Response to Reviewer #1

We thank Reviewer #1 for their valuable and helpful comments. Our responses to the comments are provided below in bold font with the reviewer's comments in italicized font.

The paper reports a comprehensive study of the climate and air quality impacts of residential solid fuel combustion. The paper reports the first assessment of the impacts of carbonaceous aerosol from residential solid fuel combustion through changes in ice nuclei as well as a comprehensive assessment of aerosol indirect effects through changes in liquid clouds.

The paper makes an important contribution to our understanding of residential solid fuel combustion on climate and is suitable for publication in ACP. Importantly the manuscript highlights that the overall climate impact of carbonaceous aerosol from residential solid fuel combustion is uncertain and that even the sign of the impact is still ambiguous. The paper certainly motivates further study of this important issue and highlights where some of the major uncertainties lie.

The manuscript is very well written and clear. I only have a few minor comments. I suggest publication in ACP after the following minor comments have been addressed.

Minor Comments:

Page 5. Have you stated the size of the emitted carbonaceous aerosol? This is important for the impacts of carbonaceous aerosol on cloud condensation nuclei and aerosol indirect effects.

Response: We have added the size range of the emitted carbonaceous aerosols in the text (Page 5, Lines 155-156): "Specifically, BC and POM from solid fuel cookstove emissions are treated in the accumulation mode, with size range of 0.058-0.27 μm (Liu et al., 2012)."

Page 12. Most models do not simulate the impacts of BC on ice nuclei. I think it would be useful to provide a brief summary of how this was treated in the Methods section of the paper.

Response: We agree with the reviewer. We have moved part of the contents from Section 3.4.4 to a new section in Methods as Section 2.5 and added the description of BC as IN following Barahona and Nenes (2008, 2009) in the text (Page 7, Lines 197-212):

2.5 Simulations: BC active as IN

"In default CAM5-Chem, BC is not treated as IN (Liu et al., 2012; Tilmes et al., 2015). IN concentrations from homogeneous nucleation are calculated as a function of vertical velocity (Liu et al., 2007). Several lab and field studies indicate that BC particles can act as IN (Cozic et al., 2008; DeMott et al., 1999; Koehler et al., 2009; Kulkarni et al., 2016). Therefore, we conduct additional simulations that treat BC as an effective IN applying the ice nucleation scheme of Barahona and Nenes (2008, 2009). The scheme estimates maximum supersaturation and ice crystal concentrations and considers competition between homogeneous and heterogeneous freezing. Homogeneous nucleation occurs in solution droplets formed on soluble aerosols (mainly sulfate), while heterogeneous nucleation occurs on IN, which here are a small subset of mineral dust and black carbon particles. The heterogeneous freezing of BC and dust is described as a generalized ice nucleation spectrum.

We perform three additional model simulations, with model configurations identical to those in Table 2, except for the treatment of BC particles as effective IN. In addition, for each model simulation, we alter the plausible maximum freezing efficiency (MFE) of BC as 0.01, 0.05 and 0.1 that provides an uncertainty range in the global climatic impact assessment."

Page 15. As stated by the authors the aerosol indirect effect is considerably larger than other studies. Ward et al. (2012) also found a large aerosol indirect effect using the CAM model (although a different version) to study carbonaceous aerosol from fires. It might be useful to point this out in the text.

Response: We have added (Page 16, Lines 489-490): "Consistent with our study, Ward et al. (2012) also found a large AIE (-1.74 to 1.00 W m^{-2}) for carbonaceous aerosols from fires using CESM CAM4-Chem."

Fig. 4 and 5. Please clarify whether these are reported at ambient conditions or at standard temperature and pressure.

Response: We have clarified the reporting units for BC and POM under standard temperature and pressure, which have been added in the caption of Fig. 4 as (Page 36, Lines 916-917) "Figure 4. Annual zonal mean BC concentrations from (a) the BASE simulation, (b) the global and (c) India solid fuel cookstove emissions. BC concentrations are calculated under standard temperature and pressure conditions (273 K, 1 atm)."

Response to Reviewer #2

We thank Reviewer #2 for their valuable and helpful comments. Our responses to the comments are provided below in bold font with the reviewer's comments in italicized font.

The manuscript by Huang et al. presents a modeling study of the radiative effects of solid fuel cookstoves, both globally and specifically in India. There is a lot of scientific and policy interest in this topic, given the potential climate co-benefits of cookstove intervention programs and the uncertainty associated with quantifying this, with several recent papers making a range of estimates that differ in sign. The contribution of this article is a welcome addition to the field, focusing particularly on details of aerosol cloud interactions, and considering the effect of BC ice nucleation, which has not been considered in such studies previously. Overall the manuscript is thus appropriate in scope for ACP. It is also generally clear, well organized, and easy to ready. I only have a few comments that are detailed below; there is some ambiguity regarding how the authors are arriving at their uncertainty estimates, and some of the motivation for the scope of their analysis (e.g., considering just India, or not considering the impact of co-emitted GHGs) could be strengthened. Addressing these would constitute minor revision.

Major Comments:

37: An important conclusion, which I mainly agree with in spirit. However, it might be stated a little bit softer for a few reasons. First, the uncertainty of up to a factor of two in estimating concentrations would seem to contribute to the overall uncertainty in the net radiate effects. Second, this study is of radiative effects, not the climate response, and the wording should reflect that. Third, it's the result of only a single model, which may not be as definitive as presented.

Response: We prefer to keep the original statement as is. First, the manuscript includes a comparative description of all previous results with several different global aerosol-climate model frameworks that supports the uncertain sign conclusion for net global radiative effect of cookstove aerosol emissions. Second, the statement clearly refers to carbonaceous aerosols only and highlights the need for improved constraints on aerosol-cloud interactions. Finally, our study quantifies the impacts of cookstove carbonaceous aerosol emissions on global average annual mean radiative effect because it is a linear predictor of

global average surface air temperature response at equilibrium. Our community is still several years away from any quantitative robust mechanistic understanding of regional climate response to regional radiative effect of aerosols (e.g. Kasoar et al., ACP, 2016). It would be fairly pedagogical to convert the global radiative effect results to global average surface air temperature response e.g. Berntsen and Fuglestvedt, PNAS, 2008.

The measurements used for comparison come from very different time periods (2010 for IMPROVE, 2009-2013 for Europe, 2000-2008 for AMS data, and 1993-2016 for AERONET). How does this impact the evaluation of modeled concentrations and AOD, given that the model uses emissions from 2010 and that there have been large changes in emissions over this time period?

Response: The measurement data availability and model configuration has necessitated using climatological observations for BC, OA and AOD with some apparent mismatches between observational years and simulations years for OA and AOD. Model simulated BC concentrations were sampled in exact correspondence to the observed temporal period. The global aerosol-climate model framework is a chemistry-climate model with specified dynamics (CCM-SD), not a chemistry transport model (CTM). CTMs exist to compare with exact measurement periods. In our case, the model output reflects a present-day climatology rather than a specific CTM run year as such. Therefore, the comparison with available climatological measurements does allow us to validate and have insights into the large-scale aerosol system dynamics and behavior. We clarify in the text (Pages 14-15, Lines 429-436): "The simulations reflect a present-day climatology forced with recycled year 2010 anthropogenic emissions. Model simulated BC concentrations were sampled in exact correspondence to the observed temporal period. In some limited cases, OA and AOD are not exactly temporally consistent with the available aerosol measurement network climatologies applied in the evaluation. For regions where carbonaceous aerosol emissions have undergone substantial changes over short periods, the model-measurement comparison may therefore introduce additional uncertainty. However, we focus the evaluation on the large-scale regional aerosol system dynamics."

215 - 220: Some previous studies have suggested that the resolution of global scale models leads to a bias that makes it difficult to match AOD from AERONET in these regions. Could this partially explain the low bias?

Response: We agree and have added (Page 9, Lines 257-259): "The model underestimate of AOD from AERONET in India may also be related to the fairly coarse global model resolution, as previously reported by Pan et al. (2015) and Zhang et al. (2015)."

Fig 6 and Section 3.4.1 (and everywhere these numbers are quoted in the text): Suddenly the results have errors associated with them (concentrations and AOD did not: : :). What is the meaning of the error estimates? Are these the standard deviations over the timeframe modeled? If so, that needs to be more clearly stated when presenting these numbers in the abstract and conclusion (that +/- is modeled temporal standard deviation). And then I wonder why similar deviations were not considered for discussion of concentrations or AOD. Further, temporal variability is very different than e.g. an estimate of uncertainty owing to sources of model error or approximations, such as the ranges provided for the RF of the simulations including BC IN that stem from uncertainty in the MFE. These ranges can't be directly compared, and yet they're presented in e.g. the abstract without distinguishing their different meanings. At present the non BC-IN ranges come across as uncertainty estimates that seem much too small (I doubt the authors believe that the aerosol RF in any single model could be that accurate).

Response: We include uncertainty estimates that are based on interannual internal climate variability (n=5 years). For consistency, we have added the uncertainty ranges based on interannual internal climate model variability to concentrations and burdens of BC and POM, and AOD where multiple years have been sampled for the comparison. For example, model BC concentrations were sampled in correspondence to the exact temporal measurement period, thus no range included for that case. This revision is reflected throughout the updated manuscript, in the abstract, Section 3.1, Section 3.2, Section 3.3 and Section 4 e.g. (Page 1, Lines 24-26): "However, the model tends to underestimate AOD over India and China by ~ 19 ± 4% but overestimate it over Africa by ~ 25 ± 11%

(uncertainty range due to interannual internal climate model variability for n=5 run years)."

