

1 **The role of dust mineral composition in atmospheric radiation and pollution in North**  
2 **China: new insights from EMIT and two-way coupled modeling**

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11  
12 **Abstract**

13 Mineral dust is a major atmospheric aerosol influencing Earth's energy balance  
14 through aerosol-radiation (ARI) and aerosol-cloud interactions (ACI). While homogeneous  
15 dust effects have been studied, the impact of mineralogical composition on regional  
16 meteorology and air quality remains underexplored, limiting accurate forecasting of dust  
17 storm impacts, especially in dust belt regions. In this study, we used a two-way coupled  
18 WRF-CHIMERE model with three mineralogical dust atlases (Nickovic et al. (2012)  
19 (N2012), Journet et al. (2014) (J2014), and a new dataset, Li et al. (2024) (L2024), from  
20 the Earth Surface Mineral Dust Source Investigation (EMIT)) to evaluate ARI effects  
21 during the March 2021 dust storm in North China. Results showed significant spatial  
22 variations in radiative forcing due to mineralogical differences. Bulk dust (without  
23 considering mineralogy) caused an average shortwave radiative forcing of  $-5.72 \text{ W/m}^2$ ,  
24 while mineral-specific forcings increased this by up to  $+0.10 \text{ W/m}^2$ . Integrating EMIT data  
25 reduced  $\text{PM}_{10}$  biases by over 15% in high-concentration regions and improved ozone  
26 predictions, with localized changes of  $-2.46$  to  $+3.52 \mu\text{g/m}^3$ . Hematite's strong absorption  
27 and quartz's reflective properties were key in altering radiative and air quality outcomes.  
28 Compared to scenarios of bulk dust, the consideration of ARI effects of mineralogical  
29 compositions can increase  $\text{PM}_{10}$  concentration by up to  $1189.48 \mu\text{g/m}^3$  in dust source  
30 regions. Future research perspectives on the utilization of high-resolution EMIT data in  
31 two-way coupled meteorology and air quality models for investigating the ACI effects of

32 mineralogical dust on cloud microphysics are proposed.

### 33 **1 Introduction**

34 Mineral dust, a dominant component of global atmospheric aerosols, primarily  
35 originates from wind erosion in arid and semi-arid regions (Schepanski, 2018; Shao et al.,  
36 2011). It can affect the Earth's energy balance through direct scattering and absorption of  
37 solar, i.e. aerosol-radiation interaction (ARI), as well as indirect effects on cloud properties  
38 by acting as cloud condensation nuclei and ice nuclei, i.e., aerosol-cloud interaction (ACI)  
39 (Choobari et al., 2014; Kok et al., 2023). By altering biogeochemical cycles, atmospheric  
40 chemistry and visibility, and air quality, transported mineral dust can exacerbate economic  
41 losses, and health risks (Adebisi et al., 2023; Cwiertny et al., 2008; Duniway et al., 2019;  
42 Maher et al., 2010; Tong et al., 2023). Accurate forecasting of mineral dust events is crucial  
43 to mitigate these adverse impacts.

44 Numerous studies have demonstrated that the magnitude of dust ARI and ACI effects  
45 is significantly influenced by its mineralogical composition. For instance, iron oxides,  
46 particularly hematite and goethite, have been identified as key components responsible for  
47 dust absorption of solar radiation, as evidenced by both observational and modeling studies  
48 (Alfaro et al., 2004; Gómez Maqueo Anaya et al., 2024; Lafon et al., 2006; Li et al., 2022;  
49 Obiso et al., 2024; Scanza et al., 2015; Song et al., 2024). Concurrently, a growing body of  
50 research has explored the impact of various dust mineral compositions, including hematite,  
51 corundum, kaolinite, mica, montmorillonite, quartz, calcite, illite, amorphous silicon,  
52 aluminum silicate, and potassium feldspar, on ice nucleation processes. Among these,  
53 potassium feldspar has emerged as a crucial component for dust nucleation activation  
54 (Harrison et al., 2016; Kumar et al., 2018). However, a notable gap exists in our  
55 understanding of how specific mineral compositions impact meteorology and air quality  
56 through ARI and ACI effects. Prior research has predominantly focused on homogeneous  
57 dust aerosols, assuming globally uniform composition and optical properties. However,  
58 this assumption introduces regional inaccuracies in estimating the impacts of dust aerosols,

59 which remain poorly understood due to uncertainties in dust composition (Ke et al., 2022;  
60 Klingmüller et al., 2019; Kok et al., 2017).

61 Many efforts have been directed to improve simulations of dust mineralogy and its  
62 representation in numerical models (Balkanski et al., 2021; Gómez Maqueo Anaya et al.,  
63 2024; Gonçalves Ageitos et al., 2023; Li et al., 2021, 2022, 2024; Li and Sokolik, 2018;  
64 Menut et al., 2020; Obiso et al., 2024; Scanza et al., 2015; Solomos et al., 2023b, a; Song  
65 et al., 2024). Most of the above are offline models, with only two studies conducting two-  
66 way feedback simulations with only WRF-Chem (Li and Sokolik, 2018) and WRF-  
67 CHIMERE (Menut et al., 2020) being applied. However, both of these studies are derived  
68 from artificially generated data and lack effective ground-based validation, as discussed in  
69 Claquin et al. (1999), Nickovic et al. (2012) (N2012 hereafter), and Journet et al. (2014)  
70 (J2014 hereafter). These validations predominantly focus on agricultural regions rather  
71 than the arid and semi-arid areas that are major sources of dust emissions (Green et al.,  
72 2020). EMIT instrument provides a new approach to invert and obtain the surface soil  
73 mineral composition and further assess the ARI and ACI effects of dust minerals (Connelly  
74 et al., 2021). To the best of our knowledge, no prior research has investigated the impact  
75 of dust on regional meteorology and air quality while considering its mineral speciation  
76 using two-way coupled models with three different mineralogical dust atlases.

77 Since the aerosol nucleation processes (ACI effects) of specific mineral components  
78 are not represented in the current two-way coupled WRF–CHIMERE framework, the  
79 present study concentrates on the ARI effects of dust minerals. This focus ensures a clear  
80 and robust assessment of how mineralogical composition influences radiative processes,  
81 without introducing additional uncertainties arising from incomplete cloud-related  
82 parameterizations. In this study, we employ a two-way coupled WRF–CHIMERE model  
83 with three mineralogical databases to investigate how dust composition influences  
84 radiation and meteorology in North China during a severe dust storm. Section 2 describes  
85 the model configuration and data sources, Section 3 presents the simulations with emphasis

86 on ARI-induced impacts on meteorology and air quality, and Section 4 summarizes the  
87 main findings.

88

## 89 **2 Methodology and data**

### 90 **2.1 Model configurations and data sources**

91 The two-way coupled WRF model version 3.7.1 and CHIMERE model version  
92 2020r3 were employed to simulate the ARI and ACI effects of mineralogical dust particles  
93 on meteorology and air quality over North China from March 12 to March 15, 2021, as  
94 shown in Fig. A1. The exchanges between meteorological and air quality variable are  
95 accomplished through the OASIS coupler (Briant et al., 2017). The simulation was  
96 conducted at a horizontal resolution of 27 km, with 165 grid cells in the east-west direction  
97 and 87 in the north-south direction, and the study domain is depicted in Fig. A1. The model  
98 has 33 vertical levels from surface to 50 hPa with 13 layers in the bottom 1km and the  
99 bottom thickness being 24.5 m. The Rapid Radiative Transfer Model for General  
100 circulation models (RRTMG) shortwave and longwave radiation schemes were employed  
101 to investigate the ARI effects (Briant et al., 2017). Additionally, the Thompson cloud  
102 microphysics scheme was utilized to assess the impacts of ACI (Tuccella et al., 2019). The  
103 initial and boundary conditions (ICs and BCs) for non-dust aerosols are prescribed by the  
104 LDMZ-INCA model, while those for dust aerosols are determined by the GOCART model.  
105 The options of other physics and chemistry schemes are presented in Table A1. The dry  
106 depositions are treated as described in Zhang et al. (2001). The parameterizations for the  
107 removal of dust particles below clouds by raindrops and snow are based on the methods  
108 proposed by Willis & Tattelman (1989) and Wang et al. (2014), respectively. Inline mineral  
109 dust emissions, incorporating mineralogy, are computed using a  $u^*$  threshold and a dust  
110 production model for saltation (Kok et al., 2014; Shao and Lu, 2000). The model accounts  
111 for the impact of soil moisture on suppressing mineral dust emissions (Fécan et al., 1998).  
112 To minimize meteorological bias, a spectral nudging approach is applied (Menut et al.,  
113 2024).

114 For the calculations of ARI effects in WRF-CHIMERE, refractive indices  
115 corresponding to these mineralogical species are provided in Table 2 of Menut et al. (2020).  
116 Concerning shortwave (SW) radiation, the aerosol optical properties, encompassing single  
117 scattering albedos and asymmetry factors at 400 and 600 nm, as well as the aerosol optical  
118 depth (AOD) at 300, 400, and 999 nm, calculated using Fast-JX, were interpolated or  
119 extrapolated to obtain values at 14 SW intervals (Briant et al., 2017; Gao et al., 2022). AOD  
120 at 16 longwave (LW) intervals ranging from 3400 to 55600 nm are directly used to  
121 calculate LW radiation.

122 To evaluate the performance of the WRF-CHIMERE model with and without  
123 mineralogical dust emissions, we compiled a comprehensive set of environmental  
124 observations. Hourly PM<sub>10</sub> and O<sub>3</sub> concentrations (132 observations) were obtained from  
125 <https://quotsoft.net/air/>, which aggregates official monitoring data from the Ministry of  
126 Ecology and Environmental Protection of China. Shortwave radiation (SSR) data (59  
127 hourly measurements) were obtained from Tang et al. (2019), with the original  
128 measurements sourced from the China Meteorological Administration. Hourly surface  
129 meteorological data (844 observations) were also obtained from the China Meteorological  
130 Administration (<https://data.cma.cn>). All datasets were subjected to quality control  
131 procedures, including checks for outliers, unit consistency, and temporal alignment, to  
132 ensure reliability and integrity.

133

## 134 **2.2 Mineral dust atlases**

135 Accurate soil composition data are essential for partitioning dust emission fluxes into  
136 contributions from individual minerals. Mineral density and refractive index data were  
137 obtained from Menut et al. (2020). Three global mineralogical composition datasets  
138 (N2012, J2014, and EMIT) provide information on 12 mineral species (Table 1) at different  
139 spatial resolutions (1 km × 1 km and 0.5° × 0.5°).

140 To ensure a consistent spatial framework and facilitate cross-dataset integration, the  
141 N2012 dataset (originally provided at 1 km × 1 km resolution and available at

142 <http://www.seevccc.rs/GMINER30>) was resampled to  $0.5^\circ \times 0.5^\circ$ . The J2014 dataset,  
143 widely employed in the WRF–CHIMERE modeling framework, includes 12 mineral  
144 species distributed across the clay and/or silt fractions (see Table 2 in [Menut et al., 2020](#)).  
145 In contrast, the EMIT dataset (<https://earth.jpl.nasa.gov/emit/data/data-products>) required  
146 additional preprocessing, as it reports only normalized spectral abundances rather than  
147 mineral mass fractions. These spectral abundances were therefore recalculated to represent  
148 the normalized mass proportions of each mineral in each substrate. Furthermore, EMIT  
149 does not include data for feldspar and quartz, necessitating additional correction procedures  
150 described below.

151 When the total mineral composition from EMIT summed to less than 100%, indicating  
152 missing mineral contributions, the residual fraction was assigned to quartz and feldspar  
153 based on their relative proportions in J2014 or N2012. Because EMIT reports illite and  
154 mica as a single category, their individual abundances were separated according to the  
155 ratios found in N2012 or J2014. For minerals that occur in both clay and silt fractions,  
156 EMIT values were partitioned following the relative contributions from N2012 or J2014.

