE-318 sunphotometers used in this study were calibrated annually,  
175 using the CARSNET calibration protocol, to verify the accuracy and reliability  
176 of the sky irradiance measurements (Holben et al., 1998; Che et al., 2009; Tao  
177 et al., 2014). The reference instruments for CARSNET were periodically  
178 calibrated at Izaña, Tenerife, Spain located at 28.31 °N, 16.50 °W (2391.0 m  
179 a.s.l.) in conjunction with the AERONET program. There are several different  
180 types of the Cimel instruments that have been used at the 50 sites in this  
181 network as follows: (1) logical type CE-318 sunphotometers (440 nm, 675 nm,  
182 870 nm, 940 nm, 1020 nm and three 870 nm at the polarization band), (2)  
183 numerical type CE-318 sunphotometers (440 nm, 675 nm, 870 nm, 940 nm,  
184 1020 nm and three polarization bands at 870 nm), (3) numerical type CE-318  
185 sunphotometers at eight wavelengths (340 nm, 380 nm, 440 nm, 500 nm, 675

186 nm, 870 nm, 940 nm, and 1020 nm), (4) and numerical type CE-318  
187 sunphotometers at nine wavelengths (340 nm, 380 nm, 440 nm, 500 nm, 675  
188 nm, 870 nm, 940 nm, 1020 nm and 1640 nm).

189 Measurements used to retrieve AODs were at 340 nm, 380 nm, 440 nm,  
190 500 nm, 675 nm, 870 nm, 1020 nm, and 1640 nm, while the total precipitable  
191 water content was obtained by using those measurements at 940 nm (Holben  
192 et al., 1998; Dubovik and King, 2000). The cloud-screened AOD data were  
193 calculated by using the ASTPwin software, and extinction Ångström exponents  
194 (EAE) were calculated from the instantaneous AODs for wavelengths of 440  
195 nm and 870 nm (Che et al., 2009, 2015). Sites with more than three daily AOD  
196 observations and more than 10 monthly AOD observation days were used to  
197 calculate the daily and monthly mean AODs and extinction Ångström  
198 exponents. The FMF is described as the fraction of fine mode particles of total  
199  $AOD_{440nm}$  ( $AOD_{fine440 nm}/AOD_{440 nm}$ ).

200

### 201 **2.3. Data processing**

202 The aerosol microphysical and optical properties, including volume size  
203 distributions ( $dV(r)/d\ln(r)$ ), the total, fine, and coarse mode aerosol effective  
204 radii ( $R_{effT}$ ,  $R_{effF}$ , and  $R_{effC}$ , respectively) single-scattering albedo (SSA),  
205 complex refractive indices, absorption AODs (AAODs), and absorption  
206 Ångström exponents (AAEs), were retrieved from the observational data from  
207 the sky scattering channel of the sunphotometers at 440 nm, 670 nm, 870 nm,  
208 1020 nm using the algorithms of Dubovik et al. (2002, 2006). In the process of  
209 retrieval, the data of surface albedo (SA) was interpolated or extrapolated to  
210 440 nm, 670 nm, 870 nm, and 1020 nm based on the daily MCD43C3 data, a  
211 product from the MODIS-Moderate Resolution Imaging Spectroradiometer  
212 surface reflectance (<https://ladsweb.modaps.eosdis.nasa.gov/>). The algorithm  
213 used to calculate aerosol volume size distributions ( $dV/\ln r$ ) was under the  
214 assumption of a homogeneous distribution of non-spherical particles following  
215 the approach of Dubovik (2006). The sphericity fraction retrieved from the

216 inversions is defined as: spherical particles/(spheroidal particles + spherical  
217 particles) (Giles et al., 2011).

218 As Dubovik et al. (2002, 2006) defined that all the particles with effective  
219 radii < 0.992 μm were considered as fine mode particles; and those > 0.992  
220 μm were considered as coarse mode particles. For the total ( $R_{\text{effT}}$ ), fine ( $R_{\text{effF}}$ )  
221 and coarse ( $R_{\text{effC}}$ ) mode aerosols, the effective radii are calculated by the  
222 equation as follows:

$$223 \quad R_{\text{eff}} = \frac{\int_{r_{\text{min}}}^{r_{\text{max}}} r^3 \frac{dN(r)}{d \ln r} d \ln r}{\int_{r_{\text{min}}}^{r_{\text{max}}} r^2 \frac{dN(r)}{d \ln r} d \ln r} \quad (1)$$

224 Where  $r_{\text{min}}$  denotes 0.05, 0.05, 0.992 μm and  $r_{\text{max}}$  denotes 15, 0.992, 15  
225 μm of the total, fine and coarse mode particles, respectively.

226 The coarse ( $PV_C$ ) and fine aerosol particle volumes distributions ( $PV_F$ ) are  
227 calculated according to a bimodal lognormal function described by Whitby  
228 (1978), Shettle and Fenn (1979) and Remer and Kaufman (1998):

$$229 \quad \frac{dV(r)}{d \ln r} = \sum_{i=1}^2 \frac{C_{v,i}}{\sqrt{2\pi}\sigma_i} \exp \left[ -\frac{(\ln r - \ln r_{v,i})^2}{2\sigma_i^2} \right] \quad (2)$$

230 where  $C_{v,i}$  means for the volume concentration;  $r_{v,i}$  means the median  
231 radius, and  $\sigma_i$  means the standard deviation.

232 The volume median radius is computed by fine and coarse modes  
233 particles as follows:

$$234 \quad \ln r_V = \frac{\int_{r_{\text{min}}}^{r_{\text{max}}} \ln r \frac{dV(r)}{d \ln r} d \ln r}{\int_{r_{\text{min}}}^{r_{\text{max}}} \frac{dV(r)}{d \ln r} d \ln r} \quad (3)$$

235 Then the standard deviation is calculated from the volume median radius:

$$236 \quad \sigma_V = \sqrt{\frac{\int_{r_{\text{min}}}^{r_{\text{max}}} (\ln r - \ln r_V)^2 \frac{dV(r)}{d \ln r} d \ln r}{\int_{r_{\text{min}}}^{r_{\text{max}}} \frac{dV(r)}{d \ln r} d \ln r}} \quad (4)$$



237 The volume concentration ( $\mu\text{m}^3/\mu\text{m}^2$ ) is speculated by the following  
 238 equation:

$$239 \quad C_V = \int_{r_{min}}^{r_{max}} \frac{dV(r)}{d \ln r} d \ln r \quad (5)$$

240 The SSA was retrieved only for  $\text{AOD}_{440\text{nm}} > 0.40$ ; this was done to avoid  
 241 the larger uncertainty inherent in the lower AOD retrieval according to Dubovik  
 242 et al. (2002, 2006). The AAOD and AAE for wavelength  $\lambda$  were calculated as  
 243 follows:

$$244 \quad \text{AAOD}(\lambda) = [1 - \text{SSA}(\lambda)] \times \text{AOD}(\lambda) \quad (6)$$

$$245 \quad \text{AAE} = \frac{-d \ln[\text{AAOD}(\lambda)]}{d \ln(\lambda)} \quad (7)$$

246 The total AODs' uncertainty was 0.01 to 0.02 according to Eck et al.  
 247 (1999). The accuracy of SSA retrieved from  $\text{AOD}_{440\text{nm}} > 0.50$  with solar zenith  
 248 angles  $> 50$  was 0.03 (Dubovik et al., 2002). The accuracy of the particle  
 249 volume size distribution was 15–25% between  $0.1 \mu\text{m} \leq r \leq 7.0 \mu\text{m}$  and 25–100%  
 250 in conditions of  $r < 0.1 \mu\text{m}$  and  $r > 7 \mu\text{m}$ .

251 Direct aerosol radiative effect (DARE in  $\text{W}/\text{m}^2$ ) was calculated by the  
 252 radiative transfer module under cloud-free conditions, which is similar to the  
 253 inversion of AERONET (García et al., 2008; 2012). The DARE at the bottom of  
 254 the atmosphere (BOA) and the top of the atmosphere (TOA) was defined as  
 255 the difference in the shortwave radiative fluxes with and without aerosol effects  
 256 as follows:

$$257 \quad \text{DARE}_{TOA} = F_{TOA}^{\uparrow 0} - F_{TOA}^{\uparrow} \quad (8)$$

$$260 \quad \text{DARE}_{BOA} = F_{BOA}^{\downarrow} - F_{BOA}^{\downarrow 0} \quad (9)$$

261  
 262 Where  $F$  and  $F^0$  denoted the broadband fluxes including and excluding

263 aerosols, respectively at the BOA and TOA. The “ $\uparrow$ ” and “ $\downarrow$ ” mean the upward  
264 fluxes and downward fluxes, respectively.

265 In the radiative transfer module, the absorption and multiple scattering  
266 effects are taken into account during flux calculations by the Discrete  
267 Ordinates (DISORT) approach (Nakajima and Tanaka, 1988; Stamnes et al.,  
268 1988). The gaseous distributions and single fixed aerosol vertical distribution  
269 (exponential to 1 km) from the multi-layered US standard 1976 atmosphere  
270 were used in the radiative flux calculations (García et al., 2008). García et al.  
271 (2008) pointed out that the error for the observed solar radiation at the surface  
272 in global was  $+2.1 \pm 3.0\%$  for an overestimation of about  $+9 \pm 12 \text{ Wm}^{-2}$ . The  
273 data used in preparing the figures for the present paper have been made  
274 available as an Appendix.

275

### 276 **3. Results and discussion**

#### 277 **3.1 Spatial distribution of aerosol microphysical properties**

278 A map showing the 50 CARSNET sampling sites and plots of the aerosol  
279 volume size distributions ( $dV/d\ln r$ ) at each of the sites is shown in Fig. 1.  
280 Generally, the annual mean effective radius of total particles ( $R_{\text{effT}}$ ) decreased  
281 from the inland northwest to the southeastern coastal areas. Furthermore, the  
282 volume concentration of total particles was found substantially higher at the  
283 urban sites. The volume of the coarse mode particles was considerably larger  
284 than that of the fine mode particles at the remotes, arid/semi-arid sites and at  
285 those sites on the CLP-Chinese Loess Plateau or nearby, indicating that those  
286 areas were most strongly affected by larger particles, most likely mineral dust  
287 as discussed below.

288 The average (arithmetic mean)  $R_{\text{effT}}$  at the remote sites was about  $0.47$   
289  $\mu\text{m}$  with the volume about  $0.05 \mu\text{m}^3/\mu\text{m}^2$  (Table 1). A large  $R_{\text{effT}}$  ( $0.64 \mu\text{m}$ ) was  
290 found at Lhasa, and the total aerosol volume there was  $0.05 \mu\text{m}^3/\mu\text{m}^2$ . These  
291 results are consistent with those reports by Li et al. (2018) who found high  
292 levels of coarse mode particles at Lhasa due to the presence of mineral dust.

293 The two other remote sites, Akedala and Shangri-La, had smaller average  $R_{\text{effT}}$   
294 values than Lhasa (0.36 and 0.39  $\mu\text{m}$ , respectively), and corresponding  
295 volumes were 0.06 and 0.03  $\mu\text{m}^3/\mu\text{m}^2$ . The average fine-mode effective radius  
296 ( $R_{\text{effF}}$ ) was 0.14  $\mu\text{m}$  at the remote sites, and fine-mode particle fractional  
297 volume ( $PV_{\text{F}}$ ) was 0.01  $\mu\text{m}^3/\mu\text{m}^2$ , while the average coarse-mode effective  
298 radii ( $R_{\text{effC}}$ ) was 2.35  $\mu\text{m}$  and the coarse-mode fractional volume ( $PV_{\text{C}}$ ) was  
299 0.03  $\mu\text{m}^3/\mu\text{m}^2$ . These findings indicated that the contribution of coarse-mode  
300 particles to the total volume of aerosol was larger at the remote sites. A study  
301 by Cong et al. (2009) at the remote Nam Co site on the Tibetan Plateau  
302 showed that dust particles mainly affected the site in spring, while  
303 anthropogenic aerosols were prevalent in the summer.

304 The average  $R_{\text{effT}}$  at the arid and semi-arid sites (0.55  $\mu\text{m}$ ) was larger than  
305 at the remote sites, and the total volume of aerosols at the arid/semi-arid sites  
306 also was large (0.14  $\mu\text{m}^3/\mu\text{m}^2$ ), nearly three times that at the remote sites.  
307 Large  $R_{\text{effT}}$  values (0.71  $\mu\text{m}$ ) were found at Tazhong, which is near the  
308 northwestern deserts, and the aerosol volume there also was high to 0.30  
309  $\mu\text{m}^3/\mu\text{m}^2$ . Large  $PV_{\text{C}}$ 's were found at the arid/semi-arid sites (0.05–0.27  
310  $\mu\text{m}^3/\mu\text{m}^2$ ). The arithmetic mean  $R_{\text{effT}}$  (0.49  $\mu\text{m}$ ) at the rural sites on or near the  
311 CLP had total aerosol volumes (0.15  $\mu\text{m}^3/\mu\text{m}^2$ ) similar to those at the  
312 arid/semi-arid sites. These results also show a major contribution to the  
313 aerosol volumes by coarse-mode particles at the sites in or near the mineral  
314 dust source regions. Bi et al. (2011) similarly found that coarse particles  
315 dominated the volume-size distribution at the Semi-Arid Climate and  
316 Environment Observatory of Lanzhou University (SACOL) on the CLP.

317 Small  $R_{\text{effT}}$  values (0.33  $\mu\text{m}$ ) were found at the rural sites in eastern China,  
318 and relatively high aerosol volumes were observed there (0.18  $\mu\text{m}^3/\mu\text{m}^2$ ). In  
319 the Yangtze River Delta (YRD) region, the  $R_{\text{effF}}$  was large range for 0.16–0.17  
320  $\mu\text{m}$ , and the  $PV_{\text{F}}$ 's were 0.12–0.13  $\mu\text{m}^3/\mu\text{m}^2$ . At the Mt. Longfeng background  
321 site in northeastern China, the total particle volume was low (0.08  $\mu\text{m}^3/\mu\text{m}^2$ ),  
322 which is consistent with minimal anthropogenic influences and low aerosol

323 loadings. Compared with the other sites, the urban areas had relatively low  
324 coarse mode aerosol concentrations, but small particles were plentiful—the  
325 average  $R_{\text{effT}}$  was  $0.37 \mu\text{m}$  and total volume was high at  $0.21 \mu\text{m}^3/\mu\text{m}^2$ . The  
326 average  $R_{\text{effF}}$  of fine-mode particles at the urban sites was  $0.16 \mu\text{m}$  with a  $PV_{\text{F}}$   
327 of  $0.10 \mu\text{m}^3/\mu\text{m}^2$  while the  $R_{\text{effC}}$  was  $2.22 \mu\text{m}$  and  $PV_{\text{C}}$  was  $0.11 \mu\text{m}^3/\mu\text{m}^2$ .

328 The effective radii and  $PV_{\text{F}}$  values showed strong relationships with  
329 population density and vehicle emissions at the urban sites. High volumes of  
330 fine mode particles occurred at the northeastern urban site of Shenyang ( $R_{\text{effT}}$   
331 =  $0.16 \mu\text{m}$ ,  $PV_{\text{F}}$  =  $0.12 \mu\text{m}^3/\mu\text{m}^2$ ); at major cities in northern China, including  
332 Shijiazhuang ( $R_{\text{effT}}$  =  $0.16 \mu\text{m}$ ,  $PV_{\text{F}}$  =  $0.12 \mu\text{m}^3/\mu\text{m}^2$ ) and Zhengzhou ( $R_{\text{effT}}$  =  
333  $0.18 \mu\text{m}$ ,  $PV_{\text{F}}$  =  $0.12 \mu\text{m}^3/\mu\text{m}^2$ ); at Chengdu, a city in the Sichuan Basin of  
334 ( $R_{\text{effT}}$  =  $0.21 \mu\text{m}$ ,  $PV_{\text{F}}$  =  $0.16 \mu\text{m}^3/\mu\text{m}^2$ ); and at the urban regions of Nanning  
335 ( $R_{\text{effT}}$  =  $0.18 \mu\text{m}$ ,  $PV_{\text{F}}$  =  $0.13 \mu\text{m}^3/\mu\text{m}^2$ ) and Panyu ( $R_{\text{effT}}$  =  $0.16 \mu\text{m}$ ,  $PV_{\text{F}}$  =  $0.10$   
336  $\mu\text{m}^3/\mu\text{m}^2$ ) in southern China. Overall, these results show that the volumes of  
337 fine-mode particles increased at the urban sites where anthropogenic  
338 influences were most apparent.

