

1 **Effects of aerosol-radiation interaction on precipitation during**  
2 **biomass-burning season in East China**

3

4 **Xin Huang<sup>1,2,3</sup>, Aijun Ding<sup>1,2,3\*</sup>, Lixia Liu<sup>1,2</sup>, Qiang Liu<sup>1,2</sup>, Ke Ding<sup>1,2</sup>, Xiaorui Niu<sup>1,2,3</sup>, Wei**  
5 **Nie<sup>1,2,3</sup>, Zheng Xu<sup>1,2,3</sup>, Xuguang Chi<sup>1,2,3</sup>, Minghuai Wang<sup>1,2,3</sup>, Jianning Sun<sup>1,2,3</sup>, Weidong**  
6 **Guo<sup>1,2,3</sup>, and Congbin Fu<sup>1,2,3</sup>**

7

8 <sup>1</sup>Joint International Research Laboratory of Atmospheric and Earth System Sciences, Nanjing  
9 University, China

10 <sup>2</sup>Institute for Climate and Global Change Research & School of Atmospheric Sciences, Nanjing  
11 University, Nanjing, 210023, China

12 <sup>3</sup>Collaborative Innovation Center of Climate Change, Jiangsu Province, China

13

14 \* Correspondence to: Aijun Ding (dingaj@nju.edu.cn)

15

## 1 **Abstract**

2 Biomass burning is a main source for primary carbonaceous particles in the atmosphere and  
3 acts as a crucial factor that alters Earth's energy budget and balance. It is also an important  
4 factor influencing air quality, regional climate and sustainability in the domain of Pan-Eurasian  
5 Experiment (PEEX). During the exceptionally intense agricultural fire season in mid-June 2012,  
6 accompanied with rapidly deteriorating air quality, a series of meteorological anomalies was  
7 observed, including a large decline in near-surface air temperature, spatial shifts and changes  
8 in precipitation in Jiangsu Province of East China. To explore the underlying processes that link  
9 air pollution to weather modification, we conducted a numerical study with parallel simulations  
10 using the fully coupled meteorology-chemistry model WRF-Chem with a high-resolution  
11 emission inventory for agricultural fires. Evaluation of the modelling results with available  
12 ground-based measurements and satellite retrievals showed that this model was able to  
13 reproduce the magnitude and spatial variations of fire-induced air pollution. During the  
14 biomass-burning event in mid-June 2012, intensive emission of absorbing aerosols trapped a  
15 considerable part of solar radiation in the atmosphere and reduced incident radiation reaching  
16 the surface on a regional scale, followed by lowered surface sensible and latent heat fluxes. The  
17 perturbed energy balance and re-allocation gave rise to substantial adjustments in vertical  
18 temperature stratification, namely surface cooling and upper-air heating. Furthermore, intimate  
19 link between temperature profile and small-scale processes like turbulent mixing and  
20 entrainment led to distinct changes in precipitation. On one hand, by stabilizing the atmosphere  
21 below and reducing the surface flux, black carbon-laden plumes tended to dissipate daytime  
22 cloud and suppress the convective precipitation over Nanjing. On the other hand, heating aloft  
23 increased upper-level convective activity and then favored convergence carrying in moist air,  
24 thereby enhancing the nocturnal precipitation in the downwind areas of the biomass burning  
25 plumes.

26

## 27 **1 Introduction**

28 Biomass burning, defined as open or quasi-open combustion of non-fossilized vegetative or  
29 organic fuel, is widely used by humans to manage and transform land cover for many purposes  
30 and has been identified as one of the most important disturbance agents in world's terrestrial  
31 ecosystems (Fearnside, 2000). It is a major source of many trace gases and particulate matters  
32 on a regional and a global scale (Andreae and Merlet, 2001; van der Werf et al., 2006; Ito et al.,

1 2007), contributing significantly to the budgets of trace gases, greenhouse gases and  
2 atmospheric aerosols (Langenfelds et al., 2002). For instance, biomass burning is estimated to  
3 be responsible for almost half of global carbon monoxide (CO) emission and more than one  
4 third of total black carbon (BC) emission (Bergamaschi et al., 2000; Bond et al., 2013). With  
5 tremendous and intensive emission of atmospheric pollutants, it has been recognized as one of  
6 the culprits of regional air pollution (Wiedinmyer et al., 2006; Ryu et al., 2007) and an important  
7 disturber of biogeochemical cycles, especially for those of carbon and nitrogen (Crutzen and  
8 Andreae, 1990; Kuhlbusch, 1998). In the Eurasian continent, i.e., the domain of Pan-Eurasian  
9 Experiment (PEEX) (Kulmala et al., 2015), biomass burning is a very important source  
10 influencing air quality, regional climate change and sustainability (Chi et al., 2013; Ding et al.,  
11 2013ab; Lappalainen et al., 2016). In the East China, the impact of biomass burning to air  
12 quality and regional climate change is particularly interesting because of the mixing of biomass  
13 burning plumes with pollutant from fossil fuel combustion sources (Ding et al., 2013a; Nie et  
14 al., 2015; Xie et al., 2015; Lappalainen et al., 2016).

15 Biomass burning, including forest fires, savanna fires, peat burning, and crop residue burning  
16 in field, generally features a high emission rate of light-absorbing carbonaceous aerosols (Reid  
17 et al., 1998; Schwarz et al., 2008). The most important one is BC, which is intensively emitted  
18 during biomass burning events due to incomplete combustions (Reid et al., 2005; Akagi et al.,  
19 2011). As the dominant absorber of solar radiation in the atmosphere, BC warms the Earth-  
20 atmospheric system and alters the partitioning of energy between the ground surface and the  
21 atmosphere, thereby modifying atmospheric thermodynamic structures and modulating  
22 hydrological cycles (Krishnan and Ramanathan, 2002; Ramanathan et al., 2005; Ding et al.,  
23 2016). These modifications induced by biomass burning have been detected in many regions,  
24 especially for those during forest fires. Surface temperature decline was extensively observed  
25 during forest fires in North America, Asia and Africa (Robock, 1988, 1991; Procopio et al.,  
26 2004; Kolusu et al., 2015). The dimming around ground surface and heating in the upper-  
27 atmosphere, especially in the upper boundary layer, could cause the suppression of daytime  
28 mixing height and result in an enhancement of surface air pollution through aerosol-boundary  
29 layer-radiation feedbacks (Ding et al., 2013a; Ding et al., 2016). This effect was named as the  
30 “Dome Effect” of BC by Ding et al. (2016). By cooling the surface and stabilizing the  
31 atmosphere, intense forest fire may lead to the inhibition of cloud formation (Andreae et al.,  
32 2004; Koren et al., 2004; Feingold et al., 2005), suppression in precipitation (Rosenfeld, 1999;  
33 Sakaeda et al., 2011), and even temporal shift in onset of monsoon (Liu et al., 2005; Lau et al.,

1 2006; Zhang et al., 2009). In one word, BC has been demonstrated to cause a significant  
2 perturbation in the radiative energy balance and has even led to regional and global climate  
3 change (Penner et al., 1992; Menon et al., 2002; Ramanathan and Carmichael, 2008).

4 Although forest and savanna fires are much less notable in China compared with tropical  
5 America, Africa and Southeast Asia (van der Werf et al., 2006), it is noteworthy that China is  
6 a large country with the world's top-ranked agricultural production, which is inevitably  
7 accompanied by a tremendous amount of crop residue. Field burning of crop residue is a  
8 common and wide-spread management practice in China during post-harvest periods for the  
9 purpose of clearing farmland and providing short-lived ash fertilization for the crop rotation  
10 (Gao et al., 2002). It is estimated that about 120 Tg crop residues were burned in field across  
11 China every year, far higher than those burned in forest fires and savanna fires (Yan et al., 2006).  
12 Previous studies have documented that field burning of crop residue led to deterioration in  
13 regional air quality during harvest season (Yang et al., 2008; Huang et al., 2012b; Li et al.,  
14 2014). What is worse, this kind of pollution occurs periodically in East China, particularly  
15 during the harvest period of wheat in June (Figure 1). However, studies regarding its effects on  
16 meteorology and climate are still limited. Ding et al. (2013a) reported that temperature and  
17 precipitation were dramatically modified during the harvest season in 2012 according to ground  
18 based measurements at a regional background station SORPES in the Yangtze River Delta  
19 region in East China (Ding et al., 2013b). However, there is a lack of a comprehensive picture  
20 of how or through which processes the biomass burning plumes influenced the air temperature  
21 and precipitation and on what scale the aerosol-weather interactions happened during this case.

22 Here we conducted numerical simulations for the biomass burning event in East China during  
23 mid-June 2012 based on the online coupled meteorology-chemistry model WRF-Chem (the  
24 Weather Research and Forecasting model coupled with Chemistry) combined with multiple  
25 ground-based measurements and remote-sensing retrievals. The rest of this paper is structured  
26 as follows: Section 2 describes the development of an emission inventory for field burning of  
27 crop residues and how the numerical simulations are configured and designed; in Section 3 we  
28 validate the modelling results using available measurements, and then analyse the perturbations  
29 in energy budget and temperature adjustments induced by crop residue burning; finally, three  
30 regions with distinct precipitation changes, located near or downwind from the burning sites,  
31 are selected to discuss in detail. Conclusions are drawn in Section 4.