157 For minerals not directly observed by EMIT (e.g., quartz and feldspar), their mass  
158 fractions were estimated using soil-type conversion methods from previous studies  
159 ([Claquin et al., 1999](#); [Journet et al., 2014](#)). The spatial distributions of clay and silt were  
160 obtained from the global SoilW texture dataset  
161 (<http://globalchange.bnu.edu.cn/research/soilw>) at 1 km resolution and resampled to  $0.5^\circ$   
162 to match EMIT data. Similarly, the J2014 and N2012 mineral datasets were resampled to  
163  $0.5^\circ$  resolution. Major minerals extracted from EMIT L3 include calcite, dolomite, chlorite,  
164 goethite, gypsum, hematite, illite+muscovite, kaolinite, montmorillonite, and vermiculite.  
165 Notably, in the official EMIT L3B dataset ([https://data.lpdaac.earthdatacloud.nasa.gov/lp-prod-protected/EMITL3ASA.001/EMIT\\_L3\\_ASA\\_001/EMIT\\_L3\\_ASA\\_001.nc](https://data.lpdaac.earthdatacloud.nasa.gov/lp-prod-protected/EMITL3ASA.001/EMIT_L3_ASA_001/EMIT_L3_ASA_001.nc)), illite  
166 and muscovite are combined because they were jointly identified during the Tetracorder  
167 analysis of L2B data using mineral groups 1 and 2 and the corresponding band depths  
168

169 ([https://github.com/nasa/EMIT-Data-Resources/blob/main/data/mineral\\_grouping\\_matrix\\_20230503.csv](https://github.com/nasa/EMIT-Data-Resources/blob/main/data/mineral_grouping_matrix_20230503.csv)).

171 The EMIT mineral fractions were normalized so that their sum at each grid point did  
172 not exceed unity. Any remaining fraction was attributed to quartz and feldspars according  
173 to their relative proportions in J2014 or N2012. To ensure consistency with the CHIMERE  
174 mineral representation, dolomite was merged into calcite, illite+muscovite was separated  
175 into illite and mica, and montmorillonite was treated as smectite. The mineral fractions  
176 were then converted to density-weighted values and renormalized at each grid point so that  
177 the total sum equaled one. Finally, each mineral was partitioned into clay and silt fractions  
178 based on the J2014 ratios, and the resulting fractions were normalized using Equations (1)–  
179 (4). The processed dataset was exported as a NetCDF file to serve as input for the  
180 CHIMERE model.

181 To ensure mineral mass balance and model consistency, a normalization and  
182 partitioning procedure was applied as follows. Equation (1) defines the total mass fraction  
183 ( $MF_j$ ) of mineral  $j$  as the sum of its contributions from the clay ( $MFC_j$ ) and silt ( $MFS_j$ )  
184 fractions:

$$185 \quad MF_j = MFC_j + MFS_j \text{ for all } j \in M_{\text{CHIMERE}} \quad (1)$$

186 Equation (2) enforces a normalization constraint so that the sum of all mineral mass  
187 fractions equals unity at each grid point.

$$188 \quad 1 = \sum_{j \in M_{\text{CHIMERE}}} MF_j \quad (2)$$

189 The normalized total fraction of each mineral ( $MF_j^*$ ) was then redistributed between  
190 clay and silt according to their relative contributions in the reference dataset (J2014 or  
191 N2012), as shown in Equations (3) and (4):

$$192 \quad MFS_j^* = MF_j^* \frac{MFS_j}{MFS_j + MFC_j} \quad (3)$$

$$193 \quad MFC_j^* = MF_j^* \frac{MFC_j}{MFS_j + MFC_j} \quad (4)$$

194 Here,  $MFS_j^*$  and  $MFC_j^*$  represent the normalized mass fractions of mineral  $j$  in the

195 silt and clay fractions, respectively. The weighting terms  $MFS_j$  and  $MFC_j$  preserve the  
 196 clay–silt distribution patterns derived from the reference datasets while maintaining the  
 197 normalized total ( $MF_j^*$ ).

198 Table 1. Mineralogical compositions in different datasets.

Mineral	Clay		Silt		EMIT
	N2012	J2014	N2012	J2014	
Smectite	✓	✓	✗	✗	✓
Illite	✓	✓	✗	✗	✓ <sup>†</sup>
Hematite	✓	✓	✓	✗	✓
Feldspar	✗	✓	✓	✓	✗
Kaolinite	✓	✓	✗	✗	✓
Calcite	✓	✓	✓	✓	✓
Quartz	✓	✓	✓	✓	✗
Gypsum	✗	✗	✓	✓	✓
Vermiculite	✗	✓	✗	✗	✓
Chlorite	✗	✓	✗	✓	✓
Goethite	✗	✗	✗	✓	✓
Mica	✗	✗	✗	✓	✓ <sup>†</sup>
Resolution	1km	0.5°	1km	0.5°	0.5°

199 <sup>†</sup> indicates the content of illite + mica

200

### 201 2.3 Scenario set up

202 Ten parallel WRF-CHIMERE simulations were performed to investigate the influence  
 203 of mineralogical dust on meteorology and air quality in China, employing three distinct  
 204 mineralogical atlases, as illustrated in Figs. 1 and A5-A7. Each simulation was conducted  
 205 both with and without enabling ARI effects, as detailed in Table 2, to isolate and compare  
 206 the effects of mineralogical dust under different modeling conditions.

207 Simulations without ARI effects (Dust\_NO, N2012\_default\_NO, N2012\_EMIT\_NO,  
 208 J2014\_default\_NO, J2014\_EMIT\_NO) were specifically designed to identify the direct  
 209 impact of mineralogical dust on meteorology and air quality, independent of the radiative

210 feedbacks induced by aerosols. These No\_ARI simulations served as a baseline for  
 211 assessing how mineralogical compositions affect meteorology and air quality in the  
 212 absence of aerosol-radiation feedback mechanisms.

213 In contrast, simulations with ARI enabled (Dust\_ARI, N2012\_default\_ARI,  
 214 N2012\_EMIT\_ARI, J2014\_default\_ARI, J2014\_EMIT\_ARI) were used to quantify the  
 215 additional effects arising from aerosol-radiation interactions. By comparing simulations  
 216 with and without ARI for each mineralogical atlas (e.g., N2012\_default\_ARI –  
 217 N2012\_default\_NO), the differential impact of ARI effects on meteorology and air quality  
 218 for various dust compositions could be identified. This approach highlights how  
 219 mineralogical properties of dust influence the strength and nature of ARI effects, thereby  
 220 modulating key atmospheric processes such as radiation balance, temperature profiles, and  
 221 pollutant distributions.

222 To evaluate the discrepancies in ARI effects among the mineralogical atlases,  
 223 differences in the ARI impacts between EMIT-derived and default dust compositions were  
 224 analyzed for both N2012 and J2014 datasets. For example, comparisons such as  
 225 (N2012\_EMIT\_ARI – N2012\_EMIT\_NO) versus (N2012\_default\_ARI –  
 226 N2012\_default\_NO) provide insight into the extent to which higher-resolution, satellite-  
 227 derived mineralogical data influence ARI effects relative to default atlas-based  
 228 representations. Similar comparisons were performed for the J2014 dataset.

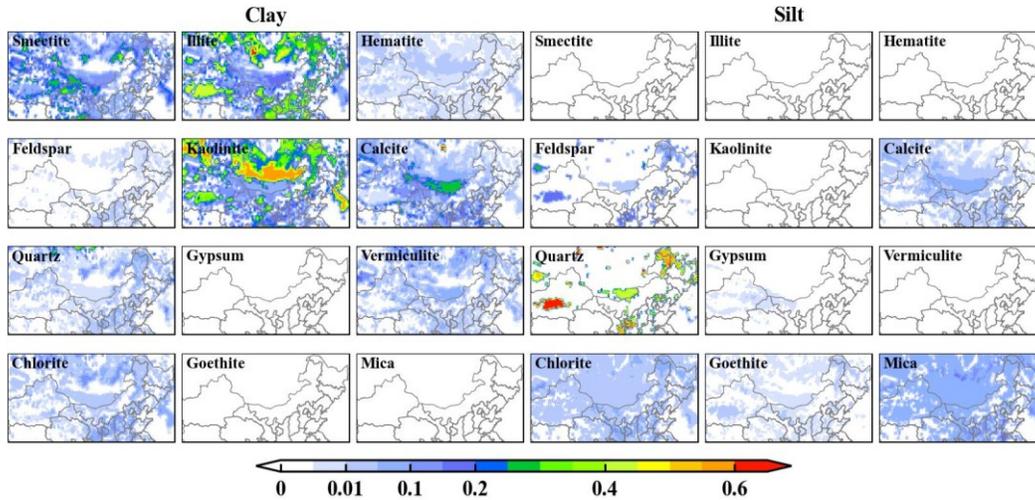
229

230 Table 2. Summary of dust emission scenarios and aerosol feedback configurations for  
 231 different simulation settings.

Scenario	Emission	Online choice	Coupling type	Aerosol feedback
Dust_NO	Bulk dust + anthropogenic emissions	[online] = 1	[cpl_case] = 1	No feedbacks
Dust_ARI		[online] = 1	[cpl_case] = 2	ARI effects
N2012_default_NO	N2012_default dust + anthropogenic emissions	[online] = 1	[cpl_case] = 1	No feedbacks
N2012_default_ARI		[online] = 1	[cpl_case] = 2	ARI effects
N2012_EMIT_NO	N2012_EMIT dust + anthropogenic emissions	[online] = 1	[cpl_case] = 1	No feedbacks
N2012_EMIT_ARI		[online] = 1	[cpl_case] = 2	ARI effects
J2014_default_NO	J2014_default dust + anthropogenic emissions	[online] = 1	[cpl_case] = 1	No feedbacks

J2014_default_ARI		[online] = 1	[cpl_case] = 2	ARI effects
J2014_EMIT_NO	J2014_EMIT dust + anthropogenic emissions	[online] = 1	[cpl_case] = 1	No feedbacks
J2014_EMIT_ARI		[online] = 1	[cpl_case] = 2	ARI effects

232



233

234 Figure 1. Spatial distribution of content for the different mineral dust species in the silt and clay fraction  
 235 of the soil for original J2014 mineralogical data.

236

### 237 3 Results and discussion

#### 238 3.1 Evaluation of meteorology and air quality

239 Table 3 presents the evaluation results for observed and simulated surface shortwave  
 240 radiation (SSR), 2-meter temperature (T2), and 10-meter wind speed (WS10) from various  
 241 scenario simulations conducted using the WRF-CHIMERE modeling system. The model  
 242 demonstrates strong overall performance, with correlation coefficients (R) between  
 243 observed and simulated values reaching approximately 0.7 for SSR and WS10, and up to  
 244 0.93 for T2. These results indicate the model’s ability to capture key atmospheric patterns  
 245 and variability across the simulation domain. Nevertheless, systematic biases are apparent,  
 246 particularly in North China, where the model tends to overestimate SSR and WS10 by  
 247 60.69%–68.92% and 17.06%–17.52%, respectively, while underestimating T2 by 0.48%–  
 248 0.58%. The overestimation of SSR likely results from uncertainties in cloud development  
 249 associated with planetary boundary layer and convection parameterizations (Alapaty et al.,  
 250 2012). The systematic overestimation of 10-m wind speed under low-wind conditions

251 commonly observed in weather models mainly stems from outdated geographic data and  
252 coarse spatial resolution (Gao et al., 2024).

253 A comparative analysis of the two configurations, N2012 and J2014, reveals that  
254 WRF-CHIMERE with N2012 generally outperforms J2014 in simulating SSR and T2,  
255 suggesting that the N2012 parameterization better captures radiative and thermodynamic  
256 processes in the region. Conversely, J2014 exhibits improved accuracy in simulating WS10,  
257 implying potential strengths in its representation of near-surface wind dynamics. These  
258 findings highlight the sensitivity of model performance to parameterization schemes and  
259 the need for tailored configurations for specific meteorological variables. The inclusion of  
260 detailed dust mineralogical compositions, while informative for certain applications,  
261 introduces additional complexities that reduce the overall accuracy of simulations.  
262 Specifically, while these compositions help mitigate the overestimation of SSR and the  
263 underestimation of T2, they exacerbate the overestimation of WS10. The integration of  
264 EMIT satellite data provides a significant boost to model performance, highlighting the  
265 value of incorporating high-resolution, real-time observational data to refine the simulation  
266 of atmospheric variables. EMIT data, with its detailed characterization of aerosol and dust  
267 properties, reduces the positive biases in SSR and WS10 while simultaneously minimizing  
268 the negative biases in T2.