339 Cheng et al. (2015) found different aerosol volume size distributions for  
340 dust and sea salt at Shanghai in the eastern China, and they showed that their  
341 relative abundances varied with season and in response to local or long-range  
342 transport. Zhao et al. (2018) also reported the effect of sea salt aerosol on the  
343 aerosol absorption and radiative effects in the coastal region over northeastern  
344 China. Especially the particles hygroscopic growth with different composition  
345 observed in special climatic conditions could affect aerosol microphysical  
346 properties by geographically variable effects (Zhang et al., 2015; Sun et al.,  
347 2010). Like in the YRD region, hygroscopic growth of fine-mode particles could  
348 lead to larger AOD and scattering enhancing reported by Sun et al. (2018) and  
349 Che et al. (2018). Xia et al. (2019) observed the aerosol hygroscopic growth on  
350 fine particle scattering coefficient in Beijing.

351

### 352 **3.2 Spatial distributions of AOD and EAE**

353 The spatial distributions of  $AOD_{440\text{ nm}}$  and  $EAE_{440-870\text{ nm}}$  are shown in Fig. 2.  
354 The  $AOD_{440\text{ nm}}$  increased from the remote/rural sites to the urban sites, and as  
355 one might expect, the remote sites were the least affected by particle  
356 emissions and had the lowest aerosol loadings. For example, the  $AOD_{440\text{ nm}}$  at  
357 the remote stations was low and had an average value of 0.12. The Lhasa and  
358 Shangri-La sites on the Tibetan Plateau had similar average  $AOD_{440\text{ nm}}$  values  
359 of 0.10. These phenomenons are similar to the study of Li et al. (2018), who  
360 showed clean air conditions at Lhasa with  $AOD < 0.1$ . Cong et al. (2007, 2009)  
361 also found a low AOD (0.05) at Nam Co, which was comparable to the  
362 background levels at other remote sites.

363 The  $AOD_{440\text{ nm}}$ 's at the arid/semi-arid sites and those on or near the Loess  
364 Plateau ranged from 0.32–0.42, which is higher than at the remote sites. The  
365 high  $AOD_{440\text{ nm}}$  at Tazhong (0.60), which is near the deserts in northwestern  
366 China was likely due to the large aerosol volume of  $0.30\ \mu\text{m}^3/\mu\text{m}^2$  (section 3.1)  
367 caused by mineral dust. Indeed, arid and semi-arid regions in northwestern  
368 China are important sources of aeolian dust on a global scale (Bi et al., 2011).  
369 Li et al. (2012) showed that the contribution of dust to the average AOD at  
370 SACOL near Lanzhou was 28.4%. Other sites that showed large  $AOD_{440\text{ nm}}$   
371 include regions with strong anthropogenic influences, such as Dengfeng (0.79)  
372 on the North China Plain, Huimin (0.83) in the YRD (0.83 to 0.87) and Huainan  
373 (0.91) in the Guanzhong Plain.

374 Compared with the sites just discussed, lower  $AOD_{440\text{ nm}}$ 's were found at  
375 the Mt. Longfeng background station of the Northeast China Plain (0.34), the  
376 semi-arid rural site as Tongyu in northeastern China (0.23), and the clean  
377 Xiyong site in southern China (0.41). Zhu et al. (2014) found a low AOD of 0.28  
378 at the North China Plain regional background site. Che et al. (2009c) have  
379 pointed out that the large AOD at Lin'an was likely affected by the high aerosol  
380 loadings in YRD Region. Among the urban sites in China, large  $AOD_{440\text{ nm}}$ 's  
381 were found in the cities with strong influences of anthropogenic activities, such  
382 as the Northeastern Plain (Shenyang 0.89), North China Plain (Zhengzhou

383 0.99), Central China (Wuhan 1.00) and Sichuan Basin (Chengdu 1.17); the  
384 average value for these sites was 0.79. Lower AOD<sub>440nm</sub>'s, that is < 0.50,  
385 occurred at remote sites in northwestern China, including Urumqi (0.42),  
386 Yinchuan (0.37); those sites are affected less by industrial activities and the  
387 population densities are lower compared with the sites in northern or eastern  
388 China.

389 It is worth noting that the particle emissions in or around the urban sites  
390 could lead to large optical extinctions due to hygroscopic aerosol growth,  
391 especially in summer when the relative humidity is often high. In a related  
392 study, Zhang et al. (2018) found a large AOD of 1.10 at Wuhan in central China  
393 and that was linked to secondary aerosol formation under the high  
394 summertime temperatures. Li et al. (2015) similarly concluded that high  
395 temperature and humidity promoted the formation of fine particles and led to  
396 hygroscopic aerosol growth at Nanjing. Qin et al. (2017) observed a high  
397 AOD<sub>500 nm</sub> of 1.04 at Shijiazhuang and related this to the hygroscopic growth of  
398 aerosol fine-mode particles during polluted days.

399 Clear spatial variability in EAE values over China is evident in Fig. 3, and  
400 at the remote sites, the average EAEs were 1.03. The EAE at Lhasa (0.77)  
401 was lower than at Akedala (EAE = 1.13), which is in an arid region of central  
402 Asia, or at Shangri-La (EAE = 1.19) in Tibet. The average coarse-mode  
403 average effective radii ( $R_{\text{effC}}$ ) at Lhasa was 2.26  $\mu\text{m}$  and the fractional volume  
404 was 0.04  $\mu\text{m}^3/\mu\text{m}^2$ , this result suggests the major components of the large  
405 mineral dust particles in aerosol populations over that region. The smaller  
406 sphericity fraction (~42.70) and lower FMF (0.66) at Lhasa indications the  
407 presence of non-spherical aerosol coarse particles compared with the  
408 spherical fine particles in the urban sites.

409 At the arid and semi-arid sites in China, the average EAE value (0.71) was  
410 relatively low and the FMF also was low (0.58). The EAE was extremely low at  
411 Tazhong (0.25), which is in the Takliman Desert in the Xinjiang Uygur  
412 Autonomous Region of northwestern China and the sphericity fraction (12.87)

413 and FMF (0.35) there were lower compared with most of the other sites. This  
414 finding indicates a strong contribution of large particles in this desert region  
415 consistent with large volume of the coarse-mode particles ( $0.27 \mu\text{m}^3/\mu\text{m}^2$ )  
416 noted in section 3.1. The average EAE reached 0.93 at the rural sites near the  
417 CLP, and the average value of FMF for those sites was 0.73. Eck et al. (2005)  
418 found especially low EAE values in March and April (0.3 and 0.4, respectively)  
419 at Yulin, China, where the dust aerosol dominated the optical column.

420 Large EAEs (1.23) were found at the sites in eastern China, and the FMFs  
421 also were large (0.89) at those sites. This result can be attributed to the strong  
422 impacts of anthropogenic in the more urbanized eastern part of the country. On  
423 the other hand, large EAE values also occurred at the clean sites in  
424 northeastern China, including Mt. Longfeng (1.38), where the sphericity  
425 fraction was 58.5 and the FMF 0.90. This shows that small particles can have  
426 stronger effects in these areas relative to some other regions of China. The  
427 EAE at Lin'an was larger than that at Shangdianzi in the Northern Plain or  
428 Longfengshan in Northeastern China for most months according to data from  
429 Che et al. (2009c). At the urban sites, large EAEs were found at sites in  
430 southern China, including Nanning (EAE = 1.36, sphericity fraction = 70.12,  
431 FMF = 0.95), Panyu (EAE = 1.43, sphericity fraction = 75.55, FMF = 0.93) and  
432 Zhuzilin (EAE = 1.45, sphericity fraction = 55.51, FMF = 0.94). This is likely  
433 because the large populations and widespread vehicle ownership in those  
434 cities led to the dominance of fine-mode particles throughout the year. Cheng  
435 et al. (2015) found a uni-modal distribution of EAE centered in 1.1–1.6 with the  
436 occurrence frequency about 72%, which indicated an abundance of fine  
437 primary particles at Shanghai in eastern China. At the urban Nanjing site,  
438 which is in east-central China, small particles were dominant, and the annual  
439 average EAE was  $1.21 \pm 0.28$  (Li et al., 2015).

440

### 441 **3.3 Spatial distribution of aerosol single-scattering albedo**

442 The spatial distribution of SSA at 440 nm of the 50 CARSNET stations is

443 shown in Fig. 4. As a frame of reference, Eck et al. (2005) reported that that  
444  $SSA_{440nm}$  from the AERONET retrievals were 0.82 to 0.98 globally. We note  
445 that  $SSA_{440nm}$  values in this range reflect slightly to strongly absorbing aerosols,  
446 and these particles originate from multitude sources (Che et al., 2018). The  
447  $SSA_{440nm}$ 's decreased from remote/rural to the urban sites and from west to  
448 east, which means that there were higher percentages of absorbing particles  
449 at the urban and eastern stations. The average  $SSA_{440nm}$  at the remote sites  
450 was about 0.91, which is indicative of particles with moderate absorption. The  
451 absorbing aerosols at the remote sites were more likely mineral dust particles  
452 because those sites are less likely to be affected by carbonaceous particles,  
453 which also are absorbing, but mainly produced by anthropogenic activities.  
454 The  $SSA_{440nm}$ 's for the arid and semi-arid sites were 0.89. The relatively high  
455 SSA at Tazhong (0.92) was probably due to slightly absorbing, coarse mode  
456 dust particles (EAE = 0.25).

457 A study by Bi et al. (2011) showed that SSAs increased slightly with  
458 wavelength when dust was present at the SACOL site. Moderately absorbing  
459 particles were found in our study on or near the Chinese Loess Plateau where  
460 the  $SSA_{440nm}$ 's were typically 0.88 to 0.89. Eck et al. (2005) concluded that the  
461 spectral SSA demonstrated effects of dust at Yulin because the SSA increased  
462 for wavelengths from 440 to 675 nm. At the rural sites in eastern China, large  
463  $SSA_{440nm}$ 's mainly occurred at sites in the YRD affected anthropogenic  
464 influences; these include Tonglu (0.93), Xiaoshan (0.93), Xiyong (0.94). Che et  
465 al. (2018) found the slightly absorbing particles came from industrial activity  
466 and anthropogenic sources at YRD region with the  $SSA_{440nm}$  between 0.91 to  
467 0.94.

468 The average value of  $SSA_{440nm}$  at the urban sites was 0.90, which  
469 indicates that particles with moderate absorption dominated the aerosol  
470 populations. Cheng et al. (2015) reported a seasonal range of SSA from 0.88  
471 to 0.91 at Shanghai, with higher values in autumn and winter compared with  
472 spring and summer. Lower  $SSA_{440nm}$ 's occurred at the urban sites and



473 industrial regions in northeastern China, such as Shenyang (0.84), Anshan  
474 (0.89), Fuhsun (0.84), which indicates that the particles were more strongly  
475 absorbing in that region. On the other hand, higher  $SSA_{440nm}$ 's were found at  
476 urban sites in southern China, including Nanning (0.92), Panyu (0.90) and  
477 Zhuzilin (0.96), and this indicates that the particles at those sites were slightly  
478 or weakly absorbing.

479 Moreover, we found that the  $SSA_{440nm}$  spatial distribution reflected the  
480 percentages of absorbing aerosols at the urban sites both in northern and  
481 eastern China. The reports of Dubovik et al. (2000, 2002, 2006) showed that  
482 SSA values vary with both particle size and composition, and Su et al. (2017)  
483 used the variations in SSA with wavelength to indicate the presence of brown  
484 carbon aerosols at Tianjin, a coastal megacity in China. Qin et al. (2017)  
485 suggested that the small SSAs found at Shijiazhuang indicated the presence  
486 of fine-mode absorbing particles, such as brown carbon. Zhuang et al. (2014)  
487 reported that the SSA at the Nanjing urban site ranged from 0.90 to 0.95, and  
488 the aerosol was more absorbing in autumn, possibly due to the biomass  
489 burning emission in the YRD. As evident in the results presented in section 3.1,  
490 one can see that the  $R_{effT}$ ,  $R_{effF}$  and  $R_{effC}$  between northeastern and southern  
491 China was very similar. For example, at Shenyang, a megacity in northeastern  
492 China, the effective radii of total, fine- and coarse-mode particles were 0.31,  
493 0.16, 2.23  $\mu m$  and the corresponding volumes were 0.22, 0.12, 0.10  $\mu m^3/\mu m^2$ ,  
494 respectively. At Hangzhou in the YRD region, the  $R_{effT}$ ,  $R_{effF}$  and  $R_{effC}$  were 0.30,  
495 0.17, 2.21  $\mu m$  with the volumes about 0.22, 0.12, 0.10  $\mu m^3/\mu m^2$ , respectively.  
496 Therefore, the different  $SSA_{440nm}$  distributions in the two regions may be  
497 attributed by the special aerosol composition related to the urban-industrial  
498 background of northeastern China (lower  $SSA_{440nm}$ ) and more  
499 anthropogenic sources in the eastern China environmental (higher  
500  $SSA_{440nm}$ ).

501 Dust aerosols with light-absorbing occur more frequently in spring in  
502 northeastern China than in more southern regions (Zhao et al., 2018).

503 Anthropogenic emissions from seasonal biomass burning and residential  
504 heating are two other main factors that affect aerosol composition between the  
505 two regions (Che et al., 2018). Especially in winter, there was high percentage  
506 of absorbing aerosols at the northeastern sites, and that was more than likely  
507 caused by emissions of carbonaceous aerosol from residential heating (Zhao  
508 et al., 2015). Climatic conditions are also the main factors affecting the  
509 absorption characteristics of aerosols in different regions of north and south  
510 China. The increased light scattering could well be due to the particles  
511 hygroscopic growth demonstrated in other studies. For example, Mai et al.  
512 (2018) found that AODs and SSAs both increased with relative humidity at  
513 Guangdong in the PRD region, which suggests that condensational growth  
514 can affect the aerosol optical properties.

515

### 516 **3.4 Spatial distributions of absorption aerosol optical depth (AAOD)**

517 The spatial distribution of AAOD at 440 nm shown as Fig. 5 indicates that  
518 overall, the  $AAOD_{440nm}$ 's increased from north to south and from remote/rural  
519 to urban sites. Lower  $AAOD_{440nm}$ 's were found at the remote stations, where  
520 the average value was 0.01. The  $AAOD_{440nm}$  at Akedala, a remote site in  
521 northwestern China, was 0.02, and that was higher than at Shangri-La or  
522 Lhasa (0.01), both of which are on the Tibetan Plateau. The low  $AAOD_{440nm}$ 's  
523 throughout that region indicates that the aerosol population was not strongly  
524 absorbing. Compared with these three sites, the average  $AAOD_{440nm}$ 's at the  
525 arid and semi-arid sites were higher (0.03); for example, an  $AAOD_{440nm}$  of  
526 0.05 was found at Tazhong, which is adjacent to the desert, and that indicates  
527 that the aerosol particles were more absorbing. As discussed in sections 3.2  
528 and 3.3, dust aerosols likely make a significant contribution to aerosol light  
529 absorption in the areas impacted by the deserts.

530 The low  $AAOD_{440nm}$  found at Xilinhot (0.02) was probably due to the low  
531 aerosol loadings ( $AOD_{440nm} = 0.21$ ) in this region. The  $AAOD_{440nm}$ 's at the Mt.  
532 Gaolan and Yulin rural sites which on or around the CLP were about 0.04 and

533 0.03, respectively, and the particles were moderately absorbing ( $SSA = 0.89$ ).  
534 The large  $AAOD_{440nm}$  at Datong (0.09) can be explained by the high  $AOD_{440nm}$   
535 (0.58) there. Indeed, large  $AAOD_{440nm}$ 's were found at rural sites in eastern  
536 China, where there were high AODs and low SSAs as noted in sections 3.2  
537 and 3.3. Of these sites, Dengfeng ( $AOD_{440nm} = 0.08$ ) and Huimin ( $AOD_{440nm} =$   
538  $0.08$ ) are located on the North China Plain, while Huainan ( $AOD_{440nm} = 0.10$ ) is  
539 on the Guanzhong Plain. Lower  $AAOD_{440nm}$ 's, from 0.02–0.03, occurred at  
540 Tongyu (0.03), which is at a semi-arid region in northeastern China, at the Mt.  
541 Longfeng (0.03) regional background site on the Northeast China Plain, at the  
542 Yushe rural site in northern China (0.03), and at the clean Xiyong site in the  
543 PRD (0.02).