(Pages 8-9, Lines 239-244): "Figure 2 shows the evaluation of simulated surface OA against observations. Over East Asia, the model slightly underestimates observed OA, with a NMB of -8.5 ± 5% (Fig. 2a). In contrast, the simulated OA concentrations overestimate the measurements by over a factor of 2 in North America, with a NMB value of 124 ± 24% (Fig. 2b). For the European sites, we find a simulated OA overestimation of measured concentrations by up to 0.9 ± 0.7 μg m^{-3}, corresponding to a NMB of +32 ± 26% (Fig. 2c)."

(Page 9, Lines 248-250): "Over India, the simulated annual mean AOD is lower than observations by about 16 ± 3% (Fig. 3a), with large bias sources mainly from the northern India regions (e.g., New Delhi and Kanpur)."

(Page 9, Lines 259-261): "A similar pattern is found over China (Fig. 3b) and the rest of Asia (Fig. 3c), with NMB values of -21 ± 4% and -15 ± 6% respectively."

(Page 9, Lines 263-265): "This directly leads to annual mean model simulated AOD values over Africa 25 ± 11% higher than observations because Saharan dust emissions dominate the AOD over North Africa (Fig. 3d)."

(Pages 9-10, Lines 270-274): "In these two regions, modeled AOD agrees with observations within a factor of 2, with NMB values -20 ± 4% and -18 ± 9% respectively. CAM5-Chem overestimates AOD over Australia (Fig. 3h) and remote sites (Fig. 3i), with NMB values of +69 ± 17% and +47 ± 12%, respectively. Globally, model simulated AOD agrees quite well with observations, with NMB values close to zero."

(Page 10, Lines 279-282): "For the control simulation, global annual mean BC burden and lifetime are 0.12 ± 0.001 Tg and 4.5 ± 0.04 days, respectively (Table 3), at the low end of the range estimated by AeroCom (Schulz et al., 2006; Textor et al., 2006)."

(Page 10, Lines 290-292): "Annual mean BC burdens from global and Indian solid fuel cookstove emissions account for about 24.2 ± 0.7% and 5.0 ± 0.0% of that in the control simulation (0.12 ± 0.001 Tg)."

(Page 10, Lines 296-298): "In our control simulation, the annual mean POM burden is 0.66 ± 0.006 Tg, and the global annual mean POM lifetime is 4.8 ± 0.04 days (Table 3)."

(Page 11, Lines 307-308): "The annual mean POM burdens from global and Indian solid fuel cookstove emissions are 0.13 ± 0.004 Tg and 0.027 ± 0.002 Tg respectively."
(Page 15, Lines 436-442): "In the control simulation, the global annual mean BC burden and lifetime are 0.12 ± 0.001 Tg and 4.5 ± 0.04 days. For POM, the burden and lifetime are 0.66 ± 0.006 Tg and 4.8 ± 0.04 days. Annual mean surface BC (POM) concentrations over Northern India, East China and sub-Saharan Africa are 1.55 ± 0.076, 0.76 ± 0.028 and 0.11 ± 0.004 μg m^{-3} (7.11 ± 0.32, 3.95 ± 0.12 and 0.48 ± 0.02 μg m^{-3}), respectively. BC and POM burdens from global solid fuel cookstove emissions are 0.029 ± 0.001 and 0.13 ± 0.004 Tg, while contributions from the Indian sector are 0.006 ± 0.000 and 0.027 ± 0.004 Tg, respectively."

We have clarified in the abstract and text to distinguish the 2 uncertainty range calculations used in the study (1) based on interannual internal climate model variability (2) based on BC maximum freezing efficiency range (Page 2, Lines 38-39):
"Here, the uncertainty range is based on sensitivity simulations that alter the maximum freezing efficiency of BC across a plausible range: 0.01, 0.05 and 0.1."
(Page 7, Lines 209-212): "We perform three additional model simulations, with model configurations identical to those in Table 2, except for the treatment of BC particles as effective IN. In addition, for each model simulation, we alter the plausible maximum freezing efficiency (MFE) of BC as 0.01, 0.05 and 0.1 that provides an uncertainty range in the global climatic impact assessment."

Introduction: I didn't get a good sense from the introduction why there is a particular interest in India as separate from the globe in this study (as opposed to China, or any other country with significant cookstove use). I'm not suggesting that the authors do more simulations for other regions, but if they wish to include the India-specific results it would make more sense to include a bit more rational for this emphasis.

Response: We have expanded the India regional focus motivation statement (Page 2, Lines 53-61):

"India contains a large concentration of solid fuel-dependent households: approximately 160 million households use solid fuels for cooking (Venkataraman et al., 2010). In India, residential biofuel combustion represents the dominant energy sector and accounts for over 50% of the total source of BC and OC emissions (Klimont et al., 2009). India has a long history of unsuccessful stove intervention programs that have sometimes focused on health benefits (Hanbar and Karve, 2002; Kanagawa and Nakata, 2007; Kishore and Ramana, 2002). Despite years of interventions, the vast majority of Indian households still rely on traditional stoves (Legros et al., 2009). The possible scope for global climate co-benefits in future Indian cookstove intervention programs warrants further examination and analysis of this region."

439 - 447: I think it's worth recognizing that there are climate impacts from GHG emissions as well. So, considering not just the aerosol emissions, these may be large enough to make the net climate images of cookstove emissions positive (Lacey 2017). It is somewhat artificial to envision a scenario wherein only the carbonaceous aerosol cookstove emissions are effected by stove replacements.

Response: We have added (Page 17, Lines 508-510) "This study does not include the greenhouse gas emission effects from the solid fuel cookstove sector, which may indeed be large enough to imply a net warming global climate impact depending on time scale (Lacey et al., 2017)."

Minor Comments:

12: Not clear – "updated" related to what? A particular previous study? Later it becomes evident what is meant (first to include BC as IN), but perhaps it could be worded differently here.

Response: We have deleted "updated".

74: Clarify whether the Butt 2016 study included just aerosols or also GHGs in the DRE.

Response: We have clarified (Page 3, Lines 86-88): "Butt et al. (2016) reported that the net DRE and AIE of aerosols from the residential emission sector (including coal) ranged from -66 to +21 mW m^{-2}, and from -52 to -16 mW m^{-2}, respectively. Their study did not include greenhouse gases."

80: Similarly, Ethiopia is the 3rd largest in Lacey 2017, but that's including GHGs in 2050, which is a slightly more specific statement than as presented here.

Response: We have revised to (Pages 3-4 Lines 89-95) "From the perspective of policy-relevant country-level assessment of cookstove burning on global climate, Lacey and Henze (2015) revealed that solid fuel cookstove aerosol emissions resulted in global air surface temperature changes ranging from 0.28 K cooling to 0.16 K warming; Lacey et al. (2017) further concluded that emissions reductions, including both aerosols and greenhouse gases, from China, India and Ethiopia contributed the most to the global surface temperature changes by 2050."

128: CAM5-Chem not CAM5-chem

Response: Corrected.

Fig 1: Sorry if I just missed it, but did the authors state how they are defining urban vs rural in their classification of measurement sites?

Response: We have added the definition of the classification of urban and rural sites in the text as (Page 5 Lines 128-129) "Here we define urban (including semi-urban) sites as the geographic locations of the measured sites locating in a city, others as rural sites."

276-279: What is the BC mass absorption coefficient (MABS) at 550 nm in this model? See e.g. Koch ACP 2009

Response: BC mass absorption cross section coefficient at 550 nm in CAM5-Chem is 14.6 m^2 g^{-1}. We have updated this in the revised text as (Page 11 Lines 319-323) "CAM5-Chem assumes that BC is internally mixed with other components in the accumulation mode and simulates enhanced absorption (BC mass absorption cross section = 14.6 m^2 g^{-1}) when BC is coated by soluble aerosol components and water vapor (Ghan et al., 2012), which results in larger estimates of the DRE than for BC alone (Bond et al., 2013; Jacobson, 2001b)."

Global radiative effects of solid fuel cookstove aerosol emissions

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Abstract. We apply the NCAR CAM5-Chem global aerosol-climate model to quantify the net global radiative effects of black and organic carbon aerosols from global and Indian solid fuel cookstove emissions for the year 2010. Our assessment accounts for the direct radiative effects, changes to cloud albedo and lifetime (aerosol indirect effect, AIE), impacts on clouds via the vertical temperature profile (semi-direct effect, SDE), and changes in the surface albedo of snow and ice (surface albedo effect). In addition, we provide the first estimate of household solid fuel black carbon emission effects on ice clouds. Anthropogenic emissions are from the IIASA GAINS ECLIPSE V5a inventory. A global dataset of black carbon (BC) and organic aerosol (OA) measurements from surface sites and aerosol optical depth (AOD) from AERONET is used to evaluate the model skill. Compared with observations, the model successfully reproduces the spatial patterns of atmospheric BC and OA concentrations, and agrees with measurements to within a factor of 2. Globally, the simulated AOD agrees well with observations, with normalized mean bias close to zero. However, the model tends to underestimate AOD over India and China by ~ 19 ± 4% but overestimate it over Africa by ~ 25 ± 11% (uncertainty range due to interannual internal climate model variability for n=5 run years). Without BC serving as ice nuclei (IN), global and Indian solid fuel cookstove aerosol emissions have a net cooling impact on global climate of -141 ± 4 mW m^{-2} and -12 ± 4 mW m^{-2}, respectively. The net radiative impacts are dominated by the AIE and SDE mechanisms, which originate from enhanced cloud condensation nuclei

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concentrations for the formation of liquid and mixed-phase clouds, and a suppression of convective transport of water vapor from the lower troposphere to the upper troposphere/lower stratosphere that in turn leads to reduced ice cloud formation. When BC is allowed to behave as a source of IN, the net global climate impacts of the global and Indian solid fuel cookstove emissions range from -275 to +154 mW m^{-2} and -33 to +24 mW m^{-2}, with globally averaged values -59 ± 215 and 0.3 ± 29 mW m^{-2} respectively. Here, the uncertainty range is based on sensitivity simulations that alter the maximum freezing efficiency of BC across a plausible range: 0.01, 0.05 and 0.1. BC-ice cloud interactions lead to substantial increases in high cloud (< 500 hPa) fractions. Thus, the net sign of the impacts of carbonaceous aerosols from solid fuel cookstoves on global climate (warming or cooling) remains ambiguous until improved constraints on BC interactions with mixed-phase and ice clouds are available.

