1	Future inhibition of ecosystem productivity by increasing wildfire pollution
2	over boreal North America
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4	Xu Yue ¹ , Susanna Strada ² , Nadine Unger ³ , Aihui Wang ⁴
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7	¹ Climate Change Research Center, Institute of Atmospheric Physics, Chinese Academy of
8	Sciences, Beijing 100029, China
9	² Laboratoire des Sciences du Climat et de l'Environnement, L'Orme des Merisiers - Bat 712,
10	91191 Gif Sur Yvette, France
11	³ College of Engineering, Mathematics and Physical Sciences, University of Exeter, Exeter,
12	EX4 4QE, UK
13	⁴ Nansen-Zhu International Research Centre, Institute of Atmospheric Physics, Chinese
14	Academy of Sciences, Beijing 100029, China
15	
15	Corresponding author
17	
18	Telenhone: 86-10-82995369
10	Email: vuvueseas@gmail.com
20	Eman: xuyueseas@gman.com
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Abstract

Biomass burning is an important source of tropospheric ozone (O_3) and aerosols. These air 30 31 pollutants can affect vegetation photosynthesis through stomatal uptake (for O_3) and light scattering and absorption (for aerosols). Wildfire area burned is projected to increase 32 33 significantly in boreal North America by the midcentury, while little is known about the 34 impacts of enhanced emissions on the terrestrial carbon budget. Here, combining site-level 35 and satellite observations and a carbon-chemistry-climate model, we estimate the impacts of fire emitted O₃ and aerosols on net primary productivity (NPP) over boreal North America. 36 37 Fire emissions are calculated based on an ensemble projection from 13 climate models. In the present day, wildfire enhances surface O₃ by 2 ppbv (7%) and aerosol optical depth (AOD) at 38 39 550 nm by 0.03 (26%) in the summer. By midcentury, area burned is predicted to increase by 40 66% in boreal North America, contributing more O_3 (13%) and aerosols (37%). Fire O_3 causes negligible impacts on NPP because ambient O₃ concentration (with fire contributions) 41 42 is below the damage threshold of 40 ppbv for 90% summer days. Fire aerosols reduce surface solar radiation but enhance atmospheric absorption, resulting in enhanced air stability and 43 44 intensified regional drought. The domain of this drying is confined to the North in the present day, but extends southward by 2050 due to increased fire emissions. Consequently, wildfire 45 aerosols enhance NPP by 72 Tg C yr⁻¹ in the present day but decrease NPP by 118 Tg C yr⁻¹ 46 in the future, mainly because of the soil moisture perturbations. Our results suggest that 47 future wildfire may accelerate boreal carbon loss, not only through direct emissions 48 increasing from 68 Tg C yr⁻¹ at present day to 130 Tg C yr⁻¹ by midcentury, but also through 49 the biophysical impacts of fire aerosols. 50

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53 1 Introduction

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55 Wildfire area burned is increasing in recent decades over North America boreal regions 56 (Stocks et al., 2002; Kasischke and Turetsky, 2006). Fire activity is closely related to weather 57 conditions and large-scale atmospheric oscillations (Gillett et al., 2004; Duffy et al., 2005), 58 and is projected to increase significantly in the future due to climatic changes (Flannigan et 59 al., 2005; Balshi et al., 2009; Groot et al., 2013; Wang et al., 2015). More area burned and the consequent fire emissions are accelerating carbon loss in boreal North America (Bond-60 Lamberty et al., 2007; Turetsky et al., 2011). Meanwhile, fire-induced air pollution, including 61 ozone (O₃) and aerosols, is predicted to increase in boreal and downwind regions by 62 midcentury (Yue et al., 2013; Yue et al., 2015). Wildfire emissions have large impacts on air 63 64 quality (Wotawa and Trainer, 2000; Morris et al., 2006), weather/climate conditions (Randerson et al., 2006; Zhao et al., 2014), and public health (Zu et al., 2016; Liu et al., 65 2017). However, little is known about how these pollutants affect ecosystem carbon 66 assimilation, and how this impact will change with the increased wildfire activity in the 67 future. 68

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70 Surface O₃ causes damages to photosynthesis through stomatal uptake (Sitch et al., 2007). In the present climate state, fire-induced O₃ enhancements are predicted to reduce net primary 71 productivity (NPP) in the Amazon forest by 230 Tg C yr⁻¹ (1 Tg = 10^{12} g), a magnitude 72 comparable to the direct release of CO_2 from fires in South America (Pacifico et al., 2015). 73 74 The aerosol effects are more uncertain because both positive and negative feedbacks occur. Appearance of aerosols increases diffuse light, which is beneficial for shaded leaves in the 75 76 lower canopy. Consequently, photosynthesis of the whole ecosystem will increase as long as the total light availability is not compromised (Kanniah et al., 2012). Rap et al. (2015) 77 estimated that biomass burning aerosols increase Amazon NPP by 78–156 Tg C yr⁻¹, which 78 79 offsets about half of the damage caused by fire O₃ (Pacifico et al., 2015). In contrast, strong light attenuation associated with high aerosol loading may decrease canopy photosynthesis 80 (Cohan et al., 2002; Oliveira et al., 2007; Cirino et al., 2014). Furthermore, the aerosol 81 radiative effects indirectly influence ecosystem productivity through concomitant 82 meteorological perturbations that are only beginning to be examined (Yue et al., 2017). 83