544 Several urban sites showed  $AAOD_{440nm}$  values greater than 0.10; these  
545 include Fushun (0.11) and Shenyang (0.14) in the northeastern China,  
546 Lanzhou (0.10) in the northwestern China, and Nanjing (0.10) and Wuhan  
547 (0.11) in the eastern and central China. Lower  $AAOD_{440nm}$ 's occurred in other  
548 urban areas, such as Yinchuan ( $AAOD_{440nm} = 0.02$ ,  $AOD_{440nm} = 0.37$ ) in the  
549 northwest and Zhuzilin ( $AAOD_{440nm} = 0.03$ ,  $AOD_{440nm} = 0.66$ ) in the PRD; both  
550 of these sites had relatively low  $AOD_{440}$ 's indicating weaker anthropogenic  
551 influences compared with metropolitan regions of some other areas. We note  
552 that there are significant uncertainties in relating aerosol absorbing properties  
553 to particle types, such as black carbon, organic matter, as well as mineral dust  
554 (Russell et al., 2010; Giles et al., 2012). Nonetheless, the information  
555 presented here on the spatial distribution of AAODs over China may be useful  
556 for the further investigations into the relationships between light absorption and  
557 particle type (Liu et al., 2017; Schuster et al., 2016a, 2016b).

558

### 559 **3.5 Spatial distribution of direct aerosol radiative effect at the Earth's** 560 **surface and top of the atmosphere**

561 The spatial distributions of the DAREs calculated for both the bottom and  
562 top of the atmosphere are shown in Fig. 6. Overall, the DARE-BOAs increased

563 from northwest to southeast and from rural to urban sites, consistent with  
564 impacts from the densely populated regions around the sites. The average  
565 DARE-BOA at the remote sites was  $-24.40 \text{ W/m}^2$ , and in comparison, a higher  
566 DARE-BOA ( $-33.65 \text{ W/m}^2$ ) occurred at Akedala, which occurred on a remote  
567 region of northwestern China. The  $\text{AOD}_{440\text{nm}}$  at Akedala was relatively low  
568 (0.17) and the SSA moderate (0.90). The moderate absorption of aerosol could  
569 lead to more strong surface cooling effects with little higher DARE-BOA than  
570 the other remote sites. The DARE-BOAs for Lhasa and Shangri-La were  
571  $-22.13$  and  $-17.43 \text{ W/m}^2$ , respectively. These results indicate weaker surface  
572 cooling effects at the remote sites relative to other regions because the aerosol  
573 loadings were relatively low, as indicated by  $\text{AOD}_{440\text{nm}}'s < 0.20$ .

574 The average DARE-BOTs at the arid and semi-arid sites of China were  
575 about  $-56.43 \text{ W/m}^2$ , and those high DARE-BOAs can be explained by the  
576 moderately absorbing particles ( $\text{SSA} = 0.89$ ) and large  $\text{AOD}_{440\text{nm}}'s$  (0.32)  
577 compared with the remote sites. A large DARE-BOA ( $-91.20 \text{ W/m}^2$ ) occurred at  
578 the Tazhong site near the northwestern deserts, and there, the high AOD (0.60)  
579 and the slight absorption of mineral dust ( $\text{SSA} = 0.92$ ) imply substantial surface  
580 cooling. The average DARE-BOA for rural sites on the Chinese Loess Plateau  
581 or surrounding was  $-74.67 \text{ W/m}^2$ , and that also implies cooling at the surface.

582 Several rural sites in northern and eastern China had large DARE-BOA  
583 values; these include Huimin ( $-111.58 \text{ W/m}^2$ ), Dengfeng ( $-104.78 \text{ W/m}^2$ ) and  
584 Huainan ( $-129.17 \text{ W/m}^2$ ), and at those sites the AODs were high, from 0.80–  
585 0.90, and the SSAs were  $\sim 0.89$ . These results show stronger surface cooling  
586 effects at sites influenced by anthropogenic emissions compared with the  
587 remote sites or those near the deserts. The large negative DARE-BOA values  
588 ( $-103.28 \text{ W/m}^2$ ) at the urban sites indicate that the combination of high  
589  $\text{AOD}_{440\text{nm}}'s$  (0.79) and moderate SSAs (0.90) can cause significant surface  
590 cooling. Indeed, anthropogenic emissions presumably led to the high  
591 DARE-BOAs at urban sites, including Shenyang ( $-144.88 \text{ W/m}^2$ ) and Fushun  
592 ( $-116.91 \text{ W/m}^2$ ) in the Northeastern Plain, Xian in the Guanzhong Plain

593 (-132.55 W/m<sup>2</sup>), Chengdu in the Sichuan Basin (-110.42 W/m<sup>2</sup>), Lanzhou in the  
594 western region (-126.17 W/m<sup>2</sup>), and Nanjing (-143.38 W/m<sup>2</sup>) and Wuhan  
595 (-171.80 W/m<sup>2</sup>) in the Central China. These results indicate that anthropogenic  
596 aerosols can cause significant direct radiative effects at urban sites.

597 The DARE-TOAs increased from north to south and from rural to urban  
598 sites, and the average DARE-TOA for the remote stations was low, about -4.79  
599 W/m<sup>2</sup> (Fig. 7). The DARE-TOAs at Lhasa and Shangri-La were -5.04 W/m<sup>2</sup>  
600 and -8.93 W/m<sup>2</sup>, respectively. A notably small DARE-TOA was found at  
601 Akedala (-0.42 W/m<sup>2</sup>), indicating that the effects of the aerosol on the  
602 temperature of earth-atmosphere system there would be weak. The average  
603 DARE-TOA at the arid and semi-arid sites was -10.17 W/m<sup>2</sup>. The large  
604 DARE-TOA found at Tazhong (-23.49 W/m<sup>2</sup>) could represent the larger  
605 contribution of slightly absorbing mineral aerosols (SSA = 0.92) and a large  
606 AOD (0.60); this indicates more cooling at surface through the absorption and  
607 scattering solar radiation compared with the less impacted sites. This is  
608 consistent with the results for Tazhong discussed in section 3.1 which showed  
609 high volumes of coarse mode particles with large radii.

610 The average DARE-TOA at rural sites on the Chinese Loess Plateau or  
611 nearby was about -14.56 W/m<sup>2</sup>. Although the SSA<sub>440nm</sub> were close to Gaolan  
612 and Yulin about 0.89, the TOAs were quite different (Mt. Gaolan -20.87 W/m<sup>2</sup>;  
613 Yulin -9.09 W/m<sup>2</sup>) which could be due to the different AOD<sub>440nm</sub> about 0.36 and  
614 0.32, respectively. In rural eastern China, the DARE-TOA was about -32.40  
615 W/m<sup>2</sup>, and to put this in context, Che et al. (2018) found that DARE-TOAs of  
616 -40 W/m<sup>2</sup> at rural sites in the YRD region, which is indicative of a relatively  
617 strong cooling effect. Low DARE-TOAs were found at the Mt. Longfeng rural  
618 site in northeastern China (DARE-TOA = -11.34, AOD<sub>440nm</sub> = 0.34, SSA = 0.89)  
619 and at the Tongyu semi-arid site in northeastern China as (DARE-TOA = -8.87,  
620 AOD<sub>440nm</sub> = 0.23, SSA = 0.88 ) where the aerosol loadings were relatively low  
621 and the absorption was moderate.

622 In the urban sites at central and eastern China, the average DARE-TOA

623 values were about  $-30.05 \text{ W/m}^2$ . Higher DARE-TOAs occurred at Anshan in  
624 the Northeastern Plain ( $-39.66 \text{ W/m}^2$ ), Chengdu in the Sichuan Basin ( $-52.21$   
625  $\text{W/m}^2$ ), Hangzhou in the YRD ( $-40.16 \text{ W/m}^2$ ), Jiaozuo ( $-39.35 \text{ W/m}^2$ ) and  
626 Zhengzhou ( $-46.18 \text{ W/m}^2$ ) in the North China Plain, and Zhuzilin ( $-40.15 \text{ W/m}^2$ )  
627 in the PRD region. The high DARE-TOA values at these urban sites imply  
628 relatively strong cooling effects due to higher aerosol loadings in the  
629 atmosphere.

630

### 631 **3.6 Spatial distributions of aerosol mixing properties**

632 The spatial distribution of aerosol mixing properties (Figure 8) was  
633 obtained by using the  $\text{SSA}_{440\text{nm}}$ , FMF, and EAE results to classify the particles  
634 based on size and absorbing properties. In previous studies by Zheng et al.  
635 (2017) and Che et al. (2018), the particles in this study were grouped into eight  
636 types as Table 2 show. Moreover, the FMF has been provided to give the  
637 particle size information in the group of the particles.

638 At the remote Akedala and Lhasa sites ( $\text{FMF} = 0.70\text{--}0.78$  and  $\text{SSA}_{440\text{nm}} =$   
639  $0.85$ ), the percentages of mixed absorbing particles (Type V) were 35-40%,  
640 while at Shangri-la ( $\text{FMF} = 0.76$ ,  $\text{SSA}_{440\text{nm}} = 0.84$ ), the percentage was slightly  
641 lower, 24.62%. The characteristics of the particles at these remote,  
642 high-altitude sites were probably affected by the rugged topography which  
643 would promote particle mixing. The proportion of coarse mode, mainly dust,  
644 particles with moderate to strong absorption (Group VII) was highest at the arid  
645 and semi-arid sites. The percent abundances of Group VII particles were 57.90%  
646 at Dunhuang ( $\text{AE} = 0.26$ ,  $\text{SSA}_{440\text{nm}} = 0.85$ ,  $\text{FMF} = 0.43$ ) and 58.52% at  
647 Tazhong ( $\text{AE} = 0.20$ ,  $\text{SSA}_{440\text{nm}} = 0.87$ ,  $\text{FMF} = 0.37$ ), respectively. Mixed  
648 absorbing particles (Type V) and strongly absorbing dust particles (Group VII)  
649 accounted for 30 to 70% of the aerosol in the rural sites on or near the CLP.  
650 The percentages of mixed absorbing particles (Type V) at Gaolan, Yulin, and  
651 Datong were 31.98%, 45.22% and 29.04%, respectively, and the average  
652 FMFs at those sites ranged from 0.70–0.76.

653 The proportions of the coarse-mode aerosols with strongly absorbing in  
654 Group VII were about 35.23% at Gaolan and 21.21% at Yulin, which was  
655 mainly dust particles with the FMFs at those sites were 0.43 and 0.48,  
656 respectively. The proportion of coarse-mode particles with strongly absorbing  
657 in Group VII and coarse-mode particles with weakly-absorbing in Group VIII at  
658 the rural sites in eastern China were < 11%. These patterns indicated that the  
659 differences in the eastern region from northwestern China because in the east,  
660 coarse-mode particles have only a minor contribution to aerosol absorption.  
661 The percentage of fine-mode particles with weakly-absorbing in Type IV and  
662 mixed absorbing particles in Type V combined about ~50% at the eastern sites.  
663 This result suggests that mixed aerosols originated from a variety of sources  
664 and that many of the sites were affected by anthropogenic emissions from  
665 megacities upwind.

666 The fine-mode particles with absorbing in Types I, II, III and V accounted  
667 for 50 to 90% at most of the urban sites. The percentages of these four particle  
668 types combined were especially large in eastern China; for example, at Panyu,  
669 particle Types I–IV composed 90.83% of the total, and the FMF there was  
670 0.90–0.94, while at Zhuzilin, the percentage of Types I–IV was 92.55%, and  
671 the FMF was 0.92–0.94. These results are another indication that fine-mode  
672 particles are important for light absorption in urban areas. In contrast, the  
673 Lanzhou and Urumqi urban sites were less affected by absorbing fine-particles  
674 because the percentages of Type I–IV particles were only 19.73% and 18.36%,  
675 respectively. The mixed absorbing Type V particles accounted for large  
676 percentages of the total at Lanzhou (48.80%, EAE = 0.88, SSA = 0.82, FMF =  
677 0.73) and at Urumqi (59.39%, EAE = 0.94, SSA = 0.84, FMF = 0.75). Different  
678 from the other urban sites, these patterns show that larger particles had  
679 significant contributions to the aerosol absorption at these two northwestern  
680 sites.

681

#### 682 **4. Conclusions**

683 Aerosol microphysical and its optical properties obtained from the  
684 ground-based sunphotometer deployed at 50 CARSNET stations were used to  
685 begin the development of their climatology characteristics and to investigate  
686 potential aerosol-climate effects over vast area of China. Direct aerosol  
687 radiative effects (DAREs) at the bottom and at the top of the atmosphere were  
688 calculated, and eight types of aerosols were classified based on the particle  
689 size and absorbing properties. The annual mean values of the  $ReffT$   
690 decreased from the arid and semi-arid sites ( $0.55 \mu\text{m}$ ) to the urban sites ( $0.37$   
691  $\mu\text{m}$ ). The aerosol volumes increased from the remote sites ( $0.05 \mu\text{m}^3/\mu\text{m}^2$ ) to  
692 the urban sites ( $0.21 \mu\text{m}^3/\mu\text{m}^2$ ). The volumes of coarse-mode particle were  
693 larger than those for the fine mode at the remote and arid/semi-arid sites—this  
694 can be explained by the greater relative abundances of mineral dust compared  
695 with pollution-derived particles at those sites. At the urban sites, where  
696 anthropogenic influences were relatively strong, the proportion of fine mode  
697 particles increased gradually with aerosol volume.

698 The  $AOD_{440\text{nm}}$  progressively increased from the remote sites (0.12) to the  
699 arid and semi-arid sites (0.32) to rural sites in eastern China (0.70) and finally  
700 to the urban sites (0.79), which were the ones most strongly affected by  
701 anthropogenic activities. The average  $EAE_{440-870 \text{ nm}}$ 's at the arid and semi-arid  
702 sites were relatively low (0.71), which indicates an important contribution of  
703 larger particles to the aerosol extinction in those regions. The consistently  
704 large  $EAE_{440-870 \text{ nm}}$ 's at the urban sites ( $> 1.20$ ) and the high FMFs that those  
705 site (0.88) are the evidence that fine mode particles are prevalent throughout  
706 year. The average  $SSA_{440\text{nm}}$ 's at the remote, rural, and urban sites were  
707 relatively similar, averaging about 0.89, and this indicates the particles were  
708 moderately absorbing.

709 Overall, dust aerosols with light-absorbing in spring and emissions came  
710 from biomass burning and residential heating during the colder months were  
711 the main factors that led to spatial differences in the percentages of absorbing  
712 aerosols over China. The  $AAOD_{440\text{nm}}$ 's increased from the remote sites (0.01)



713 to the arid and semi-arid sites (0.03) to the rural sites of eastern China (0.05)  
714 and finally to the urban sites (0.07). High  $AAOD_{440nm}$ 's were caused by  
715 light-absorbing dust aerosols at the rural sites and by the strong anthropogenic  
716 emissions in the metropolitan areas. The spatial patterns in the absorbing  
717 aerosols were not only affected by the chemical composition of aerosol, but  
718 also by physical effects imposed by topography, weather, and climate.

719 The average DARE-BOA values were  $-24.40 \text{ W/m}^2$  at the remote sites;  
720  $-56.43 \text{ W/m}^2$  at the arid and semi-arid sites;  $-74.67 \text{ W/m}^2$  at the sites on the  
721 CLP or nearby;  $-85.25 \text{ W/m}^2$  at the rural sites in eastern China; and  $-103.28$   
722  $\text{W/m}^2$  at the urban sites. The larger DARE-BOA values at the urban sites imply  
723 stronger cooling effects from anthropogenic emissions compared with those  
724 from mineral dust at the remote sites or those near the desert. Moreover, larger  
725 DARE-TOA's also occurred at the urban sites ( $-30.05 \text{ W/m}^2$ ), which indicates  
726 strong cooling effects due to the large aerosol extinctions between the  
727 earth-atmosphere system displayed the moderate to strong light absorption.  
728 Mixed-absorbing particles were the most abundant aerosol type in the remote  
729 and rural sites on or near the Chinese Loess Plateau and in eastern China.  
730 Mineral dust particles with moderate to strong absorbing were dominant in the  
731 arid and semi-arid sites while absorbing fine-mode particles accounted for 50  
732 to 90% of the aerosol at the most urban sites.

733 The results of the study have considerable value for ground truthing  
734 satellite observations and for validating aerosol models. Moreover, the results  
735 also have provided significant information on aerosol optical and radiative  
736 properties for different types of sites covering a broad expanse of China.  
737 These results also are a major step towards developing a climatology for  
738 aerosol microphysical and optical properties for China and even East Asia.