1. Introduction

Worldwide 2-3 billion people rely on solid fuels for the majority of their energy needs (Legros et al., 2009). This household biomass combustion includes burning wood fuel, agricultural residues and dung for cooking, heating and lighting. Emissions from household solid fuel combustion include greenhouse gases (carbon dioxide and methane), black carbon (BC), organic carbon (OC), and other trace gases (e.g., nitrogen oxides). Globally, BC from household solid fuel emissions accounts for approximately 25% of the total anthropogenic BC emissions (Bond et al., 2013). Among different types of cookstoves, advanced charcoal stoves show lowest BC emission factors, followed by simple charcoal, advanced biomass, rocket and simple wood stoves, respectively (Garland et al., 2017). India contains a large concentration of solid fuel-dependent households: approximately 160 million households use solid fuels for cooking (Venkataraman et al., 2010). In India, residential biofuel combustion represents the dominant energy sector and accounts for over 50% of the total source of BC and OC emissions (Klimont et al., 2009). India has a long history of unsuccessful stove intervention programs that have sometimes focused on health benefits (Hanbar and Karve, 2002; Kanagawa and Nakata, 2007; Kishore and Ramana, 2002). Despite years of interventions, the vast majority of Indian households still rely on traditional stoves (Legros et al., 2009). The possible scope for global climate co-benefits in future Indian cookstove intervention programs warrants further examination and analysis of this region. BC-rich household solid fuel emission plays an important role in affecting regional air quality (Archer-Nicholls et al.,

2016; Carter et al., 2016; Liu et al., 2016) and influencing global climate change (Bauer et al., 2010; Butt et al., 2016; Venkataraman et al., 2005). The human health consequences of solid fuel combustion are substantial (Archer-Nicholls et al., 2016; Ezzati and Kammen, 2002; Lelieveld et al., 2015). Nearly 9% of the global burden of disease is attributable to exposure to household air pollution from solid fuels, equivalent to 2.9 million premature deaths and 86 million disability adjusted life years (DALYs) annually (GBD 2015 Risk Factors Collaborators, 2016). Half of the world's population is exposed to indoor air pollution, mainly attributable to solid fuel usage for household cooking and heating (Bonjour et al., 2013; Smith et al., 2014).

Carbonaceous aerosols from solid fuel combustion interact with the Earth's radiation budget directly by absorbing and scattering solar radiation (direct radiative effect, DRE) and indirectly by changing cloud albedo and lifetime (aerosol indirect effect, AIE), modifying the vertical temperature profile (semi-direct effect, SDE), and changing the surface albedo over snow and ice (surface albedo effect, SAE) (Boucher et al., 2013; Chung, 2005; Chylek and Wong, 1995; Ghan, 2013; Ghan et al., 2012; Myhre et al., 2013). Carbonaceous aerosols affect cloud albedo and lifetimes (the AIE) by acting as cloud condensation nuclei (CCN) or ice nuclei (IN), thus modifying cloud properties and changing the top-of-atmosphere (TOA) radiative fluxes (Lohmann, 2002; Lohmann et al., 2000; Penner et al., 1992; Pierce et al., 2007; Spracklen et al., 2011b). The net climatic effect of carbonaceous aerosols from household solid fuel combustion is not well constrained and even the sign is uncertain (Bond et al., 2013). Bauer et al. (2010) estimated that the aerosol net global climate impact of residential biofuel carbonaceous aerosol emissions is -130 mW m^{-2}. Kodros et al. (2015) have estimated that net DRE of solid fuel aerosol emissions ranges from -20 to +60 mWm^{-2}, AIE from -20 to +10 mWm^{-2}, with uncertainties due to assumptions of the aerosol emission masses, size distribution, aerosol optical properties and mixing states. Butt et al. (2016) reported that the net DRE and AIE of aerosols from the residential emission sector (including coal) ranged from -66 to +21 mW m^{-2}, and from -52 to -16 mW m^{-2}, respectively. Their study did not include greenhouse gases. Moreover, neither of the latter two studies consider the aerosol cloud-lifetime effect (second indirect effect), SDE and SAE. From the perspective of policy-relevant country-level assessment of cookstove burning on global climate, Lacey and Henze (2015) revealed that solid fuel cookstove aerosol emissions resulted in global air surface temperature changes ranging from 0.28 K cooling to 0.16 K warming; Lacey et al. (2017)

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further concluded that emissions reductions, including both aerosols and greenhouse gases, from China, India and Ethiopia contributed the most to the global surface temperature changes by 2050.

None of the previous assessments have included BC-ice cloud interactions that can exert a large influence on the atmospheric radiation balance. A recent study by Kulkarni et al. (2016) showed that BC could act as IN, which was also shown by past lab and field findings (Cozic et al., 2008; DeMott et al., 1999; Koehler et al., 2009). With BC as IN, Penner et al. (2009) estimated that the total radiative forcing of anthropogenic and biomass BC emissions was -300 to -400 mW m^{-2}, with IN parameterizations following Liu and Penner (2005) and Kärcher et al. (2006). Gettelman et al. (2012) further concluded that AIE from BC emissions was -60 mW m^{-2}, with ice nucleation parameterization following Barahona and Nenes (2009). Hence, a re-assessment of the global climate change impacts of carbonaceous aerosol emissions from the solid fuel cookstove sector that newly incorporates BC as IN is urgently needed.

Here, we employ a global aerosol-climate model to quantify the impacts of solid fuel cookstove carbonaceous aerosol emissions globally and from India on global climate change. Sect. 2 presents the Methods including the evaluation measurement data sets for BC, OA and aerosol optical depth (AOD), the model description and experimental design. Sect. 3 details the results of the model evaluation and the impacts of the global and Indian solid fuel cookstove emissions on the atmospheric radiation budget and global climate. Discussion and summary are provided in Sect. 4.

2. Methods

2.1 BC and OC evaluation measurement database

Ground-based BC observations are from IMPROVE (the Interagency Monitoring of PROtected Visual Environment, http://vista.cira.colostate.edu/Improve/) for the year 2010 over North America (Malm et al., 1994), EMEP (the European Monitoring and Evaluation Programme, http://ebas.nilu.no) for 2009-2013 over Europe, and sporadic measurement campaigns for China and India. Elemental carbon (EC) concentrations are measured using Thermal Optical Reflectance (TOR) (Chow et al., 1993, 2004; EMEP/MSC-W et al., 2014). Our measurement database comprises a total of 152 sites from IMPROVE, 28 sites from EMEP, 35 sites for China, and 41 sites for India. The number of urban sites includes 8 from IMPROVE, 5 from EMEP, 17 for China,

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and 23 for India. Here we define urban (including semi-urban) sites as the geographic locations of the measured sites locating in a city, others as rural sites.

A global network of aerosol mass spectrometer (AMS) surface measurements for organic aerosol (OA) for 2000-2008 are used to compare with model simulations (Spracklen et al., 2011a; Zhang et al., 2007; Zheng et al., 2015). The AMS technique measures hydrocarbon-like OA (HOA), oxygenated OA (OOA) and total OA (HOA + OOA). HOA is a surrogate for primary OA (POA) emitted directly from fossil fuel and biomass burning, while OOA is a surrogate for secondary OA (SOA). In this study, we compare monthly mean total OA with model simulated total OA (POA + SOA). The majority of the AMS measurements in the surface concentration database were made prior to 2005.

Ground-based AOD observations from AERONET (AErosol RObtic NETwork, https://aeronet.gsfc.nasa.gov) during 1993-2016 are applied to examine model skill (Dubovikl and King, 2000; Holben et al., 1998, 2001). A climatological AOD value averaged over 1993-2016 for each site is used to compare with the model simulation. The AERONET version 2 level-2 product is used in this study.

2.2 NCAR CAM5-Chem global model description

We apply the NCAR Community Atmosphere Model version 5.3 with chemistry (CAM5-Chem) within the Community Earth System Model (CESM) version 1.2.2 (Emmons et al., 2010; Lamarque et al., 2012; Tilmes et al., 2015). The oxidant-aerosol system is fully coupled in CAM5-Chem. The horizontal resolution of CAM5-Chem is 0.9° latitude by 1.25° longitude, with 56 vertical levels from surface up to about 40 km. In the standard CAM5-Chem, aerosol microphysical processes are represented using a 3-mode scheme (MAM3; aitken, accumulation and coarse modes). MAM3 simulates both mass and number concentrations of aerosols. Aerosol size distributions in each mode are assumed to be lognormal (Liu et al., 2012). The model treats the effects of aerosol acting as CCN in liquid-phase clouds (Ghan et al., 2012). The aerosol components in MAM3 include BC, primary organic matter (POM), secondary organic aerosol (SOA), sulfate, sea salt and dust, which are assumed to be internally mixed within each lognormal mode. Specifically, BC and POM from solid fuel cookstove emissions are treated in the accumulation mode, with size range of 0.058-0.27 μm (Liu et al., 2012). Mass yields of semi-

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volatile organic gas-phase species (SOAG) from emissions of isoprene, monoterpenes, big alkanes and alkenes, as well as toluene are prescribed (Emmons et al., 2010; Liu et al., 2012; Tilmes et al., 2015). The condensable SOAG reversibly and kinetically partitions into the aerosol phase to form SOA in CAM5-Chem as described in Liu et al. (2012).