269 When comparing the ARI effects of the defaulted mineralogical compositions in  
270 N2012 and J2014 with simulations that implement EMIT satellite data, the latter shows a  
271 clear advantage. Incorporating EMIT data further reduces the positive biases in SSR and  
272 WS10, while simultaneously minimizing the negative biases in T2. This suggests that  
273 EMIT data provides a more precise representation of dust properties and atmospheric  
274 conditions, enhancing the overall reliability of the model.

275

276 Table 3. Statistics analysis of daily averaged SSR, 2-meter temperature (T2) and 10-meter wind speed  
277 (WS10) from different scenario simulations and ground observations in North China including  
278 correlation coefficient (R) and normalized mean bias (NMB).

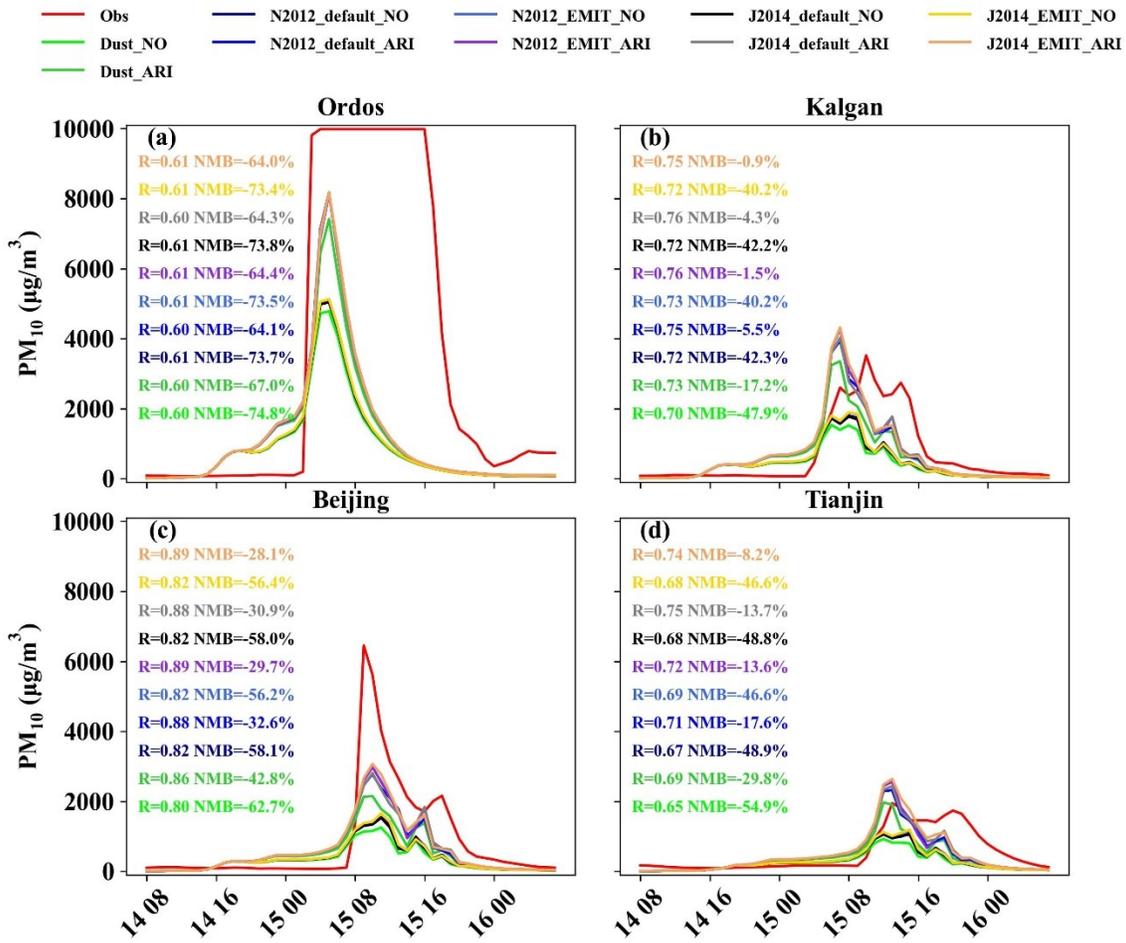
Scenario	SSR	T2	WS10
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	R	NMB (%)	R	NMB (%)	R	NMB (%)
Dust_NO	0.70	68.92	0.93	-0.58	0.71	17.06
Dust_ARI	0.72	60.69	0.94	-0.48	0.72	17.46
N2012_default_NO	0.70	68.92	0.93	-0.58	0.71	17.06
N2012_default_ARI	0.71	61.80	0.94	-0.48	0.72	17.53
N2012_EMIT_NO	0.70	68.92	0.93	-0.58	0.71	17.06
N2012_EMIT_ARI	0.72	60.88	0.94	-0.48	0.72	17.44
J2014_default_NO	0.70	68.92	0.93	-0.58	0.71	17.06
J2014_default_ARI	0.71	61.68	0.94	-0.48	0.72	17.51
J2014_EMIT_NO	0.70	68.92	0.93	-0.58	0.71	17.06
J2014_EMIT_ARI	0.72	61.22	0.94	-0.48	0.72	17.48

279

280 To assess the ability of each scenario simulation to replicate regional PM<sub>10</sub> and O<sub>3</sub>  
281 temporal patterns, Figure 2 presents hourly time series of simulated and in situ PM<sub>10</sub> and  
282 O<sub>3</sub> concentrations at four North China sites: Ordos, Kalgan, Beijing, and Tianjin. These  
283 locations represent key dust aerosol transport pathways, which play a crucial role in the  
284 region's air quality dynamics due to frequent dust storms and anthropogenic emissions. The  
285 time series plots regarding PM<sub>10</sub> and O<sub>3</sub> in Figures 2 and A2 allow for a direct comparison  
286 of model simulations with observed data, revealing important insights into model  
287 performance across different environmental conditions and geographical settings,  
288 respectively. The models exhibit relatively high correlations for PM<sub>10</sub>, with R values  
289 ranging from 0.61 to 0.89 and NMBs from -73.8% to -0.9%. In contrast, their performance  
290 for O<sub>3</sub> is notably weaker, with R values between 0.25 and 0.63 and NMBs from +10.9% to  
291 +84.2%. Among the simulations, N2012 with EMIT data and ARI effects provides the best  
292 performance for PM<sub>10</sub>, while N2012 with EMIT data without aerosol effects performs best  
293 for O<sub>3</sub> concentrations. All models accurately captured the peak PM<sub>10</sub> and O<sub>3</sub> concentrations  
294 observed during the March 12 event in North China, which was characterized by significant  
295 dust emissions and high pollutant levels. This event serves as a key test case for evaluating  
296 the models' responsiveness to extreme atmospheric conditions. However, despite the  
297 overall agreement in peak concentration timings, simulations overestimated O<sub>3</sub> and  
298 underestimated PM<sub>10</sub> at sites with high dust loads, such as Ordos and Kalgan. This

299 discrepancy highlights the challenge of simulating the complex interactions between dust  
300 aerosols, precursor gases, and photochemical reactions, particularly in regions with high  
301 dust deposition and frequent air pollution episodes. Additionally, the models tended to  
302 extend the period of elevated PM<sub>10</sub> concentrations beyond the observed time frame,  
303 suggesting that the processes controlling dust aerosol removal or dispersion were not fully  
304 captured. CHIMERE simulations using J2014 mineralogical data generally outperformed  
305 those using J2012 data, with significant reductions in PM<sub>10</sub> negative NMBs for three of the  
306 four cities, indicating the importance of accurate mineralogical characterization of dust for  
307 improving model predictions. When considering the ARI effects of bulk dust aerosols, the  
308 underestimation of PM<sub>10</sub> was alleviated, whereas the overestimation of O<sub>3</sub> was amplified,  
309 which suggests that incorporating ARI effects helps to better represent the impact of dust  
310 on local radiative forcing and air quality. Moreover, incorporating ARI effects from the  
311 default dust mineralogical atlas further enhanced these trends, underscoring the need for  
312 refined aerosol property data in enhancing model performance. Finally, using Earth-  
313 observing systems such as the EMIT satellite data led to substantial reductions in PM<sub>10</sub>  
314 negative bias at Kalgan, Beijing, and Tianjin, demonstrating the value of remote sensing  
315 data in improving model accuracy, particularly for regions with high aerosol concentrations  
316 and complex emission sources. Although considerable progress has been made in dust  
317 modeling, notable uncertainties remain. The parameterization of threshold friction velocity  
318 and soil texture in emission schemes can still result in underestimated emissions under  
319 strong winds (Zuo et al., 2024). Similarly, simplifications in coarse particle size  
320 distributions may lead to enhanced deposition and transport losses. In addition, incomplete  
321 knowledge of local soil mineralogical composition continues to limit the accurate  
322 simulation of both emission fluxes and heterogeneous chemistry (Pang et al., 2024).  
323  
324



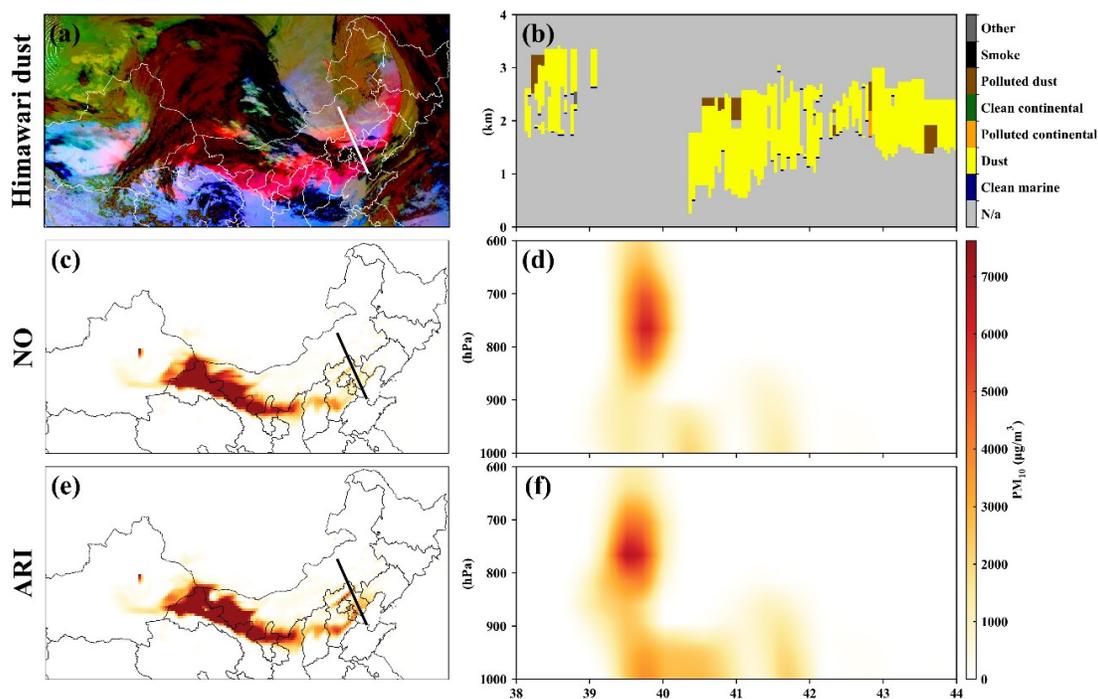
325

326 Figure 2. Statistical metrics between observed and simulated PM<sub>10</sub> concentrations by different  
 327 scenario simulations.