#### 739 **Data availability:**

740 The detailed CARSNET AOD dataset used in the study can be requested  
741 by contacting the corresponding author.

#### 742 **Competing interests.**

743 The authors declare that they have no conflict of interest.

744 **Author contribution:**

745 All authors contributed to shaping up the ideas and reviewing the paper.  
746 HC, XX and XZ designed and implemented the research, as well as prepared  
747 the manuscript; HC, HZ, YW and HW contributed to analysis of the CARSNET  
748 dataset; HC, XX, JZ, OD, BNH, PG, and ECA contributed to the CARSNET  
749 data retrieval; HC, BQ, WG, HY, RZ, LY, JC, YZ, KG, and XZ carried out the  
750 CARSNET observations; OD, BNH, PG, and ECA provided constructive  
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762 **References**

763 Bi, J.R., Huang, J.P., Fu, Q., Wang, X., Shi, J.S., Zhang, W., Huang, Z.W., and  
764 Zhang, B.D.: Toward characterization of the aerosol optical properties  
765 over Loess Plateau of Northwestern China, *Journal of Quantitative*  
766 *Spectroscopy and Radiative Transfer*, 112, 346-360,  
767 <https://doi.org/10.1016/j.jqsrt.2010.09.006>, 2011.

768 Bokoye, A.I., Royer, A., O'Neill, N.T., Cliche, P., Fedosejevs, G., Teillet, P.M.,  
769 and McArthur, L.J.B.: Characterization of atmospheric aerosols across  
770 Canada from a Ground-based sun photometer network: Aerocan, *Atmos.*  
771 *Ocean.*, 39, 429–456, <https://doi.org/10.1080/07055900.2001.9649687>,  
772 2001.

773 Breon, F., Tanré, D., and Generoso, S.: Aerosols effect on the cloud droplet  
774 size monitored from satellite, *Science*, 295, 834-838,  
775 <https://doi.org/10.1126/science.1066434>, 2002.

776 Cao, J.J., Lee, S.C., Chow, J.C., Watson, J.G., Ho, K.F., Zhang, R.J., Jin, Z.D.,  
777 Shen, Z.X., Chen, G.C., Kang, Y.M., Zou, S.C., Zhang, L.Z., Qi, S.H., Dai,  
778 M.H, Cheng, Y., and Hu, K.: Spatial and seasonal distributions of  
779 carbonaceous aerosols over China, *J. Geophys. Res.*, 112, D22S11.  
780 <http://dx.doi.org/10.1029/2006JD008205>, 2007.

781 Charlson, R. J., Schwartz, S. E., Hales, J. M., Cess, D., Coakley, J. A., and  
782 Hansen, J. E.: Climate forcing by anthropogenic aerosols, *Science*, 255,  
783 423–430, <https://doi.org/10.1126/science.255.5043.423>, 1992.

784 Che, H.Z., Shi, G.Y., Zhang, X.Y., Arimoto, R., Zhao, J.Q., Xu, L., Wang, B.,  
785 and Chen, Z.H.: Analysis of 40 years of solar radiation data from China,  
786 1961-2000, *Geophysical Research Letters*, 32, L06803,  
787 <https://doi.org/10.1029/2004GL022322>, 2005.

788 Che, H., Shi, G., Uchiyama, A., Yamazaki, A., Chen, H., Goloub, P., and Zhang,  
789 X.: Intercomparison between aerosol optical properties by a PREDE  
790 skyradiometer and CIMEL sunphotometer over Beijing, China, *Atmos.*  
791 *Chem. Phys.*, 8, 3199–3214, <http://dx.doi.org/10.5194/acp-8-3199-2008>,  
792 2008.

793 Che, H., Zhang, X., Chen, H., Damiri, B., Goloub, P., Li, Z., Zhang, X., Wei, Y.,  
794 Zhou, H., Dong, F., Li, D., and Zhou, T.: Instrument calibration and aerosol  
795 optical depth (AOD) validation of the China Aerosol Remote Sensing  
796 Network (CARSNET), *J. Geophys. Res.*, 114, D03206,  
797 <https://doi.org/10.1029/2008JD011030>, 2009a.

798 Che, H. Z., Zhang, X. Y., Alfraro, S., Chatenet, B., Gomes, L., and Zhao, J. Q.:  
799 Aerosol optical properties and its radiative forcing over Yulin, China in  
800 2001 and 2002, *Adv. Atmos. Sci.*, 26, 564–576,  
801 <https://doi.org/10.1007/s00376-009-0564-4>, 2009b.

802 Che, H. Z., Yang, Z. F., Zhang, X. Y., Zhu, C. Z., Ma, Q. L., Zhou, H. G., and

803 Wang, P.: Study on the aerosol optical properties and their relationship  
804 with aerosol chemical compositions over three regional background  
805 stations in China, *Atmospheric Environment*, 43, 1093–1099,  
806 <https://doi.org/10.1016/j.atmosenv.2008.11.010>, 2009c.

807 Che, H., Xia, X., Zhu, J., Li, Z., Dubovik, O., Holben, B., Goloub, P., Chen, H.,  
808 Estelles, V., Cuevas-Agulló, E., Blarel, L., Wang, H., Zhao, H., Zhang, X.,  
809 Wang, Y., Sun, J., Tao, R., Zhang, X., and Shi, G.: Column aerosol optical  
810 properties and aerosol radiative forcing during a serious haze-fog month  
811 over North China Plain in 2013 based on ground-based sunphotometer  
812 measurements, *Atmos. Chem. Phys.*, 14, 2125–2138,  
813 <https://doi.org/10.5194/acp-14-2125-2014>.

814 Che, H., Zhang, X.-Y., Xia, X., Goloub, P., Holben, B., Zhao, H., Wang, Y.,  
815 Zhang, X.-C., Wang, H., Blarel, L., Damiri, B., Zhang, R., Deng, X., Ma, Y.,  
816 Wang, T., Geng, F., Qi, B., Zhu, J., Yu, J., Chen, Q., and Shi, G.:  
817 Ground-based aerosol climatology of China: aerosol optical depths from  
818 the China Aerosol Remote Sensing Network (CARSNET) 2002–2013,  
819 *Atmos. Chem. Phys.*, 15, 7619–7652, [https://doi.org/10.5194/acp-](https://doi.org/10.5194/acp-15-7619-2015)  
820 [15-7619-2015](https://doi.org/10.5194/acp-15-7619-2015), 2015.

821 Che, H., Qi, B., Zhao, H., Xia, X., Eck, T.F., Goloub, P., Dubovik, O., Estelles,  
822 V., Cuevas-Agulló, E., Blarel, L., Wu, Y., Zhu, J., Du, R., Wang, Y., Wang,  
823 H., Gui, K., Yu, J., Zheng, Y., Sun, T., Chen, Q., Shi, G., and Zhang X.:  
824 Aerosol optical properties and direct radiative forcing based on  
825 measurements from the China Aerosol Remote Sensing Network  
826 (CARSNET) in eastern China, *Atmospheric Chemistry and Physics*, 18,  
827 405–425, <https://doi.org/10.5194/acp-18-405-2018>, 2018.

828 Cong, Z.Y., Kang, S.C., Liu, X.D., and Wang, G.F.: Elemental composition of  
829 aerosol in the Nam Co region, Tibetan Plateau, during summer monsoon  
830 season, *Atmospheric Environment*, 41, 1180–1187,  
831 <https://doi.org/10.1016/j.atmosenv.2006.09.046>, 2007.

832 Cong, Z.Y., Kang, S.C., Smirnov, A., and Holben, B.: Aerosol optical properties

833 at Nam Co, a remote site in central Tibetan Plateau, Atmospheric  
834 Research, 92, 42–48, <https://doi.org/10.1016/j.atmosres.2008.08.005>,  
835 2009.

836 Duan, J. and Mao, J.: Study on the distribution and variation trends of  
837 atmospheric aerosol optical depth over the Yangtze River Delta, Acta  
838 Scien. Circum., 27, 537–543,  
839 [https://doi.org/10.1007/978-1-4020-6475-3\\_126](https://doi.org/10.1007/978-1-4020-6475-3_126), 2007.

840 Dubovik, O. and King, M. D.: A flexible inversion algorithm for retrieval of  
841 aerosol optical properties from sun and sky radiance measurements, J.  
842 Geophys. Res., 105, 20673–20696,  
843 <https://doi.org/10.1029/2000JD900282>, 2000.

844 Dubovik, O., Holben, B. N., Eck, T. F., Smirnov, A., Kaufman, Y. J., King, M. D.,  
845 Tanre, D., and Slutsker, I.: Variability of absorption and optical properties  
846 of key aerosol types observed in worldwide locations, J. Atmos. Sci., 59,  
847 590–608,  
848 [https://doi.org/10.1175/1520-0469\(2002\)059<0590:VOAAOP>2.0.CO;2](https://doi.org/10.1175/1520-0469(2002)059<0590:VOAAOP>2.0.CO;2),  
849 2002.

850 Dubovik, O., Sinyuk, A., Lapyonok, T., Holben, B. N., Mishchenko, M., Yang, P.,  
851 Eck, T. F., Volten, H., Munoz, O., Veihelmann, B., van der Zande, W. J.,  
852 Leon, J. F., Sorokin, M., and Slutsker, I.: Application of spheroid models to  
853 account for aerosol particle nonsphericity in remote sensing of desert dust,  
854 J. Geophys. Res.-Atmos., 111, D11208,  
855 <https://doi.org/10.1029/2005JD006619>, 2006.

856 Dubuisson, P., Buriez, J. C., and Fouquart, Y.: High spectral resolution solar  
857 radiative transfer in absorbing and scattering media, application to the  
858 satellite simulation, J. Quant. Spectrosc. Ra., 55, 103–126,  
859 [https://doi.org/10.1016/0022-4073\(95\)00134-4](https://doi.org/10.1016/0022-4073(95)00134-4), 1996.

860 Eck, T. F., Holben, B. N., Reid, J. S., Dubovik, O., Smirnov, A., O'Neill, N. T.,  
861 Slutsker, I., and Kinne, S.: Wavelength dependence of the optical depth of  
862 biomass burning, urban, and desert dust aerosols, J. Geophys. Res., 104,

863 31333–31349, <https://doi.org/10.1029/1999jd900923>, 1999.

864 Eck, T. F., Holben, B. N., Dubovik, O., Smirnov, A., Goloub, P., Chen, H. B.,  
865 Chatenet, B., Gomes, L., Zhang, X. Y., Tsay, S. C., Ji, Q., Giles, D., and  
866 Slutsker, I.: Columnar aerosol optical properties at AERONET sites in  
867 central eastern Asia and aerosol transport to the tropical Mid-Pacific, *J.*  
868 *Geophys. Res.*, 110, D06202, <https://doi.org/10.1029/2004JD005274>,  
869 2005.

870 Estellés, V., Campanelli, M., Utrillas, M. P., Expósito, F., and Martínez-Lozano,  
871 J. A.: Comparison of AERONET and SKYRAD4.2 inversion products  
872 retrieved from a Cimel CE318 sunphotometer, *Atmos. Meas. Tech.*, 5,  
873 569–579, <https://doi.org/10.5194/amt-5-569-2012>, 2012.

874 Fan, X., Chen, H., Goloub, P., Xia, X., Zhang, W., and Chatenet, B.: Analysis of  
875 column-integrated aerosol optical thickness in Beijing from AERONET  
876 observations, *China Particuol.*, 4, 330-335, DOI:  
877 10.1016/s1672-2515(07)60285-1, 2006.

878 García, O. E., Díaz, J. P., Expósito, F. J., Díaz, A. M., Dubovik, O., Dubuisson,  
879 P., Roger, J.-C., Eck, T. F., Sinyuk, A., Derimian, Y., Dutton, E. G., Schafer,  
880 J. S., Holben, B. N., and García, C. A.: Validation of AERONET estimates  
881 of atmospheric solar fluxes and aerosol radiative forcing by groundbased  
882 broadband measurements, *J. Geophys. Res.*, 113, D21207,  
883 <https://doi.org/10.1029/2008JD010211>, 2008.

884 García, O. E., Díaz, J. P., Expósito, F. J., Díaz, A. M., Dubovik, O., and  
885 Derimian, Y.: Aerosol radiative forcing: AERONET based estimates,  
886 *climate Models*, InTech, edited by: Druyan, L., ISBN: 978- 953-51-0135-2,  
887 2012.

888 Garrett, T. J., and Zhao, C.: Increased Arctic cloud longwave emissivity  
889 associated with pollution from mid-latitudes, *Nature*, 440, 787-789,  
890 [doi:10.1038/nature04636](https://doi.org/10.1038/nature04636), 2006.

891 Gelencser, A.: *Carbonaceous Aerosol, Atmospheric and Oceanographic*  
892 *Sciences Libery.*, vol. 30. Springer, Netherland., 2004.

893 Giles, D. M., Holben, B. N., Tripathi, S. N., Eck, T. F., Newcomb, W. W.,  
894 Slutsker, I., Dickerson, R. R., Thompson, A. M., Mattoo, S., Wang, S.,  
895 Singh, R. P., Sinyuk, A., and Schafer, J. S.: Aerosol properties over the  
896 Indo-Gangetic Plain: A mesoscale perspective from the TIGERZ  
897 experiment, *J. Geophys. Res.-Atmos.*, 116(D18), DOI:  
898 10.1029/2011JD015809, 2011.

899 Giles, D. M., Holben, B. N., Eck, T. F., Sinyuk, A., Smirnov, A., Slutsker, I.,  
900 Dickerson, R. R., Thompson, A. M., and Schafer, J. S.: An analysis of  
901 AERONET aerosol absorption properties and classifications  
902 representative of aerosol source regions, *J. Geophys. Res.-Atmos.*, 117,  
903 127–135, DOI: 10.1029/2012JD018127, 2012.

904 Gobbi, G. P., Kaufman, Y. J., Koren, I., and Eck, T. F.: Classification of aerosol  
905 properties derived from AERONET direct sun data, *Atmos. Chem. Phys.*,  
906 7, 453–458, <http://dx.doi.org/10.5194/acp-7-453-2007>, 2007

907 Goloub, P., Li, Z., Dubovik, O., Blarel, L., Podvin, T., Jankowiak, I., Lecoq, R.,  
908 Deroo, C., Chatenet, B., and Morel, J. P.: PHOTONS/AERONET  
909 sunphotometer network overview: description, activities, results,  
910 Fourteenth International Symposium on Atmospheric and Ocean  
911 Optics/Atmospheric Physics, 6936, 69360V, DOI: 10.1117/12.783171,  
912 2007.

913 Haywood, J. M. and Shine, K. P.: The effect of anthropogenic sulfate and soot  
914 aerosol on the clear sky planetary radiation budget, *Geophys. Res. Lett.*,  
915 22, 603–606, <https://doi.org/10.1029/95GL00075>, 1995.

916 He, Q., Li, C., Geng, F., Yang, H., Li, P., Li, T., Liu, D., and Pei, Z.: Aerosol  
917 optical properties retrieved from Sun photometer measurements over  
918 Shanghai, China, *J. Geophys. Res.-Atmos.*, 117, D16204,  
919 <https://doi.org/10.1029/2011JD017220>, 2012.

920 Holben, B. N., Eck, T. F., Slutsker, I., Tanré, D., Buis, J. P., Setzer, A., Vermote,  
921 E., Reagan, J. A., Kaufman, Y. J., Nakajima, T., Lavenue, F., Jankowiak, I.,  
922 and Smirnov, A.: AERONET – A Federated Instrument Network and Data

923 Archive for Aerosol Characterization, Remote Sens. Environ., 66, 1–16,  
924 [https://doi.org/10.1016/S0034-4257\(98\)00031-5](https://doi.org/10.1016/S0034-4257(98)00031-5), 1998.

925 Jacobson, M. Z.: A physically based treatment of elemental carbon optics:  
926 implications for global direct forcing of aerosols, Geophys. Res. Lett., 27,  
927 217–220, DOI: 10.1029/1999gl010968, 2000.

928 Lacis, A. A. and Oinas, V.: A description of the correlated kdistribution method  
929 for modeling nongray gaseous absorption, thermal emission, and multiple  
930 scattering in vertically inhomogeneous atmospheres, J. Geophys. Res.,  
931 96, 9027–9063, DOI: 10.1029/90JD01945, 1991.

