



| 1  | Investigation of short-term effective radiative forcing of fire aerosols over North America   |
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| 2  | using nudged hindcast ensembles   |
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#### 19 Abstract

Aerosols from fire emissions can potentially have large impact on clouds and radiation. However, 20 fire aerosol sources are often intermittent and their effect on weather and climate is difficult to 21 quantify. Here we investigated the short-term effective radiative forcing of fire aerosols using the 22 global aerosol-climate model Community Atmosphere Model Version 5 (CAM5). Different from 23 previous studies, we used nudged hindcast ensembles to quantify the forcing uncertainty due to 24 25 the chaotic response to small perturbations in the atmosphere state. Daily mean emissions from three fire inventories were used to consider the uncertainty in emission strength and injection 26 heights. The simulated aerosol optical depth (AOD) and mass concentrations were evaluated 27 28 against in-situ measurements and re-analysis data. Overall, the results show the model has reasonably good predicting skills. Short (10-day) nudged ensemble simulations were then 29 performed with and without fire emissions to estimate the effective radiative forcing. Results 30 show fire aerosols have large effects on both liquid and ice clouds over the two selected regions 31 in April 2009. For the 10-day average, we found a large ensemble spread of regional mean 32 shortwave cloud radiative effect over Southern Mexico (15.6%) and the Central U.S. (64.3%), 33 34 despite that the regional mean AOD time series are almost indistinguishable during the 10-day period. Moreover, the ensemble spread is much larger when using daily averages instead of 10-35 day averages. For the case investigated here, a minimum of 9 ensemble members is necessary to 36 37 get a reasonable estimate of the ensemble mean and spread of the forcing on individual days. This demonstrates the importance of using a large ensemble of simulations to estimate the short-38 term effective aerosol radiative forcing. 39

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#### 42 1. Introduction

43 Natural and human-induced fires play an important role in the Earth system. Aerosol and gas 44 emissions from biomass burning can change the atmospheric composition and potentially affect 45 the weather and climate. Over 30% of the global total emission of black carbon (BC) comes from open burning of forests, grasslands and agricultural residues (Bond et al. 2013). For organic 46 47 aerosols, substantial increases of concentrations dominated by organic carbon enhancements are observed in regions with biomass burning events (Zeng et al. 2011; Lin et al. 2013; Brito et al. 48 49 2014; Reddington et al. 2014). As a result, biomass burning emissions have a large impact on the global and regional mean aerosol optical depth (Jacobson, 2014). 50

Through interactions with radiation and clouds, fire aerosols can significantly affect the long-51 52 term Earth's energy budget. Previous studies have investigated the global and regional radiative forcing of fire aerosols using long climatological simulations or satellite retrievals. For example, 53 Ward et al. (2012) investigated the radiative forcing of global fires in pre-industrial, present day, 54 and future periods. For the present-day condition, they estimated a direct aerosol effect (or 55 radiative forcing through aerosol-radiation interactions as defined in IPCC AR5, RFari; see 56 section 2.4) of +0.1 W m<sup>-2</sup> and an indirect effect (radiative forcing through aerosol-cloud 57 interactions, RFaci) of -1.0W m<sup>-2</sup>. Using a newer model, Jiang et al. (2016) found similar RFari 58 but slightly smaller RFaci (-0.70W m<sup>-2</sup>). Sena et al. (2013) assessed the direct impact of 59 biomass burning aerosols over the Amazon basin using satellite data. Over the 10-year studied 60 period, the estimated radiative forcing is about  $-5.6W \text{ m}^{-2}$ . 61

On short timescales, fire aerosols have even larger radiative impacts. Observed maximum daily direct aerosol radiative effects can reach -20W m<sup>-2</sup> at TOA locally in Amazonia during biomass burning seasons (Sena et al., 2013). Very large direct effects of fire aerosols were





observed during extreme fire events over Central Russia (Tarasova et al. 2004; Chubarova et al. 65 2008; Chubarova et al. 2012). Instantaneous radiative effects of emitted aerosols reached -167 66  $W m^{-2}$  and monthly mean radiative effects reached about -65  $W m^{-2}$  in the 2010 Russia 67 wildfires (Chubarova et al. 2012). Using satellite data and a radiative transfer model, Kaufman et 68 al. (2005) found a radiative effect of -9.5W m<sup>-2</sup> due to smoke aerosol-induced cloud changes over 69 Southeast Atlantic for the 3 months studied. Smoke-derived cloud albedo effect on local 70 shortwave radiative forcing is estimated to be between -2 and -4 W m<sup>-2</sup> in a day case study of 71 aircraft-measured indirect cloud effects (Zamora et al., 2016). Kolusu et al. (2015) investigated 72 73 direct radiative effect of biomass burning aerosols over tropical Southern America. By quantifying results from the first and second day of 2-day single-member forecasts in September 74 2012, they found the modeled biomass burning aerosols reduced all-sky net radiation by 8 75 W m<sup>-2</sup> at TOA and 15 W m<sup>-2</sup> at surface. 76

Previous modeling studies on the short-term fire aerosol effects mainly focused on aerosol 77 direct effects (e.g., Keil and Haywood, 2003; Chen et al., 2014; Kolusu et al., 2015), and only a 78 79 couple of studies investigated the indirect effects of fire aerosols (Lu and Sokolik, 2013). In addition, to estimate the aerosol indirect effect, long simulations (multi-years, >5 years 80 preferred) are often needed to remove the noise, because aerosol life cycle and cloud properties 81 82 are affected by strong natural variability on different timescales (Bony et al. 2006; Kooperman et al. 2012). To solve the problem, alternative methods have been proposed to help extract signals 83 with shorter simulations. For example, nudging (also called Newton relaxation method) can help 84 reduce uncertainties associated with natural variability by constraining certain meteorological 85 fields towards prescribed conditions. A robust estimate of global anthropogenic aerosol indirect 86 effects can be obtained on substantially shorter timescales (1-2 years) by implementing nudging 87





to constrain simulations with pre-industrial and present-day aerosol emissions toward identical 88 89 circulation and meteorology (Kooperman et al. 2012). When nudged towards re-analysis data, Zhang et al. (2014) found constraining only the horizontal winds is a preferred strategy to 90 estimate the aerosol indirect effect since it provides well-constrained meteorology without 91 strongly perturbing the model's mean climate state. Another example is the use of representative 92 ensembles of short simulations to replace a typical long integration. Wan et al. (2014) explored 93 94 the feasibility of this method and showed that 3-day ensembles of 20 to 50 members are able to reveal the main signals revealed by traditional 5-year simulations. 95

In this study, we performed month-long and 10-day nudged CAM5 simulations to investigate 96 97 the effects of fire aerosols on radiation and cloud processes on short time scales (less than two weeks). Horizontal winds were nudged towards reanalysis to constrain the large-scale circulation 98 and to allow for more accurate model evaluations against observations. We also used daily mean 99 100 emissions from three fire inventories to consider the uncertainty in emission strength and injection heights. Even for short simulations, small perturbations of meteorological states might 101 have large impact on the local aerosol and cloud properties, thus bring uncertainty to the aerosol 102 103 forcing estimate. Therefore, in our simulations, we also employed very weak temperature nudging in combination with ensembles to quantify the uncertainty. 104

The rest of the paper is organized as follows. Sect. 2 describes the model and data used in this study. It also introduces how the ensembles are generated in the short nudged simulations and explains how the fire aerosol forcing is estimated. Results and discussions are presented in Sect. 3 and conclusions are summarized in Sect. 4.

#### 109 2. Model, Method and Data





### 110 **2.1 Model description**

In this study, we used the Community Atmosphere Model (CAM) version 5.3 with the finite 111 112 volume dynamical core at  $1.9^{\circ}$  (latitude)  $\times 2.5^{\circ}$  (longitude) horizontal resolution with 30 vertical 113 layers. The aerosol life cycle is represented by using the modal aerosol module MAM3 (Liu et al., 2012). CAM5 links the simulated aerosol fields with cloud and radiation through interactions 114 115 of the aerosol module with the cloud microphysics and radiative transfer parameterizations. The two-moment bulk cloud microphysics scheme from Morrison and Gettelman (2008) is used to 116 117 track mass mixing ratios and number concentrations of cloud droplets and ice crystals in stratiform clouds. Representation of shallow convection is based on the work of Park and 118 Bretherton (2009). The deep convection parameterization was developed by Zhang and 119 120 McFarlane (1995) and later revised by Richter and Rasch (2008) and Neale et al. (2008). Longwave and shortwave radiative transfer are calculated with the Rapid Radiative Transfer 121 Model for GCMs (RRTMG, Malwer et al. 1997; Iacono et al. 2008). 122

### 123 **2.2 Fire Emission Inventories**

124 Three fire emission inventories were used in this study. Two of them are widely used bottomup inventories— Global Fire Emissions Database version 3.1 (GFED v3.1, van der Werf et al., 125 2010; https://daac.ornl.gov/cgi-bin/dsviewer.pl?ds\_id=1191) and GFED v4.1s (Giglio et al. 126 127 2013; Randerson et al. 2012; https://daac.ornl.gov/VEGETATION/guides/fire emissions v4.html). Another one is a top-down 128 emission inventory-Quick Fire Emissions Dataset version 2.4 (QFED v2.4). GFED v3.1 and 129 GFED v4.1s provide global monthly emissions at 0.25×0.25 degree spatial resolution from 1997 130 131 through the present. Daily emission data are obtained by disaggregating monthly emissions





based on daily temporal variability in fire emissions derived from MODIS measurements of
active fires (Mu et al. 2011). The more recent version GFED v4.1s improves by including small
fires based on active fire detections outside the burned area maps (Randerson et al., 2012).
QFED v2.4 estimates global fire emissions using the Moderate Resolution Imaging
Spectroradiometer (MODIS) measurements of fire radiative power and generates daily products
at 0.1×0.1 degree resolution.