328

329 To evaluate the model performance in simulating the horizontal distribution and  
 330 vertical profile of dust aerosol, Figure 3 presents the false RGB imagery of dust derived  
 331 from Himawari-8 thermal infrared imagery, along with CALIPSO cross sections of 532 nm  
 332 total attenuated backscatter and the vertical feature mask for the overpass of China. The  
 333 figure also includes the corresponding spatial distributions of PM<sub>10</sub> concentrations at 05:00  
 334 UTC on 15th March 2021, a time of significant dust transport in the region. This detailed  
 335 comparison allows for a comprehensive assessment of how well the model captures both  
 336 the horizontal and vertical characteristics of dust aerosol distribution. All six experiments  
 337 display similar dust distributions in the atmosphere, consistent with observations from

338 Himawari-8 and CALIPSO. This suggests that the models effectively capture the general  
 339 spatial patterns of dust transport. On March 15, 2021, the daily domain-averaged PM<sub>10</sub>  
 340 concentration was 533.81  $\mu\text{g m}^{-3}$ , with a 95% confidence interval (CI) of 0.28–5962.95  
 341 (Table S1). Specifically, the false RGB imagery from Himawari-8 clearly indicates the  
 342 presence of dust plumes in the atmosphere, with distinct thermal contrasts that help identify  
 343 the dust layers. The CALIPSO data, which provide vertical profiles of aerosol backscatter,  
 344 further validate the model’s ability to capture the vertical extent and concentration of dust  
 345 layers. These observations are critical for understanding the atmospheric processes  
 346 governing dust dispersion and their impact on air quality. The close agreement between  
 347 model simulations and satellite data across all six experiments also underscores the  
 348 robustness of the model in representing dust aerosol distribution under different simulation  
 349 conditions. This evaluation demonstrates that the models are capable of simulating the  
 350 main features of dust aerosol transport, though further refinement in capturing the fine-  
 351 scale variations and aerosol properties may still be necessary for more accurate predictions  
 352 in future studies.



353

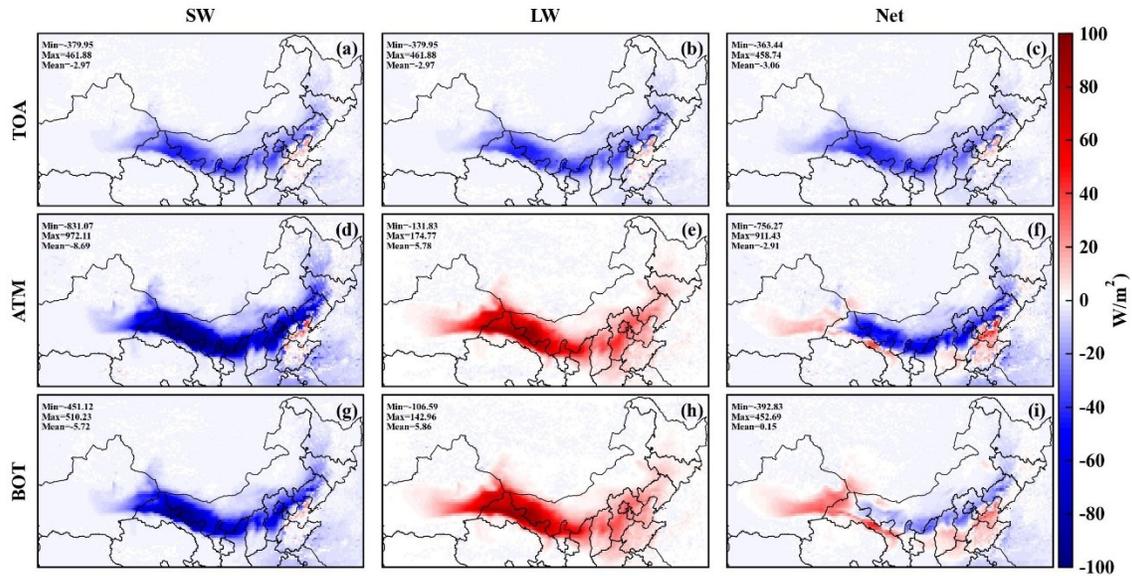
354 Figure 3. The false color imagery of dust from Himawari-8 thermal infrared imagery(a), CALIPSO

355 cross sections of 532 nm total attenuated backscatter and the vertical feature mask for the overpass of  
356 China on 13:00 (local time) 15<sup>th</sup> March 2021 (b), and corresponding horizontal (c and e) and vertical (d  
357 and f) distributions of PM<sub>10</sub> concentrations.

### 358 **3.2 Radiative effect on meteorology**

359 To further investigate the impacts of dust radiation on energy balance, the spatial  
360 distributions of the average shortwave (SW), longwave (LW), and net (NET) radiative  
361 forcing induced by bulk dust on the surface (SFC), in the atmosphere (ATM), and at the  
362 top of the atmosphere (TOA) are presented during the dust episode shown in Figure 4. The  
363 radiative forcing values provide critical insights into the energy exchanges between dust  
364 aerosols and the atmosphere, and their subsequent effects on regional climate dynamics.  
365 For SW radiation forcings, dust aerosols produced cooling effects at all three layers: the  
366 surface, the atmosphere, and the top of the atmosphere. The average SW radiative forcing  
367 was about  $-5.72 \text{ W m}^{-2}$  at the surface,  $-8.69 \text{ W m}^{-2}$  in the atmosphere, and  $-2.97 \text{ W m}^{-2}$   
368 at the TOA, highlighting the significant reduction in solar radiation reaching these layers  
369 due to the scattering and absorption properties of the dust particles. Particularly in the dust  
370 source regions, the cooling effect at the surface exceeded  $-900 \text{ W m}^{-2}$  (Figures 4a, 4d, and  
371 4g), indicating the strong influence of dust on the regional energy budget in these areas.  
372 This is a result of the large dust concentrations and their optical properties, which  
373 effectively block solar radiation from reaching the Earth's surface. In contrast, the dust-  
374 induced LW radiative forcing warmed the surface and atmosphere, with average values  
375 ranging from  $5.78$  to  $5.86 \text{ W m}^{-2}$ . This warming effect is associated with the absorption of  
376 longwave radiation by dust particles, which then re-radiate heat, contributing to local  
377 warming. However, dust particles induced negative LW radiative forcing at the TOA, with  
378 values ranging from  $-461.88$  to  $-379.95 \text{ W m}^{-2}$ , reflecting the downward flux of longwave  
379 radiation absorbed by the aerosols, which reduces the amount of energy reaching the TOA.  
380 The NET radiative forcing, which represents the combined effect of both SW and LW  
381 forcings, was positive at the surface (about  $+0.15 \text{ W m}^{-2}$ ), negative in the atmosphere  
382 (about  $-2.91 \text{ W m}^{-2}$ ), and negative at the TOA (about  $-3.06 \text{ W m}^{-2}$ ), as shown in Figures  
383 4c, 4f, and 4i. The positive NET radiative forcing at the surface suggests a slight net

384 warming effect at ground level, while the negative values in the atmosphere and at the TOA  
 385 indicate an overall cooling effect at these higher altitudes.  
 386

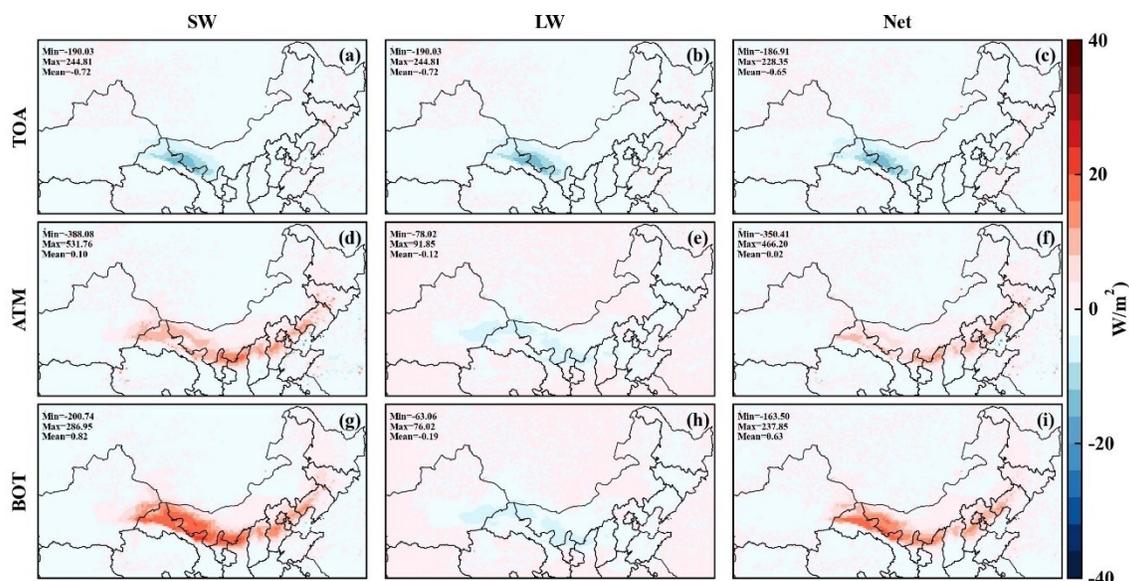


387  
 388 Figure 4. Radiation forcings due to bulk dust enabling ARI effects.  
 389

390 To assess the impact of dust mineralogical composition on radiative forcings, Figure  
 391 S6 illustrates the spatial distribution of radiative differences, considering the ARI effects  
 392 of bulk dust and comparing them to the default N2012 mineralogy atlas. This comparison  
 393 provides valuable insights into how variations in the mineralogical composition of dust  
 394 particles can influence the energy balance in the atmosphere. Compared to the ARI effects  
 395 of bulk dust, the mineralogical composition of dust aerosols can lead to increases in SW  
 396 radiation forcings at the surface and in the atmosphere, ranging from  $+0.10$  to  $+0.82 \text{ W m}^{-2}$ .  
 397 This increase reflects the different optical properties of dust mineral types, which can affect  
 398 the scattering and absorption of solar radiation. These variations in the SW radiation  
 399 forcings are particularly important for understanding how different dust types modulate the  
 400 amount of solar radiation reaching the Earth's surface and atmosphere. At the TOA,  
 401 however, the mineralogical composition resulted in a decrease of about  $-0.72 \text{ W m}^{-2}$  in  
 402 SW radiation forcing, suggesting that certain mineralogical types may be more efficient at

403 reflecting solar radiation back into space. Similar to SW radiation forcings, net radiation  
 404 forcings at the surface and in the atmosphere increased, ranging from +0.02 to +0.63 W  
 405  $m^{-2}$ , while at the TOA, net radiation forcings decreased by about  $-0.65 W m^{-2}$ . The increase  
 406 in net radiation at the surface and in the atmosphere reflects the combined effect of  
 407 increased SW absorption and the potential changes in longwave (LW) radiative properties.  
 408 For LW radiation forcings, the mineralogical composition of dust led to decreases in the  
 409 radiative forcing across different layers, ranging from  $-0.72$  to  $-0.12 W m^{-2}$ . This decrease  
 410 suggests that certain dust mineral types are more efficient at absorbing and emitting  
 411 longwave radiation, which can contribute to cooling effects in the atmosphere and at the  
 412 surface.

413



414

415 Figure 5. Difference between TOA, ATM and BOT radiation forcings with considering bulk dust and  
 416 mineralogical dust compositions (i.e., N2012\_default) enabling ARI effects.

417

418 As demonstrated in Figure A3, the selection of the soil mineralogy dataset and the  
 419 modeling approach significantly influences the calculated dust radiative forcings. When  
 420 comparing shortwave dust radiative effects (DRE) from WRF-CHIMERE simulations  
 421 using the default N2012 and J2014 mineral atlases, we observe a minor discrepancy in the  
 422 DRE amplitude, particularly for shortwave and net radiation forcing at the surface. This

423 discrepancy suggests that the choice of mineralogical dataset can influence the magnitude  
424 of radiative forcings, especially under varying atmospheric conditions. Previous research  
425 has highlighted the distinct optical properties of hematite and goethite in the shortwave  
426 spectrum (Lafon et al., 2006; Sokolik and Toon, 1999). These differences contribute to  
427 variations in the dust's radiative properties and, in turn, its effect on energy transfer in the  
428 atmosphere. Incorporating both minerals in dust production results in a flatter spectral  
429 single scattering albedo (SSA), as goethite's less pronounced dependence on shortwave  
430 wavelengths reduces the overall absorption in the shortwave spectrum (Formenti et al.,  
431 2014). This effect is particularly noticeable when comparing the radiative forcings from  
432 the different mineralogy datasets, as the presence of goethite alters the absorption and  
433 scattering characteristics of the dust particles.