2.3 Emissions

Global anthropogenic emissions are from the IIASA (International Institute for Applied System Analysis) Greenhouse Gas-Air Pollution Interactions and Synergies (GAINS) integrated assessment model ECLIPSE V5a (Evaluating the Climate and Air Quality Impacts of Short-lived Pollutants version 5a) for the year 2010 (Amann et al., 2011, 2013; Klimont et al., 2017; Stohl et al., 2015). Species in ECLIPSE V5a include BC, POM, sulfur dioxide, nitrogen oxides, carbon monoxide, volatile organic compounds, and ammonium, with their annual global budgets for the year 2010 shown in Table 1. ECLIPSE V5a emissions available at 0.5° latitude by 0.5° longitude spatial resolutions are re-gridded to the model spatial resolution. ECLIPSE V5a does not include shipping or wildfire biomass burning emissions, which are instead obtained from the IPCC AR5 RCP8.5 scenario for the year 2010 (Riahi et al., 2011).

2.4 Simulations: BC not active as IN

Atmosphere-only simulations are performed in specified dynamics (SD) mode with offline meteorological fields from the Goddard Earth Observing System model version 5 (GEOS-5). In this SD mode configuration, the internally derived meteorological fields (e.g., horizontal wind component, air temperature and latent heat flux) are nudged by 10% towards reanalysis fields from GEOS-5 for every model time step. The nudging technique in CAM5-Chem has been evaluated to quantify the aerosol indirect effect in order to reduce the influence of natural variability (Kooperman et al., 2012). Sea surface temperature and sea ice in the model are prescribed from the Climatological/Slab-Ocean Data Model (DOCN) and Climatological Ice Model (DICE) respectively, with monthly-varying decadal mean averaged over 1981-2010.

We perform three sets of model simulations using the model configurations shown in Table 2. The first set of simulations represents the control with anthropogenic emissions following ECLIPSE V5a, as described above (hereafter referred to as BASE). The second set of simulations are

identical to the BASE simulation except the global solid fuel cookstove emissions for aerosols and gas-phase aerosol and ozone precursors are set to zero (termed as GBLSF_OFF). The third set of simulations is identical to BASE except the solid fuel cookstove emissions are set to zero over the Indian sub-continent (termed as INDSF_OFF). We run all the above simulations for 6 years from 2005 to 2010, with the first year discarded as spin-up and the last five years averaged for output analysis. The differences between BASE and GBLSF_OFF isolate the impacts of the global solid fuel cookstove sector aerosol emissions, and the differences between BASE and INDSF_OFF isolate the impacts of the Indian solid fuel cookstove sector aerosol emissions. Top-of-the-atmosphere (TOA) aerosol shortwave (SW) and longwave (LW) radiative effects are calculated using the Rapid Radiative Transfer Model for GCMs (RRTMG) that is coupled to CAM5-Chem (Ghan, 2013; Ghan et al., 2012).

2.5 Simulations: BC active as IN

In default CAM5-Chem, BC is not treated as IN (Liu et al., 2012; Tilmes et al., 2015). IN concentrations from homogeneous nucleation are calculated as a function of vertical velocity (Liu et al., 2007). Several lab and field studies indicate that BC particles can act as IN (Cozic et al., 2008; DeMott et al., 1999; Koehler et al., 2009; Kulkarni et al., 2016). Therefore, we conduct additional simulations that treat BC as an effective IN applying the ice nucleation scheme of Barahona and Nenes (2008, 2009). The scheme estimates maximum supersaturation and ice crystal concentrations and considers competition between homogeneous and heterogeneous freezing. Homogeneous nucleation occurs in solution droplets formed on soluble aerosols (mainly sulfate), while heterogeneous nucleation occurs on IN, which here are a small subset of mineral dust and black carbon particles. The heterogeneous freezing of BC and dust is described as a generalized ice nucleation spectrum.

We perform three additional model simulations, with model configurations identical to those in Table 2, except for the treatment of BC particles as effective IN. In addition, for each model simulation, we alter the plausible maximum freezing efficiency (MFE) of BC as 0.01, 0.05 and 0.1 that provides an uncertainty range in the global climatic impact assessment.

3 Results

3.1 Evaluation of surface BC and OA concentrations

Surface observation networks from IMPROVE, EMEP, and various campaigns in China and India are employed to compare with model simulations, as shown in Figure 1. We diagnose the normalized mean bias (NMB) for each source region, calculated as

$$\mathrm{NMB} = \left(\frac{\sum_i (M_i - O_i)}{\sum_i O_i}\right) \times 100\% \quad (1)$$

where M and O represent monthly mean model simulated and observational concentrations at site *i* respectively, and $\sum$ is the sum over all the sites within a source region.

In general, the model simulated surface BC concentrations agree with observations to within a factor of 2, consistent with previous studies (Huang et al., 2013; Wang et al., 2011, 2014a, 2014b). A total of 41 surface BC observational sites are used to evaluate the model simulation over India (Fig. 1a). On average, the model underestimates surface BC concentrations by approximately 45% and 34% over urban and rural sites respectively, with a total NMB -41% (Fig. 1a), which implies a marked underestimation of the BC emissions in India. Previous modeling studies have also reported large underestimates of BC surface concentrations over India against observations (Gadhavi et al., 2015; He et al., 2014; Zhang et al., 2015). Part of the model/measurement discrepancy is related to a sampling bias because the majority of the observations are located over urban or heavily polluted regions. For China sites, the NMB value is -16% (Fig. 1b). Similar to India, the model substantially underestimates the surface BC concentrations over urban sites with a NMB of -30%. However, the model performs relatively well over rural areas, with a NMB close to zero. For IMPROVE, the NMB values for rural and urban sites are -15% and -43%, respectively, with a total NMB -28% (Fig. 1c). Over Europe, the model simulated surface BC concentrations agree quite well with observations, with a NMB value of -8%, although two urban sites show substantial model underestimation (Fig. 1d).

The 40 AMS surface OA measurements are grouped into three categories: East Asia (8 sites), North America (17 sites) and Europe (15 sites) (Spracklen et al., 2011a; Zhang et al., 2007; Zheng et al., 2015). Figure 2 shows the evaluation of simulated surface OA against observations. Over East Asia, the model slightly underestimates observed OA, with a NMB of -8.5 ± 5% (Fig. 2a). In contrast, the simulated OA concentrations overestimate the measurements by over a factor of 2 in North America, with a NMB value of 124 ± 24% (Fig. 2b). For the European sites, we find a

simulated OA overestimation of measured concentrations by up to 0.9 ± 0.7 μg m^{-3}, corresponding to a NMB of +32 ± 26% (Fig. 2c).

3.2 Evaluation of model AOD

Figure 3 compares simulated AOD values against observations over nine regions across the globe, including India, China, Rest of Asia (excluding China and India), Africa, South America, North America, Europe, Australia and remote regions. Over India, the simulated annual mean AOD is lower than observations by about 16 ± 3% (Fig. 3a), with large bias sources mainly from the northern India regions (e.g., New Delhi and Kanpur). This is consistent with Quennehen et al. (2016) who also reported that model simulated AOD values were generally lower than satellite-derived AOD over northern India, using the same emission inventory as our study. As discussed in Sect. 3.1, model simulated surface BC concentrations over India are also underestimated (by up to 41%), therefore, the low bias of model simulated AOD can be attributed, in part, to the underestimation of Indian BC emissions from ECLIPSE V5a emission inventory (Stohl et al., 2015), although global anthropogenic BC budgets in ECLIPSE V5a lie in the high end compared with previous studies (Bond et al., 2004, 2013; Janssens-Maenhout et al., 2015). The model underestimate of AOD from AERONET in India may also be related to the fairly coarse global model resolution, as previously reported by Pan et al. (2015) and Zhang et al. (2015). A similar pattern is found over China (Fig. 3b) and the rest of Asia (Fig. 3c), with NMB values of -21 ± 4% and -15 ± 6% respectively. Model simulated AOD values from several sites in West Asia (Fig. 3c) are higher than observations, which is probably caused by the model overestimation of dust emissions (He and Zhang, 2014). This directly leads to annual mean model simulated AOD values over Africa 25 ± 11% higher than observations because Saharan dust emissions dominate the AOD over North Africa (Fig. 3d). For South America, the model generally agrees quite well with observations (Fig. 3e), except for a few sites where model simulated AOD values are lower than observations by more than a factor of 2. This is probably due to the model underestimation of biomass burning emissions there (Reddington et al., 2016). AOD values over North America (Fig. 3f) and Europe (Fig. 3g) are relatively lower (with values generally < 0.3), due to lower anthropogenic emissions. In these two regions, modeled AOD agrees with observations within a factor of 2, with NMB values -20 ± 4% and -18 ± 9% respectively. CAM5-Chem overestimates AOD over Australia (Fig. 3h) and remote sites (Fig. 3i), with NMB values of +69 ± 17% and +47

± 12%, respectively. Globally, model simulated AOD agrees quite well with observations, with NMB values close to zero.