32

## 1 2 Data and Methodology

### 2 2.1 Emission inventory

3 Modelling aerosols' radiative effects during this biomass burning event first requires accurate  
4 quantification and meticulous characterization of emission from field burning of crop residue.  
5 Here, emission intensities of trace gases and particulate matters, specifically including carbon  
6 dioxide (CO<sub>2</sub>), CO, methane (CH<sub>4</sub>), Non-Methane Organic Compounds (NMOCs), nitrogen  
7 oxides (NO<sub>x</sub>), ammonia (NH<sub>3</sub>), sulfur dioxide (SO<sub>2</sub>), BC, organic carbon (OC), and particulate  
8 matter (PM<sub>2.5</sub> and PM<sub>10</sub> are particles with aerodynamic diameter less than 2.5 and 10 microns,  
9 respectively), were estimated based on a bottom-up method. According to the farming season  
10 (available at [zzys.agri.gov.cn](http://zzys.agri.gov.cn)) and province-level statistics on crop cultivation (NBSC, 2013),  
11 we can deduce that the intensive agricultural fires in June were mainly related to wheat straw  
12 burning as a consequence of the extensively spreading cultivation mode of "winter wheat-  
13 summer corn/rice" in East China. Burned biomass at province-level was calculated based on  
14 statistical data of crop productions, residue-to-production ratios, percentages of crop residues  
15 burned in the field. Emissions of various pollutants were derived from the product of burned  
16 mass and experiment results on crop-specific combustion efficiencies and pollutant-specific  
17 emission factors. The detailed methods and involved datasets are described in our previous  
18 work (Huang et al., 2012b).

19 To determine the locations and time of crop residue fires, MODIS (Moderate Resolution  
20 Imaging Spectroradiometer) Thermal Anomalies/Fire Daily L3 Global Product  
21 (MOD/MYD14A1) combined with burned area product (MCD45A1) were introduced for the  
22 purpose of emission spatiotemporal allocations (Giglio et al., 2003; Boschetti et al., 2009).  
23 MOD/MYD14A1 provides fire identification by examining the brightness temperature relative  
24 to neighbouring pixels. MCD45A1 was also incorporated in this work because its bidirectional  
25 reflectance model-based change detection approach has been proved to be capable of presenting  
26 a more accurate mapping of smaller fragments of burn scars (Roy and Boschetti, 2009). Global  
27 Land Cover (GLC) product with a spatial resolution of 1 km was used in this study to identify  
28 the burning of different biomass. Only fire detections that occurred on farmland, that is land  
29 cover classes defined as "Farm" and "Mosaic of cropping", was identified as field burning of  
30 crop residue. Emission at province level estimated using the aforementioned method were then  
31 allocated equally to each fire spot.

1 The fire emission estimation developed in this work was compared with the FINN fire emission  
2 dataset. Spatially, these two emission inventories generally were consistent with each other  
3 because the locations for the fires in both inventories are based on MODIS Thermal Anomalies  
4 Product (Figure S1). Some inconsistencies, such as the density of fire in central Jiangsu, are  
5 attributed to the different land cover dataset applied for the identification of underlying biomass  
6 type. FINN fire emission estimation used MODIS Collection 5 Land Cover Type data  
7 (Wiedinmyer et al., 2011), while we employed Global Land Cover data. This inventory differs  
8 slightly from FINN estimation in magnitude. Taking CO emission in the inner model domain  
9 for instance, we estimate that 4.5 Tg CO was emitted while FINN gives the value of 7.5 Tg  
10 during the first half of June 2012. It might be attributed to different methods to estimate burned  
11 biomass. FINN used MODIS Vegetation Continuous Fields to assign the burned mass. The fuel  
12 loading of farmland was assumed to be 0.5 kg/m<sup>2</sup> (Wiedinmyer et al., 2006). However, in China,  
13 crop straw is used in multiple ways that differ regionally, like biofuel, biogas production and  
14 animal feed supply, which is highly dependent on crop species. We estimated the emission  
15 using a “bottom-up” method by fully considering crop yields, crop-specific straw usage and  
16 combustion efficiency.

17 During this agricultural fire event, the spatial pattern of fire detections in Figure 2a indicates  
18 that open burning of straw mostly concentrated in northern parts of Anhui and Jiangsu province  
19 and got extremely severe on 9 and 13 June, as displayed in Figure 2b. Burning of crop residues  
20 dominated local emissions of atmospheric pollutants when compared with corresponding  
21 anthropogenic emissions. Taking BC for instance (Figure 2c and d), emission rate from field  
22 burning of crop residues far outweighed that from industry, power plant, residential activity and  
23 transportation combined (Li et al., 2015).

## 24 **2.2 Numerical simulation**

25 The numerical simulations in this study were conducted using WRF-Chem version 3.6.1, which  
26 is an online-coupled chemical transport model considering multiple physical and chemical  
27 processes, including emission and deposition of pollutants, advection and diffusion, gaseous  
28 and aqueous chemical transformation, aerosol chemistry and dynamics (Grell, G. et al., 2011).  
29 The model has been widely utilized to investigate aerosol-radiation-cloud interactions and  
30 aerosol-boundary layer feedback (Grell, G. et al., 2011; Zhao, C. et al., 2013; Fan et al., 2015;  
31 Huang et al., 2015; Ding et al., 2016; Gao et al., 2016). In the present work, we adopted two  
32 nested model domains centred at 115.0°E, 33.0°N (Figure 1a). The parent domain with a grid

1 resolution of 20 km covered the eastern China and its surrounding areas to get synoptic forcing.  
2 The fine resolution of 4 km for the inner one allowed better characterization of small-scale  
3 physical processes, especially those linked to convective motions, cloud formation and rainfall  
4 onset. There were 31 vertical layers from the ground level to the top pressure of 50 hPa, 20 of  
5 which were placed below 4 km to achieve finer vertical resolution within the boundary layer.  
6 The initial and boundary conditions of meteorological fields were updated from the 6-hour  
7 NCEP (National Centres for Environmental Prediction) global final analysis (FNL) data with a  
8  $1^\circ \times 1^\circ$  spatial resolution. To investigate the aerosols' radiative effects on 10 June 2012 when  
9 the precipitation was substantially modified, the simulations were conducted for the time period  
10 from 20 May to 15 June. The meteorological initializing date for 10 June was 12:00 UTC on 9  
11 June. Each run covered 60 hours and the last 48-hour modelling results were kept. The chemical  
12 outputs from the preceding run were used as the initial conditions for the following run. The  
13 first 20 days were regarded as the model spin-up period for atmospheric chemistry, so as to  
14 better characterize aerosol distributions and minimize the influences of initial conditions and  
15 allow the model to reach a state of statistical equilibrium under the applied forcing (Berge et  
16 al., 2001; Lo et al., 2008).

17 Key parameterization options for the WRF-Chem modelling were the Noah land surface  
18 scheme to describe the land-atmosphere interactions (Ek et al., 2003), the YSU boundary layer  
19 scheme (Hong, 2010), and the RRTMG short- and long-wave radiation scheme (Mlawer et al.,  
20 1997). The Lin microphysics scheme that accounts for six forms of hydrometeor (Lin et al., 1983)  
21 together with the Grell cumulus parameterization was applied to reproduce the cloud and  
22 precipitation processes (Grell, G. A. and Devenyi, 2002) for the coarse domain. Cumulus  
23 parameterization was switched off for the inner domain. For the numerical representation of  
24 atmospheric chemistry, we used the CBMZ (Carbon-Bond Mechanism version Z)  
25 photochemical mechanism combined with MOSAIC (Model for Simulating Aerosol  
26 Interactions and Chemistry) aerosol model (Zaveri and Peters, 1999; Zaveri et al., 2008).  
27 Aerosols were assumed to be spherical particles. The size distribution was divided into four  
28 discrete size bins defined by their lower and upper dry particle diameters (0.039–0.156, 0.156–  
29 0.625, 0.625–2.5, and 2.5–10.0  $\mu\text{m}$ ). Aerosols in each size bin were assumed to be internally  
30 mixed and their optical properties, including extinction coefficient, single-scattering albedo  
31 (SSA) and asymmetry factor, were computed based on Mie theory (Fast et al., 2006) using  
32 volume averaged refractive indices (Barnard et al., 2010). Similar model configurations and  
33 settings have achieved good performance in our previous simulations over the eastern China

1 (Huang et al., 2015; Ding et al., 2016). Detailed configurations and domain settings are listed  
2 in Table 1.

3 Both natural and anthropogenic emissions were included for the regional WRF-Chem  
4 modelling in the present work. Typical anthropogenic emissions were obtained from the Multi-  
5 resolution Emission Inventory for China (MEIC) database (Li et al., 2015), in which emissions  
6 sources were classified into five main sectors: power plants, residential combustion, industrial  
7 processes, on-road mobile sources, and agricultural activities. This database covers most of  
8 anthropogenic pollutants, such as SO<sub>2</sub>, NO<sub>x</sub>, CO, volatile organic compounds (VOCs), PM,  
9 BC, and OC. NH<sub>3</sub> emission over China was derived from Huang et al. (2012a). VOCs emitted  
10 from typical anthropogenic activities and aforementioned crop residue burning were speciated  
11 into model-ready lumped species using profiles for Carbon-Bond Mechanism (Hsu et al., 2006).  
12 The biogenic VOC and NO emissions were calculated online by using the Model of Emissions  
13 of Gases and Aerosols from Nature (MEGAN) that embedded in WRF-Chem (Guenther et al.,  
14 2006). More than 20 biogenic species, including isoprene, monoterpenes (e.g.,  $\alpha$ -pinene and  $\beta$ -  
15 pinene) and sesquiterpenes, were considered and then involved in the photochemistry  
16 calculation. In China, crop residues are usually burned in piles, which is characterized by short-  
17 lived and small-scale smoldering. Consequently, the plume rise of biomass burning plumes was  
18 not considered in this study, and the straw fire emission was placed in the lowest two levels  
19 from the surface to around 50 meter in this simulation.