434 As depicted in Figure 6, the distinct day-night variations in shortwave radiation  
435 forcing (SWRF) induced by ARI effects have been thoroughly demonstrated when  
436 considering different mineralogical atlases compared to bulk dust. These variations reflect  
437 the different impacts that dust aerosols have on solar radiation during the day and night,  
438 with a clear difference in the magnitude of the effects between the two periods. Notably,  
439 SWRF variations were more pronounced during the daytime than at night, which can be  
440 attributed to the stronger interaction between dust aerosols and incoming solar radiation  
441 during daylight hours. The presence of dust aerosols alters the reflection, absorption, and  
442 scattering of sunlight, leading to significant changes in the radiation balance, especially  
443 during the day when solar energy is at its peak.

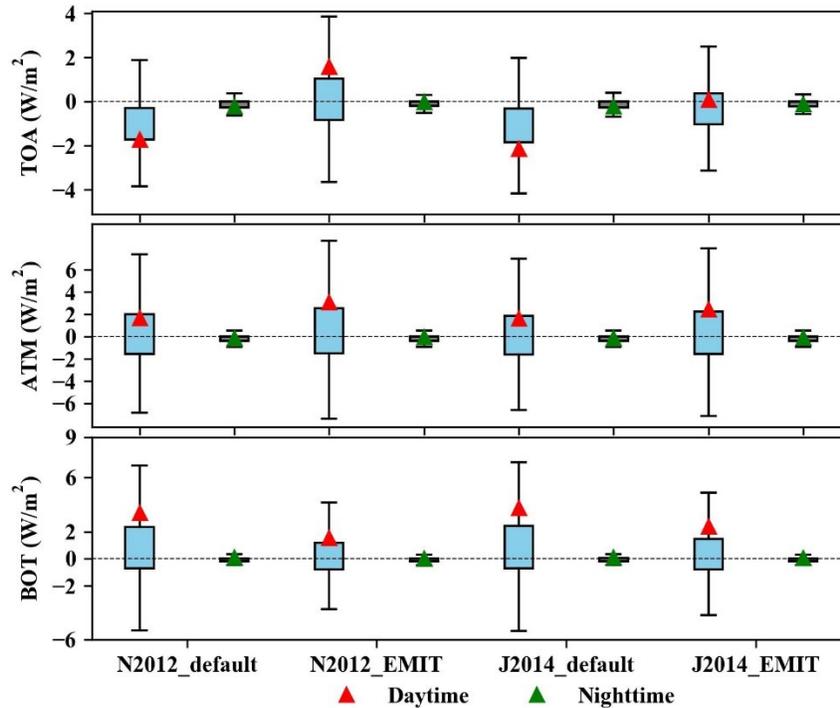
444 Incorporating default dust mineralogical compositions into the simulations led to an  
445 increase in daytime SWRF at the surface and within the atmosphere, ranging from 1.60 to  
446  $3.74 \text{ W m}^{-2}$ . This increase suggests that the specific mineralogy of dust aerosols contributes  
447 to greater absorption and scattering of solar radiation, amplifying the cooling effect at the  
448 surface and the atmosphere. However, at the top of the atmosphere (TOA), the SWRF  
449 decreased by approximately  $2.00 \text{ W m}^{-2}$ , which could be indicative of increased reflection

450 of shortwave radiation back into space due to the dust particles' optical properties. This  
451 shift in radiative forcing at the TOA highlights the role of dust in altering the energy fluxes  
452 across different atmospheric layers.

453       When comparing simulations using default dust mineralogical compositions to those  
454 employing Earth-observing EMIT satellite data within the WRF-CHIMERE model,  
455 notable differences in SWRF were observed. Daytime SWRF at the surface was reduced  
456 for the N2012 mineralogy dataset ( $-1.88 \text{ W m}^{-2}$ ) and J2014 mineralogy dataset ( $-1.37 \text{ W}$   
457  $\text{m}^{-2}$ ) when using EMIT data, compared to the default dust mineralogy compositions. This  
458 reduction could be due to more accurate mineralogical characterization, which alters the  
459 dust's optical properties and reduces its ability to absorb and scatter sunlight. Conversely,  
460 SWRF was enhanced in the atmosphere (N2012:  $+1.44 \text{ W m}^{-2}$ , J2014:  $+0.84 \text{ W m}^{-2}$ ) when  
461 using the EMIT data, indicating that the updated mineralogical information leads to a  
462 different interaction with solar radiation in the atmospheric layer, possibly due to changes  
463 in dust composition that affect scattering and absorption properties at higher altitudes.

464       Furthermore, SWRF at the TOA transitioned from negative to positive in simulations  
465 using the EMIT data. For the N2012 dataset, the SWRF varied from  $-1.73$  to  $+1.59 \text{ W m}^{-2}$ ,  
466 and for the J2014 dataset, it ranged from  $-2.14$  to  $+0.07 \text{ W m}^{-2}$ . This shift suggests that  
467 more accurate dust mineralogy data, particularly from satellite observations, can have a  
468 significant impact on the amount of solar radiation reflected back to space, thereby  
469 influencing the radiative balance at the TOA. The transition from negative to positive  
470 forcing at the TOA emphasizes the importance of incorporating realistic mineralogical  
471 information to enhance the accuracy of dust-related radiative forcing calculations and  
472 better understand their role in climate systems.

473



474  
 475 Figure 6. Day-night changes of TOA, ATM and BOT shortwave radiation forcings from simulations  
 476 using different composition atlases (N2012\_default, N2012\_EMIT, J2014\_default and J2014\_EMIT)  
 477 compared to bulk dust.

478  
 479 **3.3 Radiative effect on air quality**

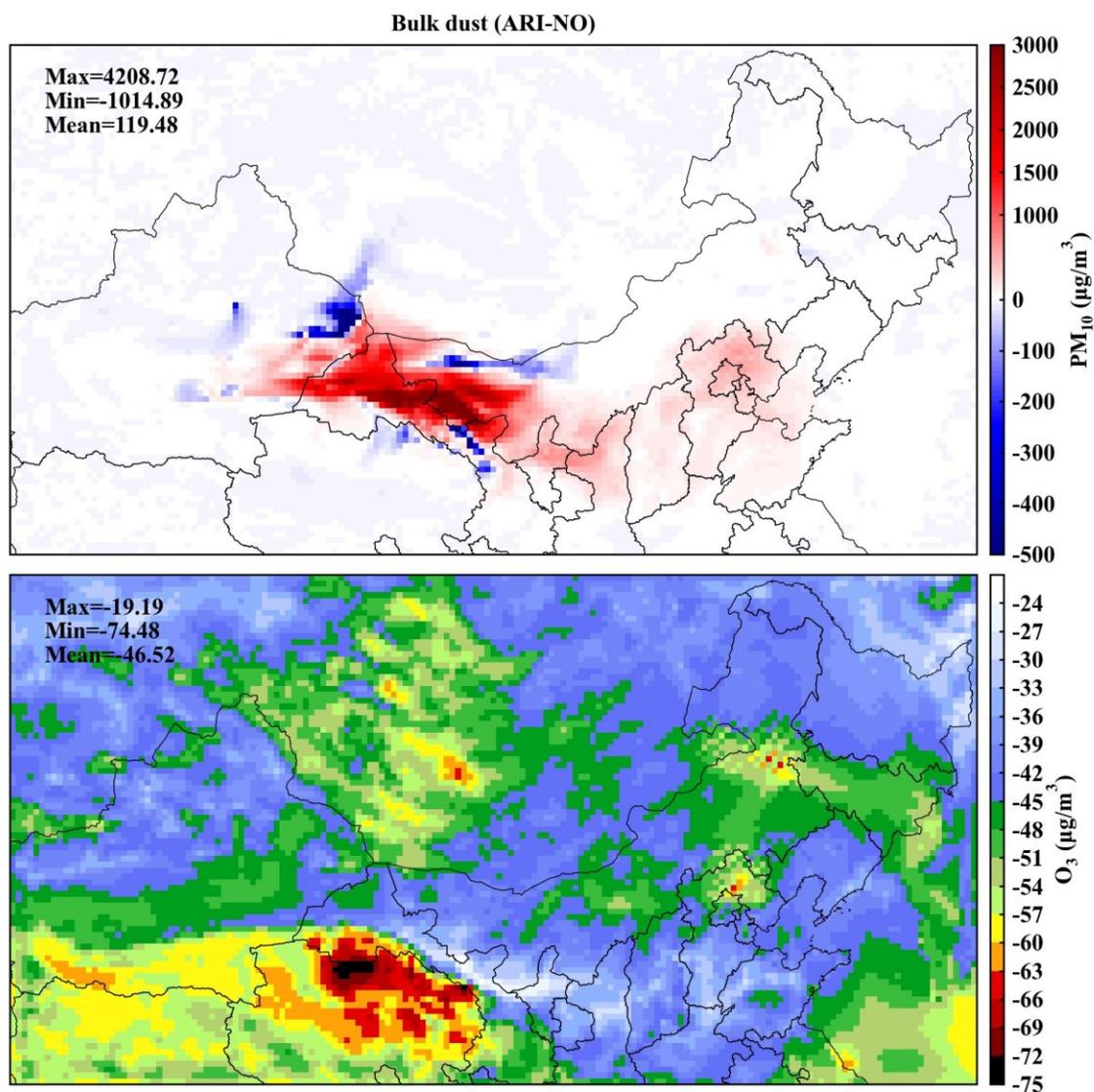
480 Aerosol effects not only gave rise to changes in meteorological variables but also had  
 481 a significant impact on air quality. As shown in Figure 7 and Table S2, the inclusion of bulk  
 482 dust aerosol feedbacks in the WRF-CHIMERE model resulted in substantial increases in  
 483 PM<sub>10</sub> concentrations, with an average increase of 119.48  $\mu\text{g m}^{-3}$  with a 95% CI of  $-27.63$   
 484 to 1408.39  $\mu\text{g m}^{-3}$ . This rise in particulate matter highlights the important role of dust  
 485 aerosols in contributing to local and regional air pollution, especially in regions that are  
 486 susceptible to dust storms. Along with these increases in PM<sub>10</sub> concentrations, O<sub>3</sub>  
 487 concentrations slightly decreased, with an average reduction of  $-46.52 \mu\text{g m}^{-3}$ . This  
 488 reduction in ozone can be attributed to the complex interaction between dust particles and  
 489 ozone precursor gases, where dust aerosols can act as both a sink for ozone and influence  
 490 the photochemical processes that govern its formation and degradation. These reactions  
 491 would be related to the adsorption and catalytic decomposition of ozone on the surface of

492 mineral dust particles, as well as the potential for dust to alter the concentration of reactive  
493 species in the atmosphere through heterogeneous chemistry (Cwiertny et al., 2008). For  
494 example, the presence of adsorbed water on dust particles can compete with ozone for  
495 reactive sites, reducing the overall uptake and decomposition of ozone (Usher et al., 2003).  
496 Additionally, the photochemical reactions involving dust particles, such as the photolysis  
497 of nitrate ions, can produce reactive radicals that further influence the atmospheric  
498 chemistry of ozone (Ma et al., 2021).

499 The most pronounced increases in PM<sub>10</sub> concentrations occurred in the Badain Jaran  
500 Desert, a well-known dust source region, where peak values exceeded 1200 µg m<sup>-3</sup>. This  
501 reflects the large dust emissions typical of desert environments, where strong winds  
502 mobilize vast quantities of particulate matter. Downwind regions, including Ningxia,  
503 Shaanxi, and Beijing, also experienced significant PM<sub>10</sub> elevations, with concentration  
504 differences reaching approximately 600 µg m<sup>-3</sup> compared to baseline levels. The inclusion  
505 of speciated dust influences long-range transport and can substantially affect PM<sub>10</sub>  
506 concentrations. Comparison of the subfigures in Fig. 8 reveals pronounced regional  
507 differences in PM<sub>10</sub> predictions arising from the use of different mineralogical databases.  
508 Incorporating detailed mineralogical data enhances the accuracy of dust composition  
509 representation and its associated effects on PM<sub>10</sub>, highlighting the critical role of mineral  
510 speciation in dust modeling and regional air quality assessment.