3.3 Contribution of solid fuel cookstove sector emissions to atmospheric BC and POM

3.3.1 BC

Annual BC emissions and budgets are reported in Table 3 based on the anthropogenic inventory from ECLIPSE V5a. Annual BC emissions from the global and Indian solid fuel cookstove emissions are 2.31 and 0.36 Tg yr^{-1}, accounting for 23.7% and 3.7% of the total BC emissions. For the control simulation, global annual mean BC burden and lifetime are 0.12 ± 0.001 Tg and 4.5 ± 0.04 days, respectively (Table 3), at the low end of the range estimated by AeroCom (Schulz et al., 2006; Textor et al., 2006).

Figure 4 shows the zonal mean BC concentrations from the control simulation (Fig. 4a), global (Fig. 4b) and Indian (Fig. 4c) solid fuel cookstove emissions respectively. For the control simulation, in general, the highest BC concentrations (by up to 0.40 μg m^{-3}) occur at the surface over the emission source regions in the mid-latitudes (e.g., China and India). In the tropics and mid-latitudes, zonal mean BC concentrations decrease with increasing altitude, due to wet removal and deposition, as found in Huang et al. (2013). A similar vertical distribution is observed for the impacts from global and Indian solid fuel cookstove emissions, although the magnitude is smaller, compared with the control simulation. Annual mean BC burdens from global and Indian solid fuel cookstove emissions account for about 24.2 ± 0.7% and 5.0 ± 0.0% of that in the control simulation (0.12 ± 0.001 Tg).

3.3.2 POM

Global POM emissions are mainly from biomass burning (31 Tg yr^{-1}) and anthropogenic emissions (18.9 Tg yr^{-1}), with global and Indian solid fuel cookstove emissions accounting for, 21% and 3.4% respectively, of the total POM emissions (Table 3). In our control simulation, the annual mean POM burden is 0.66 ± 0.006 Tg, and the global annual mean POM lifetime is 4.8 ± 0.04 days (Table 3).

In Figure 5, we show the annual zonal mean POM concentrations for the control simulation (Fig. 5a) and for global (Fig. 5b) and Indian (Fig. 5c) solid fuel cookstove emissions. There are two maxima in the annual zonal mean POM concentrations near the surface. One is located in the tropics due to the large biomass burning emissions there, and the other is located over mid-latitude regions and originates mainly from anthropogenic emissions (Chung and Seinfeld, 2002; Huang et al., 2013). For POM concentrations from global solid fuel cookstove emissions, a single maximum is evident in the Northern Hemisphere (NH) subtropics at the surface (Fig. 5b). The surface maximum for the Indian solid fuel cookstove emissions reaches a maximum in the NH subtropics. The annual mean POM burdens from global and Indian solid fuel cookstove emissions are 0.13 ± 0.004 Tg and 0.027 ± 0.002 Tg respectively.

3.4 Impacts of solid fuel cookstove aerosol emissions on global climate change

3.4.1 Direct radiative effect (DRE)

The DRE impacts of the global and Indian solid fuel cookstove emissions are shown in Figure 6. For the global solid fuel cookstove sector, the globally averaged DRE from aerosol emissions is +70 ± 3 mW m^{-2} without treating BC as IN, which is a warming effect. The positive DRE from global solid fuel cookstove emissions shows large spatial variability, with the largest impacts located over western Africa, followed by India and China (figure not shown). The contributions of BC and POM to DRE are +105 ± 4 (warming) and -14 ± 1 (cooling) mW m^{-2}, respectively. In other words, the warming effect of BC is partially offset by the cooling effect from POM. Additional cooling effects may come from sulfate and SOA. CAM5-Chem assumes that BC is internally mixed with other components in the accumulation mode and simulates enhanced absorption (BC mass absorption cross section = 14.6 m^2 g^{-1}) when BC is coated by soluble aerosol components and water vapor (Ghan et al., 2012), which results in larger estimates of the DRE than for BC alone (Bond et al., 2013; Jacobson, 2001b).

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The DRE from Indian solid fuel cookstove emissions also corresponds to a net warming effect (Fig. 6), with a global annual mean value of +11 ± 1 mW m^{-2}. Large impacts are found over continental India, the Tibetan Plateau and southeastern China. On a global annual basis, DRE values from BC and POM emissions from the Indian solid fuel cookstove sector are +18 ± 1 and -3 ± 0.2 mW m^{-2}, respectively.

3.4.2 Aerosol indirect, semi-direct and surface albedo effects: BC not active as IN

Global annual mean AIE and SAE values from global and Indian solid fuel cookstove aerosol emissions are shown in Figure 6. In our study, AIE includes the first (albedo) and second (lifetime) indirect effects, as well as the semi-direct effect. Annually averaged AIE from the global solid fuel cookstove sector is -226 ± 5 mW m^{-2} (Fig. 6), with annual mean shortwave (SW) AIE -122 ± 22 mW m^{-2} and longwave (LW) AIE -104 ± 17 mW m^{-2}, without treating BC as IN. Both the annual mean SW and LW AIE thus yield cooling effects. The cooling signals of SW AIE mainly occur over the western coast of South America, west and east coasts of Africa, South China and Himalaya regions (figure not shown). This is directly linked to the contribution of global solid fuel cookstove aerosol emissions to CCN (Pierce et al., 2007), which increases the cloud droplet number concentrations (CDNC) and cloud liquid water path (CLWP). Figure 7 shows the global vertically-integrated distribution of CLWP from the contribution of global solid fuel cookstove aerosol emissions. The higher CLWP is due to the enhanced lifetime of liquid and mixed-phase clouds, which therefore reflect more solar radiation, leading to cooling effect. For the LW AIE, the largest cooling effect is found over tropical regions, especially over southern India and the Indian Ocean. In order to investigate the causes of the LW AIE cooling effect, we analyze the cloud fraction change over a defined region (Latitude:0-20ºN; Longitude:60-90ºE) due to the effect from the global solid fuel cookstove sector. As shown in Figure 8a, cloud fraction in the lower troposphere increases. However, in the middle and upper troposphere cloud fraction decreases by up to 0.6%, with the strongest decrease found at ~150 hPa. We further analyze the changes in shallow and deep convective mass fluxes of moisture over the same domain. As shown in Figure 8b, moist shallow convective mass flux generally shows increases in the lower troposphere, which means that solid fuel cookstove aerosol emissions enhance the convective transport of water vapor within the boundary layer. By contrast, the deep convective mass flux demonstrates decreases from surface up to the middle troposphere (Fig. 8c). This indicates that solid fuel cookstove aerosol emissions may stabilize the boundary layer and inhibit the transport of water vapor from the surface to the upper troposphere/lower stratosphere, which leads to decreases in ice cloud formation, thus reducing cloud cover in the upper troposphere and lower stratosphere (UTLS) region at around 200 hPa (Fig. 8a) and a LW AIE cooling effect.

The global annual mean AIE from Indian solid fuel cookstove aerosol emissions accounts for approximately 10% (-22 ± 3 mW m^{-2}) relative to the value of AIE from the global solid fuel cookstove sector (Fig. 6), with globally averaged SW and LW AIE values of -3 ± 11 and -19 ± 11 mW m^{-2} respectively.

Global annual mean SAE values from global and Indian solid fuel cookstove sector are relatively small: +15 ± 3 and -2 ± 3 mW m^{-2}, respectively (Fig. 6). The warming effect is mainly due to the deposition of BC on the surface of snow and sea ice (Flanner et al., 2007; Ghan, 2013; Ghan et al., 2012).

3.4.3 Total radiative effect: BC not active as IN

The net total radiative effect of global and Indian solid fuel cookstove aerosol emissions are both cooling, with the global annual mean estimated to be -141 ± 4 and -12 ± 4 mW m^{-2} respectively (Fig. 6). This suggests that if we remove solid fuel cookstove aerosol emissions, it will result in warming and thus slightly increased global surface air temperature. That being said, this is likely to be quite sensitive to model representation of aerosol mixing state (Fierce et al., 2017).