20 Previous studies have shown that, under highly polluted conditions, the ARI dominated over  
21 the aerosol-cloud interaction (ACI) that is related to aerosols' ability to act as CCN (e.g.,  
22 Rosenfeld et al., 2008; Fan et al., 2015). We also conducted another numerical experiment  
23 which included both ACI and ARI. The ACI-induced radiative perturbations were much less  
24 notable than those caused by ARI both at the surface and in the atmosphere (Figure S2),  
25 implying the dominant role of ARI during this fire event. Since that the focus of this study is  
26 on ARI and ACI's effect was not that significant, the prognosed aerosol was disabled to act as  
27 cloud condensation nuclei (CCN) or ice nuclei (IN) in the simulations and therefore the effects  
28 from ACI were not accounted for in the following analysis. Accordingly, wet scavenging of  
29 aerosol was disabled too. In order to disentangle aerosols' role in radiative transfer and  
30 subsequent effects on cloud and precipitation during this biomass-burning event in the mid-  
31 June of 2012, we designed three parallel numerical experiments. Domain settings and model  
32 configurations for these simulations were exactly the same as mentioned before. The control



1 experiment (CTL) did not include aerosol's effects on either longwave or shortwave radiation  
2 transfer. On the contrary, the other two took account of aerosols' perturbations on radiation  
3 transfer: ARI-A with anthropogenic emissions (anthropogenic activities refer to power  
4 generation, transportation, industrial and residential activities hereafter) and ARI-AB that  
5 included both anthropogenic activities and biomass burning emissions.

### 6 **3 Results and discussions**

#### 7 **3.1 Fire-induced pollution and observed anomalies in meteorology**

8 As demonstrated by existing studies (Andreae et al., 1988; Huang et al., 2012c; Ding et al.,  
9 2013a), air quality was dramatically deteriorated and the visibility was impeded during biomass  
10 burning events. We compare the simulated daily averaged PM<sub>10</sub> concentration with the  
11 corresponding measurements derived from Air Pollution Index (API) in Figure 3 (If not  
12 mentioned specially, the simulation refers to ARI-AB experiment hereafter). Both observations  
13 and simulations manifested the fact that intensive agricultural fires led to the severe pollution  
14 in mid-June. Since 9 June when the detected fire spots became intense and extensive, PM<sub>10</sub>  
15 concentrations in northern Anhui and northwest Jiangsu province began to increase, especially  
16 for those regions near the fire location. For instance, the observed daily mean PM<sub>10</sub>  
17 concentrations reached up to around 250 µg/m<sup>3</sup> at Fuyang (FY) and Xuzhou (XZ) and even  
18 exceeded 400 µg/m<sup>3</sup> at Bengbu (BB) on 9 June (the locations of cities mentioned in this article  
19 are labelled in Figure 2). Although the simulated temporal variations agree with observations,  
20 model-predicted PM<sub>10</sub> concentration at FY and BB were 196 and 168 µg/m<sup>3</sup>, respectively. The  
21 underestimation might be due to that rapid formation of secondary aerosol like sulfate and  
22 secondary organic matters is not so well described in current atmospheric chemical transport  
23 models (Capes et al., 2008; Xie et al., 2015). XZ and BB suffered from the second-round fire  
24 smoke two days later, with a maximum daily mean concentration of 548 µg/m<sup>3</sup> observed at BB.  
25 Figure 4 illustrates the satellite-retrieved 660-nm aerosol optical depth (AOD) and SSA from  
26 MODIS Aerosol Product MOD04\_L2 (daily level 2 data produced at the spatial resolution of  
27 10 km, collection 6) around 11:00 local time (LT) on 9 June when the first-round of extensive  
28 fire pollution broke out. Their comparisons with ARI-AB modelled spatial distributions of  
29 PM<sub>2.5</sub> and BC column-integrated mass loadings further confirm model's ability to reproduce  
30 atmospheric pollution for this event. The AOD observation shows that high aerosol loadings  
31 were concentrating in northeast Anhui and the north-central Jiangsu, shaping a belt of pollution  
32 from the fire sites to the downwind areas. The similar pattern was also simulated by the model.

1 The PM<sub>2.5</sub> mass loading was found to exceed 200 mg/m<sup>2</sup> near BB, NJ and most parts of central  
2 Jiangsu. This strap-shaped pollution was particularly obvious in terms of BC column  
3 concentrations, which was also consistent with a relatively lower SSA along BB, Yangzhou  
4 (YZ) and Taizhou (TZ). While solely including anthropogenic emissions, ARI-A experiment  
5 failed to represent the spatial pattern of high AOD in the northern Anhui and Jiangsu and the  
6 low SSA value near BB (Fig.4 a, d).

7 Along with the severe air pollution and poor visibility, anomalies in meteorology occurred on  
8 9-10, June. Ding et al. (2013a) found that, during these two days, a sharp decline existed in the  
9 observed air temperature in NJ and YZ, compared with weather forecast results and NCEP FNL  
10 data, but the simulations and observations showed a good agreement when the heavy air  
11 pollution was not present before 8 June. At YZ the temperature difference was as high as 5.9  
12 and 9.2 °C on 9 and 10 June, respectively. Simultaneously, measured solar radiation intensity  
13 and sensible heat flux showed very low values on 10 June in comparison with non-episode days.  
14 Moreover, local meteorological agency forecasted a convective rainfall to occur in NJ and  
15 surrounding areas in the afternoon of 10 June, with the rainfall centre passing by NJ around  
16 14:00 LT. However, this forecasted rainfall never happened that day.

17 On the basis of ground-based measurements, vertical sounding data, remote-sensing images  
18 and their comparisons with numerical simulations, we found that agricultural fires worsen  
19 regional air quality to a large extent and caused a series of anomalies in temperature and  
20 precipitation in the mid-June of 2012. How the biomass burning plumes influenced the air  
21 temperature and precipitation will be the main issue to be addressed in the following discussions.

### 22 **3.2 Perturbations in energy budget and temperature responses**

23 To better understand aerosols' role in the energy re-allocation on 10 June when precipitation  
24 was evidently modified, radiative forcing in the atmosphere and at the ground surface was  
25 estimated by differentiating the CTL, ARI-A and ARI-AB simulations (Figure 5). At the surface,  
26 daily mean incident short-wave radiation was weakened by 45.5 W/m<sup>2</sup> (averaged over the inner  
27 domain) as the extinction of aerosol was quite large with a satellite-observed 660-nm AOD  
28 exceeding 2.0 (Figure 4b). Meanwhile, about 60.4 W/m<sup>2</sup> shortwave energy was blocked in the  
29 atmosphere over the inner domain due to the fact that absorbing aerosols were accumulated on  
30 that day. A positive domain-averaged radiative forcing of +14.9 W/m<sup>2</sup> was simulated at the top  
31 of the atmosphere (TOA) on 10 June. Comparatively, radiative perturbations caused by

1 agricultural fires (ARI-AB minus ARI-A) were more substantial than those due to  
2 anthropogenic emissions (ARI-A minus CTL) in magnitude, particularly in the atmosphere, as  
3 presented in Figure 5. Spatially, radiative effects due to anthropogenic activities concentrated  
4 in the economically developed Yangtze River Delta region while agricultural fires exerted  
5 significant impact on radiation balance in northern and central Jiangsu and north part of Anhui.  
6 Table 2 compares the radiative perturbations caused by anthropogenic activities and biomass  
7 burning emissions over three regions with distinct precipitation changes (marked in Figure 8).  
8 As shown, both of them tended to heat the atmosphere and cool the ground surface. Fire plumes  
9 dominated the radiative effect in terms of atmospheric warming. Radiation measurements  
10 collected at Heifei (HF) and sensible and latent heat flux recorded at Lishui (in South Nanjing)  
11 are compared with the diurnal variations of corresponding simulations in Figure 6, which  
12 supports that significant radiative perturbations took place at NJ and HF. Substantially  
13 weakened daytime solar irradiance was observed on 10 June, when the peak value of  
14 downwelling shortwave radiation was  $618.3 \text{ W/m}^2$  at HF and was only  $309.7 \text{ W/m}^2$  at NJ.  
15 Taking aerosol's effect on radiation into account tended to predict lower downwards solar  
16 radiation, which was closer to observation for both cities. Reduction in shortwave energy hitting  
17 the surface in turn decreased outgoing heat fluxes, and therefore simulated sensible and latent  
18 heat fluxes at 12:00 LT on 10 June in ARI-AB experiment decreased by 89.3 and 76.1  $\text{W/m}^2$ ,  
19 respectively, compared with CTL experiment.

20 Overall, the magnitude of the radiative forcing on 10 June was comparable in northern Anhui  
21 and central Jiangsu, differing from the distribution pattern of fire-induced air pollution that  
22 remarkably concentrated in northern Anhui. As revealed in our previous estimation, among all  
23 components of the ambient aerosols, BC is the most important disturber of shortwave radiation  
24 transfer at the surface and in the atmosphere as well (Huang et al., 2015; Ding et al., 2016).  
25 Although fire emission mostly concentrated in the northern Anhui and resulted in a high BC  
26 concentration of  $20 \mu\text{g/m}^3$  there, high-altitude BC was spread much more broadly. At an altitude  
27 of 2 km, BC concentration around  $5 \mu\text{g/m}^3$  stretched from northern Anhui to central Jiangsu  
28 (Figure S3). Such distinct distributions between two layers were partly attributed to the stagnant  
29 condition near the surface and stronger horizontal transport in the upper level. It is emphasized  
30 that upper-level BC has higher absorbing efficiency (Ding et al., 2016). That is why the  
31 distributions of both positive radiative forcing in the atmosphere and negative forcing at the  
32 surface generally consisted with BC's spatial pattern in the upper air.