To drive CAM5 simulations, fire emission data were regridded to the model resolution and distributed vertically. For the GFED v3.1 and QFED v2.4 emission data we adopted the same injection heights (from surface to 6 km) as used in the standard CAM5 model. While for GFEDv4.1s, in this study the injection heights were estimated using a fire plume model and scaled to the 6-hourly interval.

The fire emission inventories were first analyzed to select appropriate time periods and 143 144 regions for our study before being used to drive model simulations. Fig.1 shows the multi-year mean biomass burning emissions from GFED v4.1 over North America. The emission manifests 145 significant seasonality with large dry matter consumption during March to April and June to 146 147 September. The summer and autumn burning covers Pacific Northwest and part of Canada and is mainly associated with forest fires, while the spring burning occurs in more densely populated 148 regions like Mexico and central and eastern United States with a large contribution of 149 150 agricultural fires in croplands (Korontzi et al., 2006; Magi et al., 2012). Similar features are also captured in GFED v3.1 and OFED v2.4 with differences in the magnitude. We chose to analyze 151 the simulated fire aerosol effect in April, the peak month of spring burning, when there are 152 153 extreme fire activities over Mexico (10 N to 25N, 100W to 80W) and occasionally large fires in the Central U.S. (35 N to 45N, 100W to 85W). For the U.S., extended fire period is rare, making 154





it necessary to perform short-term evaluation. Fire aerosols formed from these two regions are 155 156 often transported to the Eastern and Southeastern U.S., where they mix with aerosols from anthropogenic sources and potentially have significant impact on clouds and radiation over these 157 areas. Time series of regional mean fire emissions in April during 2003-2014 shows that 158 relatively large fires occur in both regions in 2009 (Fig.S1). Values of fire emissions in 2009 are 159 larger than the multi-year April mean by a factor of 1.9 in the Central U.S. and 1.5 in Southern 160 Mexico. Thus, in the following model simulations, we focused on analyzing the aerosol 161 properties and radiative effects over the two selected regions (denoted by the red boxes in Fig.1) 162 in April 2009. 163

Fire emitted BC from different emission inventories in April 2009 is shown is Fig.2. Although GFED v4.1s includes the contributions of small fires (Randerson et al., 2012), the emitted BC in GFED v4.1 shows no substantial increase compared to GFED v3.1during the selected period. Only an increase by 1.75 is seen over Southern Mexico. In the Central U.S., the BC emission is even slightly weaker in GFED v4.1. QFED v2.4 shows a much larger BC emission than the GFED inventories. Values of emitted BC in QFED v2.4 are larger than those in GFED v4.1s by a factor of 9.7 in the Central U.S. and a factor of 2.7 in Southern Mexico.

#### 171 2.3 Simulations

Two groups of simulations were conducted (Table1) using the same greenhouse gas concentrations, sea surface conditions and anthropogenic emissions of aerosols and precursors. Each group includes four simulations, performed either without fire emission or with daily fire emissions from one of the three fire emission inventories introduced in section 2.2. The emitted species include BC, OC, and SO<sub>2</sub>. Horizontal winds were nudged to 6-hourly ERA-Interim reanalysis (Dee et al., 2011) as described in Zhang et al. (2014) in both groups.





Simulations in Group A are month-long single-member nudged simulations. These simulations were performed to provide longer time series for model evaluation and generate initial condition files for simulations in Group B. They started from January 1, 2009 and were integrated for four months with 3-month spin-up. Initial condition files were generated on April 1 at 00 UTC for simulations in group B.

Simulations in group B are 10-day ensemble simulations. Unlike the traditional way of 183 184 perturbing initial conditions, in this study we constructed the ensembles by implementing a very weak temperature nudging and perturbing the nudging time scale. This is because under the 185 influence of horizontal-wind nudging, ensemble differences generated by perturbing initial 186 187 conditions would fade away during the integration. In contrast, our method can consider the influence of small temperature perturbations during the entire simulation period, as nudging is 188 applied at every time step. On the other hand, the large-scale circulation patterns simulated in the 189 190 different ensemble members are very similar (not shown), so the noises caused by the chaotic system can be constrained and the effective fire aerosol forcing signal can be easily identified. 191

Each ensemble in group B includes 10 members. The only difference between the members is the relaxation time scale of temperature, which varies from 10 to 11 days at an interval of 0.1 day. All simulations started on April 1, 2009 and were integrated for 10 days. For each simulation (e.g. E\_QF), the initial condition was generated by combining the meteorological fields from initial condition outputs in the S\_NF simulation with aerosol and precursor concentrations from initial condition outputs in the single-member simulation forced by the corresponding fire emission (S\_QF).

## 199 2.4 Calculation of fire aerosol RF





The IPCC AR5 report provides a more useful characterization of aerosol forcing by allowing for 200 201 rapid tropospheric adjustments (Boucher et al., 2013) compared to the original definition of 202 aerosol forcing. It quantifies aerosol radiative effects in terms of Effective Radiative Forcing 203 from aerosol-radiation interactions (ERFari) and Effective Radiative Forcing from aerosol-cloud interactions (ERFaci). ERFari refers to the combined effect of instantaneous radiative forcing 204 from direct scattering and absorption of sunlight (aerosol direct effect) and related subsequent 205 206 rapid adjustments of atmospheric state variables and cloudiness (aerosol semi-direct effect). 207 ERFaci refers to the indirect forcing resulting from aerosol induced changes in cloud albedo (cloud albedo effect) and subsequent changes in cloud lifetime as rapid adjustments (second 208 209 aerosol indirect effect) via microphysical interactions.

To allow for a straightforward comparison with previous studies in the literature, we followed the IPCC concept of including rapid adjustments (effective aerosol radiative forcing), but continued to decompose the aerosol effect in the conventional terms as aerosol direct radiative effect (DRE), aerosol cloud radiative effect (CRE) and surface albedo effect. Note that as nudging timescale determines the degree to which model physics are constrained (Kooperman et al., 2012), the use of a 6-hour relaxation time scale for horizontal wind nudging means only very fast adjustments are considered in the simulations.

Similar to Jiang et al. (2016), our calculations of fire aerosol DRE, CRE and surface albedo effect are based on the work of Ghan et al. (2012) and Ghan (2013). They were calculated as the radiative flux differences between simulations with and without fire emissions (denoted by  $\Delta$ ). In each simulation, aerosol (direct) forcing was defined as the difference between all-sky and cleansky TOA radiative fluxes (F – F<sub>clean</sub>). Aerosol induced cloud forcing change was defined as the difference between all-sky and clear sky TOA radiative fluxes under clean-sky





223 conditions ( $F_{clean} - F_{clean,clear}$ ). The rest were related to surface albedo forcing ( $F_{clean,clear}$ ). 224 Thus fire aerosol DRE, CRE, and surface albedo effect were expressed as  $\Delta(F - F_{clean})$ , 225  $\Delta(F_{clean} - F_{clean,clear})$ , and  $\Delta F_{clean,clear}$ , respectively. More details about the method can be 226 found in section 2 of Ghan (2013). CRE includes contributions of both aerosol indirect effect and 227 aerosol semi-direct effect but was analyzed as a single term (i.e., the sum).

### 228 2.5 Observational Data

In this study, we used two sets of AOD reanalysis and the AERONET data (Holben et al. 229 230 1998) to evaluate the modeled AOD. The two AOD reanalysis datasets are the Naval Research Laboratory (NRL) reanalysis (Rubin et al. 2015) and the Monitoring Atmospheric Composition 231 and Climate (MACC) reanalysis (Eskes et al. 2015). Both are generated by assimilating AOD 232 233 retrievals from MODIS (Zhang et al., 2008; Benedetti et al., 2009) with forecast fields. The NRL 234 reanalysis provides 6-hourly AOD at 1°horizontal resolution. The MACC dataset provides 3hourly AOD at 1.125° horizontal resolution. Daily averages in April, 2009 were used for model 235 236 evaluation in this study. AERONET retrievals of AOD from April 1 to April 30 in 2009 were used for model evaluation. Two sites are available in the selected regions: Cart Site (36°N, 237 97°W) and Mexico City (19°N, 99°W). LEV 2.0 cloud-screened all points AOD at 500 nm and 238 675 nm was used to generate hourly AOD at 550nm. 239

In addition, the simulated BC and primary organic matter (POM) concentrations were compared with observations from the Interagency Monitoring of Protected Visual Environments (IMPROVE) (Malm et al. 2004). IMPROVE aerosol data are only available over the Central U.S. A total of fifteen sites were selected and marked in Fig 2, which include the sites west of 94°W near the source region (asterisks) and sites east of 94°W in the downwind region (dots).