932 Lee, K. H., Li, Z., Cribb, M. C., Liu, J., Wang, L., Zheng, Y., Xia, X., Chen, H.,  
933 and Li, B.: Aerosol optical depth measurements in eastern China and a  
934 new calibration method, J. Geophys. Res., 115, 4038–4044,  
935 <https://doi.org/10.1029/2009JD012812>, 2010.

936 Li, C.C., Mao, J.T., Lau, A.K.H., Chen, J.C., Yuan, Z.B., Liu, X.Y., Zhu, A.H.,  
937 and Liu, G.Q.: Characteristics of distribution and seasonal variation of  
938 aerosol optical depth in eastern China with MODIS products, Chinese  
939 Science Bulletin., 48, 2488-2495, DOI: 10.1360/03wd0224, 2003.

940 Li, Z., Chen, H., Cribb, M., Dickerson, R., Holban, B., Li, C., Lu, D., Luo, Y.,  
941 Maring, H., Shi, G., Tsay, S.C., Wang, P., Wang, Y., Xia, X., and Zhao, F.:  
942 Overview of the East Asian Studies on tropospheric aerosols, an  
943 international regional experiment(EAST-AIRE), J. Geophys. Res., 112,  
944 D22S00. <http://dx.doi.org/10.1029/2007JD008853>, 2007.

945 Li, X., and Zhang, L.: Analysis of aerosol sources and optical properties based  
946 on backward trajectory method over SACOL, Acta Physica Sin., 61, 1-9,  
947 DOI: 10.7498/aps.61.023402, 2012.

948 Li, Z. Q., Eck, T., Zhang, Y., Zhang, Y. H., Li, D. H., Li, L., Xu, H., Hou, W. Z., Lv,  
949 Y., Goloub, P., and Gu, X. F.: Observations of residual submicron fine  
950 aerosol particles related to cloud and fog processing during a major  
951 pollution event in Beijing, Atmos. Environ., 86, 187–192, DOI:  
952 10.1016/j.atmosenv.2013.12.044, 2014.



953 Li, Z., Lau, W. K.-M., Ramanathan, V., Wu, G., Ding, Y., Manoj, M. G., Liu, J.,  
954 Qian, Y., Li, J., Zhou, T., Fan, J., Rosenfeld, D., Ming, Y., Wang, Y., Huang,  
955 J., Wang, B., Xu, X., Lee, S.- S., Cribb, M., Zhang, F., Yang, X., Takemura,  
956 T., Wang, K., Xia, X., Yin, Y., Zhang, H., Guo, J., Zhai, P. M., Sugimoto, N.,  
957 Babu, S. S., and Brasseur, G. P.: Aerosol and monsoon climate  
958 interactions over Asia, *Rev. Geophys.*, 54, 866–929,  
959 <https://doi.org/10.1002/2015RG000500>, 2016.

960 Li, Z.Q., Xu, H., Li, K. T., Li, D. H., Xie, Y. S., Li, L., Zhang, Y., Gu, X.F., Zhao,  
961 W., Tian, Q.J., Deng, R.R., Su, X.L., Huang, B., Qiao, Y.L., Cui, W.Y., Hu,  
962 Y., Gong, C.L., Wang, Y.Q., Wang, X.F., Wang, J.P., Du, W.B., Pan, Z.Q.,  
963 Li, Z.Z., and Bu, D.: Comprehensive Study of Optical, Physical, Chemical,  
964 and Radiative Properties of Total Columnar Atmospheric Aerosols over  
965 China: An Overview of Sun–Sky Radiometer Observation Network  
966 (SONET) Measurements, *Bulletin of the American Meteorological Society.*,  
967 99(4), 739–755, <https://doi.org/10.1175/BAMS-D-17-0133.1>, 2018.

968 Luo, Y., Lu, D., Zhou, X., and Li, W.: Analyses on the spatial distribution of  
969 aerosol optical depth over china in recent 30 years, *Chinese Journal of*  
970 *Atmospheric Sciences.*, 26, 721-730, DOI: 10.1002/mop.10502, 2002.

971 Mai, B., Deng, X., Xia, X., Che, H., Guo, J., Liu, X., Zhu, J., and Ling, C.:  
972 Column-integrated aerosol optical properties of coarse- and fine-mode  
973 particles over the Pearl River Delta region in China, *Sci. Total. Environ.*,  
974 622–623, 481-492, DOI: 10.1016/j.scitotenv.2017.11.348, 2018.

975 Myhre, G.: Consistency between satellite-derived and modeled estimates of  
976 the direct aerosol effect, *Science*, 325, 187-190, DOI:  
977 10.1126/science.1174461, 2009.

978 Nakajima, T. and Tanaka, M.: Algorithms for radiative intensity calculations in  
979 moderately thick atmospheres using a truncation approximation, *J. Quant.*  
980 *Spectrosc. Ra.*, 40, 51–69,  
981 [https://doi.org/10.1016/0022-4073\(88\)90031-3](https://doi.org/10.1016/0022-4073(88)90031-3), 1988.

982 Pappalardo, G., Amodeo, A., Apituley, A., Comeron, A., Freudenthaler, V.,

983 Linné, H., Ansmann, A., Bösenberg, J., D'Amico, G., Mattis, I., Mona, L.,  
984 Wandinger, U., Amiridis, V., AladosArboledas, L., Nicolae, D., and  
985 Wiegner, M.: EARLINET: towards an advanced sustainable European  
986 aerosol lidar network, *Atmos. Meas. Tech.*, 7, 2389–2409,  
987 <https://doi.org/10.5194/amt-7-2389-2014>, 2014.

988 Prats, N., Cachorro, V. E., Berjón, A., Toledano, C., and De Frutos, A. M.:  
989 Column-integrated aerosol microphysical properties from AERONET Sun  
990 photometer over southwestern Spain, *Atmos. Chem. Phys.*, 11, 12535–  
991 12547, DOI:10.5194/acp-11-12535-2011, 2011.

992 Qin, K., Wang, L.Y., Wu, L.X., Xu, J., Rao, L.L., Letu, H., Shi, T.W., Wang, R.F.,  
993 2017. A campaign for investigating aerosol optical properties during winter  
994 hazes over Shijiazhuang, China, *Atmos. Res.*, 198, 113-122,  
995 <https://doi.org/10.1016/j.atmosres.2017.08.018>, 2017.

996 Ramanathan, V., Crutzen, P. J., Kiehl, J. T., and Rosenfeld, D.: Aerosol,  
997 climate, and hydrological cycle, *Science* 294, 2119–2124,  
998 <https://doi.org/10.1126/science.1064034>, 2001.

999 Remer, L. A., and Kaufman, Y. J.: Dynamic aerosol model: Urban/industrial  
1000 aerosol, *J. Geophys. Res.*, 103, 13859–13871,  
1001 <https://doi.org/10.1029/98jd00994>, 1998.

1002 Roger, J. C., Mallet, M., Dubuisson, P., Cachier, H., Vermote, E., Dubovik, O.,  
1003 and Despiiau, S.: A synergetic approach for estimating the local direct  
1004 aerosol forcing: applications to an urban zone during the ESCOMPTE  
1005 experiment, *J. Geophys. Res.*, 111, D13208,  
1006 <https://doi.org/10.1029/2005JD006361>, 2006.

1007 Russell, P. B., Bergstrom, R. W., Shinozuka, Y., Clarke, A. D., DeCarlo, P. F.,  
1008 Jimenez, J. L., Livingston, J. M., Redemann, J., Dubovik, O., and Strawa,  
1009 A.: Absorption Angstrom Exponent in AERONET and related data as an  
1010 indicator of aerosol composition, *Atmos. Chem. Phys.*, 10, 1155–1169,  
1011 <https://doi.org/10.5194/acp-10-1155-2010>, 2010.

1012 Shen, L., Zhao, C., Ma, Z., Li, Z., Li, J., and Wang, K.: Observed decrease of

1013 summer sea-land breeze in Shanghai from 1994 to 2014 and its  
1014 association with urbanization, *Atmos. Res.*,  
1015 doi:10.1016/j.atmosres.2019.05.007, 2019.

1016 Shettle, E. P., and Fenn, R. W.: Models for the aerosols of the lower  
1017 atmosphere and the effects of humidity variations on their optical  
1018 properties. AFCRL Tech. Rep. 79 0214, Air Force Cambridge Research  
1019 Laboratory, Hanscom Air Force Base, MA, 100 pp, 1979.

1020 Stamnes, K., Tsay, S. C., Wiscombe, W., and Jayaweera, K.: Numerically  
1021 stable algorithm for discrete-ordinate-method radiative transfer in multiple  
1022 scattering and emitting layered media, *Appl. Optics.*, 27, 2502–2509,  
1023 <https://doi.org/10.1364/ao.27.002502>, 1988.

1024 Su, X. L., Wang, Q., Li, Z. Q., Calvello, M., Esposito, F., Pavese, G., Lin, M. J.,  
1025 Cao, J. J., Zhou, C. Y., Li, D. H., Xu, H.: Regional transport of  
1026 anthropogenic pollution and dust aerosols in spring to Tianjin — A coastal  
1027 megacity in China, *Sci. Total Environ.*, 584-585, 381-392,  
1028 <https://doi.org/10.1016/j.scitotenv.2017.01.016>, 2017.

1029 Sun, J. Y., Zhang, Q., Canagaratna, M. R., Zhang, Y. M., Ng, N. L., Sun, Y. L.,  
1030 Jayne, J. T., Zhang, X. C., Zhang, X. Y., and Worsnop, D. R.: Highly time-  
1031 and size-resolved characterization of submicron aerosol particles in  
1032 Beijing using an Aerodyne Aerosol Mass Spectrometer, *Atmos. Environ.*,  
1033 44, 131–140, 2010.

1034 Sun, T., Che, H., Qi, B., Wang, Y., Dong, Y., Xia, X., Wang, H., Gui, K., Zheng,  
1035 Y., Zhao, H., Ma, Q., Du, R., and Zhang, X.: Aerosol optical characteristics  
1036 and their vertical distributions under enhanced haze pollution events:  
1037 effect of the regional transport of different aerosol types over eastern  
1038 China, *Atmos. Chem. Phys.*, 18, 2949–2971,  
1039 <https://doi.org/10.5194/acp-18-2949-2018>, 2018.

1040 Takamura, T., and Nakajima, T.: Overview of SKYNET and its activities, *Opt.*  
1041 *Pura Apl.*, 37, 3303–3308, 2004.

1042 Tao, R., Che, H. Z., Chen, Q. L., Wang, Y. Q., Sun, J. Y., Zhang, X. C., Lu, S.,

1043 Guo, J. P., Wang, H., and Zhang, X.Y.: Development of an integrating  
1044 sphere calibration method for Cimel sunphotometers in China aerosol  
1045 remote sensing network, *Particuology* 13, 88–99.  
1046 <https://doi.org/10.1016/j.partic.2013.04.009>, 2014.

1047 Twomey, S. A., Piepgrass, M., and Wolfe, T. L.: An assessment of the impact of  
1048 pollution on the global cloud albedo, *Tellus*, 36B, 356–366,  
1049 <https://doi.org/10.1111/j.1600-0889.1984.tb00254.x>, 1984.

1050 Wan, X., Kang, S.C., Wang, Y.S., Xin, J.Y., Liu, B., Guo, Y.H., Wen, T.X., Zhang,  
1051 G.S., and Cong, Z.Y.: Size distribution of carbonaceous aerosols at a  
1052 high-altitude site on the central Tibetan Plateau (Nam Co Station, 4730m  
1053 a.s.l.), *Atmos. Res.*, 153, 155–164,  
1054 <https://doi.org/10.1016/j.atmosres.2014.08.008>, 2015.

1055 Wang, X., Huang, J.P., Ji, M.X., and Higuchi, K.: Variability of east Asia dust  
1056 events and their long-term trend, *Atmos. Environ.*, 42, 3156–3165,  
1057 <http://dx.doi.org/10.1016/j.atmosenv.2007.07.046>, 2007.

1058 Wang, L. C., Gong, W., Xia, X. A., Zhu, J., Li, J., and Zhu, Z. M.: Long-term  
1059 observations of aerosol optical properties at Wuhan, an urban site in  
1060 Central China, *Atmos. Environ.*, 101, 94–102, DOI:  
1061 [10.1016/j.atmosenv.2014.11.021](https://doi.org/10.1016/j.atmosenv.2014.11.021), 2015.

1062 Wang, P., Che, H. Z., Zhang, X. C., Song, Q. L., Wang, Y. Q., Zhang, Z. H., Dai,  
1063 X., and Yu, D. J.: Aerosol optical properties of regional background  
1064 atmosphere in Northeast China, *Atmos. Environ.*, 44, 4404–4412, DOI:  
1065 [10.1016/j.atmosenv.2010.07.043](https://doi.org/10.1016/j.atmosenv.2010.07.043) , 2010.

1066 Wang, Z., Liu, D., Wang, Y., Wang, Z., and Shi, G.: Diurnal aerosol variations  
1067 do affect daily averaged radiative forcing under heavy aerosol loading  
1068 observed in Hefei, China, *Atmos. Meas. Tech.*, 8, 2901–2907,  
1069 <https://doi.org/10.5194/amt-8-2901-2015>, 2015.

1070 Wehrli, C.: Calibration of filter radiometers for the GAW Aerosol Optical Depth  
1071 network at Jungfraujoch and Mauna Loa, in: *Proceedings of ARJ*  
1072 *Workshop, SANW Congress, Davos, Switzerland*, 70–71, 2002.

1073 Whitey, K. T.: The physical characteristics of sulfur aerosols. *Atmos. Environ.*,  
1074 41(supp-S), 25-49, DOI: 10.1016/j.atmosenv.2007.10.057, 1978.

1075 Wu, Y.F., Zhang, R.J., Pu, Y.F., Zhang, L.M., Ho, K.F., and Fu, C.B.: Aerosol  
1076 optical properties observed at a semi-arid rural site in Northeastern China,  
1077 *Aerosol. Air. Qual. Res.*, 12,503–514, DOI: 10.4209/aaqr.2011.11.0202 ,  
1078 2012.

1079 Xia, C., Sun, J. Y., Qi, X. F., Shen, X. J., Zhong, J. T., Zhang, X. Y., Wang, Y. Q.,  
1080 Zhang, Y. M., and Hu, X. Y.: Observational study of aerosol hygroscopic  
1081 growth on scattering coefficient in Beijing: A case study in March of 2018,  
1082 *Sci. Total Environ.*, 685, 239-247, 2019.

1083 Xia, X.: A closer looking at dimming and brightening in China during 1961–  
1084 2005, *Ann. Geophys.*, 28, 1121–1132, doi:10.5194/angeo-28-1121-2010,  
1085 2010.

1086 Xia, X., Chen, H., Goloub, P., Zong, X., Zhang, W., and Wang, P.:  
1087 Climatological aspects of aerosol optical properties in North China Plain  
1088 based on ground and satellite remote-sensing data, *J. Quant. Spectrosc.*  
1089 *Ra.*, 127, 12–23, DOI: 10.1016/j.jqsrt.2013.06.024, 2013.

1090 Xie, C., Nishizawa, T., Sugimoto, N., Matsui, I., and Wang, Z.: Characteristics  
1091 of aerosol optical properties in pollution and Asian dust episodes over  
1092 Beijing, China. *Appl. Opt.*, 47, 4945-4951, DOI: 10.1364/AO.47.004945,  
1093 2008.

1094 Xie, S., Liu, X., Zhao, C., and Zhang, Y.: Sensitivity of CAM5-Simulated Arctic  
1095 Clouds and Radiation to Ice Nucleation Parameterization, *J. Climate.*, 26,  
1096 5981-5999, doi:10.1175/jcli-d-12-00517.1, 2013.

1097 Xin, J., Wang Y, Li, Z.Q., Wang, P.C., Hao, W.M., Nordgren, B.L., Wang, S.G.,  
1098 Liu, G.R., Wang, L.L., Wen, T.X., Sun, Y., and Hu, B.: Aerosol optical  
1099 depth (AOD) and Ångström exponent of aerosols observed by the  
1100 Chinese Sun Hazemeter Network from August 2004 to September 2005, *J.*  
1101 *Geophys. Res.*, 112, D05203, DOI: 10.1029/2006JD007075, 2007

1102 Xin, J.Y., Wang, L.L., Wang, Y.S., Li, Z., and Wang, P.: Trends in aerosol

1103 optical properties over the Bohai Rim in Northeast China from 2004 to  
1104 2010, *Atmos. Environ.*, **45**, 6317–6325, DOI:  
1105 10.1016/j.atmosenv.2011.08.052, 2011.