1 The perturbations in the energy budget and the following re-allocation gave rise to substantial  
2 modulation in vertical stratification of air temperature. In comparison with CTL experiment,  
3 ARI-AB experiment predicted an obvious decline in near-surface temperature by considering  
4 the effects of aerosol-radiation interaction. Hourly observed 2-m air temperature was compared  
5 with corresponding simulations by two experiments during the time period from 8 to 15 June.  
6 Model-performance statistics including mean bias (MB), mean error (ME) and root mean  
7 square error (RMSE) are presented in Table 3. As shown, CTL simulation had a systematic  
8 positive bias in 2-m temperature and ARI predicted lower temperature for both areas near fire  
9 locations (BB and XZ) and downwind regions (NJ and SY). The decreases in temperature were  
10 pronounced in BB and XZ with a large difference of approximately 1.2 °C, which notably  
11 narrowed the gaps with observations. On 10 June when the fire-induced pollution became  
12 intensive, the magnitude of surface cooling was remarkably high near the fire sites.  
13 Temperature response in Figure 5e support this cooling effect. For instance, compared to CTL,  
14 simulated near-surface temperature by ARI-A and ARI-AB experiment at XZ was cooled by  
15 almost 1.2 and 8.0 °C at 20:00 LT on 10 June (Figure 7b). In addition to the cooling tendency  
16 of near-surface temperature, aerosols' radiative effects also increased air temperature at a higher  
17 altitude, which were more apparent over the downwind areas (Figure 5f). According to the  
18 comparisons between simulated temperature profiles by the three parallel experiments in Figure  
19 7, the warming of air temperature was particularly evident around an altitude of 2 km at SY  
20 with a maximum of 3.0 °C and biomass burning aerosol played a leading role.

21 The different temperature responses over the source region of fire emission and downwind areas  
22 could be partially interpreted by the fact that near the fire locations, pronounced surface cooling  
23 counteracted part of the atmospheric warming, which would otherwise elevate upper-air  
24 temperature, through vertical mixing; while for the downwind area where the surface was less  
25 radiatively cooled, the atmosphere was prone to being warmed. As a result of surface cooling  
26 and atmospheric heating, vertical convective motions were weakened, triggering perturbations  
27 in pressure and wind fields (Figure 5e and f). It is obvious that suppressed convection was  
28 generally along with the resultant wind convergence around 2 km and surface divergence,  
29 which may further play a significant role in water vapor transport, entrainment and also cloud  
30 formation.

### 1 3.3 Effects on cloud and precipitation

2 In addition to the attenuation of solar radiation and the modulation in temperature gradients,  
3 precipitation also showed many disparities between CTL and ARI-AB simulations. The satellite  
4 observation from Tropical Rainfall Measuring Mission (TRMM) Multisatellite Precipitation  
5 Analysis product (3B42), which provides merged-infrared precipitation information at a  
6  $0.25 \times 0.25^\circ$  spatial resolution and has been demonstrated to perform well in East China  
7 (Simpson et al., 1988; Zhao and Yatagai, 2014), was used to evaluate the simulated precipitation.  
8 As demonstrated in Figure 8, ARI-AB experiment agrees better with TRMM observations than  
9 CTL experiment in terms of precipitation intensities and also spatial pattern on 10 June.  
10 Specifically, CTL and ARI-A simulation suggested a convective rain in Zone 1 (NJ and its  
11 adjacent areas) around 14:00 LT (the locations of Zone 1-3 are marked in Figure 8), however  
12 the ARI-AB simulation did not show any precipitation then, consistent with the TRMM  
13 observations. Besides, ARI-AB displayed enhanced precipitation in northern Jiangsu province.  
14 A precipitation with the intensity of 3 and 5 mm/h was predicted by ARI-AB experiment in  
15 Zone 2 (XZ and its adjacent areas) and Zone 3 (SY and its adjacent areas), which, however,  
16 never occurred in CTL and ARI-A experiment. Concerning temporal variations, 3-hour  
17 precipitation rates for these three zones derived from TRMM 3B42 retrievals are plotted in  
18 Figure 9. Compared to CTL and ARI-A experiment, ARI-AB experiment which considered  
19 radiative effects of aerosol from both anthropogenic and biomass burning emissions succeeded  
20 in capturing the approximate onset time for all the three regions.

#### 21 3.3.1 Suppressed daytime precipitation

22 Over Zone 1, CTL and ARI-A simulation produced a convective rainfall event in the afternoon  
23 that actually did not happen, while ARI-AB simulation with no precipitation was closer to the  
24 observations. According to the energy budget and radiation flux calculation (Figure 5), on 10  
25 June more than  $6 \text{ MJ/m}^2$  solar radiation that supposed to reach the surface was blocked in the  
26 atmosphere over Zone 1, most of which was caused by biomass burning aerosol. The presence  
27 of light-absorbing aerosols reduced sensible heat flux and evapotranspiration at the surface  
28 (Figure 6). Large-eddy simulation for biomass burning regions of Brazil deduced that the peak  
29 reductions in sensible and latent heat flux were  $60$  and  $70 \text{ W/m}^2$  (Feingold et al., 2005), which  
30 are quantitatively similar to those near NJ estimated in this work. It was shown that reduced  
31 surface flux alone was sufficient to explain the observed cloud dissipation during the biomass  
32 burning event in Brazil. For this case, this convective rain got disappeared merely by nudging

1 2-m temperature in the WRF modelling run by Ding et al. (2013a), highlighting the importance  
2 of surface flux modification in the development of these convective clouds.

3 To figure out the role of vertical thermal behaviors in Zone 1, temporal variations of zone-  
4 averaged differences in temperature, relative humidity (RH) profiles between ARI-AB and CTL  
5 experiments are illustrated in Figure 10a and b. From 9:00 LT in the morning, a 1-km-thick belt  
6 with BC-laden smoke approached Zone 1 and covered over the boundary layer top. The  
7 radiative extinction by the elevated smoke layer led to a cooling effect at the surface, which  
8 reduced the boundary layer height and decreased the air temperature in the boundary layer.  
9 Simultaneously, relatively strong warming effect between the altitudes of 1-3 km increased the  
10 air temperature above the boundary layer. The cooling at the lower altitude and warming at the  
11 upper altitude made the stability significantly increased, especially near the top of the boundary  
12 layer, which further suppressed the development of boundary layer. For the perturbations in  
13 humidity, the enhanced atmospheric stability reduced the boundary layer height and hindered  
14 the upward transport of water vapor to a higher altitude, while the heating aloft decreased RH  
15 by increasing the air temperature there. These led to a resultant decrease of more than 20% in  
16 RH above the boundary layer. A more stable and shallower boundary layer in ARI-AB  
17 experiment had a tendency to reduce convective mixing and effectively cut off the cloud layer  
18 from its source of moisture, subsequently desiccating the cloud layer, and leading to  
19 substantially weakened vertical motions. Accordingly, ARI-AB simulated updraft velocity  
20 above 1 km was only one-tenth that of CTL experiment in the afternoon of 10 June, as  
21 demonstrated in Figure 10f. Even though anthropogenic aerosol also weakened convective  
22 motions in ARI-A experiment, the potential temperature profile was hardly changed and the  
23 weakening effect of convection was not comparable with that caused by biomass burning  
24 aerosols. Therefore, compared with CTL and ARI-A experiment, much less moist static energy  
25 (MSE) was carried upwards and the excess MSE accumulated in a shallower boundary layer  
26 due to much weaker convection in ARI-AB experiment (Figure 10e).

27 In addition to Zone 1, this warmed belt was also blanketing a wider range from 116 to 120 °E  
28 at the moment when the CTL-predicted rainfall started (Figure 9a shows that the rainfall  
29 occurred around 14:00 LT), as shown in the longitude-height cross sections of temperature  
30 difference between CTL and ARI-AB experiment in Figure 10c. In CTL run, cumulus cloud  
31 layer appeared above the inversion capping the boundary layer (Figure 10d). However, the  
32 absorbing aerosol in ARI-AB run heated the atmosphere aloft and stabilized the sub-cloud layer.

1 The decrease in specific humidity was collocated with warmed upper air since that atmospheric  
2 heating and surface cooling weakened vertical convection and further reduced the vertical  
3 transport of water vapor. Lower entrainment rate together with higher saturation pressure  
4 resulted in daytime decoupling and thinning of the cloud layer all along the longitude from 116  
5 to 120 °E. This effect might be further strengthened by a positive feedback loop as described  
6 by Jacobson (2002) in which cloud loss leads to an increasing opportunity for BC's light  
7 absorption.