Observed organic carbon concentrations were multiplied by 1.4 for comparison with simulated 245 246 POM. Detailed descriptions about the data and sites available are at http://vista.cira.colostate.edu/improve/. The IMPROVE network collect 24-hour aerosol data on 247 every third day. Daily averages during April, 2009 are compared on IMPROVE observation days 248 249 only.

#### 250 **3. Results**

In this part, the model performance is first evaluated based on the simulations in group A. Next, we present the simulated short-term effective fire aerosol forcing on 10-day and daily timescales based on the results from group B simulations. We will demonstrate the importance of using ensemble simulations in estimating the short-term aerosol effective forcing and give a quantitative estimate of how many ensemble members are needed for the case selected in this study.

### 257 **3.1 Model Evaluation**

258 Model simulated AOD are evaluated against the NRL and MACC reanalysis data (Fig. 3). The simulated temporal variation of regional mean AOD over the central U.S. is consistent with 259 that in the reanalysis, but the magnitudes of simulated AOD are lower (Fig. 3). A better 260 261 agreement is found between the model and the NRL data, despite the horizontal winds in the simulation are nudged towards a reanalysis that is very similar to the data used to derive MACC. 262 Temporal correlation coefficients (TCC) between the modeled AOD and the NRL reanalysis are 263 0.87 and 0.82 for S QF and S GF4 simulations, respectively, but are lower (0.67 and 0.78) 264 between the modeled AOD and the MACC reanalysis. The corresponding root mean square 265





errors rise from 0.13 (S\_QF) and 0.1 (S\_GF4) to 0.23 and 0.21. Generally, AOD is underestimated by a factor of 2-4 in all simulations compared to the reanalysis, especially in simulations with GFED emissions. Previous studies have suggested the need to scale up GFED emissions by a factor of 1-3 to match the observed AOD (Tosca et al., 2013). This is consistent with the large negative bias in the simulation S\_GF3 and S\_GF4. Simulated AOD in these two simulations are almost indistinguishable due to the small difference in the total fire emission in the region.

273 Over Mexico, different simulations produce similar temporal variations in AOD, but the 274 magnitude is smaller in the GFED simulations. Large discrepancies are found between model 275 results and reanalysis data during Apr. 17-20. An increase of AOD is captured by both reanalysis datasets, while model results display a decrease of AOD compared to earlier days in the 276 simulation period. Note that the two sets of reanalysis data also have some differences 277 occasionally. For example, during Apr. 10-12, NRL data displays an increase of AOD, while 278 MACC data show the opposite. These discrepancies may partly result from the large internal 279 variability in this tropical region, where the simulated atmosphere state and its influence on 280 281 aerosol transport are more likely to disagree between the model and the reanalysis. Generally speaking, the model forced with different fire emissions is capable of capturing daily variation of 282 AOD in both regions, especially during Apr. 1-10. This period was selected for further 283 284 investigation of the short-term fire aerosol effect.

Model simulated AOD are also evaluated against AERONET retrievals (Fig. 4). At Cart Site ( $36^{\circ}N$ , 97°W), with the QFED emission (S\_QF) the model performs well in simulating both the temporal variation (TCC=0.62) and magnitude of AOD. Simulations with GFED emissions also reproduce the temporal evolution well (TCC = 0.58 for S\_GF3 and 0.55 for S\_GF4), but with





significantly low bias (mean bias by a factor of 2). The simulated difference in AOD magnitude 289 290 is similar to that found by Zhang et al. (2014) over the northern sub-Saharan African. Using the 291 QFEDv2.4 fire emission, the simulated regional mean AOD is a factor of 1.5 higher than that 292 using the GFEDv3.1 emission in their study. Relatively good performance of S\_QF is also seen over Mexico. The simulated time evolution agrees well with AERONET retrievals except for 293 small discrepancies (e.g. during Apr.17 -19). A better agreement with the AERONET retrievals 294 is found for the NRL data than MACC reanalysis at both sites. Consistent with the evaluation 295 296 using reanalysis, the simulated temporal evolution of AOD during Apr. 1-10 agrees well with 297 both reanalysis data and AERONET retrievals in selected regions. This gives us further 298 confidence in choosing this period for further investigation.

299 The model is further evaluated against the IMPROVE data for BC and POM mass concentrations (Fig. 5). In the downwind region, the simulated mass concentrations in simulation 300 301 S QF lie within a factor of 2 of the observed values at most sites. However, the magnitude is generally underestimated in simulations with the GFED emissions (S\_GF3 and S\_GF4), 302 especially in S GF3. BC and POM concentrations in the downwind regions are affected by 303 304 transport of aerosols from Southern Mexico (Fig. S3). A larger amount of fire emission in Southern Mexico would result in a higher BC (POM) concentration in the downwind region. 305 This explains the slightly higher concentrations in the simulation S GF4 than S GF3, as BC and 306 307 POM emissions over Southern Mexico are higher in GFED v4.1 due to the inclusion of small 308 fires (Randerson et al., 2012). The good agreement between S OF and observations suggests that the QFED data have a reasonable total emission rate. However, in the source region, the 309 310 simulation S\_QF displays large positive bias with a large majority of the values fall out of the afactor-of-2 band. Given the reasonable total emission rate in QFED and a good agreement of 311





AOD with AERONET retrievals at Cart Site, this might result from the discrepancies in the 312 313 vertical distribution of the fire emissions. Fire-emitted BC and POM in simulations S OF and S GF3 reach maximum values in the lowest level and decrease sharply to the next level, while 314 low-level fire emissions in S\_GF4 distribute in a more uniform way (Fig. S4). As the sampling 315 was done on the lowest model level at most sites to compare with the IMPROVE data, this 316 explains the strong overestimation in S\_QF. Although the same impact from vertical distribution 317 of fire emission also appears in simulation S\_GF3, it is partly offset by its negative bias in the 318 total emission rate. 319

#### 320 3.2 10-day Mean Results

Given the good model performance during April 1-10, we proceed to analyze the short-term effects of fire aerosols during this period with nudged ensemble simulations. We define "fire AOD" as the AOD difference between the simulations with and without fire emissions.

# 324 3.2.1 Fire Aerosol Distribution

Fig. 6 shows the spatial distributions of 10-day average ensemble mean fire AOD. For 325 reference, the total AOD in the simulation without fire emissions is shown in Fig. S2. During the 326 period, regional mean AOD increases by 6.4% (E\_GF3), 6.4% (E\_GF4) and 70.2% (E\_QF) in 327 the central U.S. and 10.4% (E\_GF3), 13.3% (E\_GF4), and 49.6% (E\_QF) in Southern Mexico 328 when fire emissions are included. In E QF, high fire AOD covers almost the entire selected 329 330 region and extends further north. Maximum values of fire AOD stay above 0.2 around the Yucatan Peninsula. Over the Central U.S, significant fire AOD ranging between 0.04 and 0.1 331 appears in the southwest part of the selected region. Apart from the significant AOD difference 332 333 in selected regions, large fire AOD also appears near the eastern coast as a result of local fire





emission and the eastward transport of fire aerosols from both regions. Overall, the modeled fire

AOD is much smaller in simulations with GFED emissions.

### 336 3.2.2 Fire Aerosol Radiative Effect

337 As described in Sect. 2.4, fire aerosol radiative effect can be decomposed into three items 338 including fire aerosol DRE, fire aerosol CRE and fire aerosol surface albedo effect. Fig.7 shows 339 the spatial distributions of shortwave direct effect (SDRE) and shortwave cloud radiative effect (SCRE). They are major contributors to the total fire aerosol forcing in the selected regions. For 340 reference, total aerosol forcing and total shortwave cloud forcing in the simulation without fire 341 emissions are shown in Fig. S2. The spatial distribution of SDRE and SCRE are similar for the 342 343 three cases, but with different magnitudes and statistical significant regions for simulations with QFED and GFED fire emissions. In the Central U.S., fire aerosol SDRE is negligible in GFED 344 forced simulations due to small fire AOD. Although the fire AOD is larger in simulation E QF, 345 the compensation between warming effect of fire BC and cooling effect of fire POM still results 346 a weak forcing of about -0.1W m<sup>-2</sup>. Over southern Mexico, all simulations produce significant 347 cooling by fire aerosol SCRE with maximum values three times as large as those of 348 corresponding SDRE. For both SDRE and SCRE, the largest fire aerosol effects appear in the 349 350 E QF simulation while the E GF3 yields the weakest forcing, which is consistent with the modeled fire AOD in these simulations. 351

In the following analysis, we will focus on the results from the E\_QF simulation. Both SDRE and SCRE spread outside the two selected regions and extend eastward reaching coast regions. A stronger fire aerosol effect is seen in the Southern Mexico region. Strong SDRE appears over the Yucatan Peninsula where fire AOD peaks (Fig. 6). Regional mean 10-day average of SDRE and SCRE reach -1.66W m<sup>-2</sup> and -3.02W m<sup>-2</sup> respectively. In the central U.S, despite moderate fire





aerosol SDRE, SCRE near fire source region is weaker than -4 W  $m^{-2}$ , which is comparable to

that in the extended regions.