1106 Xin J, Wang Y, Pan Y, Ji, D., Liu, Z., Wen, T.X., Wang, Y.H., Li, X.R., Sun, Y.,  
1107 Sun, J., Wang, P.C., Wang, G.H., Wang, X.M., Cong, Z.Y., Song, T., Hu, B.,  
1108 Wang, L.L., Tang, G.Q., Gao, W.K., Guo, Y.H., Miao, H.Y., Tian, S.L., and  
1109 Wang, L.: The campaign on atmospheric aerosol research network of  
1110 China: CARE-China, *Bulletin of the American Meteorological Society.*, **96**,  
1111 1137-1155, <https://doi.org/10.1175/BAMS-D-14-00039.1>, 2015.

1112 Yang, X., Zhao, C., Zhou, L., Li, Z., Cribb, M., and Yang, S.: Wintertime cooling  
1113 and a potential connection with transported aerosols in Hong Kong during  
1114 recent decades, *Atmos. Res.*, **211**, 52–61,  
1115 doi:10.1016/j.atmosres.2018.04.029, 2018.

1116 Yang, X., Zhao, C., Zhou, L., Wang, Y., and Liu, X.: Distinct impact of different  
1117 types of aerosols on surface solar radiation in China, *J. Geophys.*  
1118 *Res.-Atmospheres*, **121**, 6459–6471. DOI:  
1119 <https://doi.org/10.1002/2016jd024938>, 2016.

1120 Yang, Y., Zhao, C., Dong, X., Fan, G., Zhou, Y., Wang, Y., Zhao, L., Lv, F., and  
1121 Yan, F.: Toward understanding the process-level impacts of aerosols on  
1122 microphysical properties of shallow cumulus cloud using aircraft  
1123 observations, *Atmos. Res.*, doi:10.1016/j.atmosres.2019.01.027, 2019.

1124 Yang, Y., Zhao, C., Sun, L., and Wei, J.: Improved aerosol retrievals over  
1125 complex regions using NPP Visible Infrared Imaging Radiometer Suite  
1126 observations, *Earth. Space. Sci.*, doi:10.1029/2019ea000574, 2019.

1127 Yuan, Y., Shuai, Y., Li, X.W., Liu, B., and Tan, H.P.: Using a new aerosol  
1128 relative optical thickness concept to identify aerosol particle species,  
1129 *Atmos. Res.*, **150**, 1–11, DOI: 10.1016/j.atmosres.2014.07.007, 2014.

1130 Zhang, L., Sun, J. Y., Shen, X. J., Zhang, Y. M., Che, H., Ma, Q. L., Zhang, Y.  
1131 W., Zhang, X. Y., and Ogren, J. A.: Observations of relative humidity  
1132 effects on aerosol light scattering in the Yangtze River Delta of China,

1133 Atmos. Chem. Phys., 15, 8439-8454,  
1134 <https://doi.org/10.5194/acp-15-8439-2015>, 2015.

1135 Zhang, M., Ma, Y., Gong, W., Liu, B., Shi, Y., and Chen, Z.: Aerosol optical  
1136 properties and radiative effects: Assessment of urban aerosols in central  
1137 China using 10-year observations, *Atmos. Environ.*, 182, 275-285, DOI:  
1138 10.1016/j.atmosenv.2018.03.040, 2018

1139 Zhang, R., Jing, J., Tao, J., Hsu, S.-C., Wang, G., Cao, J., Lee, C.S.L., Zhu, L.,  
1140 Chen, Z., Zhao, Y., and Shen, Z.: Chemical characterization and source  
1141 apportionment of PM<sub>2.5</sub> in Beijing: seasonal perspective, *Atmos. Chem.*  
1142 *Phys.*, 13, 7053–7074, <http://dx.doi.org/10.5194/acp-13-7053-2013>, 2013

1143 Zhao, C., and Garrett, T. J.: Effects of Arctic haze on surface cloud radiative  
1144 forcing, *Geophys. Res. Lett.*, 42, 557–564. doi:10.1002/2014gl062015,  
1145 2015.

1146 Zhao, C., Lin, Y., Wu, F., Wang, Y., Li, Z., Rosenfeld, D., and Wang, Y.:  
1147 Enlarging Rainfall Area of Tropical Cyclones by Atmospheric Aerosols,  
1148 *Geophys. Res. Lett.*, doi:10.1029/2018gl079427, 2018.

1149 Zhao, C., Wang, Y., Shi, X., Zhang, D., Wang, C., Jiang, J. H., Zhang, Q., and  
1150 Fan, H.: Estimating the contribution of local primary emissions to  
1151 particulate pollution using high-density station observations, *J. Geophys.*  
1152 *Res.-Atmospheres*, doi:10.1029/2018jd028888, 2019.

1153 Zhao, H., Che, H., Ma, Y., Xia, X., Wang, Y., Wang, P., and Wu, X.: Temporal  
1154 variability of the visibility, particulate matter mass concentration and  
1155 aerosol optical properties over an urban site in Northeast China, *Atmos.*  
1156 *Res.*, 166, 204–212, DOI: 10.1016/j.atmosres.2015.07.003, 2015.

1157 Zhao H., Che H., Wang Y., Wang H., Ma Y., and Wang Y.: Investigation of the  
1158 Optical Properties of Aerosols over the Coastal Region at Dalian,  
1159 Northeast China, *Atmosphere.*, 7(8), 103, DOI: 10.3390/atmos7080103,  
1160 2016

1161 Zhao, H., Che, H., Xia, X., Wang, Y., Wang, H., Wang, P., Ma, Y., Yang, H., Liu,  
1162 Y., Wang, Y., Gui, K., Sun, T., Zheng, Y., and Zhang, X.: Multi-year

1163 ground-based measurements of aerosol optical properties and direct  
1164 radiative effect over different surface types in northeastern China, *J.*  
1165 *Geophys. Res.-Atmos.*, 123, 13,887–13,916,  
1166 <https://doi.org/10.1029/2018JD029141>, 2018

1167 Zhang, K., Zhao, C., Fan, H., Yang, Y., and Sun, Y.: Toward Understanding the  
1168 Differences of PM<sub>2.5</sub> Characteristics Among Five China Urban Cities,  
1169 *Asia-Pac. J. Atmos. Sci.*, doi:10.1007/s13143-019-00125-w, 2019.

1170 Zheng, C., Zhao, C., Zhu, Y., Wang, Y., Shi, X., Wu, X., Chen, T., Wu, F., and  
1171 Qiu, Y.: Analysis of influential factors for the relationship between PM<sub>2.5</sub>  
1172 and AOD in Beijing, *Atmos. Chem. Phys.*, 17, 13473–13489, doi:  
1173 10.5194/acp-17-13473-2017, 2017.

1174 Zheng, Y., Che, H., Xia, X., Wang, Y., Wang, H., Wu, Y., Tao, J., Zhao, H., An,  
1175 L., Li, L., Gui, K., Sun, T., Li, X., Sheng, Z., Liu, C., Yang, X., Liang, Y.,  
1176 Zhang, L., Kuang, X., Luo, S., and You, Y.: Five-year observation of  
1177 aerosol optical properties and its radiative effects to planetary boundary  
1178 layer during air pollution episodes in North China: Intercomparison of a  
1179 plain site and a mountainous site in Beijing, *Sci. Total. Environ.*, 674,  
1180 140-158, 2019.

1181 Zhu, J., Che, H., Xia, X., Chen, H. B., Goloub, P., and Zhang, W.:  
1182 Column-integrated aerosol optical and physical properties at a regional  
1183 background atmosphere in North China Plain, *Atmos. Environ.*, 84, 54–64,  
1184 DOI: 10.1016/j.atmosenv.2013.11.019, 2014.

1185 Zhuang, B., Wang, T., Li, S., Liu, J., Talbot, R., Mao, H., Yang, X., Fu, C., Yin,  
1186 C., Zhu, J., Che, H., and Zhang, X.: Optical properties and radiative  
1187 forcing of urban aerosols in Nanjing over China, *Atmos. Environ.*, 83, 43–  
1188 52, DOI: 10.1016/j.atmosenv.2013.10.052, 2014.

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1193 **Figure captions**

1194 **Figure 1. Annual spatial distribution of aerosol volume-size distributions**  
1195 **at the CARSNET sites**

1196 **Figure 2. Annual spatial distribution of aerosol optical depth (AOD) at 440**  
1197 **nm at the CARSNET sites**

1198 **Figure 3. Annual spatial distribution of extinction Ångström exponent**  
1199 **(AE) 440-870 nm at the CARSNET sites**

1200 **Figure 4. Annual spatial distribution of fine mode fraction at the**  
1201 **CARSNET sites**

1202 **Figure 5. Annual spatial distribution of the single scattering albedo (SSA)**  
1203 **at 440 nm at the CARSNET sites**

1204 **Figure 6. Annual spatial distribution of absorption aerosol optical depth**  
1205 **(AAOD) at 440 nm at the CARSNET sites**

1206 **Figure 7. Annual spatial distribution of direct aerosol radiative effect at**  
1207 **the bottom of the atmosphere at the CARSNET sites**

1208 **Figure 8. Annual spatial distribution of direct aerosol radiative effect at**  
1209 **the top of the atmosphere at the CARSNET sites**

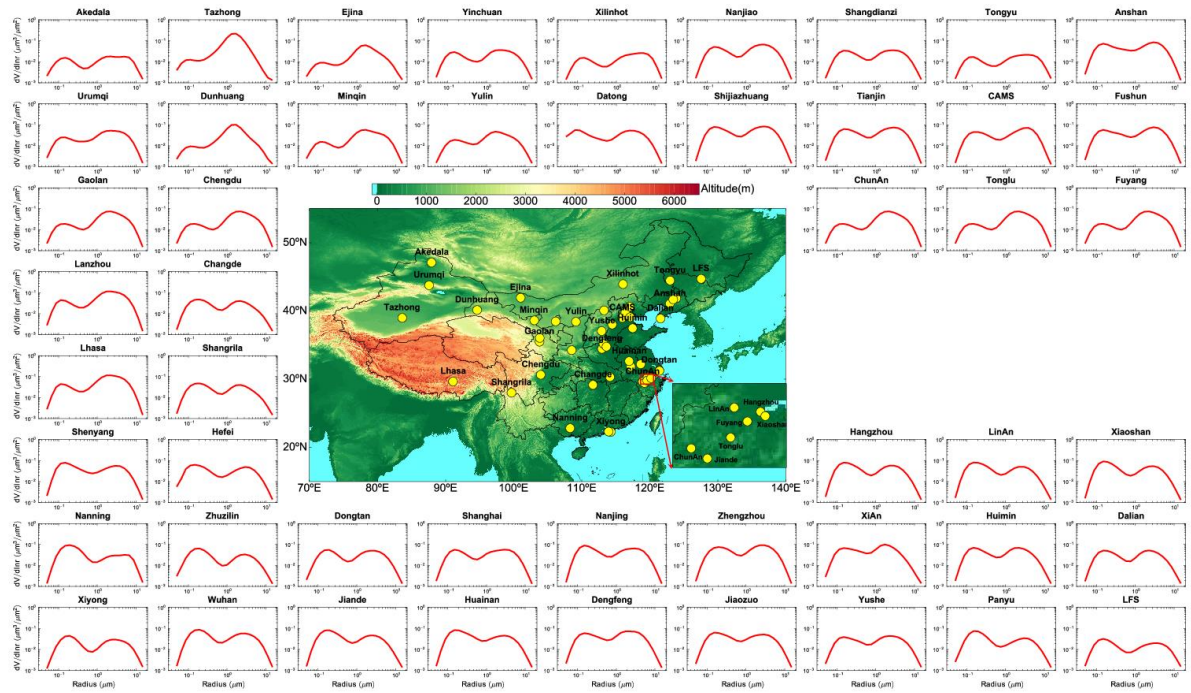
1210 **Figure 9. Annual spatial distribution of the aerosol type classification of**  
1211 **types I–VII at the CARSNET sites**

1212 **Table 1 The aerosol type classification based on the optical properties**

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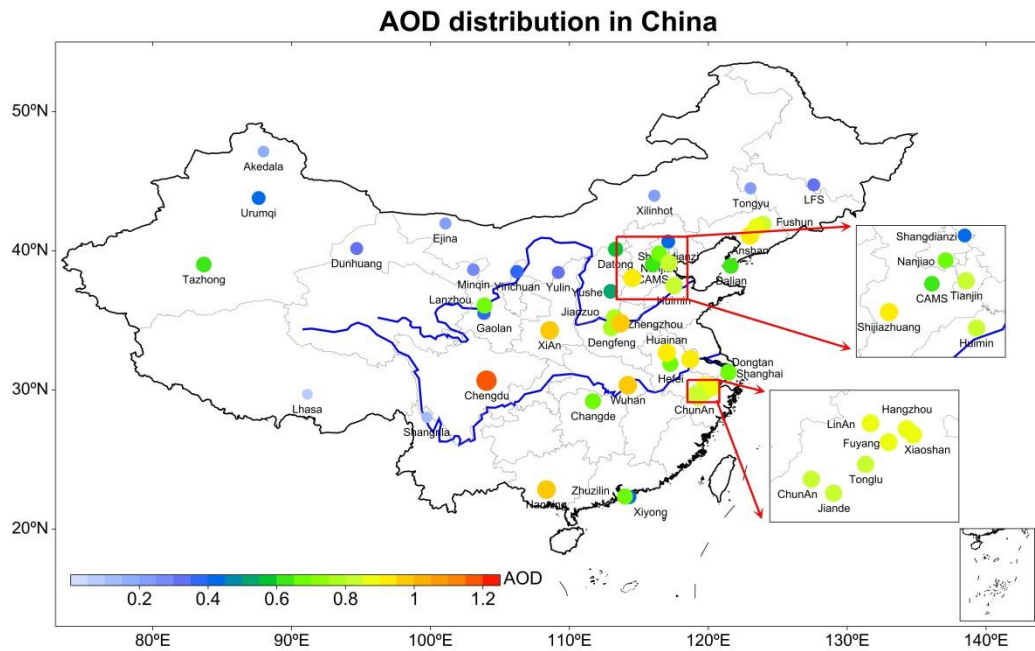
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**Figure 1. Annual spatial distribution of aerosol volume-size distributions at the CARSNET sites**



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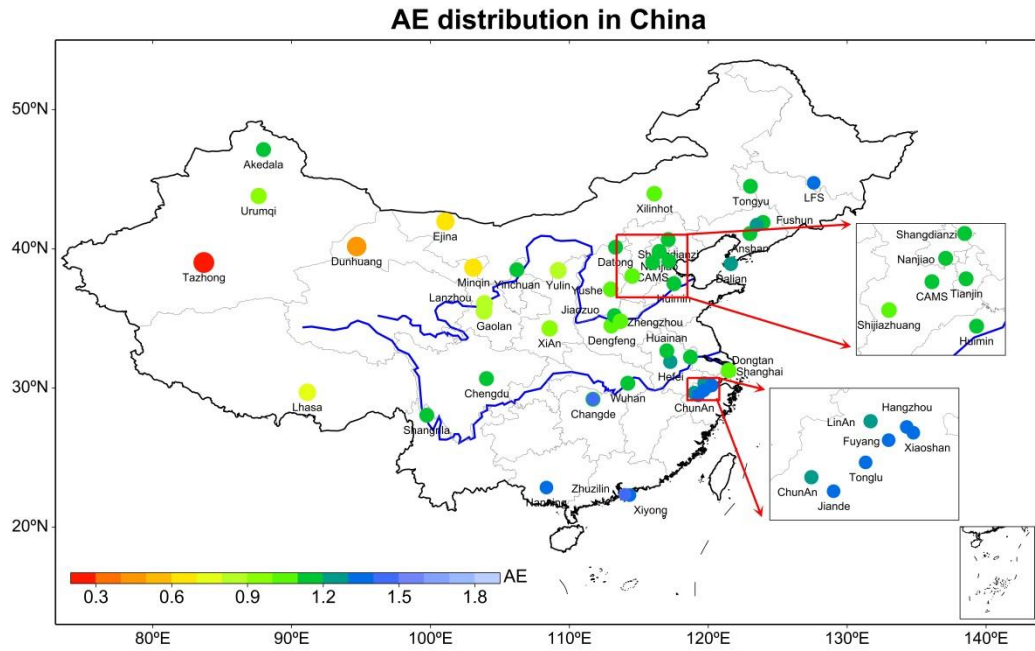
1219 **Figure 2. Annual spatial distribution of aerosol optical depth (AOD) at 440**  
1220 **nm at the CARSNET sites**  
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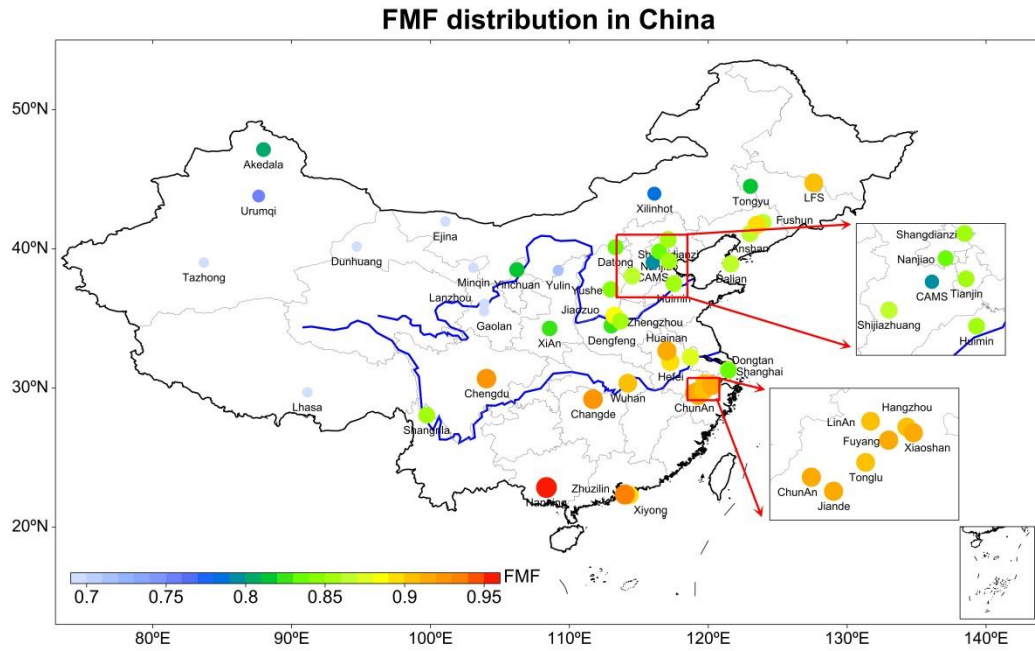
1224 **Figure 3. Annual spatial distribution of extinction Ångström exponent**  
1225 **(AE) 440-870 nm at the CARSNET sites**