511 Ozone changes along transport pathways were generally smaller than the surrounding  
512 concentrations, typically ranging from -60 to -40 µg m<sup>-3</sup> with a mean value of -46.52 µg  
513 m<sup>-3</sup> (95% CI: -63.38 to -31.74) as shown in Table S2. These smaller changes in O<sub>3</sub>  
514 concentrations reflect the fact that dust aerosols have a more localized and complex effect  
515 on ozone formation and destruction, with significant variability depending on the regional  
516 and temporal context. In particular, dust-induced reductions in ozone are likely to be  
517 influenced by the local presence of other atmospheric constituents. The photochemical  
518 reactions involving dust particles, such as the photolysis of nitrate ions, can produce

519 reactive radicals that further influence the atmospheric chemistry of ozone (Ma et al., 2021).  
520  
521



522  
523 Figure 7. Changes in PM<sub>10</sub> and O<sub>3</sub> concentrations resulting from bulk dust-induced ARI effects,  
524 compared to the scenario without aerosol feedbacks.

525

526 The spatial differences in PM<sub>10</sub> and O<sub>3</sub> concentrations simulated by WRF-CHIMERE  
527 with different mineralogy atlases compared to bulk dust, enabling ARI effects, are depicted  
528 in Figures 8 and A4. These comparisons reveal substantial changes in the PM<sub>10</sub> and O<sub>3</sub>  
529 concentrations across the different mineralogical compositions, including N2012\_Default,

530 N2012\_EMIT, J2014\_Default, J2014\_EMIT, and bulk dust. This suggests that the  
531 normalization of the 12 minerals from these atlases significantly modifies meteorological  
532 conditions, further influencing the relative abundances of dust particles and their  
533 subsequent effects on air quality and atmospheric composition.

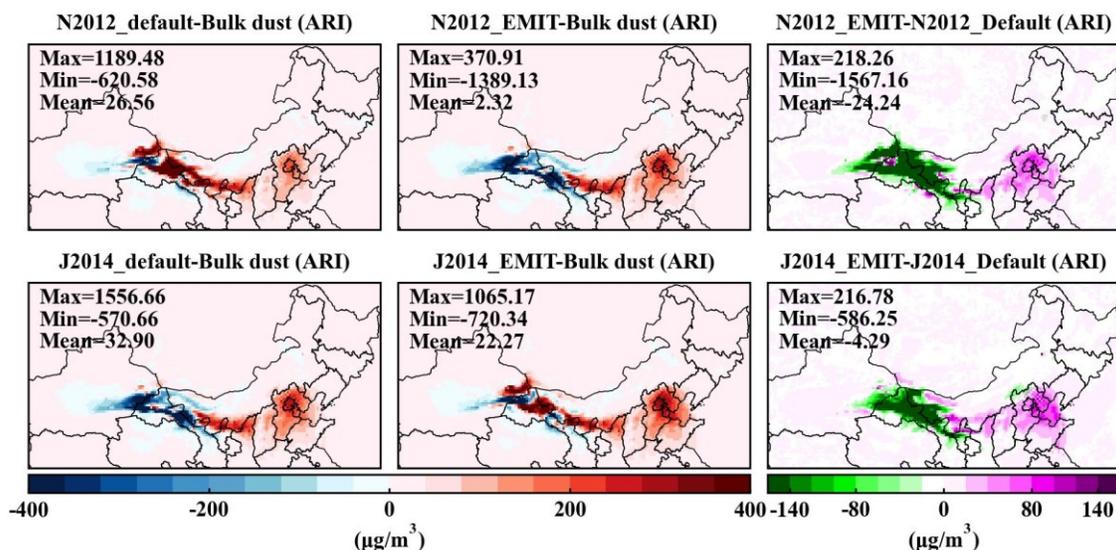
534 When compared to bulk dust, reduced PM<sub>10</sub> concentrations were primarily observed  
535 in the Taklimakan Desert, with decreases of around 60 μg m<sup>-3</sup>, while increases in PM<sub>10</sub>  
536 concentrations occurred in the Badain Jaran Desert and its downwind regions, with  
537 concentrations rising up to 1000 μg m<sup>-3</sup>. These regional variations indicate that different  
538 dust mineralogical compositions can impact the emission and transport of dust, with certain  
539 mineral types leading to more efficient scattering or absorption of radiation, which may  
540 alter the local meteorological conditions and dust dispersion patterns.

541 For O<sub>3</sub> concentrations, reductions and enhancements were mainly observed in the  
542 Horqin sandy land and North China Plain, with changes up to 4 μg m<sup>-3</sup>, respectively. This  
543 highlights the complex interaction between dust aerosols and ozone chemistry, where dust  
544 can either enhance or reduce ozone concentrations depending on the region. Dust aerosols  
545 can influence ozone levels by acting as a surface for heterogeneous chemical reactions or  
546 by modifying the photochemical processes that control ozone formation and destruction.

547 When considering the EMIT data, PM<sub>10</sub> concentrations were reduced in dust source  
548 regions and enhanced in downwind regions, with reductions of up to -1567.16 μg m<sup>-3</sup> and  
549 increases of +218.26 μg m<sup>-3</sup>. This suggests that more accurate mineralogical data can  
550 influence dust transport patterns, leading to greater reductions in PM<sub>10</sub> at the source regions  
551 and increased dust concentrations in the downwind areas. These findings further emphasize  
552 the role of mineralogical composition in modulating dust aerosol behavior and distribution.

553 For O<sub>3</sub>, enhancements appeared in source regions, while reductions were observed in  
554 downwind regions, with changes ranging from -2.46 to +3.52 μg m<sup>-3</sup>. These trends suggest  
555 that more accurate dust speciation can influence regional ozone levels in different ways,  
556 with possible implications for local air quality and atmospheric chemistry. Notably, the

557 impacts on PM<sub>10</sub> concentrations from N2012\_EMIT compared to N2012\_Default were  
 558 larger than those observed from J2014\_EMIT versus J2014\_Default, while the impacts on  
 559 O<sub>3</sub> concentrations followed the opposite trends. This indicates that the choice of dust  
 560 mineralogical dataset has a differential impact on PM<sub>10</sub> and O<sub>3</sub> concentrations,  
 561 underscoring the importance of considering mineral composition in aerosol modeling to  
 562 more accurately predict air quality and climate effects.  
 563



564  
 565 Figure 8. Difference in PM<sub>10</sub> concentrations considering bulk dust and various dust mineralogy atlases  
 566 that enable ARI effects.  
 567

568 Figure 9 shows the percentage changes in surface concentrations of mineral dust with  
 569 and without considering ARI effects. These results provide valuable insight into how the  
 570 inclusion of ARI effects modifies the composition and radiative properties of dust aerosols,  
 571 depending on the mineralogical dataset used. For the N2012\_default and N2012\_EMIT  
 572 data, quartz and feldspar accounted for a substantial portion of the total dust, ranging from  
 573 approximately 51.7% to 57.4% for quartz and 18.6% to 19.8% for feldspar. This indicates  
 574 that quartz and feldspar are the dominant mineral components in the dust modeled with the  
 575 N2012 dataset.

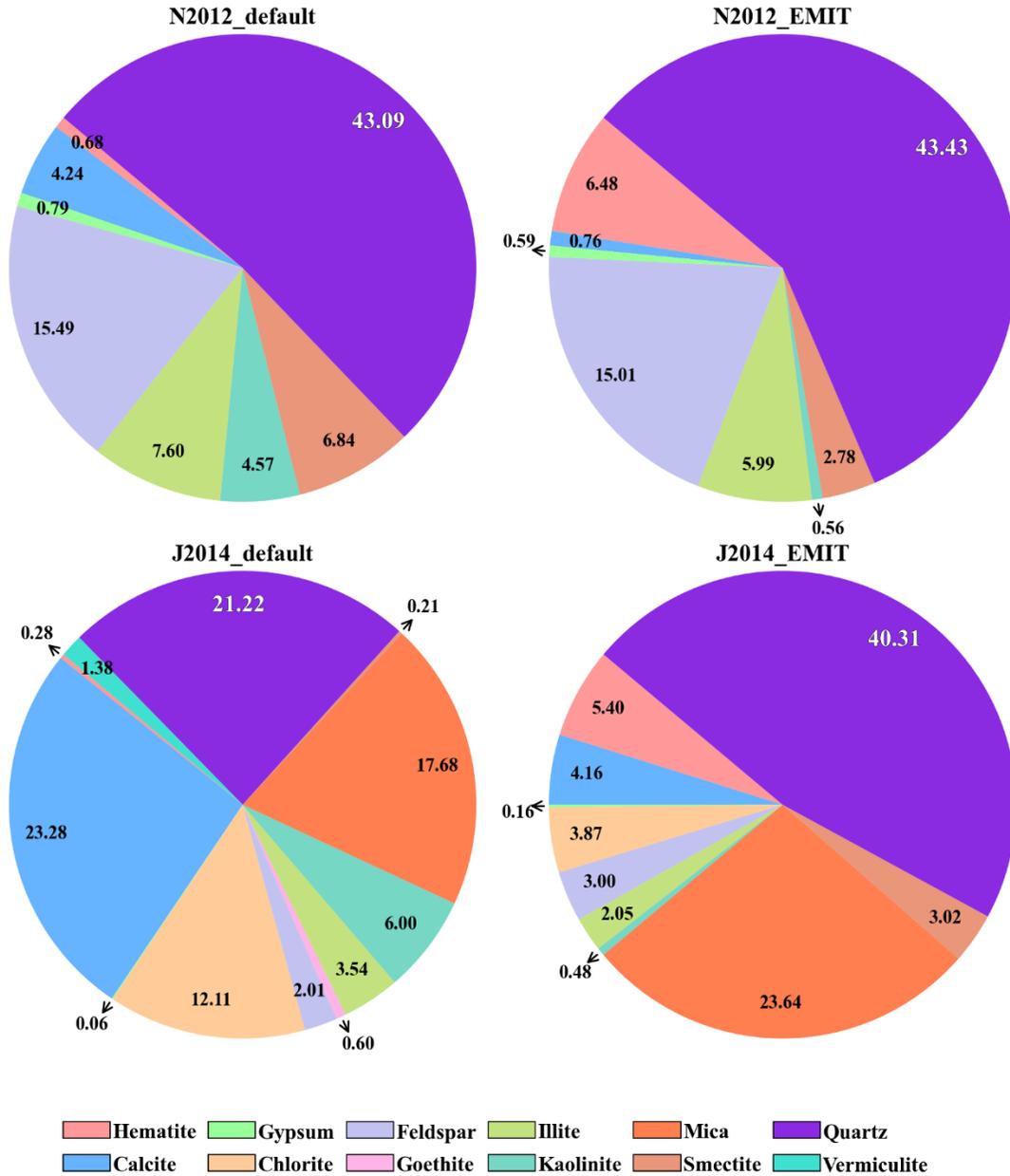
576 In contrast, for the J2014\_default dataset, the mineral composition was more

577 diversified, with calcite, quartz, and mica contributing about 26.3%, 24.0%, and 20.0%,  
578 respectively, to the total dust composition. This shift in mineral proportions reflects the  
579 differences in the mineralogical characterization between the N2012 and J2014 datasets,  
580 with J2014 incorporating a broader range of dust minerals. For J2014\_EMIT, the mineral  
581 composition shifted further, with quartz and mica making up approximately 46.8% and  
582 27.5% of the dust, respectively. This highlights the importance of using accurate  
583 mineralogical data, such as that from EMIT satellite observations, to better represent the  
584 composition of dust aerosols in simulations.