### 8 3.3.2 Enhanced nocturnal precipitation

9 A precipitation rate of over 2.5 mm/h was observed around 19:00-20:00 LT on 10 June in XZ  
10 and its surrounding areas (Zone 2). Only ARI-AB simulation captured this precipitation event.  
11 As shown in Figure 11a, there existed two layers with a high BC concentration of up to 10  
12  $\mu\text{g}/\text{m}^3$  during daytime over Zone 2. One was near the surface and peaked around 18:00 LT,  
13 which could be linked to local fire emissions. The other one was lying over the boundary layer  
14 top, which was apparent at an altitude of 0.8 km before the boundary layer developed and at 2  
15 km after 15:00 LT. It was very likely to be associated with the transport of upstream fire  
16 pollution. Owing to strong radiative heating effect of BC, a warmer layer was formed above  
17 1 km during daytime with temperature increase over 1.0 °C. On the contrary, near-surface  
18 temperature kept decreasing. The decline reached its maximum around 20:00 LT. It was also  
19 supported by Figure 7b in which the near-surface temperature decreased by almost 8.0 °C at  
20 XZ. Until 16:00 LT, the upper-air warming due to radiative absorption was gradually  
21 compensated by cooling from the surface through vertical mixing. Changes in RH were almost  
22 opposite of those in air temperature. Around 18:00 LT, RH at 3-km altitude started to increase  
23 and then a precipitating cloud formed there.

24 To get a better insight on the dynamical processes that contribute to precipitation change,  
25 longitude-height cross section of zonal mean responses of temperature, water vapor and wind  
26 profile just before the onset time of precipitation are demonstrated in Figure 11c and d.  
27 Noteworthy is that warmed upper air between 117 to 119 °E led to less condensation there.  
28 More water vapor accumulated below 1 km and was then transported toward Zone 2 by the  
29 prevailing east wind near the surface, leading to an excess water vapor over Zone 2 in ARI-AB  
30 experiment (Figure 11e). Simultaneously, radiatively heated air parcel with a temperature  
31 increase of 0.5 °C was found around 2 km over Zone 2. The warmer layer around 2-3 km  
32 combined with large drops in temperature beneath resulted in a buoyancy-driven lifting force.

1 Moreover, horizontal heterogeneity in atmospheric heating provided the low-level convergence  
2 for maintaining convection in a conditionally unstable atmosphere around 3 km. The zone-  
3 averaged updraft velocity in ARI-AB experiment tripled that predicted by CTL and ARI-A at  
4 the altitude of 3 km when the precipitation began (Figure 11f). Understandably, what made the  
5 precipitating cloud formed around 3 km over Zone 2 were the accumulated MSE near the  
6 surface and anomalous updraft of the air that favored the vertical uplift of MSE. The release of  
7 latent heat may increase the upper-air instability and in turn enhance the precipitation.

8 For the downwind region Zone 3, the warming effect caused by aerosol-radiation interaction  
9 was evident for the air column above 0.5 km all day long on 10 June (Figure 12a). The warming  
10 pattern was coincident with the distribution of BC concentration since BC is the predominant  
11 light-absorbing aerosol specie in the atmosphere. As a result of increased air temperature, RH  
12 decreased substantially during daytime. At late night, an extra precipitating cloud formed above  
13 2 km over Zone 3 in ARI-AB simulation, leading to a nocturnal precipitation with a strength of  
14 approximately 6 mm/h at 01:00 LT on 11 June. What triggered this rainfall event is a bit  
15 complicated than that over Zone 2. First, the whole air column was getting cooled at the moment  
16 when the precipitation took place, inevitably raising RH value. The RH increase was quite  
17 apparent at the altitude of 3-4 km. Second, daytime radiative absorption by BC-laden plumes  
18 around 2 km heated the surrounding air. Relatively warmer layer at the altitude of ~ 2 km  
19 generated a positive buoyant updraft (Figure 12f), hence air parcel there was displaced upwards  
20 along with enhanced convergence carrying in moist air. This effect has been proposed by Fan  
21 et al. (2015) as part of termed “enhanced conditional instability”, by which absorbing aerosols  
22 escalate convection downwind of a heavily polluted area and promote precipitation.  
23 Comparatively, radiative heating of biomass burning aerosol was the main contributor to the  
24 significant enhancement of upper-level updraft. Last but not the least, spatially heterogeneous  
25 aerosol-related heating was associated with greater horizontal temperature lapse, resulting in a  
26 convergence flow above 3 km with an additional onshore wind (Figure 12d). Zone 3 is only  
27 about 20 km from the Yellow sea. It is plausible that more water vapor-saturated air masses  
28 originating from the ocean brought in excess water vapor and consequently elevated the  
29 humidity above 3 km. More MSE accumulated above 3 km in ARI-AB experiment compared  
30 with those simulated by CTL and ARI-A experiment before the precipitation also support this  
31 view (Figure 12e). We suggest that these precipitating clouds formed because of instability at  
32 the top of the smoke layer, driven by the strong radiation absorption that warmed the



1 surrounding air. Therefore, the heated BC-laden air was ascended and cooled, leading to the  
2 formation of clouds preferentially in the conditionally unstable zone in the upper air.

### 3 **3.4 Uncertainties**

4 Though the modelling work here characterized cloud and precipitation anomalies during the  
5 biomass burning event, we may also question to what extent the modelling reproduced the  
6 relevant processes in the real world. As widely acknowledged, accurate simulation of smoke  
7 plume and prediction of clouds are both challenging for regional/global models. One  
8 contributor to the uncertainties is the characterization of fire emission. The magnitude was  
9 determined by statistical information and laboratory experiment data, whose accuracy and  
10 representativeness may introduce some uncertainties. The spatiotemporal distribution of fire  
11 emission was allocated based on MODIS retrievals. Loss of information due to cloud coverage  
12 and poor detection efficiency of short-lived or small-scale fires are major limitations (Giglio et  
13 al., 2003). Another challenge is quantification of heat release from biomass burning and  
14 subsequent effects on local and regional meteorology. Furthermore, much emphasis has been  
15 paid to the vertical distribution of absorbing aerosol, to which the cloud response is highly  
16 sensitive (Koch and Del Genio, 2010). The vertical profile of absorbing aerosol in this  
17 simulation underwent little constrain due to limited observation at that time. The regional model  
18 is hardly capable of precisely presenting turbulent flows and vertical transport, thus introducing  
19 uncertainties in three dimensional distributions of BC. It also should be noted that BC is co-  
20 emitted with other components such as OC and sulfur dioxide that oxidizes to sulfate (Xie et  
21 al., 2015). Mixing with other scattering aerosol would considerably amplify the absorbing  
22 efficiency of BC. Model's ability to account for the evolution of mixing state and how to  
23 quantify its amplification also affect the simulated radiative behaviors. Besides, poorly  
24 recognized secondary organic carbon (SOC) formation processes and its light absorption makes  
25 it imperative to reassess and redefine the chemical mechanism and optical properties of OC in  
26 models (Saleh et al., 2014). The large uncertainty in simulating clouds and further aerosol-cloud  
27 interaction is another limitation (e.g., Wang et al., 2011; Tao et al., 2012). To improve the model  
28 performance in all these chemical and physical processes, more comprehensive measurements  
29 and modelling efforts are needed in the future.

30

## 1    **4    Conclusions**

2    To investigate radiative effects of aerosol-radiation interaction on cloud and precipitation  
3    modifications during the exceptionally active agricultural fire season in June 2012, a bottom-  
4    up emission inventory of crop open burning was developed and then the fully coupled online  
5    WRF-Chem model was applied in this work. The evaluation of simulation through ground-  
6    based observations and satellite retrievals showed that the model generally captured spatial  
7    patterns and temporal variations of fire pollution, which was predominantly concentrating over  
8    northern Anhui and central-north Jiangsu. It is evident that post-harvest burning of crop residues  
9    emitted a tremendous amount of atmospheric pollutants and deteriorated regional air quality to  
10   a large extent in East China. Elevated concentration of aerosols, particularly light-absorbing  
11   BC, would heat the atmosphere and cool the ground surface through both direct solar radiation  
12   attenuation (direct radiative forcing) and cloud redistribution (semi-direct radiative forcing).  
13   This radiative cooling (heating) effects were distinct close to (downwind from) the source  
14   regions of fire sites. Adjusted temperature stratification was intimately linked to small-scale  
15   processes such as turbulent mixing, entrainment and the evolution of the boundary layer.  
16   Subsequently, over Nanjing and its adjacent regions, absorbing aerosols immediately above the  
17   boundary layer top increased the inversion beneath, reducing available moisture and leading to  
18   a burn-off effect of cloud. Meanwhile, fire plumes played an enhancement role in nocturnal  
19   precipitation over northern Jiangsu by increasing up-level convective activity and fostering  
20   low-level convergence that carries in more moist air. Overall, aerosols' radiative effect on  
21   precipitation modification is therefore likely to depend to a large extent on local meteorological  
22   conditions like atmospheric instability and humidity.

23

## 24    **Acknowledgements**

25    This work was supported by the National Natural Science Foundation of China  
26    (D0512/91544231, D0512/41422504, and D0510/41505109). Part of this work was supported  
27    by the Jiangsu Provincial Science Fund for Distinguished Young Scholars awarded to A. J.  
28    Ding (No. BK20140021).

29

30

1 Table 1. WRF-Chem modelling configuration options and settings.

Domain setting		
	Domain 1	Domain 2
Horizontal grid	130×130	160×160
Grid spacing	20 km	4 km
Vertical layers	31	31
Configuration options		
Long-wave radiation	RRTMG	
Short-wave radiation	RRTMG	
Land-surface	Noah	
Boundary layer	YSU	
Microphysics	Lin et al.	
Cumulus parameterization	Grell–Deveny (only for domain 1)	
Photolysis	Fast-J	
Gas-phase chemistry	CBMZ	
Aerosol scheme	MOSAIC	

2

3

1 Table 2. Radiative perturbations at the surface (SUR) and in the atmosphere (ATM) caused by  
 2 anthropogenic activities and agricultural fires for three zones with distinct precipitation changes.