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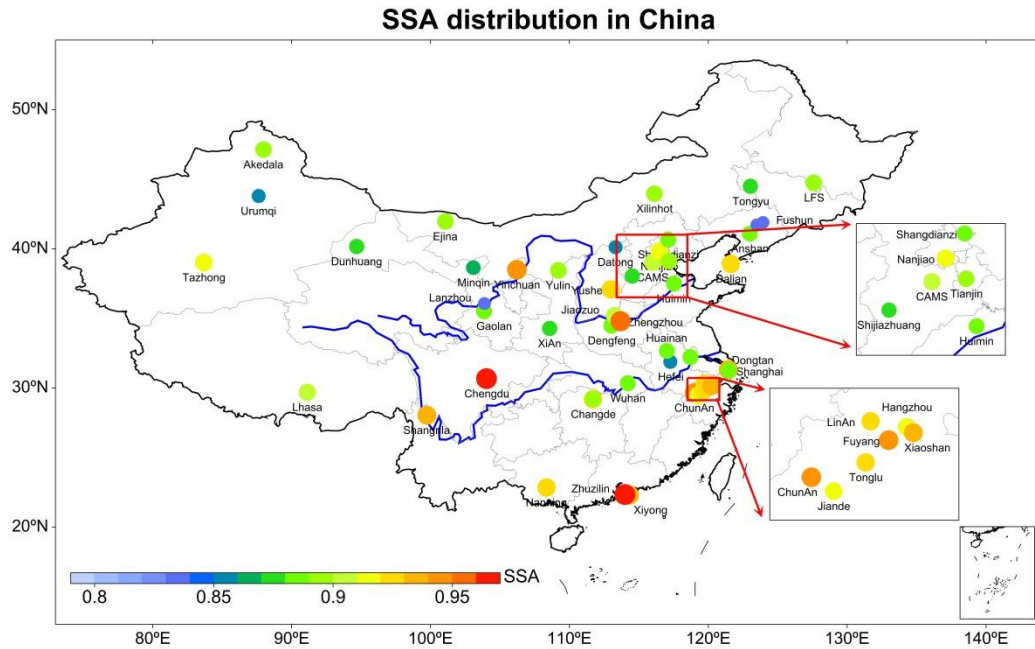
1229 **Figure 4. Annual spatial distribution of fine mode fraction at the**  
1230 **CARSNET sites**  
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1234 **Figure 5. Annual spatial distribution of the single scattering albedo (SSA)**  
1235 **at 440 nm at the CARSNET sites**

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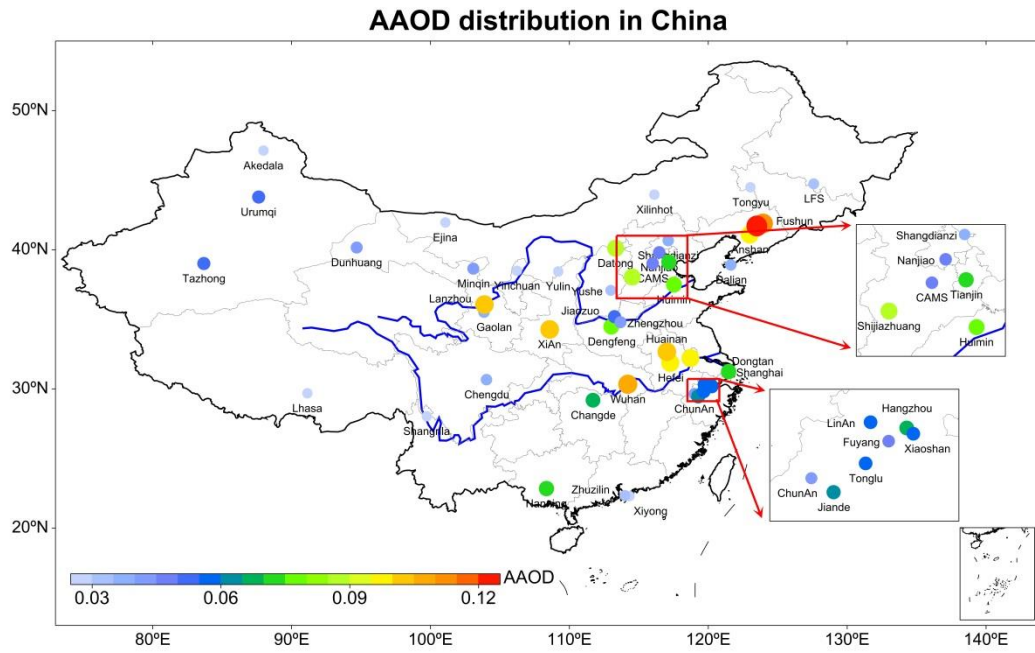
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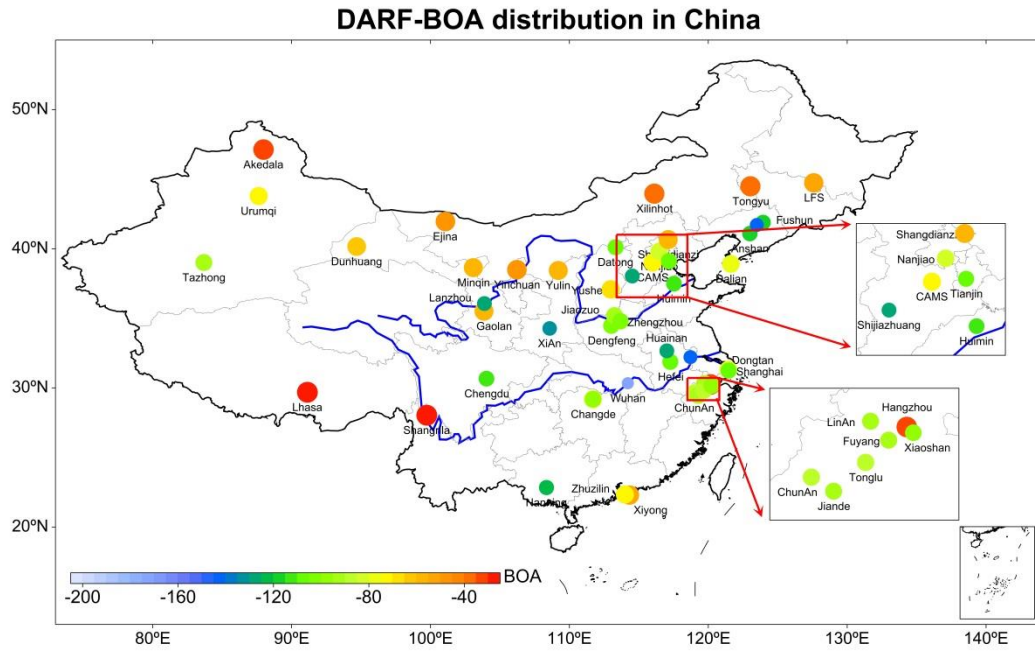
1241 **Figure 6. Annual spatial distribution of absorption aerosol optical depth**  
1242 **(AAOD) at 440 nm at the CARSNET sites**  
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1246 **Figure 7. Annual spatial distribution of direct aerosol radiative effect at**  
1247 **the bottom of the atmosphere at the CARSNET sites**

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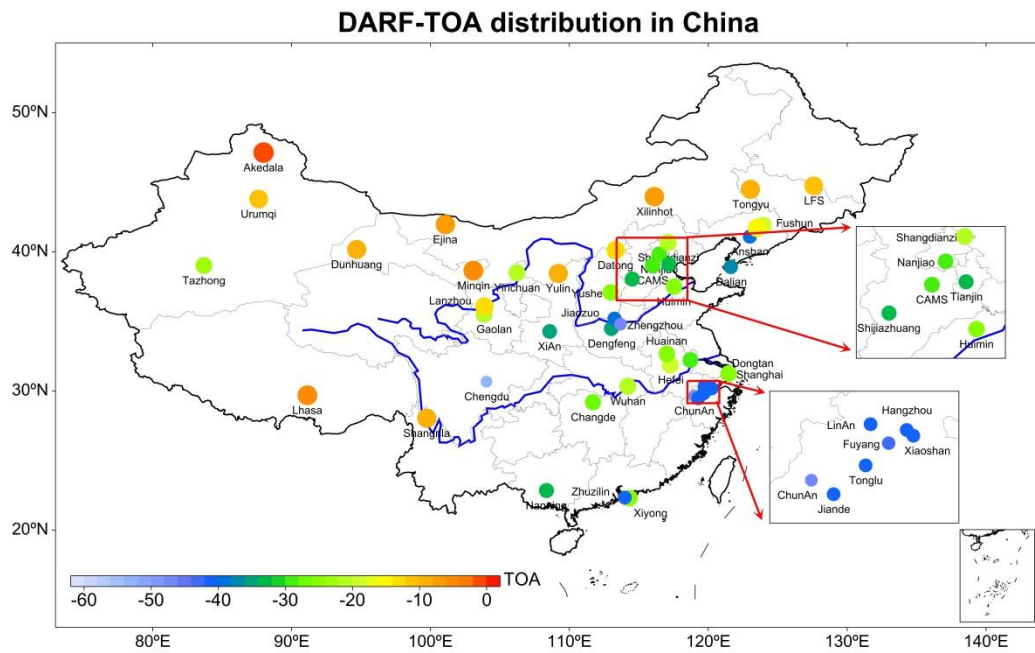


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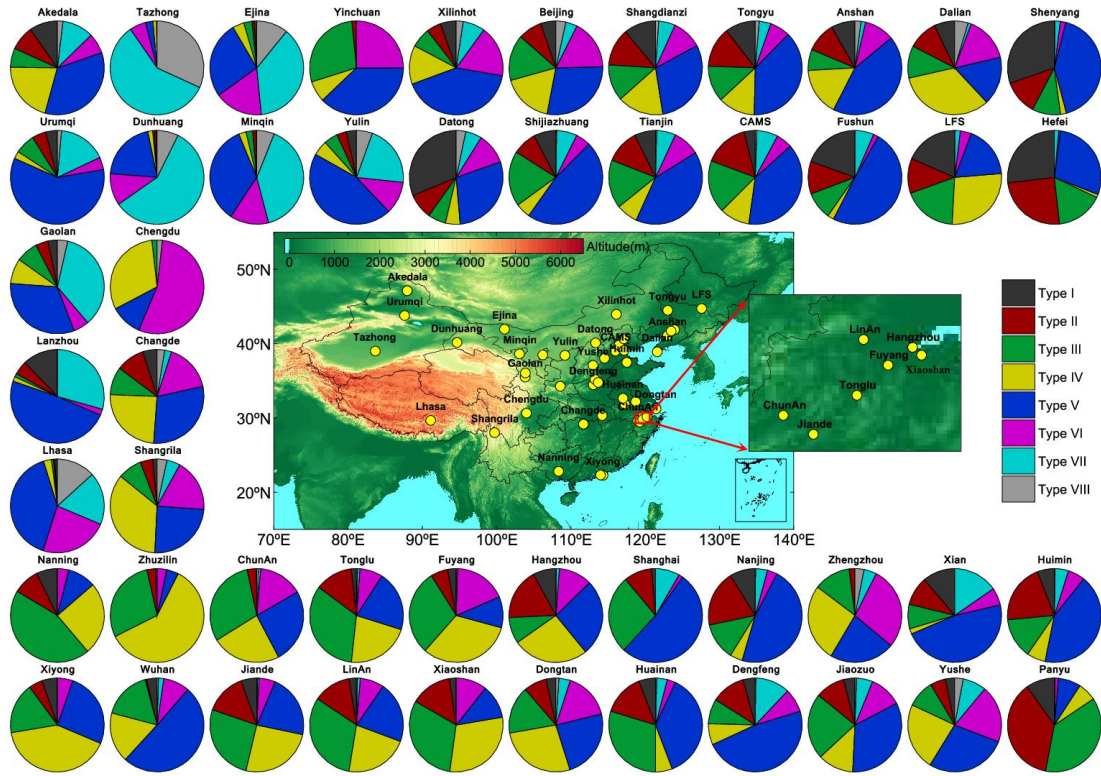
1251 **Figure 8. Annual spatial distribution of direct aerosol radiative effect at**  
1252 **the top of the atmosphere at the CARSNET sites**  
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1256 **Figure 9. Annual spatial distribution of the aerosol type classification of**  
 1257 **types I–VII at the CARSNET sites**

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1261 **Table 1. The aerosol type classification based on the optical properties.**

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Type	EAE	SSA	absorbing properties and particle size
I	$EAE > 1.20$	$SSA_{440nm} \leq 0.85$	fine-mode particles with highly-absorbing
II	$EAE > 1.20$	$0.85 \leq SSA_{440nm} < 0.90$	fine-mode particles with moderately-absorbing
III	$EAE > 1.20$	$0.90 \leq SSA_{440nm} < 0.95$	fine-mode particles with slightly-absorbing
IV	$EAE > 1.20$	$SSA_{440nm} > 0.95$	fine-mode particles with weakly-absorbing
V	$0.60 \leq EAE < 1.20$	$SSA_{440nm} \leq 0.95$	mixed-absorbing particles
VI	$0.60 \leq EAE < 1.20$	$SSA_{440nm} > 0.95$	mixed-slightly absorbing particles
VII	$EAE \leq 0.60$	$SSA_{440nm} \leq 0.95$	coarse mode particles with strongly absorbing (mainly dust)
VIII	$AE \leq 0.60$	$SSA_{440nm} > 0.95$	coarse-mode particles with weakly-absorbing

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1280 **Appendix**1281 **Table 1. Site information for the 50 CARSNET sites used in this study**