359 Given the largely insignificant change in cloud fraction (Fig. 8), fire aerosol SCRE in both regions are mainly induced by changes in liquid water path (LWP) and droplet number 360 concentrations (CDNC). Changes in ice water path (IWP) and ice crystal number concentration 361 (ICNC) can also significantly affect SCRE, but with an opposite sign and mostly in the central 362 363 U.S. Fire aerosol SCRE in the central U.S is associated with significant increases in both column-integrated droplet number concentration (smaller droplet effective radius) and LWP, 364 indicating important contributions of both the aerosol first and second indirect effects. Increased 365 366 CDNC enhances cloud albedo by decreasing droplet sizes (Twomey, 1977) and allows more liquid water to accumulate by decreasing precipitation efficiency (Albrecht, 1989; Ghan et al., 367 2012). Note that although LWP and CDNC over southern Mexico change in a smaller magnitude 368 369 than those in central U.S., fire aerosol SCRE is stronger over Southern Mexico. This is mainly due to the reductions in IWP and ICNC over the Central U.S. These changes, which possibly 370 caused by fire aerosol-induced changes in the circulation (Ten Hoeve et al, 2012), lead to a 371 372 positive SCRE that partly offsets the negative SCRE caused by changes in warm clouds. In the northeast of the extended coastal regions, a more significant change of LWP comparable to that 373 in the central U.S appears, while a more significant change of CDNC comparable to that in 374 375 Southern Mexico occurs in the southwest. The combined effect leads to the total fire aerosol 376 effect in the extended regions.

The ensemble method provides another effective way to distinguish fire aerosol radiative effect by comparing the radiative forcing distribution of ensemble members between simulations with and without fire emission. A significant difference in the distribution of total aerosol (cloud)





forcing indicates a significant fire aerosol direct (cloud) effect. As shown in Fig. 9, a shift towards stronger magnitude occurs to the total aerosol forcing when fire aerosols are considered. Simulation E\_QF has a larger percentage of grid cells with SDRE below -4.2W m<sup>-2</sup>, while more grid cells exceed -4.2W m<sup>-2</sup> in E\_NF, which indicates a significant negative fire aerosol direct effect. Same shift also appears to the total cloud forcing with more grid cells having cloud forcing below -30W m<sup>-2</sup> in the simulation E\_QF. Regional mean total aerosol and cloud forcing in southern Mexico become more negative (-0.86 and -3.0 W m<sup>-2</sup>) with fire aerosols.

Fig. 10 illustrates ensemble behavior of 10-day average regional mean total aerosol and cloud 387 forcing from all simulations as well as resulted fire aerosol SDRE and SCRE. The GFED forced 388 389 simulations not only resemble in ensemble mean, but also have small difference in ensemble member distribution. Although members in the E\_QF simulation capture stronger aerosol 390 forcing, thus stronger fire aerosol SDRE than those in E\_GF3 and E\_GF4, the ensemble spread 391 392 (as indicated by the maximum and minimum values) in the three simulations is similar. Moreover, the E\_QF simulation yields a smaller spread of SCRF compared with the GFED 393 forced simulations despite a stronger ensemble mean SCRF. In each fire simulation, ensemble 394 395 mean fire aerosol SCRE has a much larger magnitude than SDRE. So is the corresponding ensemble spread. Taking results from E\_QF simulation as an example, ensemble spread of 396 SCRE reaches 0.47 W m<sup>-2</sup>, accounting for 15.6% of the corresponding ensemble mean, while 397 ensemble spread of SDRE is 0.03W m<sup>-2</sup> accounting for 3.5% of the corresponding ensemble 398 399 mean.

### 400 3.3 Daily RF

401 The fire aerosol effect is also investigated for individual days. The spatial distributions of
402 SDRE and SCRE on April 7 are shown in Fig 11, when relatively high fire emissions appear in





both regions. Negative fire aerosol SDRE appears in the central U.S. biomass-burning region 403 404 indicating the dominant role of POM scattering. Fire aerosol SDRE over Southern Mexico shows a contrast of warming effect in land region and cooling effect in adjacent ocean despite similar 405 aerosol loading in the two regions. However, they do have nearly equal clear-sky BC absorption 406 and POM scattering (Fig. 12). Difference in the low-level cloud distributions between two 407 regions leads to different signs of the simulated all-sky SDRE. Over land, when clouds appear 408 409 under elevated aerosol layers, more solar radiation is reflected back to space and this leads to amplified BC absorption and more positive direct aerosol forcing (Keil and Haywood, 2003; 410 Zhang et al., 2016; Jiang et al., 2016). In contrast, neither absorption nor scattering changes 411 412 significantly from clear-sky to all-sky condition over adjacent areas over the ocean, since the cloud fraction is small. Same enhanced absorption of above-cloud aerosols is also found over the 413 west Atlantic Ocean. Fire aerosols produce remarkable negative SCRE up to -16W m<sup>-2</sup> over 414 Southern Mexico land in response to the increase in CDNC and LWP. 415

### 416 **3.4 Discussion about the Simulation Strategy**

Fig. 13 shows the daily variation of the regional mean total (direct) aerosol forcing and cloud 417 forcing. Both the ensemble mean and spread are investigated here. The total aerosol forcing 418 419 exhibits considerable diversity across ensemble members within each simulation, even though the simulated AOD is nearly indistinguishable (Fig. 3). Taking results from simulation E OF as 420 an example, maximum values of difference between members exceed 0.4 W m<sup>-2</sup> for aerosol 421 forcing and 5 W m<sup>-2</sup> for cloud forcing, which are approximate 10% of the corresponding 422 423 ensemble mean values. The large spread of total aerosol forcing and cloud forcing will lead to 424 uncertainties in the estimation of fire aerosol effect. This points out the importance of conducting ensemble simulations in order to get a more comprehensive estimate of the daily fire aerosol 425





effect. The minimum ensemble size required for this case is investigated in terms of the ensemble mean and spread estimate. Simulated ensemble mean fire aerosol SDRE remains nearly unchanged regardless of the ensemble size (Fig. 14a). However, discrepancies in the ensemble mean fire aerosol SCRE (Fig. 14b) are substantial when the number of ensemble members is smaller than 8. The same is true for the ensemble spread of fire aerosol SCRE (Fig. S5). Overall, the time evolution and magnitude of ensemble mean and spread tend to converge when the number of ensemble members reaches about 9 for different days we investigated here.

Fire aerosol sources are often intermittent and height-dependent and there is a need to 433 estimate the short-term effective aerosol forcing. Although nudging helps to constrain large-scale 434 435 features, the simulated cloud properties (e.g. cloud fraction and LWP) and their response to aerosol changes can still be sensitive to small perturbations in the atmospheric state. Therefore, 436 for investigating the short-term aerosol effect, a single simulation might not be sufficient to tell 437 whether the aerosol effect is significant. The use of ensembles provides an effective way to 438 estimate the uncertainty. Previous investigations of short-term fire aerosol effect are mainly 439 based on single-member simulations (Wu et al., 2011; Sena et al., 2013; Kolusu et al., 2015). 440 441 While this might be less a problem for SDRE, one should be more careful when investigating the aerosol indirect effect and conduct ensemble simulations to see whether the estimated fire 442 aerosol effects are robust. 443

#### 444 **4.** Summary

In this study, we investigated the short-term effect of fire aerosols on cloud and radiation using CAM5 simulations. Month-long single-member simulations and 10-day ensemble simulations were conducted in April 2009. In order to help extract signals on short time scales,





448 we used nudging to constrain horizontal winds in all simulations. Our investigation focused on 449 Southern Mexico where there were constant intensive fire activities and the Central U.S. with 450 occasionally large fires. Apart from the local effect, fire emissions from the two regions are 451 shown to affect downwind coastal regions through transport.