	SUR		ATM	
	Anthropogenic activities	Agricultural fires	Anthropogenic activities	Agricultural fires
Zone 1	-27.3	-35.1	36.3	41.1
Zone 2	-33.3	-41.7	31.9	45.6
Zone 3	-23.8	-14.8	27.7	21.1

3  
4  
5  
6

7 Table 3. Statistical analyses of the simulated 2-m temperature and the corresponding  
 8 observations at four different cities.

	MB <sup>a</sup>		ME <sup>a</sup>		RMSE <sup>a</sup>	
	CTL	ARI	CTL	ARI	CTL	ARI
NJ	0.85	0.37	1.70	1.66	2.39	2.15
BB	2.19	0.98	2.51	1.65	3.27	2.16
XZ	1.67	0.51	2.37	2.19	3.32	2.89
SY	-0.28	-0.46	1.97	1.65	2.52	2.03

9 <sup>a</sup>MB, ME and RMSE refer to mean bias, mean error and root-mean-square error respectively.  
 10

## 1 **References**

- 2 Akagi, S. K., Yokelson, R. J., Wiedinmyer, C., Alvarado, M. J., Reid, J. S., Karl, T., Crounse,  
3 J. D., and Wennberg, P. O.: Emission factors for open and domestic biomass burning for use in  
4 atmospheric models, *Atmos. Chem. Phys.*, 11, 4039-4072, 10.5194/acp-11-4039-2011, 2011.
- 5 Andreae, M. O., Browell, E. V., Garstang, M., Gregory, G. L., Harriss, R. C., Hill, G. F., Jacob,  
6 D. J., Pereira, M. C., Sachse, G. W., Setzer, A. W., Dias, P. L. S., Talbot, R. W., Torres, A. L.,  
7 and Wofsy, S. C.: Biomass-Burning Emissions and Associated Haze Layers over Amazonia, *J.*  
8 *Geophys. Res. Atmos.*, 93, 1509-1527, doi:10.1029/Jd093id02p01509, 1988.
- 9 Andreae, M. O., and Merlet, P.: Emission of trace gases and aerosols from biomass burning,  
10 *Global Biogeochem. Cy.*, 15, 955-966, doi:10.1029/2000gb001382, 2001.
- 11 Andreae, M. O., Rosenfeld, D., Artaxo, P., Costa, A. A., Frank, G. P., Longo, K. M., and Silva-  
12 Dias, M. A. F.: Smoking rain clouds over the Amazon, *Science*, 303, 1337-1342,  
13 doi:10.1126/science.1092779, 2004.
- 14 Barnard, J. C., Fast, J. D., Paredes-Miranda, G., Arnott, W. P., and Laskin, A.: Technical Note:  
15 Evaluation of the WRF-Chem "Aerosol Chemical to Aerosol Optical Properties" Module using  
16 data from the MILAGRO campaign, *Atmos. Chem. Phys.*, 10, 7325-7340, 10.5194/acp-10-  
17 7325-2010, 2010.
- 18 Bergamaschi, P., Hein, R., Heimann, M., and Crutzen, P. J.: Inverse modeling of the global CO  
19 cycle 1. Inversion of CO mixing ratios, *J. Geophys. Res. Atmos.*, 105, 1909-1927,  
20 doi:10.1029/1999jd900818, 2000.
- 21 Berge, E., Huang, H. C., Chang, J., and Liu, T. H.: A study of the importance of initial  
22 conditions for photochemical oxidant modeling, *J. Geophys. Res. Atmos.*, 106, 1347-1363,  
23 doi:10.1029/2000jd900227, 2001.
- 24 Bond, T. C., Doherty, S. J., Fahey, D. W., Forster, P. M., Berntsen, T., DeAngelo, B. J., Flanner,  
25 M. G., Ghan, S., Karcher, B., Koch, D., Kinne, S., Kondo, Y., Quinn, P. K., Sarofim, M. C.,  
26 Schultz, M. G., Schulz, M., Venkataraman, C., Zhang, H., Zhang, S., Bellouin, N., Guttikunda,  
27 S. K., Hopke, P. K., Jacobson, M. Z., Kaiser, J. W., Klimont, Z., Lohmann, U., Schwarz, J. P.,  
28 Shindell, D., Storelvmo, T., Warren, S. G., and Zender, C. S.: Bounding the role of black carbon  
29 in the climate system: A scientific assessment, *J. Geophys. Res. Atmos.*, 118, 5380-5552,  
30 10.1002/jgrd.50171, 2013.
- 31 Boschetti, L., Roy, D., and Hoffmann, A.: MODIS Collection 5 Burned Area Product-MCD45,  
32 User's Guide, Ver, 2, 2009.
- 33 Capes, G., Johnson, B., McFiggans, G., Williams, P., Haywood, J., and Coe, H.: Aging of  
34 biomass burning aerosols over West Africa: Aircraft measurements of chemical composition,  
35 microphysical properties, and emission ratios, *J. Geophys. Res. Atmos.*, 113, 2008.
- 36 Chi, X., Winderlich, J., Mayer, J.-C., Panov, A. V., Heimann, M., Birmili, W., Heintzenberg,  
37 J., Cheng, Y., and Andreae, M. O.: Long-term measurements of aerosol and carbon monoxide  
38 at the ZOTTO tall tower to characterize polluted and pristine air in the Siberian taiga, *Atmos.*  
39 *Chem. Phys.*, 13, 12271-12298, doi:10.5194/acp-13-12271-2013, 2013.
- 40 Crutzen, P. J., and Andreae, M. O.: Biomass Burning in the Tropics - Impact on Atmospheric  
41 Chemistry and Biogeochemical Cycles, *Science*, 250, 1669-1678, doi:10.1126/science.250.  
42 4988.1669, 1990.

1 Ding, A. J., Fu, C. B., Yang, X. Q., Sun, J. N., Petaja, T., Kerminen, V. M., Wang, T., Xie, Y.,  
2 Herrmann, E., Zheng, L. F., Nie, W., Liu, Q., Wei, X. L., and Kulmala, M.: Intense atmospheric  
3 pollution modifies weather: a case of mixed biomass burning with fossil fuel combustion  
4 pollution in eastern China, *Atmos. Chem. Phys.*, 13, 10545-10554, 10.5194/acp-13-10545-  
5 2013, 2013a.

6 Ding, A. J., Fu, C. B., Yang, X. Q., Sun, J. N., Zheng, L. F., Xie, Y. N., Herrmann, E., Nie, W.,  
7 Petaja, T., Kerminen, V. M., and Kulmala, M.: Ozone and fine particle in the western Yangtze  
8 River Delta: an overview of 1 yr data at the SORPES station, *Atmos. Chem. Phys.*, 13, 5813-  
9 5830, 10.5194/acp-13-5813-2013, 2013b.

10 Ding, A. J., Huang, X., Nie, W., Sun, J. N., Kerminen, V. M., Petäjä, T., Su, H., Cheng, Y. F.,  
11 Yang, X. Q., Wang, M. H., Chi, X. G., Wang, J. P., Virkkula, A., Guo, W. D., Yuan, J., Wang,  
12 S. Y., Zhang, R. J., Wu, Y. F., Song, Y., Zhu, T., Zilitinkevich, S., Kulmala, M., and Fu, C. B.:  
13 Enhanced haze pollution by black carbon in megacities in China, *Geophys. Res. Lett.*,  
14 doi:10.1002/2016GL067745, 2016.

15 Draxler, R. R., and Rolph, G.: HYSPLIT (HYbrid Single-Particle Lagrangian Integrated  
16 Trajectory) model access via NOAA ARL READY website (<http://www.arl.noaa.gov/ready/hysplit4.html>). NOAA Air Resources Laboratory, Silver Spring, in, Md, 2003.

18 Ek, M. B., Mitchell, K. E., Lin, Y., Rogers, E., Grunmann, P., Koren, V., Gayno, G., and  
19 Tarpley, J. D.: Implementation of Noah land surface model advances in the National Centers  
20 for Environmental Prediction operational mesoscale Eta model, *J. Geophys. Res. Atmos.*, 108,  
21 Artn 8851, doi:10.1029/2002jd003296, 2003.

22 Fan, J. W., Rosenfeld, D., Yang, Y., Zhao, C., Leung, L. R., and Li, Z. Q.: Substantial  
23 contribution of anthropogenic air pollution to catastrophic floods in Southwest China, *Geophys.*  
24 *Res. Lett.*, 42, 6066-6075, 10.1002/2015GL064479, 2015.

25 Fast, J. D., Gustafson, W. I., Easter, R. C., Zaveri, R. A., Barnard, J. C., Chapman, E. G., Grell,  
26 G. A., and Peckham, S. E.: Evolution of ozone, particulates, and aerosol direct radiative forcing  
27 in the vicinity of Houston using a fully coupled meteorology-chemistry-aerosol model, *J.*  
28 *Geophys. Res. Atmos.*, 111, Artn D21305, doi:10.1029/2005jd006721, 2006.

29 Fearnside, P. M.: Global warming and tropical land-use change: greenhouse gas emissions from  
30 biomass burning, decomposition and soils in forest conversion, shifting cultivation and  
31 secondary vegetation, *Climatic change*, 46, 115-158, 2000.

32 Feingold, G., Jiang, H. L., and Harrington, J. Y.: On smoke suppression of clouds in Amazonia,  
33 *Geophys. Res. Lett.*, 32, Artn L02804, doi:10.1029/2004gl021369, 2005.