585         The inclusion of EMIT data led to an increase in the absorption percentage of hematite  
586 by about 8% for N2012 and 6% for J2014. Hematite is a highly absorbing mineral,  
587 especially in the shortwave spectrum, and its increased presence enhances the dust's ability  
588 to absorb solar radiation, thereby affecting the DRE in the shortwave spectrum. This  
589 increase in hematite absorption is significant, as it directly impacts the radiative effects of  
590 dust, potentially contributing to a greater cooling effect on the atmosphere by modifying  
591 the shortwave radiation balance.

592         While quartz constitutes the largest portion of the dust in both the N2012 and J2014  
593 datasets, its DRF effects are relatively limited, as noted in Li et al. (2021). Quartz is known  
594 for its high reflectivity in the shortwave spectrum, and while it makes up a large fraction  
595 of the total dust mass, it has a less pronounced effect on radiative forcing compared to more  
596 absorbing minerals like hematite or mica. This suggests that, despite its dominance in dust  
597 composition, quartz plays a smaller role in modifying the energy balance of the atmosphere  
598 through direct radiative effects.

599  
600



601  
 602 Figure 9. Contributions of different mineralogical compositions using N2012\_default, N2012\_EMIT,  
 603 J2014\_default, and J2014\_EMIT, considering ARI effects, compared to the scenario without enabling  
 604 aerosol feedbacks.

605  
 606 **3.4 Limitations and uncertainties for aerosol feedbacks of mineralogical dust**

607 The accuracy of simulated dust emission is intricately linked to soil properties, such  
 608 as soil texture and moisture, which primarily influence the threshold friction velocity  
 609 required for dust particle mobilization (Kim and Choi, 2015; Su and Fung, 2015). These

610 factors play a critical role in determining the magnitude and spatial distribution of dust  
611 emissions, underscoring the need for precise and high-resolution soil data in dust modeling.  
612 While the current EMIT L3 data offers a spatial resolution of  $0.5^\circ \times 0.5^\circ$ , real-time higher  
613 spatial resolution datasets, such as the  $60 \text{ m} \times 60 \text{ m}$  EMIT L2B mineral atlas  
614 ([https://earth.jpl.nasa.gov/emit/internal\\_resources/282](https://earth.jpl.nasa.gov/emit/internal_resources/282)), can provide a more detailed  
615 representation of soil mineralogy, thereby enhancing the fidelity of dust emission  
616 simulations.

617 The uncertainty associated with dust mineralogical datasets is being actively  
618 addressed by NASA's EMIT. This initiative has deployed a hyperspectral imaging  
619 spectrometer aboard the International Space Station to deliver global retrievals of soil  
620 mineral compositions with unprecedented spatial detail. The spectrometer captures spectral  
621 absorption features within the UV to near-infrared range ( $0.38\text{--}2.5 \mu\text{m}$ ), offering critical  
622 insights into the distribution and variability of soil minerals (Castellanos et al., 2024;  
623 Connelly et al., 2021). In addition to satellite-based observations, ground-based stations  
624 play a vital role in measuring dust mineralogical compositions using stationary instruments,  
625 which provide localized and highly accurate data. Complementing these measurements,  
626 aircraft-based instruments offer the capability to sample dust particles along specific flight  
627 tracks, providing valuable vertical and spatial profiles of speciated dust properties (e.g.,  
628 size and mixing state, Panta et al., 2023; Ryder et al., 2015). Together, these observational  
629 platforms form a robust foundation for validating and improving dust models.

630 Aerosol-cloud interactions involving speciated dust are another critical aspect of dust-  
631 climate interactions that require further investigation, especially for feldspar and quartz  
632 (Atkinson et al., 2013; Chatziparaschos et al., 2023). Incorporating these interactions into  
633 two-way coupled WRF-CHIMERE models can provide a more comprehensive  
634 understanding of the feedback mechanisms between dust aerosols and cloud microphysics.  
635 Such implementations are currently a focus of ongoing work, aiming to refine the  
636 representation of dust-induced radiative and microphysical effects in regional and global

637 models. These efforts will not only improve model accuracy but also enhance our ability  
638 to predict the impacts of dust on weather, air quality, and climate.

639

#### 640 **4 Conclusion**

641 Dust mineral composition plays a vital role in regulating atmospheric radiation and  
642 air quality, yet its effects remain poorly constrained in current atmospheric models.  
643 Understanding these impacts is particularly important for North China, where severe dust  
644 storms frequently affect regional climate and pollution. This study investigates how  
645 variations in mineral composition influence aerosol–radiation interactions and their  
646 implications for meteorology and air quality during a major dust storm event.

647 The findings revealed significant spatial variations in radiative forcing due to  
648 differences in dust mineralogy. Compared to the ARI effects of bulk dust, the mineralogical  
649 composition of dust aerosols can increase SW radiation forcing at the surface and in the  
650 atmosphere by +0.10 to +0.82 W m<sup>-2</sup>, while simultaneously causing a decrease of  
651 approximately -0.72 W m<sup>-2</sup> in SW radiation forcing at the TOA. Integrating EMIT data  
652 into the model reduced PM<sub>10</sub> biases by over 15% in high-concentration regions and  
653 improved ozone predictions, with localized changes ranging from -2.46 to +3.52 μg m<sup>-3</sup>.  
654 Specifically, the ARI effects of these mineralogical compositions led to a notable increase  
655 in PM<sub>10</sub> levels, reaching up to 1189.48 μg m<sup>-3</sup> in dust source regions, when compared to  
656 bulk dust scenarios.

657 These findings highlight the importance of incorporating dust mineralogical data to  
658 improve simulations of radiative forcing and air quality impacts. Within the scope of this  
659 study, the results indicate that overall dust mineralogical composition, rather than dust mass  
660 alone, plays a decisive role in ARI effects, with hematite exerting a dominant influence  
661 despite its minor abundance, although the radiative effects of individual mineral species  
662 were not separately quantified. Systematic biases in surface radiation, near-surface winds,  
663 and temperature persist, reflecting challenges in simulating dust–atmosphere interactions  
664 and uncertainties in mineralogical datasets. Incorporating meteorological spectral nudging

665 in future simulations could provide a more realistic representation of ARI-induced dust  
666 perturbations under different mineralogical compositions. Coupling this approach with  
667 higher-resolution soil and satellite data, as well as additional observational constraints,  
668 would further refine dust emission simulations and reduce model biases, particularly in  
669 regions frequently affected by severe dust events.

670

### 671 **Data and software availability**

672 The meteorological ICs and BCs, Chemical ICs and BCs and emission data used for  
673 WRF–CHIMERE and all data used to create figures and tables in this study are provided  
674 in an open repository on Zenodo (<https://doi.org/10.5281/zenodo.14728874>, Gao et al.,  
675 2025a). Himawari and CALIPSO satellite data are available at  
676 <ftp://ftp.ptree.jaxa.jp/jma/netcdf> and <https://subset.larc.nasa.gov/calipso>, respectively.

677 The source codes of the two-way coupled WRFv3.7.1–CHIMERE v2020r3 models  
678 are obtained from <https://www.lmd.polytechnique.fr/chimere>. The related source codes,  
679 configuration information, namelist files and automated run scripts of these three two-way  
680 coupled models are archived at Zenodo with the following associated DOI:  
681 <https://doi.org/10.5281/zenodo.14729124> (Gao et al., 2025b).

682

### 683 **Author contributions**

684 CG, XZ, HY and LH carried out the data collection, related analysis, figure plotting,  
685 and paper writing. HZ, SZ, and AX were involved with the original research plan and made  
686 suggestions for the paper writing.

### 687 **Competing interests**

688 The contact author has declared that neither they nor their co-authors have any  
689 competing interests.

690

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700

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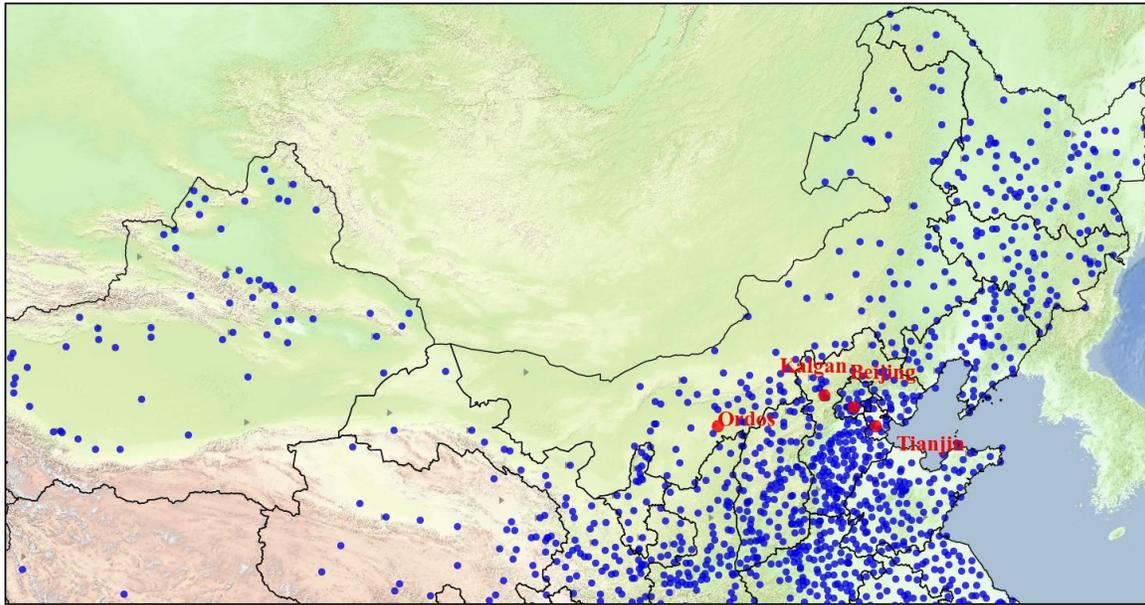
905 **Appendix**

906 Table A1. Model setups and inputs for the WRF-CHIMERE model.

		WRF-CHIMERE
Domain configuration	Horizontal grid spacing	27 km (165 × 87)
	Vertical resolution	33 levels
Physics parameterization	Shortwave radiation	RRTMG
	Longwave radiation	RRTMG
	Cloud microphysics	Thompson
	PBL	YSU
	Cumulus	Grell-Freitas
	Surface	Monin-Obukhov
	Land surface	Noah LSM
Chemistry scheme	Icloud	Xu-Randall method
	Aerosol mechanism	SAM
	Aerosol size distribution	Sectional (10 bins)
	Aerosol mixing state	Core-Shell
	Gas-phase chemistry	MELCHIOR2
Emission	Photolysis	Fast-JX with cloud effects
	Dust emission	Kok
Input data	Meteorological ICs and BCs	FNL
	Chemical ICs and BCs	LMDZ-INCA