Modeled AOD and mass concentrations (BC and POM) were evaluated against observations. 452 In general, all simulations with fire emissions reproduce the observed temporal variation of daily 453 mean AOD well, although the simulated magnitude is smaller. The model performance is better 454 when QFEDv2.4 is used, which has larger fire emissions. Modeled regional mean AOD values in 455 simulations using two versions of GFED fire emission data are barely distinguishable, despite the 456 457 inclusion of small fires and changed injection heights in GFEDv4.1 used in this study. Both of them simulate about a factor of 1.5 smaller AOD than that in the simulation using the QFED fire 458 emissions. At sites in the downwind region, the modeled BC and POM mass concentrations in 459 460 the simulation with QFEDv2.4 emission (S QF) agree well with the IMPROVE data. In contrast, simulations with the other two fire emission datasets (S\_GF3 and S\_GF4) have a low bias. The 461 simulated AOD in the source region in S OF also agrees well with the AERONET data (Cart 462 463 Site). If there is no large compensating error in the model, QFEDv2.4 seems more reasonable in terms of the total (vertically-integrated) emission rate. On the other hand, S\_QF strongly 464 overestimates BC and POM concentrations in the source region. Considering that the source-465 466 region AOD and the downwind surface mass concentrations are well simulated, the overestimation suggests that the actual emission peak might appear at higher levels compared to 467 the height-dependent injection rates applied in the S\_QF simulation. 468

Based on the evaluation, we chose the first 10 days as the simulation period and focused on
the simulation with QFEDv2.4 fire emission in our ensemble nudged simulations. In our method,





the nudged ensembles are generated by adding a very weak temperature nudging along with 471 horizontal-wind nudging and perturbing the nudging time scale of temperature gently. In this 472 way, small temperature perturbations are added to the simulation at each time step, while the 473 large-scale circulation features are very similar between individual members. We first 474 investigated the 10-day mean effective fire aerosol forcing. Decomposition of total aerosol 475 radiative forcing shows that fire aerosol effects in the two selected regions are dominated by the 476 477 shortwave cloud radiative effect SCRE. All fire simulations show similar spatial distribution of SDRE and SCRE, but with different magnitudes and statistically significant regions. The 478 479 similarity in the spatial distribution is expected since the three emission datasets differ mainly in 480 the emission magnitude and not much in spatial distribution in the focused regions of this study. Fire aerosol effects in simulations with GFED emissions (E GF3 and E GF4) are weaker than 481 that with QFEDv2.4 emission (E\_QF) by a factor of 1.5 for SCRE and a factor of more than 4 482 for SDRE. Generally speaking, the difference in simulated AOD and fire aerosol indirect 483 radiative effects between simulations is smaller compared to the difference between fire 484 emissions, consistent with the findings in sub-Saharan African biomass-burning region (Zhang et 485 486 al. 2014).

Fire aerosols produce a negative direct effect of  $-0.1 \text{ W m}^{-2}$  in the Central U.S. and -0.86W m<sup>-2</sup> in Southern Mexico in E\_QF during the 10-day period. Within each region, negative fire aerosol SDRE peaks where fire AOD reaches maximum. Unlike the limited area affected by significant fire aerosol SDRE, fire aerosol SCRE from selected regions spreads eastward and northward, affecting remote coastal regions. Maximum SCRE stays below  $-4 \text{ W m}^{-2}$  in the central U.S. and  $-10 \text{ W m}^{-2}$  in Southern Mexico in response to significantly increased LWP and CDNC. Decreases of IWP and ICNC also contribute to fire aerosol SCRE in the Central U.S. but





with an opposite sign. The offset effect of the positive forcing induced by changes in cloud ice
properties explains the smaller SCRE in the central U.S. despite the larger changes in cloud
droplet properties.

497 We also investigated fire aerosol effects on the daily time scale, where the variation in the simulated fire aerosol effect can be large among the ensemble members. The large ensemble 498 spread of total aerosol and cloud forcing indicates large uncertainties in estimating daily fire 499 500 aerosol effects, despite similar AOD across ensemble members. Further investigations show that 501 the simulated ensemble mean and spread with less than 7 members differs considerably to those 502 with more members. A minimum of 9 members is necessary to achieve a steady estimate of the 503 magnitude and temporal variation of SCRE in this case. Our results suggest that for short-term 504 simulations of aerosol and cloud processes, even small perturbations might result in large difference across members despite constrained large scale features. In order to obtain a robust 505 506 estimate of the effective fire aerosol forcing during a short period, it is important to conduct ensemble simulations with sufficient ensemble members. 507

508

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#### 520 **References:**

- 521 Albrecht, B. A.: Aerosols, cloud microphysics, and fractional cloudiness, Science, 245, 1227-
- 522 1231, 1989.
- 523 Benedetti, A., Morcrette, J. J., Boucher, O., Dethof, A., Engelen, R., Fisher, M., Flentje, H.,
- 524 Huneeus, N., Jones, L., and Kaiser, J.: Aerosol analysis and forecast in the European centre for
- 525 medium-range weather forecasts integrated forecast system: 2. Data assimilation, Journal of
- 526 Geophysical Research: Atmospheres, 114, 2009.
- 527 Bond, T. C., Doherty, S. J., Fahey, D., Forster, P., Berntsen, T., DeAngelo, B., Flanner, M.,
- 528 Ghan, S., Kärcher, B., and Koch, D.: Bounding the role of black carbon in the climate system: A
- scientific assessment, Journal of Geophysical Research: Atmospheres, 118, 5380-5552, 2013.
- 530 Bony, S., Colman, R., Kattsov, V. M., Allan, R. P., Bretherton, C. S., Dufresne, J.-L., Hall, A.,
- 531 Hallegatte, S., Holland, M. M., and Ingram, W.: How well do we understand and evaluate
- climate change feedback processes?, Journal of Climate, 19, 3445-3482, 2006.
- 533 Boucher, O., Randall, D., Artaxo, P., Bretherton, C., Feingold, G., Forster, P., Kerminen, V.-M.,
- 534 Kondo, Y., Liao, H., and Lohmann, U.: Clouds and aerosols, in: Climate change 2013: the
- 535 physical science basis. Contribution of Working Group I to the Fifth Assessment Report of the
- 536 Intergovernmental Panel on Climate Change, Cambridge University Press, 571-657, 2013.
- 537 Brito, J., Rizzo, L. V., Morgan, W. T., Coe, H., Johnson, B., Haywood, J., Longo, K., Freitas, S.,
- 538 Andreae, M. O., and Artaxo, P.: Ground-based aerosol characterization during the South
- 539 American Biomass Burning Analysis (SAMBBA) field experiment, Atmospheric Chemistry and
- 540 Physics, 14, 12069-12083, 2014.





- 541 Chen, D., Liu, Z., Schwartz, C. S., Lin, H.-C., Cetola, J. D., Gu, Y., and Xue, L.: The impact of
- 542 aerosol optical depth assimilation on aerosol forecasts and radiative effects during a wild fire
- event over the United States, Geoscientific Model Development, 7, 2709-2715, 2014.
- 544 Chubarova, N., Nezval, Y., Sviridenkov, I., Smirnov, A., and Slutsker, I.: Smoke aerosol and its
- radiative effects during extreme fire event over Central Russia in summer 2010, Atmospheric
- 546 Measurement Techniques, 5, 557-568, 2012.
- 547 Chubarova, N. Y., Prilepsky, N. G., Rublev, A. N., and Riebau, A. R.: A Mega-Fire event in
- central Russia: fire weather, radiative, and optical properties of the atmosphere, and
  consequences for subboreal forest plants, Developments in environmental science, 8, 247-264,
  2008.
- 551 Dee, D., Uppala, S., Simmons, A., Berrisford, P., Poli, P., Kobayashi, S., Andrae, U., Balmaseda,
- M., Balsamo, G., and Bauer, P.: The ERA-Interim reanalysis: Configuration and performance of
  the data assimilation system, Quarterly Journal of the royal meteorological society, 137, 553597, 2011.
- Eskes, H., Huijnen, V., Arola, A., Benedictow, A., Blechschmidt, A.-M., Botek, E., Boucher, O.,
  Bouarar, I., Chabrillat, S., and Cuevas, E.: Validation of reactive gases and aerosols in the
  MACC global analysis and forecast system, Geoscientific model development, 8, 3523-3543,
  2015.
- Ghan, S. J., Liu, X., Easter, R. C., Zaveri, R., Rasch, P. J., Yoon, J.-H., and Eaton, B.: Toward a
  minimal representation of aerosols in climate models: Comparative decomposition of aerosol
  direct, semidirect, and indirect radiative forcing, Journal of Climate, 25, 6461-6476, 2012.
- 562 Ghan, S. J.: Technical Note: Estimating aerosol effects on cloud radiative forcing, Atmos. Chem.
- 563 Phys., 13, 9971-9974, doi:10.5194/acp-13-9971-2013, 2013.