34 Gao, M., Carmichael, G., Wang, Y., Saide, P., Yu, M., Xin, J., Liu, Z., and Wang, Z.: Modeling  
35 study of the 2010 regional haze event in the North China Plain, *Atmospheric Chemistry &*  
36 *Physics*, 16, 2016.

37 Gao, X., Ma, W., Ma, C., Zhang, F., and Wang, Y.: Analysis on the current status of utilization  
38 of crop straw in China (in Chinese), *Journal of Huazhong Agricultural University*, 21, 242-247,  
39 2002.

40 Giglio, L., Descloitres, J., Justice, C. O., and Kaufman, Y. J.: An enhanced contextual fire  
41 detection algorithm for MODIS, *Remote. Sens. Environ.*, 87, 273-282, 10.1016/S0034-  
42 4257(03)00184-6, 2003.

1 Grell, G., Freitas, S. R., Stuefer, M., and Fast, J.: Inclusion of biomass burning in WRF-Chem:  
2 impact of wildfires on weather forecasts, *Atmos. Chem. Phys.*, 11, 5289-5303, 10.5194/acp-  
3 11-5289-2011, 2011.

4 Grell, G. A., and Devenyi, D.: A generalized approach to parameterizing convection combining  
5 ensemble and data assimilation techniques, *Geophys. Res. Lett.*, 29, Artn 1693, doi:  
6 10.1029/2002gl015311, 2002.

7 Guenther, A., Karl, T., Harley, P., Wiedinmyer, C., Palmer, P. I., and Geron, C.: Estimates of  
8 global terrestrial isoprene emissions using MEGAN (Model of Emissions of Gases and  
9 Aerosols from Nature), *Atmos. Chem. Phys.*, 6, 3181-3210, 2006.

10 Hong, S. Y.: A new stable boundary-layer mixing scheme and its impact on the simulated East  
11 Asian summer monsoon, *Q. J. Roy. Meteor. Soc.*, 136, 1481-1496, doi:10.1002/Qj.665, 2010.

12 Hsu, Y., Strait, R., Roe, S., and Holoman, D. S.: 4.0. Speciation Database Development  
13 Documentation. Final Report. EPA contract. Nos, EP-D-06.001, work assignment Numbers 0-  
14 03 and 68-D-02-063, WA 4-04 and WA 5-05. EPA/600/R-06/16. [http://www.epa.  
15 gov/ttn/chief/software/speciate/speciate4/documentation/speciatedoc\\_1206.pdf](http://www.epa.gov/ttn/chief/software/speciate/speciate4/documentation/speciatedoc_1206.pdf), 2006.

16 Huang, X., Song, Y., Li, M., Li, J., Huo, Q., Cai, X., Zhu, T., Hu, M., and Zhang, H.: A high -  
17 resolution ammonia emission inventory in China, *Global Biogeochem. Cy.*, 26, 2012a.

18 Huang, X., Li, M. M., Li, J. F., and Song, Y.: A high-resolution emission inventory of crop  
19 burning in fields in China based on MODIS Thermal Anomalies/Fire products, *Atmos. Environ.*,  
20 50, 9-15, 10.1016/j.atmosenv.2012.01.017, 2012b.

21 Huang, X., Song, Y., Li, M. M., Li, J. F., and Zhu, T.: Harvest season, high polluted season in  
22 East China, *Environ. Res. Lett.*, 7, Artn 044033, doi:10.1088/1748-9326/7/4/044033, 2012c.

23 Huang, X., Song, Y., Zhao, C., Cai, X. H., Zhang, H. S., and Zhu, T.: Direct Radiative Effect  
24 by Multicomponent Aerosol over China, *J. Climate*, 28, 3472-3495, 10.1175/Jcli-D-14-00365.1,  
25 2015.

26 Ito, A., Ito, A., and Akimoto, H.: Seasonal and interannual variations in CO and BC emissions  
27 from open biomass burning in Southern Africa during 1998-2005, *Global Biogeochem. Cy.*, 21,  
28 Artn Gb2011, doi:10.1029/2006gb002848, 2007.

29 Jacobson, M. Z.: Control of fossil-fuel particulate black carbon and organic matter, possibly the  
30 most effective method of slowing global warming, *J. Geophys. Res. Atmos.*, 107, Artn 4410,  
31 doi:10.1029/2001jd001376, 2002.

32 Koch, D., and Del Genio, A. D.: Black carbon semi-direct effects on cloud cover: review and  
33 synthesis, *Atmos. Chem. Phys.*, 10, 7685-7696, 10.5194/acp-10-7685-2010, 2010.

34 Kolusu, S. R., Marsham, J. H., Mulcahy, J., Johnson, B., Dunning, C., Bush, M., and Spracklen,  
35 D. V.: Impacts of Amazonia biomass burning aerosols assessed from short-range weather  
36 forecasts, *Atmos. Chem. Phys.*, 15, 12251-12266, 10.5194/acp-15-12251-2015, 2015.

37 Koren, I., Kaufman, Y. J., Remer, L. A., and Martins, J. V.: Measurement of the effect of  
38 Amazon smoke on inhibition of cloud formation, *Science*, 303, 1342-1345, doi: 10.1126  
39 /science.1089424, 2004.

40 Krishnan, R., and Ramanathan, V.: Evidence of surface cooling from absorbing aerosols,  
41 *Geophys. Res. Lett.*, 29, Artn 1340, doi:10.1029/2002gl014687, 2002.

42 Kuhlbusch, T. A. J.: Black carbon and the carbon cycle, *Science*, 280, 1903-1904,  
43 doi:10.1126/science.280.5371.1903, 1998.

- 1 Kulmala, M., Lappalainen, H.K., Petäjä, T., Kurten, T., Kerminen, V-M., Viisanen, Y., Hari,  
2 P., Bondur, V., Kasimov, N., Kotlyakov, V., Matvienko, G., Baklanov, A., Guo, H., Ding, A.,  
3 Hansson, H-C., and Zilitinkevich, S., 2015. Introduction: The Pan-Eurasian Experiment (PEEX)  
4 – multi-disciplinary, multi-scale and multi-component researchn and capacity building  
5 initiative, *Atmos. Chem. Phys.*, 15, 13085-13096, doi:10.5194/acp-15-13085-2015, 2015.
- 6 Langenfelds, R. L., Francey, R. J., Pak, B. C., Steele, L. P., Lloyd, J., Trudinger, C. M., and  
7 Allison, C. E.: Interannual growth rate variations of atmospheric CO<sub>2</sub> and its delta C-13, H-2,  
8 CH<sub>4</sub>, and CO between 1992 and 1999 linked to biomass burning, *Global Biogeochem. Cy.*, 16,  
9 Artn 1048, doi:10.1029/2001gb001466, 2002.
- 10 Lappalainen, H.K. et al., Pan-Eurasian Experiment (PEEX): Towards holistic understanding of  
11 the feedbacks and interactions in the land-atmosphere-ocean-society continuum in the Northern  
12 Eurasian region, submitted to *Atmos. Chem. Phys.*, 2016.
- 13 Lau, K. M., Kim, M. K., and Kim, K. M.: Asian summer monsoon anomalies induced by aerosol  
14 direct forcing: the role of the Tibetan Plateau, *Clim. Dynam.*, 26, 855-864, doi:10.1007/s00382-  
15 006-0114-z, 2006.
- 16 Li, J. F., Song, Y., Mao, Y., Mao, Z. C., Wu, Y. S., Li, M. M., Huang, X., He, Q. C., and Hu,  
17 M.: Chemical characteristics and source apportionment of PM<sub>2.5</sub> during the harvest season in  
18 eastern China's agricultural regions, *Atmos. Environ.*, 92, 442-448, 10.1016/j.atmosenv.  
19 2014.04.058, 2014.
- 20 Li, M., Zhang, Q., Kurokawa, J., Woo, J., He, K., Lu, Z., Ohara, T., Song, Y., Sreets, D., and  
21 Carmichael, G.: MIX: a mosaic Asian anthropogenic emission inventory for the MICS-Asia  
22 and the HTAP projects, *Atmos. Phys. Chem. Discuss.*, 34813-34869, doi:10.5194/acpd-15-  
23 34813-2015, 2015.
- 24 Lin, Y. L., Farley, R. D., and Orville, H. D.: Bulk Parameterization of the Snow Field in a Cloud  
25 Model, *J. Clim. Appl. Meteorol.*, 22, 1065-1092, doi: 10.1175/1520-0450(1983)022<1065:  
26 Bpotsf>2.0.Co;2, 1983.
- 27 Liu, Y., Fu, R., and Dickinson, R.: Smoke aerosols altering South American monsoon, *B. Am.*  
28 *Meteorol. Soc.*, 86, 1062-1063, 2005.
- 29 Lo, J. C. F., Yang, Z. L., and Pielke, R. A.: Assessment of three dynamical climate downscaling  
30 methods using the Weather Research and Forecasting (WRF) model, *J. Geophys. Res. Atmos.*,  
31 113, Artn D09112, doi:10.1029/2007jd009216, 2008.
- 32 Menon, S., Hansen, J., Nazarenko, L., and Luo, Y. F.: Climate effects of black carbon aerosols  
33 in China and India, *Science*, 297, 2250-2253, doi:10.1126/science.1075159, 2002.
- 34 Mlawer, E. J., Taubman, S. J., Brown, P. D., Iacono, M. J., and Clough, S. A.: Radiative transfer  
35 for inhomogeneous atmospheres: RRTM, a validated correlated-k model for the longwave, *J.*  
36 *Geophys. Res. Atmos.*, 102, 16663-16682, doi:10.1029/97jd00237, 1997.
- 37 NBSC: China Statistical Yearbook National Bureau of Statistics of China ed., China Statistics  
38 Press, Beijing, 2013.
- 39 Penner, J. E., Dickinson, R. E., and Oneill, C. A.: Effects of Aerosol from Biomass Burning on  
40 the Global Radiation Budget, *Science*, 256, 1432-1434, doi:10.1126/science.256. 5062.1432,  
41 1992.
- 42 Procopio, A. S., Artaxo, P., Kaufman, Y. J., Remer, L. A., Schafer, J. S., and Holben, B. N.:  
43 Multiyear analysis of amazonian biomass burning smoke radiative forcing of climate, *Geophys.*  
44 *Res. Lett.*, 31, Artn L03108, doi:10.1029/2003gl018646, 2004.