3.4.4 Total radiative effect: BC active as IN

For the radiative effect of global solid fuel cookstove emissions with BC as IN, global annual mean DRE is 105 ± 13 mW m^{-2}, ranging from +90 to +115 mW m^{-2}, which is 29-64% higher than the DRE values from the default scheme (Fig. 6). Intriguingly, large globally averaged negative SW AIE (-1.36 ± 0.63 W m^{-2}) and positive LW AIE (+1.18 ± 0.44 W m^{-2}) for global solid fuel cookstove aerosol emissions are found, with annual mean values for the SW AIE ranging from -1.83 to -0.64 W m^{-2} and from +0.67 to +1.45 W m^{-2} for the LW AIE. This results in a rather uncertain net AIE, with a global annual mean AIE of -177 ± 223 mW m^{-2} (Fig. 6). The reason for the large global annual average negative SW AIE and positive LW AIE is a substantial increase in high cloud (< 500 hPa) fractions when BC acts as an efficient IN. For instance, with MFE = 0.1, large increases (by up to 9%) in high cloud fractions from global solid fuel cookstove aerosol emissions are found over subtropical regions, especially over the southern Atlantic Ocean (Fig. 9). With BC particles active as IN, ice particle sizes become smaller, leading to a slower settling velocity for ice particles and thus an increase in the lifetime of ice clouds. Increases in high clouds

Deleted: In default CAM5-Chem, BC is not treated as IN (Liu et al., 2012; Tilmes et al., 2015). However, several lab and field studies have shown that BC particles could act as IN (Cozic et al., 2008; DeMott et al., 1999; Koehler et al., 2009; Kulkarni et al., 2016), as discussed in Section 1. Therefore, we conduct sensitivity studies in our model simulations by treating BC as an effective IN, with the ice nucleation scheme by Barahona and Nenes (2008, 2009). We run three additional model simulations, with model configurations identical to those in Table 2, except for the treatment of BC particles as effective IN. In addition, for each model simulation, we alter the plausible maximum freezing efficiency (MFE) of BC as 0.01, 0.05 and 0.1, from which the uncertainty ranges of the climatic impacts from global and Indian solid fuel cookstove aerosol emissions with BC as IN are quantified.¶

not only reflect more solar radiation back to space, but also trap more LW radiation within the troposphere. For SAE, the global annual mean value is +12 ± 10 mW m^{-2} (Fig. 6). As a result, the net total radiative effect of global solid fuel cookstove aerosol emissions ranges from -275 to + 154 mW m^{-2}, with a global annual mean of -59 ± 215 mW m^{-2} (Fig. 6). Again, the source of the large uncertainty of the total radiative effect is due to the choice of MFE values. With MFE = 0.01, the global mean LW AIE (+672 mW m^{-2}) outweighs SW AIE (-638 mW m^{-2}), and therefore results in a net warming effect. For other MFE values (0.05 and 0.1), the absolute global annual mean SW AIE values are always higher than the LW AIE, leading to a net negative (i.e., cooling) total radiative effect.

For the Indian solid fuel cookstove sector, the global annual mean net total radiative effect is 0.3 ± 29 mW m^{-2}, with an AIE of -18 ± 37 and a SAE of +1 ± 8 mW m^{-2}, respectively.

4 Discussion and Summary

In this study, we employ the atmospheric component of a global 3-D climate model CESM v1.2.2, CAM5.3-Chem, to investigate the impacts of solid fuel cookstove emissions on global climate change. We update the default anthropogenic emission inventory using IIASA ECLIPSE V5a for the year 2010. We focus our analysis on the radiative effects of global and Indian solid fuel cookstove aerosol emissions. Model performance is evaluated against a global dataset of BC and OA measurements from surface sites and AOD from AERONET. Compared with observations, the model successfully reproduces the spatial patterns of atmospheric BC and OA concentrations, and generally agrees with measurements to within a factor of 2. Globally, the simulated AOD agrees quite well with observations, with NMB values close to zero. Nevertheless, the model tends to underestimate AOD values over source regions (except for Africa) and overestimate AOD over remote regions. The underestimates of AOD over India and China indicate that anthropogenic emissions of carbonaceous aerosols and sulfate precursors in ECLIPSE V5a are underestimated because carbonaceous aerosols and sulfate account for over 60% of the AOD over these two countries (Lu et al., 2011; Streets et al., 2009), which may introduce uncertainties for our climate estimates. The simulations reflect a present-day climatology forced with recycled year 2010 anthropogenic emissions. Model simulated BC concentrations were sampled in exact correspondence to the observed temporal period. In some limited cases, OA and AOD are not

exactly temporally consistent with the available aerosol measurement network climatologies applied in the evaluation. For regions where carbonaceous aerosol emissions have undergone substantial changes over short periods in the past few years, the model-measurement comparison may therefore introduce additional uncertainty. However, we focus the evaluation on the large-scale regional aerosol system dynamics. In the control simulation, the global annual mean BC burden and lifetime are 0.12 ± 0.001 Tg and 4.5 ± 0.04 days. For POM, the burden and lifetime are 0.66 ± 0.006 Tg and 4.8 ± 0.04 days. Annual mean surface BC (POM) concentrations over Northern India, East China and sub-Saharan Africa are 1.55 ± 0.076, 0.76 ± 0.028 and 0.11 ± 0.004 μg m^{-3} (7.11 ± 0.32, 3.95 ± 0.12 and 0.48 ± 0.02 μg m^{-3}), respectively. BC and POM burdens from global solid fuel cookstove emissions are 0.029 ± 0.001 and 0.13 ± 0.004 Tg, while contributions from the Indian sector are 0.006 ± 0.000 and 0.027 ± 0.004 Tg, respectively.

In the default CESM simulations without treating BC as IN, globally averaged DRE values from global and Indian solid fuel cookstove emissions are +70 ± 3 and +11 ± 1 mW m^{-2}, respectively. The contributions of BC and POM from global solid fuel cookstove emissions to the DRE are +105 ± 4 and -14 ± 1 mW m^{-2}. Global annual mean SW and LW AIE values from global solid fuel cookstove emissions are -122 ± 22 and -104 ± 17 mW m^{-2}, with contributions from India yielding -3 ± 11 mW m^{-2} for the SW AIE and -19 ± 11 mW m^{-2} for the LW AIE, respectively. The cooling effect of the SW AIE is associated with the increases of CCN and CDNC, whereas the negative effects of LW AIE are caused by the suppression of convection that transports water vapor from lower troposphere to upper troposphere/stratosphere, thus reducing ice cloud cover. The CAM5-Chem also computes the SAE, with global and Indian solid fuel cookstove emissions contributing +15 ± 3 and -2 ± 3 mW m^{-2}, respectively. As a result, the net total radiative effects of global and Indian solid fuel cookstove emissions are -141 ± 4 and -12 ± 4 mW m^{-2}, respectively, both producing a net cooling effect.

Sensitivity studies are carried out to examine the impacts of global and Indian solid fuel cookstove emissions on climate by treating BC as an effective IN, with MFE as 0.01, 0.05 and 0.1, respectively. For the climate impacts of global solid fuel cookstove emissions, global annual mean DRE is +105 ± 13 mW m^{-2}, which is ~ 50% higher than the default model scheme in which BC particles are not treated as IN (Fig. 6). This is driven by the increases of BC burden (due to prolonged BC lifetimes) from global solid fuel cookstove emissions by up to 17% with BC as IN.

Because the BC absorption effect dominates the DRE, increases in BC burden enhance the magnitude of annual mean DRE (Jacobson, 2001a). Compared with the default model scheme, significant changes in globally averaged SW AIE are found, with a global annual mean of -1.36 ± 0.63 W m^{-2}, which is about an order of magnitude higher than that from the default scheme. Moreover, in contrast to the cooling effect found in the default scheme, annual mean positive LW AIE is simulated here (+1.18 ± 0.44 W m^{-2}). The above changes in cookstove emission induced SW and LW AIE are caused by the substantial increases in high cloud (< 500 hPa) fractions with BC particles acting as IN by up to 9% due to the effect of solid fuel cookstove emissions. Large increases in high cloud fractions are found mainly over tropical regions, especially over southern Africa. For the SAE, similar to the model default scheme, the global annual mean value is +12 ± 10 mW m^{-2}. Summing up the DRE, the AIE and the SAE, the net total radiative effect of global solid fuel cookstove emissions is -59 ± 215 mW m^{-2}. For the Indian sector, the global mean total radiative effect is 0.3 ± 29 mW m^{-2}, with a net AIE -18 ± 37 and a SAE +1 ± 8 mW m^{-2}, respectively.

We compare our simulation results with previous studies as shown in Figure 10. The globally averaged DRE in our control simulation is more than four times higher than that from the baseline simulation of Kodros et al. (2015), which assumes homogeneous particle mixing state (Fig. 10). Annual emissions of BC from global solid fuel cookstove sector in our study (2.3 Tg C yr^{-1}) is approximately 44% higher than that from global biofuel emissions (1.6 Tg C yr^{-1}) in Kodros et al. (2015), which, to some extent, leads to differences in annual mean DRE values together with different optical calculations. The annual mean DRE value from another study by Butt et al. (2016) differs from ours in magnitude and sign, and concluded that annually averaged DRE from residential combustion sources was -5 mW m^{-2} (Fig. 10). The negative effect of DRE in Butt et al. (2016) is partially driven by the inclusion of SO_2 emissions (8.9 Tg SO_2 yr^{-1}) from commercial coal combustion in the residential sector, leading to the cooling effect of sulfate and organic aerosols outweighing the warming from BC. For AIE, our control simulation is 38 times higher than that from Kodros et al. (2015) and over an order of magnitude higher than that from Butt et al. (2016). Consistent with our study, Ward et al. (2012) also found a large AIE (-1.74 to 1.00 W m^{-2}) for carbonaceous aerosols from fires using CESM CAM4-Chem. Both Kodros et al. (2015) and Butt et al. (2016) used offline radiative models to calculate AIE and only considered the first (albedo) aerosol indirect effect, which may partially explain the AIE differences. As mentioned

earlier, the AIE in our study includes aerosol first and second indirect effects as well as the semi-direct effect. Lacey and Henze (2015) estimated that the global surface air temperature changes due to solid wood fuel removal ranged from -0.28 K (cooling) to +0.16 K (warming), with a central estimate of -0.06 K (cooling). This cooling estimate is opposite to our study. However, we acknowledge that there are fundamental differences in calculating the radiative effect between our study and Lacey and Henze (2015), which employed absolute regional temperature potentials to quantify the climate responses.