- 564 Giglio, L., Randerson, J. T., and van der Werf, G. R. (2013), Analysis of daily, monthly, and
- 565 annual burned area using the fourth-generation global fire emissions database (GFED4) J.
- 566 Geophys. Res. Biogeosci., 118, 317–328, doi:10.1002/jgrg.20042.
- 567 Holben, B. N., Eck, T., Slutsker, I., Tanre, D., Buis, J., Setzer, A., Vermote, E., Reagan, J.,
- 568 Kaufman, Y., and Nakajima, T.: AERONET—A federated instrument network and data archive
- for aerosol characterization, Remote sensing of environment, 66, 1-16, 1998.
- 570 Iacono, M. J., Delamere, J. S., Mlawer, E. J., Shephard, M. W., Clough, S. A., and Collins, W.
- 571 D.: Radiative forcing by long-lived greenhouse gases: Calculations with the AER radiative
- transfer models, Journal of Geophysical Research: Atmospheres, 113, 2008.
- 573 Jacobson, M. Z.: Effects of biomass burning on climate, accounting for heat and moisture fluxes,
- 574 black and brown carbon, and cloud absorption effects, Journal of Geophysical Research:
- 575 Atmospheres, 119, 8980-9002, 2014.
- 576 Jiang, Y., Lu, Z., Liu, X., Qian, Y., Zhang, K., Wang, Y., and Yang, X.-Q.: Impacts of global
- 577 open-fire aerosols on direct radiative, cloud and surface-albedo effects simulated with CAM5,
- 578 Atmospheric Chemistry and Physics (Online), 16, 2016.
- 579 Kaufman, Y. J., Koren, I., Remer, L. A., Rosenfeld, D., and Rudich, Y.: The effect of smoke,
- 580 dust, and pollution aerosol on shallow cloud development over the Atlantic Ocean, Proceedings
- of the National Academy of Sciences of the United States of America, 102, 11207-11212, 2005.
- 582 Keil, A., and Haywood, J. M.: Solar radiative forcing by biomass burning aerosol particles
- 583 during SAFARI 2000: A case study based on measured aerosol and cloud properties, Journal of
- 584 Geophysical Research: Atmospheres, 108, 2003.





- 585 Kolusu, S., Marsham, J., Mulcahy, J., Johnson, B., Dunning, C., Bush, M., and Spracklen, D.:
- 586 Impacts of Amazonia biomass burning aerosols assessed from short-range weather forecasts,
- 587 Atmospheric Chemistry and Physics, 15, 12251-12266, 2015.
- 588 Kooperman, G. J., Pritchard, M. S., Ghan, S. J., Wang, M., Somerville, R. C., and Russell, L. M.:
- 589 Constraining the influence of natural variability to improve estimates of global aerosol indirect
- 590 effects in a nudged version of the Community Atmosphere Model 5, Journal of Geophysical
- 591 Research: Atmospheres, 117, 2012.
- 592 Korontzi, S., McCarty, J., Loboda, T., Kumar, S., and Justice, C.: Global distribution of
- <sup>593</sup> agricultural fires in croplands from 3 years of Moderate Resolution Imaging Spectroradiometer
- 594 (MODIS) data, Global Biogeochemical Cycles, 20, 2006.
- 595 Lin, N.-H., Tsay, S.-C., Maring, H. B., Yen, M.-C., Sheu, G.-R., Wang, S.-H., Chi, K. H.,
- 596 Chuang, M.-T., Ou-Yang, C.-F., and Fu, J. S.: An overview of regional experiments on biomass
- 597 burning aerosols and related pollutants in Southeast Asia: From BASE-ASIA and the Dongsha
- 598 Experiment to 7-SEAS, Atmospheric Environment, 78, 1-19, 2013.
- Liu, X., Easter, R. C., Ghan, S. J., Zaveri, R., Rasch, P., Shi, X., Lamarque, J.-F., Gettelman, A.,
- 600 Morrison, H., and Vitt, F.: Toward a minimal representation of aerosols in climate models:
- 601 Description and evaluation in the Community Atmosphere Model CAM5, Geoscientific Model
- 602 Development, 5, 709, 2012. Jiang, Y.: Impacts of global open-fire aerosols on direct radiative,
- cloud and surface-albedo effects simulated with CAM5, Atmos. Chem. Phys, 1680, 7324, 2016.
- 604 Lu, Z., and Sokolik, I. N.: The effect of smoke emission amount on changes in cloud properties
- and precipitation: A case study of Canadian boreal wildfires of 2007, Journal of Geophysical
- 606 Research: Atmospheres, 118, 2013.





- 607 Magi, B., Rabin, S., Shevliakova, E., and Pacala, S.: Separating agricultural and non-agricultural
- fire seasonality at regional scales, Biogeosciences, 9, 3003, 2012.
- 609 Malm, W. C., Schichtel, B. A., Pitchford, M. L., Ashbaugh, L. L., and Eldred, R. A.: Spatial and
- 610 monthly trends in speciated fine particle concentration in the United States, Journal of
- 611 Geophysical Research: Atmospheres, 109, 2004.
- 612 Mlawer, E. J., Taubman, S. J., Brown, P. D., Iacono, M. J., and Clough, S. A.: Radiative transfer
- 613 for inhomogeneous atmospheres: RRTM, a validated correlated-k model for the longwave,
- Journal of Geophysical Research: Atmospheres, 102, 16663-16682, 1997.
- 615 Morrison, H., and Gettelman, A.: A new two-moment bulk stratiform cloud microphysics
- scheme in the Community Atmosphere Model, version 3 (CAM3). Part I: Description and
- numerical tests, Journal of Climate, 21, 3642-3659, 2008.
- 618 Mu, M., Randerson, J., van der Werf, G., Giglio, L., Kasibhatla, P., Morton, D., Collatz, G.,
- 619 DeFries, R., Hyer, E., and Prins, E.: Daily and hourly variability in global fire emissions and
- 620 consequences for atmospheric model predictions of carbon monoxide, 2011.
- 621 Neale, R. B., Richter, J. H., and Jochum, M.: The impact of convection on ENSO: From a
- delayed oscillator to a series of events, Journal of Climate, 21, 5904-5924, 2008.
- 623 Park, S., and Bretherton, C. S.: The University of Washington shallow convection and moist
- turbulence schemes and their impact on climate simulations with the Community Atmosphere
- 625 Model, Journal of Climate, 22, 3449-3469, 2009.
- 626 Randerson, J., Chen, Y., Werf, G., Rogers, B., and Morton, D.: Global burned area and biomass
- 627 burning emissions from small fires, Journal of Geophysical Research: Biogeosciences, 117,
- **628** 2012.





- 629 Reddington, C., Yoshioka, M., Balasubramanian, R., Ridley, D., Toh, Y., Arnold, S., and
- 630 Spracklen, D.: Contribution of vegetation and peat fires to particulate air pollution in Southeast
- Asia, Environmental Research Letters, 9, 094006, 2014.
- 632 Richter, J. H., and Rasch, P. J.: Effects of convective momentum transport on the atmospheric
- circulation in the Community Atmosphere Model, version 3, Journal of Climate, 21, 1487-1499,
- 634 2008.
- Rubin, J. I., Reid, J. S., Hansen, J. A., Anderson, J. L., Hoar, T. J., Reynolds, C. A., Sessions, W.
- 636 R., and Westphal, D. L.: Development of the Ensemble Navy Aerosol Analysis Prediction
- 637 System (ENAAPS) and its application of the Data Assimilation Research Testbed (DART) in
- support of aerosol forecasting, Atmospheric Chemistry and Physics, 16, 3927, 2016.
- 639 Sena, E., Artaxo, P., and Correia, A.: Spatial variability of the direct radiative forcing of biomass
- 640 burning aerosols and the effects of land use change in Amazonia, Atmospheric Chemistry and
- 641 Physics, 13, 1261-1275, 2013.
- 642 Stier, P., Schutgens, N., Bellouin, N., Bian, H., Boucher, O., Chin, M., Ghan, S., Huneeus, N.,
- Kinne, S., and Lin, G.: Host model uncertainties in aerosol radiative forcing estimates: results
  from the AeroCom Prescribed intercomparison study, Atmospheric Chemistry and Physics, 13,
- 645 3245-3270, 2013.
- Tarasova, T., Gorchakova, I., Sviridenkov, M., Anikin, P., and Romashova, E.: Estimation of the
  radiative forcing of smoke aerosol from radiation measurements at the Zvenigorod scientific
  station in the summer of 2002, Izvestiya Atmospheric and Oceanic Physics, 40, 454-463, 2004.
- 649 Ten Hoeve, J. E., Jacobson, M. Z., and Remer, L. A.: Comparing results from a physical model
- 650 with satellite and in situ observations to determine whether biomass burning aerosols over the
- Amazon brighten or burn off clouds, Journal of Geophysical Research: Atmospheres, 117, 2012.





- 652 Tosca, M., Randerson, J., and Zender, C.: Global impact of smoke aerosols from landscape fires
- on climate and the Hadley circulation, Atmospheric Chemistry and Physics, 13, 5227-5241,
- 654 2013.
- Twomey, S.: The influence of pollution on the shortwave albedo of clouds, Journal of the atmospheric sciences, 34, 1149-1152, 1977.
- van der Werf, G. R., Randerson, J. T., Giglio, L., Collatz, G. J., Mu, M., Kasibhatla, P. S.,
- 658 Morton, D. C., DeFries, R. S., Jin, Y., and van Leeuwen, T. T.: Global fire emissions and the
- contribution of deforestation, savanna, forest, agricultural, and peat fires (1997–2009), Atmos.
- 660 Chem. Phys., 10, 11707-11735, doi:10.5194/acp-10-11707-2010, 2010.
- 661 Ward, D., Kloster, S., Mahowald, N., Rogers, B., Randerson, J., and Hess, P.: The changing
- radiative forcing of fires: global model estimates for past, present and future, AtmosphericChemistry and Physics, 12, 2012.
- 664 Wan, H., Rasch, P. J., Zhang, K., Qian, Y., Yan, H., and Zhao, C.: Short ensembles: an efficient
- 665 method for discerning climate-relevant sensitivities in atmospheric general circulation models,
- 666 Geoscientific Model Development, 7, 1961-1977, 2014.
- 667 Wu, L., Su, H., and Jiang, J. H.: Regional simulations of deep convection and biomass burning
- over South America: 2. Biomass burning aerosol effects on clouds and precipitation, Journal of
- 669 Geophysical Research: Atmospheres, 116, 2011.
- 670 Zamora, L. M., Kahn, R., Cubison, M. J., Diskin, G., Jimenez, J., Kondo, Y., McFarquhar, G.,
- 671 Nenes, A., Thornhill, K., and Wisthaler, A.: Aircraft-measured indirect cloud effects from
- biomass burning smoke in the Arctic and subarctic, Atmospheric Chemistry and Physics, 16,
- 673 715-738, 2016.