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No.	Site Name	Long.	Lat.	Alt.	Site information	Obs. Num	Period
Remote sites (three sites)							
1	Akedala	47.12	87.97	562.0	55 km west of Fuhai county, Xinjiang province, and 250–300 km southeast of Kazakistan.	947	2010-2017
2	Lhasa	29.67	91.13	3663.0	In the center of Lhasa city, Qinghai-Xizang Plateau.	437	2012-2017
3	Shangri-La	28.02	99.73	3583.0	12 km northeast of Shangri-La county, Diqing area, Yunnan province	325	2013-2017
Arid and semi-arid sites (six sites)							
4	Dunhuang	40.15	94.68	1139.0	1.5 km northeast of Dunhuang city, Gansu province; near Kumutage Desert of China	2030	2012-2017
5	Ejina	41.95	101.07	940.5	West of Inner-Mongolia Province, near Mongolia and Badanjilin desert	1970	2013-2017
6	Minqin	38.63	103.08	1367.0	In Minqin county, east to Tenggeli desert and north to Badanjilin Desert, Gansu Province	481	2013-2017
7	Tazhong	39.00	83.67	1099.4	In the middle of Taklamakan Desert, Xinjiang Province	1279	2013-2017
8	Xilinhot	43.95	116.12	1003.0	5 km southeast of Xilinhot City, near Hunshandake sand-land, Inner-Mongolia Province,	1464	2013-2017
9	Tongyu	44.42	122.87	151.0	In Tonyu city, west of Jilin Province	817	2010-2011
Rural sites on the Chinese Loess Plateau or nearby (three sites)							
10	Mt.Gaolan	36.00	103.85	2161.6	5 km north of Lanzhou city in Gansu province	769	2015-2016
11	Yulin	38.43	109.20	1135.0	10 km north of Yulin city in Shaanxi province	716	2010-2016
12	Datong	40.10	113.33	1067.3	9 km of Datong City, but within area of rapid urbanization, Shanxi Province	914	2014-2017
Rural sites in eastern China (15 sites)							
13	Changde	29.17	111.70	565.0	18 km northwest from Changde city, Hunan province.	344	2013-2016
14	Dongtan	31.52	121.96	10.0	In the Chongmin Island, 30km east of Shanghai city	986	2012-2016
15	ChunAn	29.61	119.05	171.4	151 km southwest from Hangzhou city, Zhejiang province.	1286	2011-2015
16	Huimin	37.48	117.53	11.7	100 km Northeast of Jinan City, Shandong Province	2243	2009-2017
17	Lin'an	30.30	119.73	138.6	150 km northeast of Shanghai, and 50 km west of Hangzhou city, Zhejiang province	1834	2011-2015
18	Mt.Longfeng	44.73	127.60	330.5	In Wuchang county, 175 km northeast of Harbin city, Heilongjiang Province	1515	2012-2016
19	Fuyang	30.07	119.95	17.0	44.1 km southwest from Hangzhou city, Zhejiang province.	710	2014-2015
20	Shangdianzi	40.65	117.12	293.0	In Miyun county, 150 km northeast to Beijing city.	1520	2014-2017
21	Yushe	37.07	112.98	1041.5	1.5 km east of Yushe city in Shanxi Province	1479	2013-2017
22	Dengfeng	34.46	113.02	350.0	75 km Southwest of Zhengzhou City, Henan Province	712	2013
23	Huainan	32.65	117.02	52.0	In the central of Hefei City, Anhui Province	794	2014-2015
24	Jiande	29.45	119.28	89.0	In the southwest from Hangzhou city, Zhejiang province.	1550	2011-2015
25	Tonglu	29.80	119.64	46.1	100 km northwest from Hangzhou city, Zhejiang province.	1717	2011-2015
26	Xiaoshan	30.16	120.25	14.0	In the south of Hangzhou city, Zhejiang province.	600	2014-2015
27	Xiyong	22.28	114.33	155.2	In the eastern of Shenzhen city, Guangdong province.	189	2016
Urban sites (23 sites)							
28	Anshan	41.08	123.00	23.0	In Anshan city, central Liaoning province	193	2009-2013
29	Beijing-Nanjiao	39.80	116.47	31.3	In the southeast Beijing at city margin	1732	2014-2017
30	Beijing-CAMS	39.93	116.32	106.0	Chinese Academy of Meteorological Sciences in Beijing	1113	2012-2018
31	Chengdu	30.65	104.03	496.0	In Chengdu city, Sichuan province.	55	2014-2015
32	Dalian	38.90	121.63	91.5	Southeast coastal city in Liaoning Province	736	2012-2015
33	Fushun	41.88	123.95	80.0	In Fushun city, central Liaoning province.	231	2009-2013
34	Hangzhou	30.23	120.17	42.0	In Hangzhou city, Zhejiang province.	1663	2011-2015

35	Hefei	31.98	116.38	92.0	In Hefei city, Anhui province.	197	2016
36	Jiaozuo	35.18	113.25	113.0	In the center of Jiaozuo city, Henan province.	981	2016-2017
37	Lanzhou	36.05	103.88	1517.3	In Lanzhou city, Gansu province.	1493	2013-2017
38	Nanjing	32.05	118.77	99.3	In Nanjing city, Jiangsu province	1258	2007-2015
39	Nanning	22.82	108.35	172.0	In Nanning city, Guangxi province	286	2013-2017
40	Panyu	23	113.35	145.0	In district of Guangzhou city, Guangdong Province	436	2012-2016
41	Shanghai	31.22	121.55	14.0	In Pudong district of Shanghai city	144	2016
42	Shenyang	41.77	123.50	60.0	In Shenyang city, central Liaoning province.	541	2009-2013
43	Tianjin	39.10	117.17	3.3	Northern coastal city in North China Plain	1705	2013-2017
44	Urumqi	43.78	87.62	935.0	In Urumqi city, Xijiang province	1411	2012-2017
45	Xi'an	34.43	108.97	363.0	20 km north of center of Xian city, but within Jing RiverIndustrial District, Shaanxi province	652	2012-2016
46	Yinchuan	38.48	106.22	1111.5	In Yinchuan city, Ningxia province.	124	2017
47	Zhengzhou	34.78	113.68	99.0	In Zhengzhou city, Henan province.	1485	2013-2017
48	Shijiazhuang	38.03	114.53	75.0	In the center of Shijiazhuang city, Hebei province.	1178	2015-2017
49	Wuhan	30.32	114.21	30	In the center of Wuhan city, Hubei province	220	2008
50	Zhuzilin	22.32	114.00	63.0	In the central of Shenzhen city, Guangdong province.	915	2010-2017

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**Table 2. Annual data for aerosol microphysical properties, optical and direct radiative parameters**

No.	Site	<sup>a</sup> ReffT	<sup>a</sup> ReffF	<sup>a</sup> ReffC	<sup>a</sup> VolT	<sup>a</sup> VolF	<sup>a</sup> VolC	<sup>a</sup> AODt	<sup>b</sup> Alpha	<sup>a</sup> FMF	<sup>a</sup> SSAT	<sup>a</sup> Image	<sup>a</sup> Real	<sup>a</sup> AAOD	<sup>a</sup> BOA	<sup>a</sup> TOA
Remote sites (3 sites)																
1	Akedala	0.36	0.14	2.45	0.06	0.02	0.04	0.17	1.13	0.81	0.90	0.0117	1.4540	0.02	-33.65	-0.42
2	Lhasa	0.64	0.13	2.26	0.05	0.01	0.04	0.10	0.77	0.66	0.90	0.0106	1.5541	0.01	-22.13	-5.04
3	Shangri-La	0.39	0.14	2.33	0.03	0.01	0.02	0.10	1.19	0.85	0.93	0.0086	1.4626	0.01	-17.43	-8.93
	Average	0.47	0.14	2.35	0.05	0.01	0.03	0.12	1.03	0.77	0.91	0.0103	1.4902	0.01	-24.40	-4.79
Arid and semi-arid sites (6 sites)																
4	Dunhuang	0.62	0.14	1.52	0.15	0.02	0.13	0.33	0.48	0.44	0.88	0.0103	1.5491	0.04	-63.61	-8.96
5	Ejina	0.56	0.14	1.78	0.11	0.02	0.09	0.24	0.64	0.52	0.89	0.0116	1.5265	0.03	-47.66	-7.20
6	Minqin	0.56	0.13	1.87	0.13	0.02	0.11	0.30	0.68	0.59	0.86	0.0145	1.5430	0.04	-59.83	-5.01
7	Tazhong	0.71	0.14	1.38	0.30	0.03	0.27	0.60	0.25	0.35	0.92	0.0054	1.5257	0.05	-91.20	-23.49
8	Xilinhot	0.48	0.13	2.45	0.08	0.02	0.05	0.21	1.03	0.78	0.89	0.0139	1.5183	0.02	-37.14	-7.47
9	Tongyu	0.39	0.13	2.36	0.07	0.02	0.05	0.23	1.16	0.82	0.88	0.0179	1.5377	0.03	-39.13	-8.87
	Average	0.55	0.14	1.89	0.14	0.02	0.12	0.32	0.71	0.58	0.89	0.0123	1.5334	0.03	-56.43	-10.17
Rural sites on the Chinese Loess Plateau or nearby (3 sites)																
10	Mt.Gaolan	0.58	0.14	2.03	0.16	0.03	0.13	0.36	0.81	0.64	0.89	0.0108	1.5154	0.04	-59.36	-20.87
11	Yulin	0.53	0.15	2.05	0.11	0.03	0.08	0.32	0.84	0.72	0.89	0.0122	1.5070	0.03	-56.81	-9.09
12	Datong	0.35	0.13	2.15	0.19	0.09	0.10	0.58	1.15	0.83	0.86	0.0171	1.4905	0.09	-107.86	-13.71
	Average	0.49	0.14	2.08	0.15	0.05	0.10	0.42	0.93	0.73	0.88	0.0134	1.5043	0.05	-74.67	-14.56
Rural sites in eastern China (15 sites)																
13	Changde	0.32	0.16	2.18	0.14	0.07	0.07	0.58	1.15	0.88	0.93	0.0101	1.4619	0.04	-75.33	-31.44
14	Dongtan	0.37	0.16	2.12	0.17	0.08	0.09	0.62	1.21	0.86	0.93	0.0080	1.4624	0.04	-79.41	-33.18
15	ChunAn	0.30	0.18	2.30	0.19	0.12	0.08	0.81	1.22	0.92	0.94	0.0066	1.4095	0.04	-86.49	-46.48
16	Huimin	0.36	0.15	2.07	0.22	0.10	0.12	0.83	1.14	0.86	0.89	0.0147	1.4852	0.08	-111.58	-25.49

17	Lin'an	0.29	0.17	2.24	0.21	0.12	0.09	0.87	1.29	0.91	0.93	0.0089	1.4172	0.06	-93.09	-41.73
18	Mt.Longfeng	0.28	0.15	2.44	0.08	0.04	0.04	0.34	1.38	0.90	0.89	0.0165	1.4647	0.03	-51.17	-11.34
19	Fuyang	0.29	0.17	2.28	0.21	0.13	0.09	0.89	1.31	0.92	0.94	0.0070	1.4147	0.05	-91.69	-42.29
20	Shangdianzi	0.40	0.15	2.33	0.12	0.05	0.07	0.43	1.17	0.86	0.89	0.0148	1.4840	0.04	-59.99	-20.58
21	Yushe	0.41	0.15	2.18	0.14	0.06	0.08	0.50	1.07	0.84	0.92	0.0090	1.4878	0.03	-66.72	-25.99
22	Dengfeng	0.39	0.15	2.03	0.23	0.09	0.13	0.79	1.02	0.83	0.89	0.0131	1.4782	0.08	-104.78	-35.84
23	Huainan	0.30	0.17	2.25	0.21	0.13	0.08	0.91	1.17	0.92	0.88	0.0166	1.4308	0.10	-129.17	-24.44
24	Jiande	0.29	0.17	2.18	0.20	0.12	0.08	0.84	1.34	0.91	0.92	0.0099	1.4085	0.06	-91.06	-40.07
25	Tonglu	0.29	0.17	2.20	0.20	0.12	0.08	0.83	1.31	0.91	0.93	0.0091	1.4269	0.06	-89.82	-41.28
26	Xiaoshan	0.28	0.17	2.24	0.22	0.13	0.09	0.87	1.35	0.91	0.93	0.0082	1.4134	0.06	-95.23	-40.39
27	Xiyong	0.33	0.16	2.43	0.11	0.06	0.05	0.41	1.32	0.89	0.94	0.0074	1.4072	0.02	-53.18	-25.45
	Average	0.33	0.16	2.23	0.18	0.09	0.08	0.70	1.23	0.89	0.92	0.0107	1.4435	0.05	-85.25	-32.40
Urban sites (23 sites)																
28	Anshan	0.36	0.17	2.24	0.26	0.12	0.14	0.94	1.12	0.86	0.89	0.0158	1.4759	0.10	-117.99	-39.66
29	Beijing-Nanjiao	0.45	0.15	2.33	0.19	0.07	0.12	0.65	1.12	0.84	0.92	0.0100	1.4939	0.05	-82.06	-29.43
30	Beijing-CAMS	0.50	0.16	2.37	0.19	0.07	0.12	0.65	1.12	0.79	0.90	0.0115	1.5108	0.05	-72.66	-29.10
31	Chengdu	0.34	0.21	2.26	0.26	0.16	0.10	1.17	1.12	0.92	0.97	0.0033	1.4116	0.04	-110.42	-52.21
32	Dalian	0.35	0.16	2.24	0.16	0.08	0.09	0.62	1.22	0.87	0.93	0.0095	1.4584	0.04	-75.50	-37.42
33	Fushun	0.38	0.17	2.34	0.22	0.09	0.12	0.80	1.12	0.87	0.84	0.0244	1.4954	0.11	-116.91	-19.59
34	Hangzhou	0.30	0.17	2.21	0.22	0.12	0.10	0.87	1.30	0.90	0.91	0.0109	1.4337	0.07	-31.57	-40.16
35	Hefei	0.29	0.15	2.37	0.18	0.10	0.08	0.69	1.28	0.90	0.85	0.0195	1.4253	0.10	-105.83	-19.22
36	Jiaozuo	0.35	0.16	2.17	0.20	0.10	0.10	0.76	1.14	0.88	0.91	0.0105	1.4722	0.05	-92.29	-39.35
37	Lanzhou	0.54	0.14	2.04	0.28	0.06	0.22	0.66	0.81	0.66	0.83	0.0197	1.5193	0.10	-126.17	-13.81
38	Nanjing	0.33	0.16	2.16	0.25	0.12	0.12	0.94	1.13	0.88	0.88	0.0154	1.4446	0.10	-143.38	-28.29
39	Nanning	0.30	0.18	2.53	0.20	0.13	0.06	0.97	1.36	0.95	0.92	0.0107	1.4272	0.07	-121.92	-33.35
40	Panyu	0.26	0.16	2.29	0.16	0.10	0.06	0.69	1.43	0.93	0.90	0.0137	1.4155	0.07	-96.03	-26.56

41	Shanghai	0.40	0.15	1.93	0.19	0.08	0.11	0.68	1.10	0.84	0.88	0.0142	1.4814	0.07	-106.89	-24.34
42	Shenyang	0.31	0.16	2.23	0.22	0.12	0.10	0.89	1.20	0.90	0.84	0.0253	1.4589	0.14	-144.88	-15.02
43	Tianjin	0.42	0.16	2.26	0.23	0.10	0.13	0.83	1.11	0.86	0.89	0.0134	1.4957	0.07	-108.09	-33.26
44	Urumqi	0.48	0.14	2.14	0.15	0.04	0.10	0.42	0.93	0.75	0.85	0.0192	1.5371	0.05	-70.55	-11.74
45	Xi'an	0.37	0.16	1.85	0.26	0.11	0.15	0.98	0.98	0.82	0.88	0.0150	1.4888	0.10	-132.55	-35.93
46	Yinchuan	0.38	0.14	2.02	0.11	0.04	0.07	0.37	1.12	0.81	0.94	0.0054	1.4930	0.02	-48.67	-21.89
47	Zhengzhou	0.43	0.18	2.22	0.28	0.12	0.16	0.99	1.10	0.86	0.95	0.0045	1.4626	0.04	-101.10	-46.18
48	Shijiazhuang	0.40	0.16	2.28	0.26	0.12	0.14	0.95	1.09	0.87	0.88	0.0154	1.4754	0.09	-125.05	-33.66
49	Wuhan	0.34	0.17	2.22	0.22	0.12	0.10	1.00	1.16	0.91	0.88	0.0196	1.4779	0.11	-171.80	-20.40
50	Zhuzilin	0.27	0.17	2.45	0.15	0.09	0.05	0.66	1.45	0.94	0.96	0.0049	1.4438	0.03	-73.16	-40.65
	Average	0.37	0.16	2.22	0.21	0.10	0.11	0.79	1.15	0.86	0.90	0.0136	1.4695	0.07	-103.28	-30.05

1307 Table 1 (Continued)

1308 <sup>a</sup> Optical parameters at a wavelength of 440 nm.

1309 <sup>b</sup> Angström exponents between 440 and 870 nm.