Cookstove intervention programs have been implemented in developing countries, such as China, India and some African countries, to improve air quality and human health and to mitigate climate change (Anenberg et al., 2017; Aung et al., 2016; Carter et al., 2016). Our results suggest that large-scale efforts to replace inefficient cookstoves in developing countries with advanced technologies is not likely to reduce global warming through aerosol reductions, and may even lead to increased global warming when aerosol-cloud interactions are taken into account. Therefore, without improved constraints on BC interactions with clouds, especially mixed-phase and ice clouds, the net sign of the impacts of carbonaceous aerosols from solid fuel cookstoves on global climate (warming or cooling) remains ambiguous. This study does not include the greenhouse gas emission effects from the solid fuel cookstove sector, which may indeed be large enough to imply a net warming global climate impact depending on time scale (Lacey et al., 2017).

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Table 1. Annual budget for various species for the BASE, GBLSF_OFF and INDSF_OFF simulations for the year 2010.

Specie	ECLIPSE V5a (BASE)[a]	GBLSF_OFF[a]	INDSF_OFF[a]
BC	7.23	4.92	6.87
POM	18.9	8.53	17.2
SO_2	98.5	97.1	98.37
NO_x	120.5	118	119.8
VOC	81.1	52.4	76.6
CO	548	358	516
NH_3	54.9	54.6	54.87

[a]Units are Tg specie/yr.

Table 2. Model experiments setup.

Experiments	Anthropogenic emission scenario
BASE	ECLIPSE V5a
GBLSF_OFF	ECLIPSE V5a excluding global solid fuel cookstove emissions
INDSF_OFF	ECLIPSE V5a excluding Indian solid fuel cookstove emissions

878 **Table 3. Global budgets, burden and lifetime of BC and POM from model control**
879 **simulations.**

Specie	BC	POM
Sources (Tg specie/yr)	9.73	49.9
fossil fuel and biofuel	7.23	18.9
biomass burning emissions	2.5	31
Sinks (Tg specie/yr)	9.72	49.8
Dry Deposition	1.8	8.14
Wet Deposition	7.92	41.7
Burden (Tg)[a]	0.12 ± 0.001	0.66 ± 0.006
Lifetime (days)[a]	4.5 ± 0.04	4.8 ± 0.04

880 [a]standard deviation represents the uncertainty error owing to temporal variability in the
881 model.

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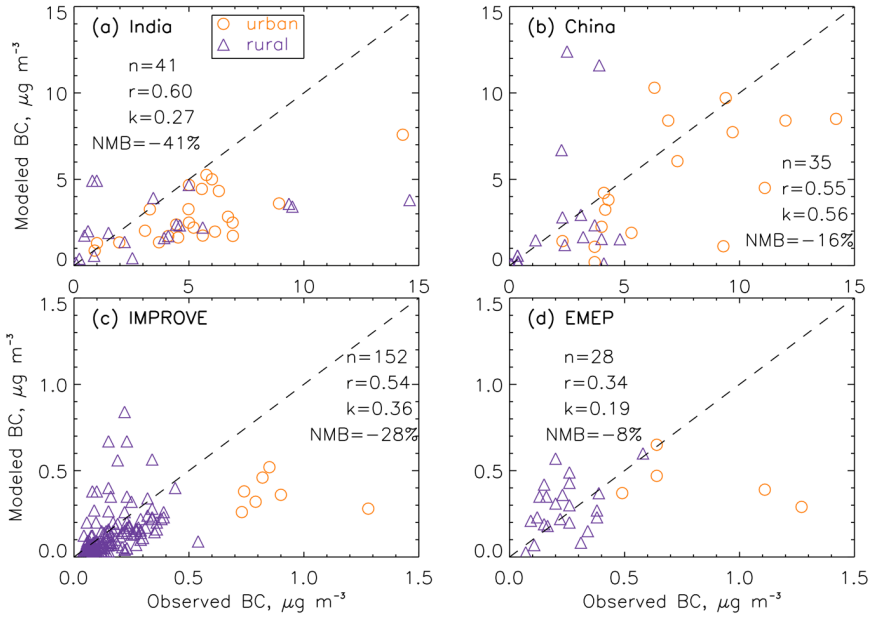

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892 **Figure 1.** Comparisons of observational and model simulated annual mean surface BC
893 concentrations from (a) India, (b) China, (3) IMPROVE, and (d) EMEP. Urban and rural sites are
894 shown in orange circles and blue triangles for each region. For each panel, the total number of
895 observational sites (n), model-to-observation regression slopes (k), correlation coefficient (r) and
896 NMB values are included. The dashed line in each panel represents the 1:1 ratio.

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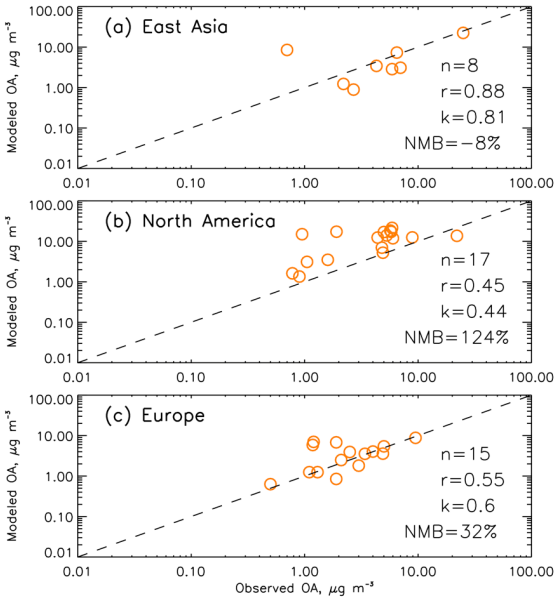

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901 **Figure 2.** Comparisons of observational and model simulated surface OA concentrations from (a)
902 East Asia, (b) North America, and (3) Europe. For each panel, the total number of observational
903 sites (n), model-to-observation regression slopes (k), correlation coefficient (r) and NMB values
904 are included. The dashed line in each panel represents the 1:1 ratio.

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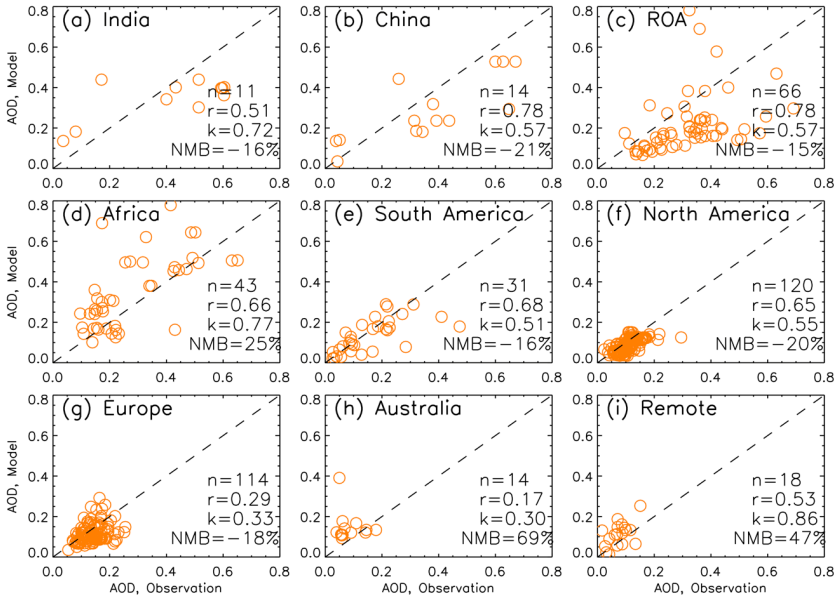

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908 **Figure 3.** Scatter plots of AOD between model simulation and observations over (a) India, (b)
909 China, (c) Rest of Asia (ROA), excluding China and India, (d) Africa, (e) South America, (f) North
910 America, (g) Europe, (h) Australia and (i) Remote. For each panel, the total number of
911 observational sites (n), model-to-observation regression slopes (k), correlation coefficient (r) and
912 NMB are included.

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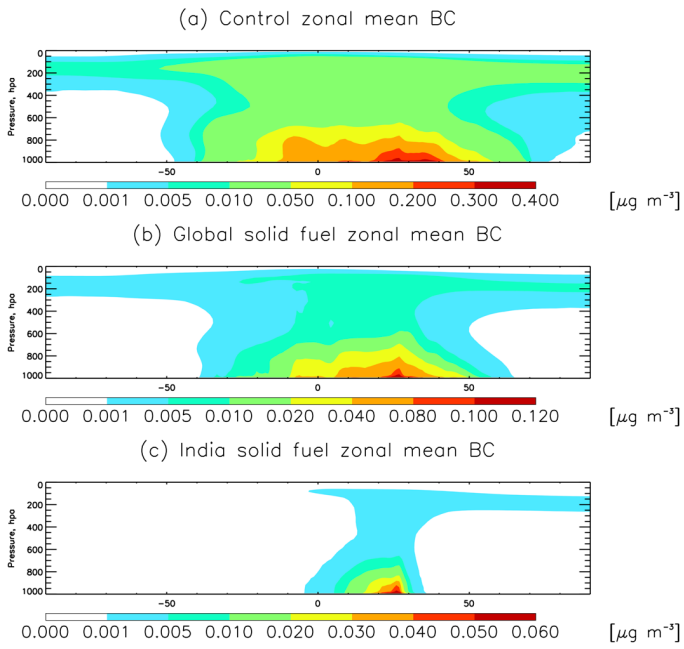

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915 **Figure 4.** Annual zonal mean BC concentrations from (a) the BASE simulation, (b) the global and
916 (c) India solid fuel cookstove emissions. BC concentrations are calculated under standard
917 temperature and pressure conditions (273 K, 1 atm).