- 674 Zhang, F., Wang, J., Ichoku, C., Hyer, E. J., Yang, Z., Ge, C., Su, S., Zhang, X., Kondragunta,
- 675 S., and Kaiser, J. W.: Sensitivity of mesoscale modeling of smoke direct radiative effect to the
- 676 emission inventory: a case study in northern sub-Saharan African region, Environmental
- 677 Research Letters, 9, 075002, 2014.
- 678 Zhang, G. J., and McFarlane, N. A.: Sensitivity of climate simulations to the parameterization of
- 679 cumulus convection in the Canadian Climate Centre general circulation model, Atmosphere-
- 680 ocean, 33, 407-446, 1995.
- 681 Zhang, J., Reid, J. S., Westphal, D. L., Baker, N. L., and Hyer, E. J.: A system for operational
- aerosol optical depth data assimilation over global oceans, Journal of Geophysical Research:Atmospheres, 113, 2008.
- K., Wan, H., Liu, X., Ghan, S. J., Kooperman, G., Ma, P.-L., Rasch, P. J., Neubauer, D.,
- and Lohmann, U.: Technical Note: On the use of nudging for aerosol-climate model
  intercomparison studies, Atmospheric Chemistry and Physics, 14, 8631-8645, 2014.
- 687 Zhang, Z., Meyer, K., Yu, H., Platnick, S., Colarco, P., Liu, Z., and Oreopoulos, L.: Shortwave
- 688 direct radiative effects of above-cloud aerosols over global oceans derived from 8 years of
- 689 CALIOP and MODIS observations, Atmospheric Chemistry and Physics, 16, 2877-2900, 2016.
- 690

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# Table 1. List of CAM5 simulations.

| Name                               | Fire emission | Simulation period    | Member | Nudging          |  |  |
|------------------------------------|---------------|----------------------|--------|------------------|--|--|
| Group A: Single member simulations |               |                      |        |                  |  |  |
| S_NF                               | No            |                      |        |                  |  |  |
| S_GF3                              | GFED v3       |                      |        |                  |  |  |
| S_GF4                              | GFED v4.1     | January 1- April 30, | 1      | Horizontal winds |  |  |
| S_QF                               | QFED v2.4     | 2009                 | -      | (6h)             |  |  |
| Group B: Ensemble simulations      |               |                      |        |                  |  |  |
| E_NF                               | No            |                      |        | Horizontal winds |  |  |
| E_GF3                              | GFED v3       | April 1 - April 10,  | 10     | (6h) and         |  |  |
| E_GF4                              | GFED v4.1     |                      |        | temperature      |  |  |
| E_QF                               | QFED v2.4     | 2009                 |        | (~10d)*          |  |  |

\* See section 2.3 for details about ensembles





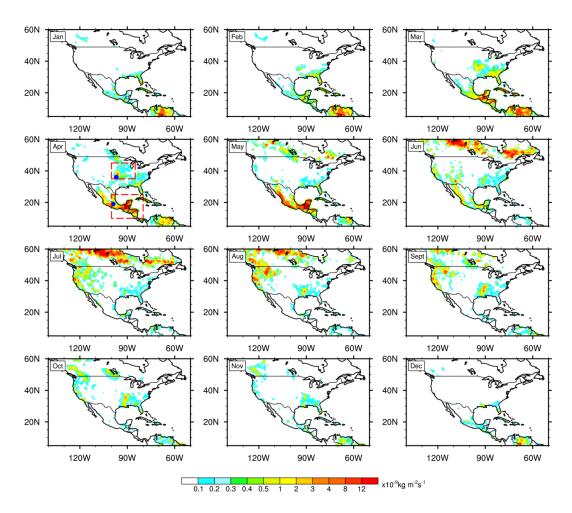


Figure 1. Spatial distributions of multi-year monthly mean biomass burning consumed dry matter over North America during 2003-2014 from GFEDv4.1. Boxes denote selected regions: central U.S (35 - 45°N, 85 - 100°W) and Southern Mexico (10 - 25°N, 80 - 100°W). Dots denote locations of AERONET sites: Cart Site (36°N, 97°W) and Mexico City (19°N, 99°W)





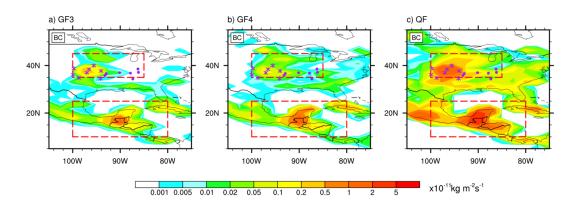


Figure 2. Spatial distributions of monthly mean BC emissions from three emission inventories in April, 2009. IMPROVE data sites are shown as asterisks for sites near the source region and as dots for sites in the region downwind of the fire source.





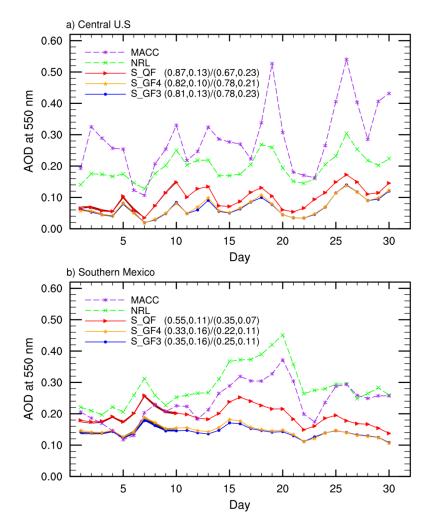


Figure 3. Time series of daily regional mean AOD in April, 2009 in simulations and reanalysis data. Numbers in parenthesis denote TCC and RMSE between each simulation in group A and reanalysis data (left: NRL; right: MACC). Individual lines indicate group A simulations. Shaded areas (very narrow) in slightly darker colors during April 1-10 illustrate maximum and minimum values of daily mean AOD among ensemble members in group B simulations.





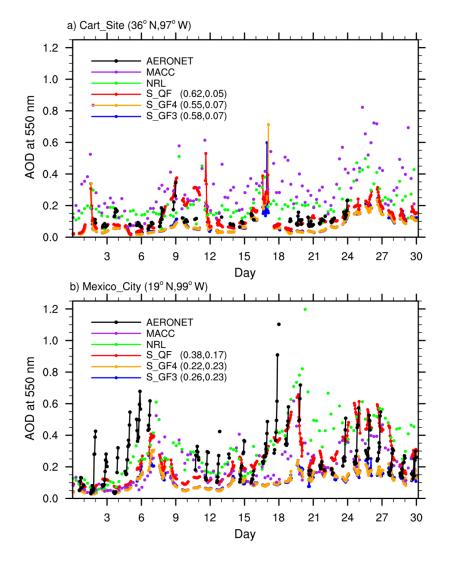


Figure 4. Time series of hourly regional mean AOD in April, 2009 from group A simulations, reanalysis data and AERONET retrievals at AERONET sites. Numbers in parenthesis denote TCC (left) and RMSE (right) between each simulation and AERONET AOD.





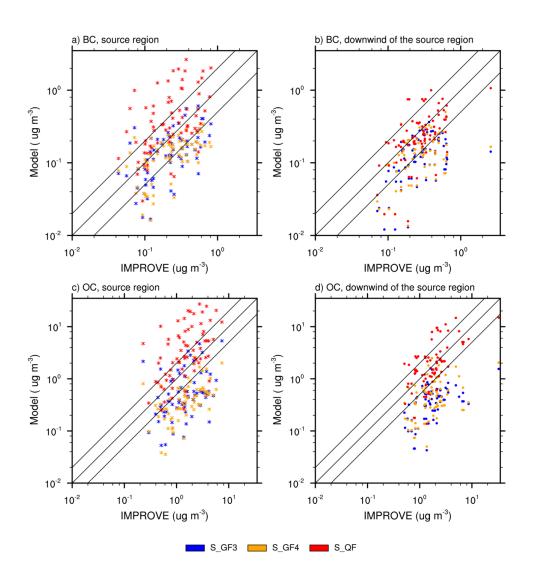


Figure 5. Evaluation of the simulated BC (up) and POM (bottom) concentrations in group A simulations against the IMPROVE data at sites near the source and downwind the source region. Locations of these sites are marked with the same symbol as in Fig. 2.