- 1 Ramanathan, V., Chung, C., Kim, D., Bettge, T., Buja, L., Kiehl, J. T., Washington, W. M., Fu,  
2 Q., Sikka, D. R., and Wild, M.: Atmospheric brown clouds: Impacts on South Asian climate  
3 and hydrological cycle, *P. Natl. Acad. Sci. USA*, 102, 5326-5333, 10.1073/pnas.0500656102,  
4 2005.
- 5 Ramanathan, V., and Carmichael, G.: Global and regional climate changes due to black carbon,  
6 *Nat Geosci*, 1, 221-227, 10.1038/ngeo156, 2008.
- 7 Reid, J. S., Hobbs, P. V., Liousse, C., Martins, J. V., Weiss, R. E., and Eck, T. F.: Comparisons  
8 of techniques for measuring shortwave absorption and black carbon content of aerosols from  
9 biomass burning in Brazil, *J. Geophys. Res. Atmos.*, 103, 32031-32040, doi:10.  
10 1029/98jd00773, 1998.
- 11 Reid, J. S., Koppmann, R., Eck, T. F., and Eleuterio, D. P.: A review of biomass burning  
12 emissions part II: intensive physical properties of biomass burning particles, *Atmos. Chem.*  
13 *Phys.*, 5, 799-825, 2005.
- 14 Robock, A.: Enhancement of Surface Cooling Due to Forest Fire Smoke, *Science*, 242, 911-  
15 913, 1988.
- 16 Robock, A.: Surface Cooling Due to Forest-Fire Smoke, *J. Geophys. Res. Atmos.*, 96, 20869-  
17 20878, doi:10.1029/91jd02043, 1991.
- 18 Rosenfeld, D.: TRMM observed first direct evidence of smoke from forest fires inhibiting  
19 rainfall, *Geophys. Res. Lett.*, 26, 3105-3108, doi:10.1029/1999gl006066, 1999.
- 20 Rosenfeld, D., Lohmann, U., Raga, G. B., O'Dowd, C. D., Kulmala, M., Fuzzi, S., Reissell, A.,  
21 and Andreae, M. O.: Flood or drought: How do aerosols affect precipitation?, *Science*, 321,  
22 1309-1313, 10.1126/science.1160606, 2008.
- 23 Roy, D. P., and Boschetti, L.: Southern Africa Validation of the MODIS, L3JRC, and  
24 GlobCarbon Burned-Area Products, *IEEE T. Geosci. Remote*, 47, 1032-1044,  
25 10.1109/Tgrs.2008.2009000, 2009.
- 26 Ryu, S. Y., Kwon, B. G., Kim, Y. J., Kim, H. H., and Chun, K. J.: Characteristics of biomass  
27 burning aerosol and its impact on regional air quality in the summer of 2003 at Gwangju, Korea,  
28 *Atmos. Res.*, 84, 362-373, 10.1016/j.atmosres.2006.09.007, 2007.
- 29 Sakaeda, N., Wood, R., and Rasch, P. J.: Direct and semidirect aerosol effects of southern  
30 African biomass burning aerosol, *J. Geophys. Res. Atmos.*, 116, Artn D12205, doi:  
31 10.1029/2010jd015540, 2011.
- 32 Saleh, R., Robinson, E. S., Tkacik, D. S., Ahern, A. T., Liu, S., Aiken, A. C., Sullivan, R. C.,  
33 Presto, A. A., Dubey, M. K., Yokelson, R. J., Donahue, N. M., and Robinson, A. L.: Brownness  
34 of organics in aerosols from biomass burning linked to their black carbon content, *Nat. Geosci.*,  
35 7, 647-650, 10.1038/NNGEO2220, 2014.
- 36 Schwarz, J. P., Gao, R. S., Spackman, J. R., Watts, L. A., Thomson, D. S., Fahey, D. W.,  
37 Ryerson, T. B., Peischl, J., Holloway, J. S., Trainer, M., Frost, G. J., Baynard, T., Lack, D. A.,  
38 de Gouw, J. A., Warneke, C., and Del Negro, L. A.: Measurement of the mixing state, mass,  
39 and optical size of individual black carbon particles in urban and biomass burning emissions,  
40 *Geophys. Res. Lett.*, 35, Artn L13810, doi:10.1029/2008gl033968, 2008.
- 41 Simpson, J., Adler, R. F., and North, G. R.: A Proposed Tropical Rainfall Measuring Mission  
42 (Trmm) Satellite, *B. Am. Meteorol. Soc.*, 69, 278-295, doi:10.1175/1520-0477(1988)069  
43 <0278:Aptrmm>2.0.Co;2, 1988.

- 1 Tao, W. K., J. P. Chen, Z. Q. Li, C. Wang, and C. D. Zhang, Impact of Aerosols on  
2 Convective Clouds and Precipitation, *Reviews of Geophysics*, 50, doi:10.1029/2011rg000369,  
3 2012.
- 4 van der Werf, G. R., Randerson, J. T., Giglio, L., Collatz, G. J., Kasibhatla, P. S., and Arellano,  
5 A. F.: Interannual variability in global biomass burning emissions from 1997 to 2004, *Atmos.*  
6 *Chem. Phys.*, 6, 3423-3441, 2006.
- 7 Wang, M., S. Ghan, M. Ovchinnikov, X. Liu, R. Easter, E. Kassianov, Y. Qian, and H. Morrison,  
8 Aerosol indirect effects in a multi-scale aerosol-climate model PNNL-MMF, *Atmos. Chem.*  
9 *Phys.*, 11(11), 5431-5455, doi:10.5194/Acp-11-5431-2011, 2011.
- 10 Wiedinmyer, C., Quayle, B., Geron, C., Belote, A., McKenzie, D., Zhang, X. Y., O'Neill, S.,  
11 and Wynne, K. K.: Estimating emissions from fires in North America for air quality modeling,  
12 *Atmos. Environ.*, 40, 3419-3432, 10.1016/j.atmosenv.2006.02.010, 2006.
- 13 Wiedinmyer, C., Akagi, S. K., Yokelson, R. J., Emmons, L. K., Al-Saadi, J. A., Orlando, J. J.,  
14 and Soja, A. J.: The Fire INventory from NCAR (FINN): a high resolution global model to  
15 estimate the emissions from open burning, *Geosci. Model Dev.*, 4, 625-641, doi:10.5194/gmd-  
16 4-625-2011, 2011.
- 17 Xie, Y. N., Ding, A. J., Nie, W., Mao, H. T., Qi, X. M., Huang, X., Xu, Z., Kerminen, V. M.,  
18 Petaja, T., Chi, X. G., Virkkula, A., Boy, M., Xue, L. K., Guo, J., Sun, J. N., Yang, X. Q.,  
19 Kulmala, M., and Fu, C. B.: Enhanced sulfate formation by nitrogen dioxide: Implications from  
20 in situ observations at the SORPES station, *J. Geophys. Res. Atmos.*, 120, 12679-12694,  
21 10.1002/2015JD023607, 2015.
- 22 Yan, X. Y., Ohara, T., and Akimoto, H.: Bottom-up estimate of biomass burning in mainland  
23 China, *Atmos. Environ.*, 40, 5262-5273, 10.1016/j.atmosenv.2006.04.040, 2006.
- 24 Yang, S. J., He, H. P., Lu, S. L., Chen, D., and Zhu, J. X.: Quantification of crop residue burning  
25 in the field and its influence on ambient air quality in Suqian, China, *Atmos. Environ.*, 42, 1961-  
26 1969, 10.1016/j.atmosenv.2007.12.007, 2008.
- 27 Zaveri, R. A., and Peters, L. K.: A new lumped structure photochemical mechanism for large-  
28 scale applications, *J. Geophys. Res. Atmos.*, 104, 30387-30415, 1999.
- 29 Zaveri, R. A., Easter, R. C., Fast, J. D., and Peters, L. K.: Model for simulating aerosol  
30 interactions and chemistry (MOSAIC), *J. Geophys. Res. Atmos.*, 113, D13204, 2008.
- 31 Zhang, Y., Fu, R., Yu, H., Qian, Y., Dickinson, R., Silva Dias, M. A. F., da Silva Dias, P. L.,  
32 and Fernandes, K.: Impact of biomass burning aerosol on the monsoon circulation transition  
33 over Amazonia, *Geophys. Res. Lett.*, 36, 2009.
- 34 Zhao, C., Leung, L. R., Easter, R., Hand, J., and Avise, J.: Characterization of speciated aerosol  
35 direct radiative forcing over California, *J. Geophys. Res. Atmos.*, 118, 2372-2388,  
36 10.1029/2012JD018364, 2013.
- 37 Zhao, T., and Yatagai, A.: Evaluation of TRMM 3B42 product using a new gauge - based  
38 analysis of daily precipitation over China, *Int. J. Climatol.* 34, 2749-2762, 2014.