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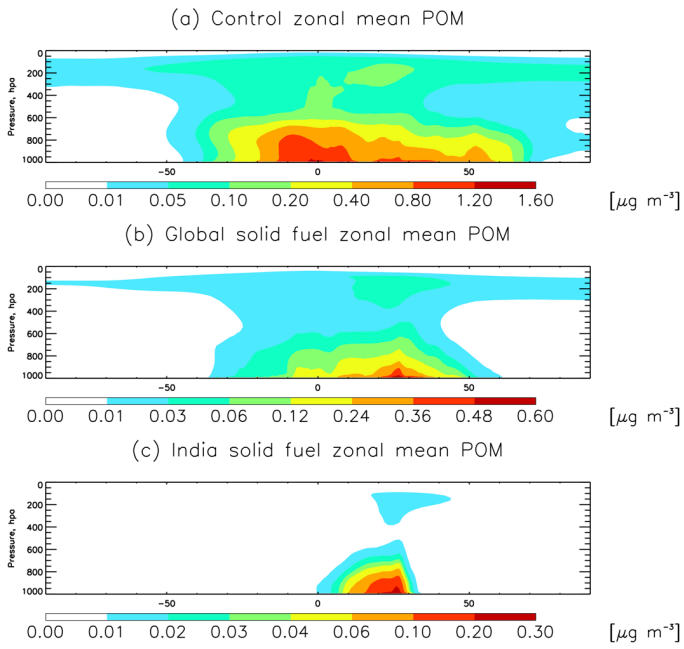


Figure 5. Same as Fig. 4 but for POM.

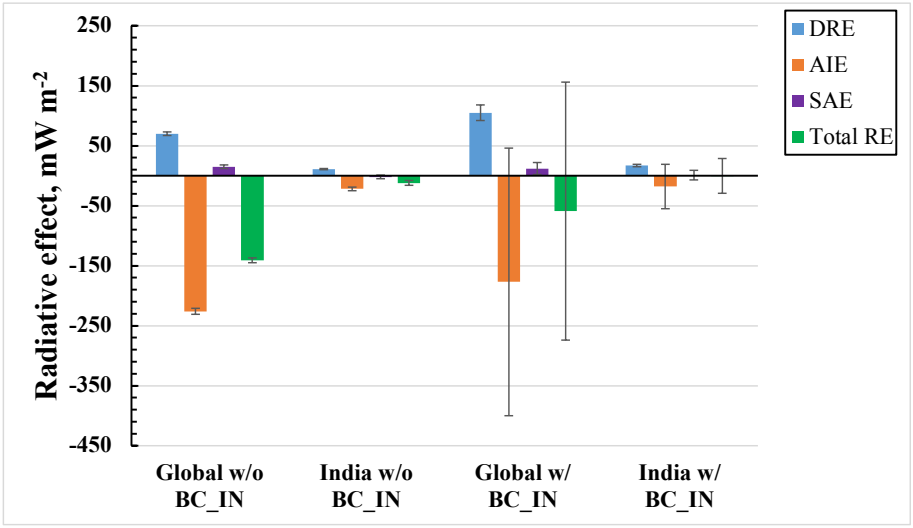
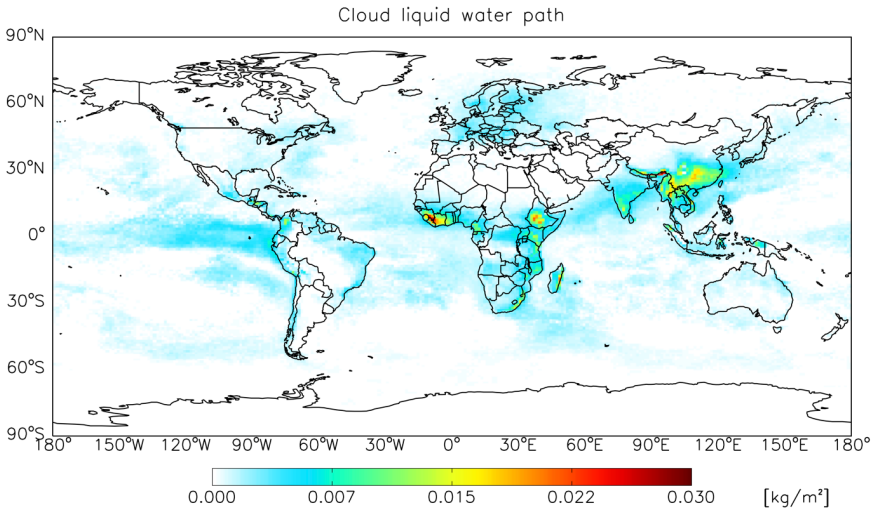


Figure 6. Radiative effect (RE) for global and Indian solid fuel cookstove aerosol emissions with BC not serving as IN (w/o BC_IN) and BC as IN (BC_IN), with DRE (blue), AIE (orange), SAE (purple) and total RE (green). Error bars represent one standard deviation for each RE. For BC as IN, standard deviations of RE are solely based on the choices of maximum freezing efficiency of BC as 0.01, 0.05 and 0.1 respectively.



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934 **Figure 7.** Global vertically-integrated cloud liquid water path from the global solid fuel cookstove
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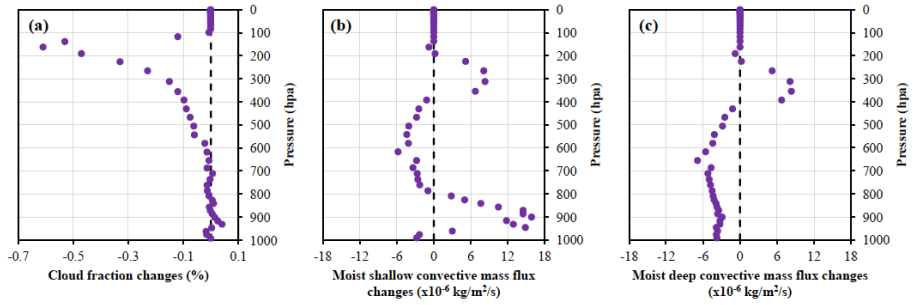
941 

942 **Figure 8.** Changes in vertical cloud fractions (a), shallow (b) and deep (c) convective mass flux
943 within the India and Indian Ocean domain from global solid fuel cookstove emissions.

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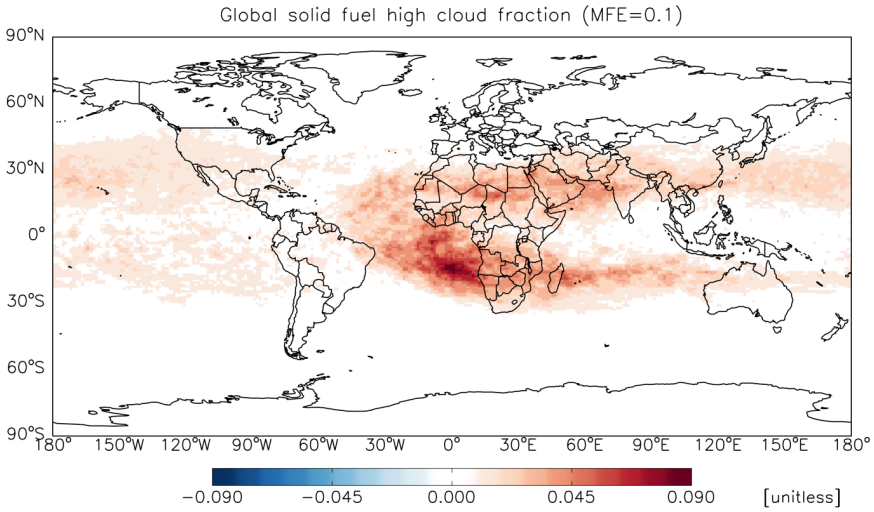

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950 **Figure 9.** Global distribution of high cloud fraction due to solid fuel cookstove aerosol emissions
951 with BC as IN and MFE=0.1.

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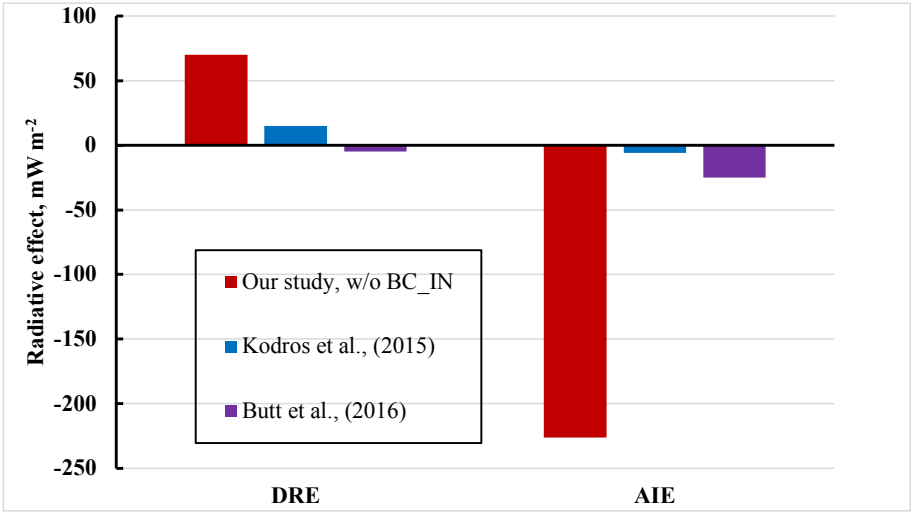

Figure 10. Comparisons of DRE (left) and AIE (right) radiative effects from global solid fuel cookstove emissions in our control simulation with Kodros et al. (2015) and Butt et al. (2016).