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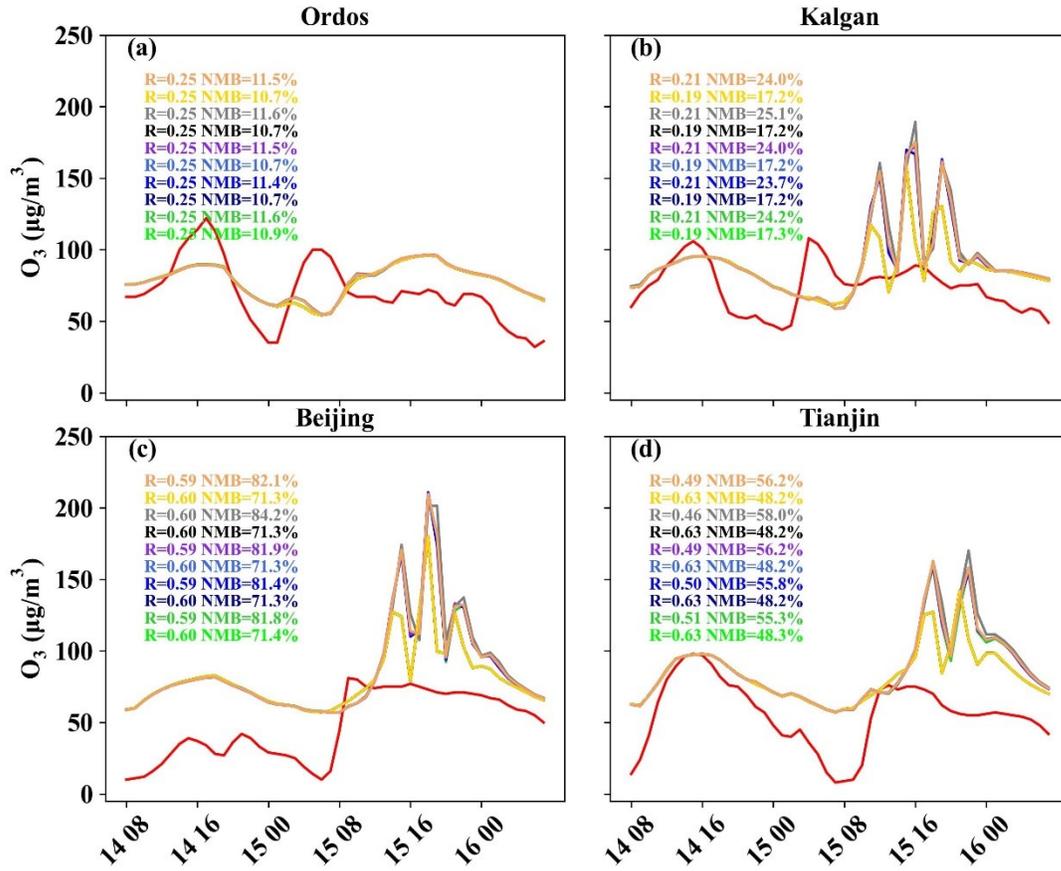
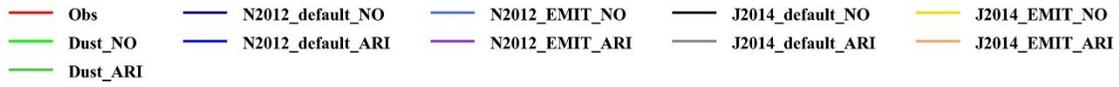


- ▶ Solar radiation monitoring station
- Meteorological monitoring station
- Air quality monitoring station

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910 Figure A1. Simulation domain and locations of meteorology and air quality monitoring stations.

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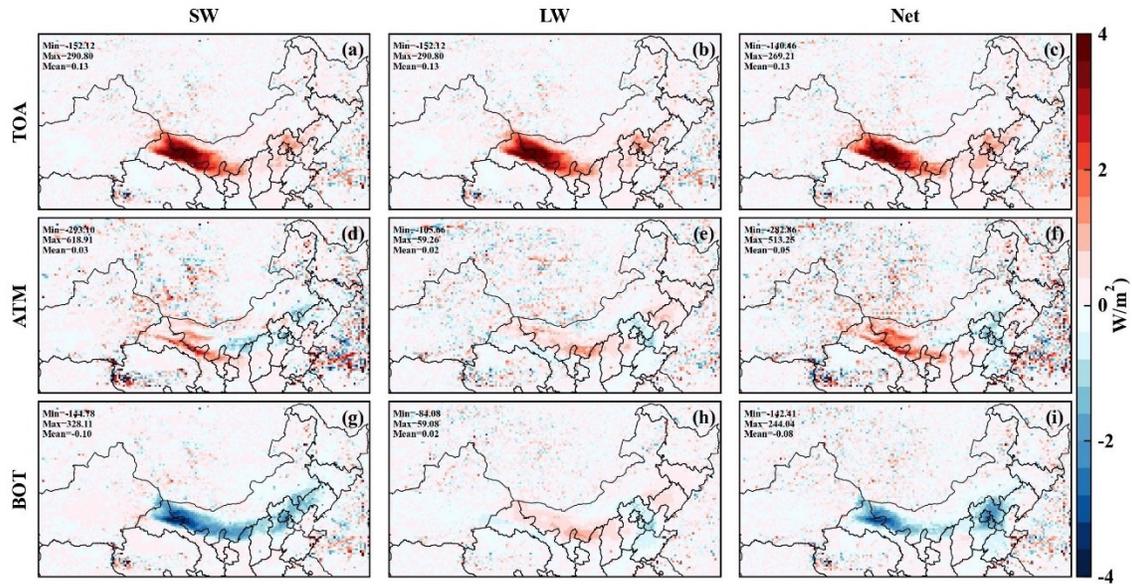


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913 Figure A2. Statistical metrics between observed and simulated O<sub>3</sub> concentrations by different  
 914 scenario simulations.

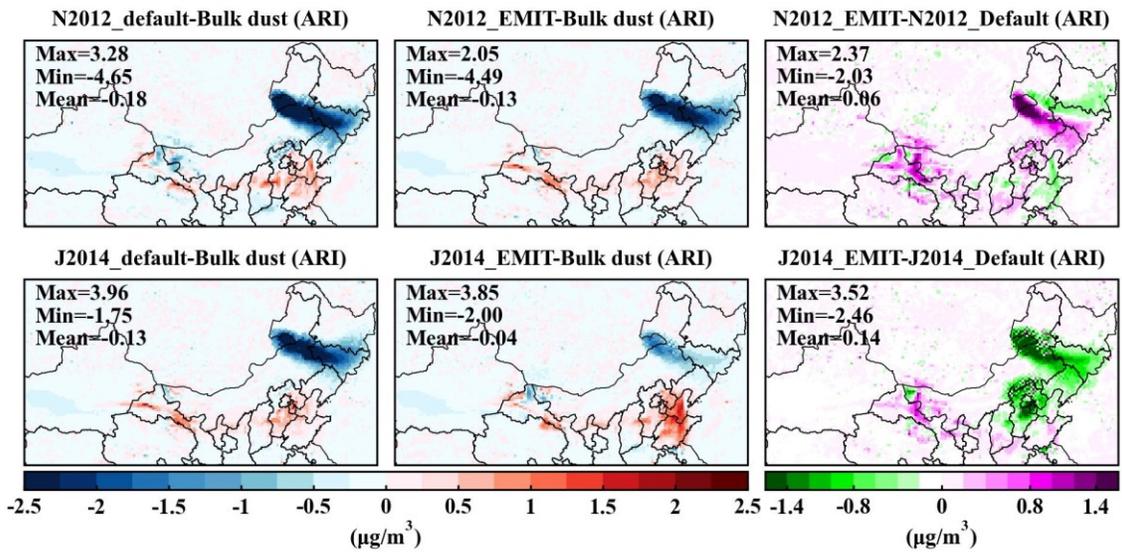
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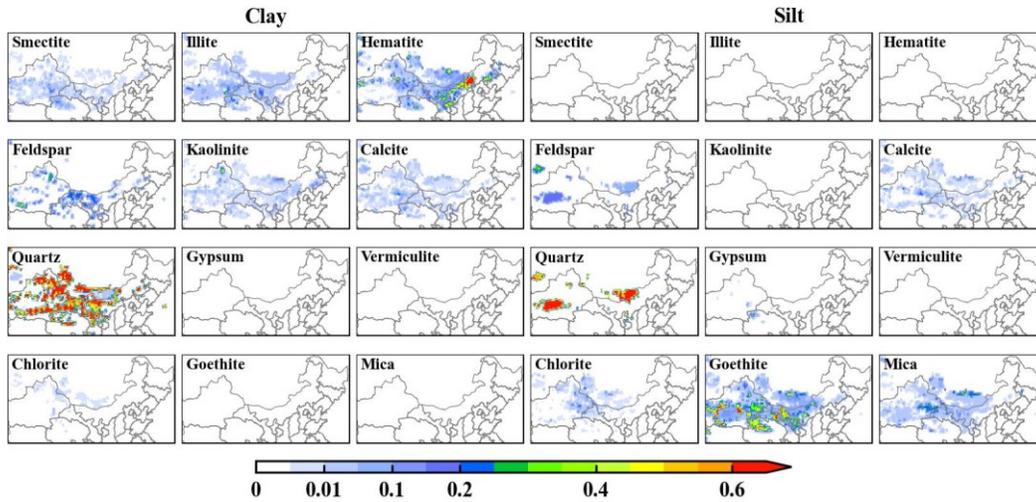
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Figure A3. Difference between TOA, ATM and BOT radiation forcings with considering J2014 and N2012 mineralogical dust compositions (i.e., N2012\_default) enabling ARI effects.



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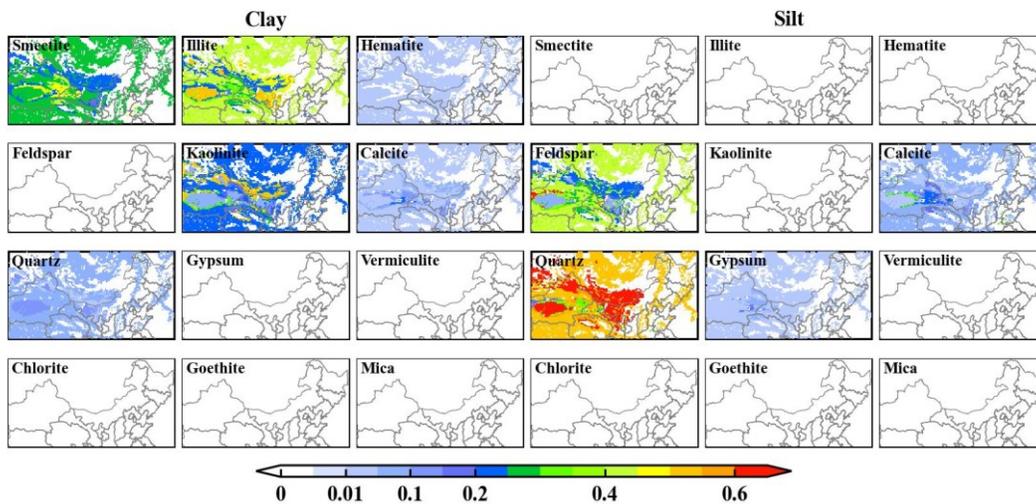
Figure A4. Difference in O<sub>3</sub> concentrations considering bulk dust and various dust mineralogy atlases that enable ARI effects.



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927 Figure A5. Spatial distribution of content for the different mineral dust species in the silt and  
 928 clay fraction of the soil for J2014 with EMIT satellite data.

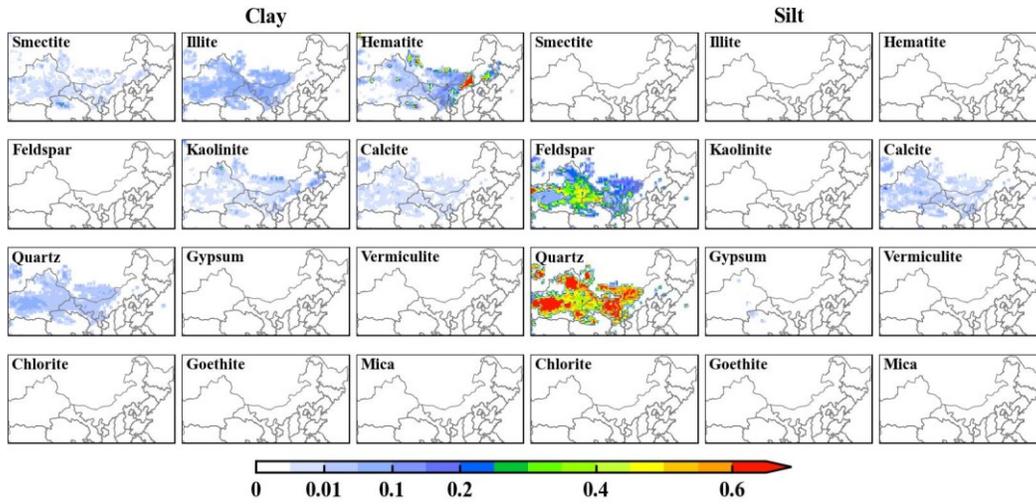
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931 Figure A6. Spatial distribution of content for the different mineral dust species in the silt and clay  
 932 fraction of the soil for original N2012 mineralogical data.

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934

935 Figure A7. Spatial distribution of content for the different mineral dust species in the silt and clay  
 936 fraction of the soil for N2012 with EMIT satellite data.

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