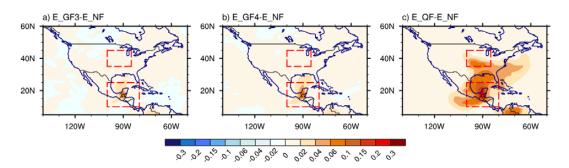


Figure 6. Spatial distributions of 10-day average (Apr. 1-10) ensemble mean AOD differences between simulations with (E\_GF3, E\_GF4, and E\_QF) and without fire emission (E\_NF).





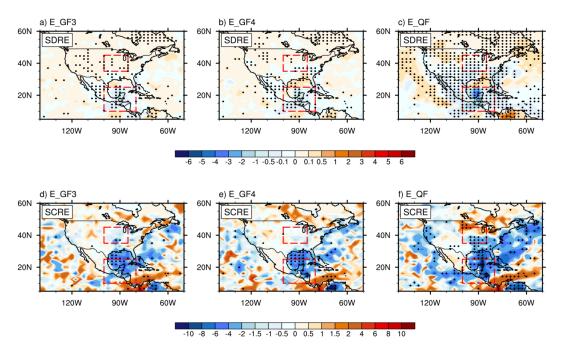


Figure 7. Spatial distributions of 10-day average (Apr. 1-10) ensemble mean fire aerosol shortwave direct radiative effect (SDRE) and shortwave cloud radiative effect (SCRE) ( $W m^{-2}$ ) in group B simulations. Dots denote regions where SDRE is statistically significant at the 95% confidence level based on the KS test.





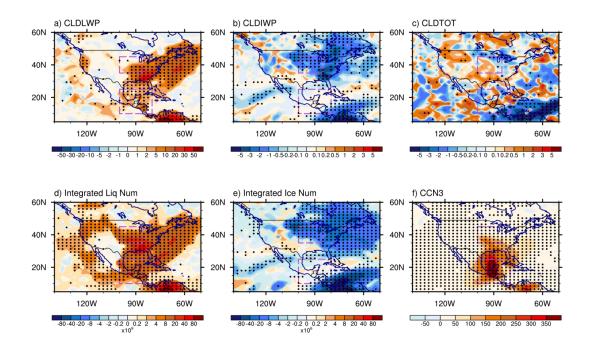


Figure 8. Difference of 10-day average (Apr.1-10) ensemble mean between simulations E\_NF and E\_QF: a) cloud liquid water path (  $g m^{-2}$  ), b) cloud ice water path (  $g m^{-2}$  ), c) total cloud fraction (%), d) column-integrated droplet number concentration (  $m^{-2}$  ), e) column-integrated ice number concentration ( $m^{-2}$  ), and f) cloud condensation nuclei at 0.1% supersaturation near 900 hPa. Dots denote regions where the difference is statistically significant at the 95% confidence level based on the KS test.





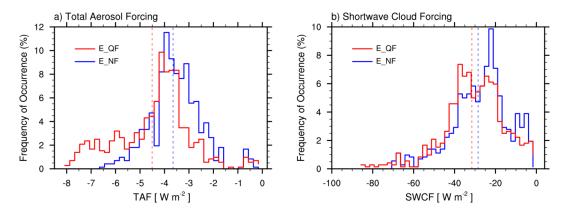


Figure 9. Probability distributions of 10-day average (Apr.1-10) a) total aerosol forcing and b) cloud forcing over Southern Mexico in simulations E\_NF and E\_QF sampled from grid values of ensemble members (72x10 for each case). Dashed lines indicate the mean of the distribution.





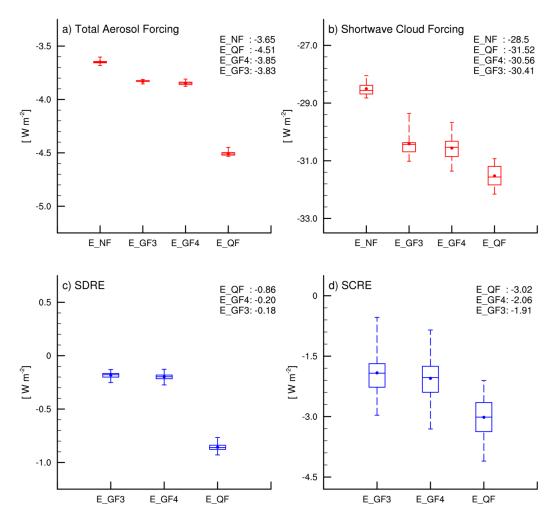


Figure 10. 10-day average (Apr. 1-10) regional mean a) total aerosol forcing, b) total shortwave cloud forcing and fire aerosol, c) SDRE, and d) SCRE in Southern Mexico in group B simulations. Box denotes the  $25^{th}$  and  $75^{th}$  percentiles. Bars outside the box indicate minimum and maximum. Bar within the box denotes the  $50^{th}$  percentile. Total aerosol and cloud forcing are sampled from different ensemble members (10 for each case). Fire aerosol SDRE and SCRF are sampled by calculating the difference between members in simulations E\_QF (E\_GF3/E\_GF4) and E\_NF (10x10 for each case).





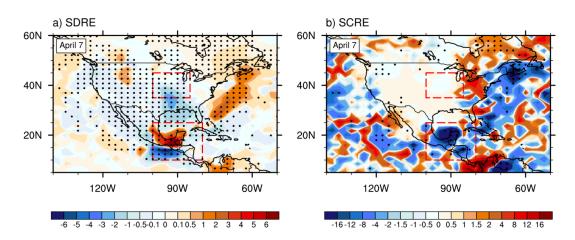


Figure 11. Spatial distributions of ensemble mean fire aerosol a) SDRE and b) SCRE ( $W m^{-2}$ ) on April 7 in the E\_QF simulation. Dots denote grids where fire aerosol effect is statistically significant at the 95% confidence level based on the KS test.





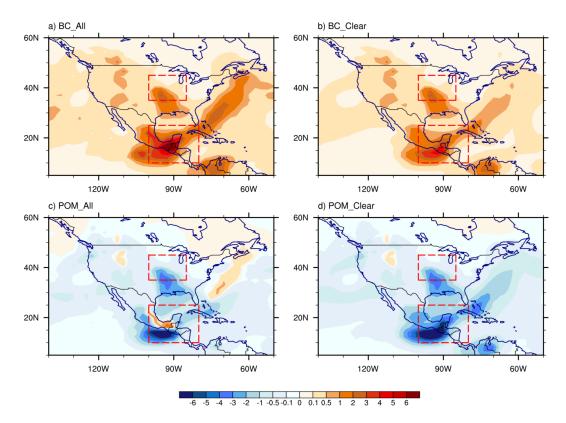


Figure 12. Spatial distributions of fire BC SDRE and fire POM SDRE (  $W m^{-2}$  ) on all-sky and clear-sky conditions on April 7 in the E\_QF simulation.





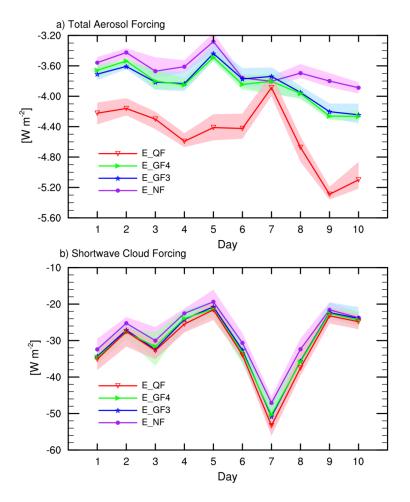


Figure 13. Time series of daily regional mean total a) aerosol forcing and b) cloud forcing in Southern Mexico during Apr.1-10, 2009 in group B simulations. Individual lines indicate ensemble mean values. Shaded areas illustrate the ensemble spread (from minimum to maximum).





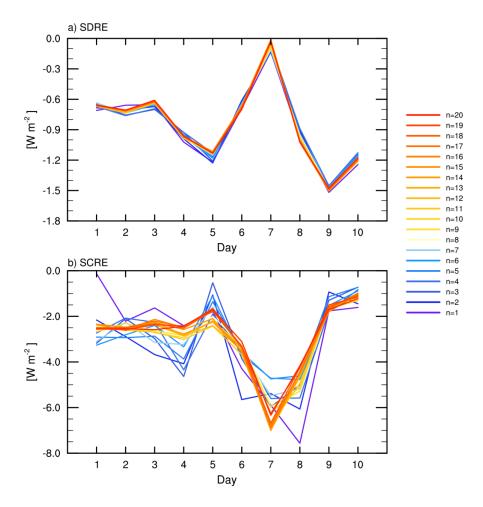


Figure 14. Time series of daily ensemble mean fire aerosol a) SDRE and b) SCRE averaged over Southern Mexico during Apr. 1-10, 2009 in QFED forced ensemble simulations with varying the total number of ensemble members (n=1-20).