85 Future wildfire activity is projected to increase over boreal North America but with large uncertainties (Flannigan et al., 2005; Tymstra et al., 2007; Girardin and Mudelsee, 2008; 86 Nitschke and Innes, 2008; Amiro et al., 2009; Balshi et al., 2009; Bergeron et al., 2010; 87 Wotton et al., 2010; de Groot et al., 2013; Wang et al., 2016). For example, Amiro et al. 88 89 (2009) predicted an increase of 34% in Canadian area burned for a 2×CO₂ scenario (2040-2060) relative to a 1×CO₂ condition (1975-1995), using the Canadian Fire Weather Index 90 91 (CFWI) and output from Canadian global climate model (CGCM) version 1. Balshi et al. (2009) projected that area burned in boreal North America would double by the year 2045-92 2050 relative to 1991-2000, using the Multivariate Adaptive Regression Splines (MARS) 93 94 approach and meteorological output from CGCM version 2. The increasing rate in Balshi et al. (2009) is higher than that in Amiro et al. (2009), indicating substantial uncertainties in fire 95 projections originating from both fire models and simulated future climate. However, even 96 97 with the same fire models and climate change scenario, large uncertainties (in both magnitude and signs) are found in the projection of area burned among individual climate 98 models (Moritz et al., 2012; Yue et al., 2013). The multi-model ensemble approach has 99 shown superior predictability over single models in historical climate simulations (Flato et al., 100 2013) and near-term climate predictions (Kirtman et al., 2014), and has been used as a 101 102 standard technique to assess changes of climate variables in the long-term projections 103 (Collins et al., 2013). Following this strategy, Yue et al. (2015) used output from 13 climate 104 models to drive fire regression models and predicted an average increase of 66% in boreal area burned at 2046-2065 relative to 1981-2000 under the IPCC A1B scenario (Solomon et 105 106 al., 2007). Yue et al. (2015) further calculated that the wildfire emission increase by the 2050s would increase mean summertime surface O₃ by 5 ppbv in Alaska and 3 ppbv in 107 108 Canada. The study found regional maximum O₃ enhancements as high as 15 ppbv, suggesting 109 the potential for possible vegetation damage and land carbon loss due to the enhanced boreal 110 fire-related air pollution. Wildfire aerosols are also expected to increase significantly but not predicted in Yue et al. (2015). 111

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In this study, we quantify the impacts of O_3 and aerosols emitted from boreal wildfires on the land carbon uptake in North America in the present climate state and in the future world at 2050, taking advantage of the ensemble projection of future wildfire emissions by Yue et al. (2015). The major chain we investigate includes i) generation of aerosols and surface ozone from wildfire emissions and ii) impact of fire-emitted aerosols and ozone on plant

photosynthesis through physical and biogeochemical processes (Fig. 1). We first analyze 118 relationships between gross primary production (GPP) and aerosol optical depth (AOD) at 119 120 550 nm over the boreal regions based on observations. We then perform a suite of Earth 121 system model simulations using NASA GISS ModelE2 that embeds the Yale Interactive 122 Terrestrial Biosphere model (YIBs), a framework known as ModelE2-YIBs (Yue and Unger, 123 2015). Future projections of wildfire emissions from Yue et al. (2015) are applied as input to 124 ModelE2-YIBs model to project fire-induced O₃ and aerosol concentrations in the 2010s and 2050s. The impacts of the boreal fire O₃ on forest photosynthesis are predicted using the flux-125 based damage algorithm proposed by Sitch et al. (2007), which has been fully evaluated 126 127 against available O₃ damage sensitivity measurements globally and over North America (Yue and Unger, 2014; Yue et al., 2016; Yue et al., 2017). Fire aerosols induce perturbations to 128 129 radiation, meteorology, and hydrology, leading to multiple influences on the land carbon uptake. Sensitivity experiments are performed using the YIBs model in offline mode to 130 131 isolate the contributions of changes in the individual meteorological drivers.

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- 134 2 Materials and methods
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- 2.1 Observed GPP-AOD relationships
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Following the approach by Strada et al. (2015), we investigate the GPP sensitivity to diffuse 138 139 radiation and AOD variability in boreal regions. First, we identify study sites in Canada and Alaska from the AmeriFlux (AMF) network (http://ameriflux.lbl.gov/). There are much fewer 140 141 boreal sites than those in temperate regions. We select AMF sites providing hourly (or half-142 hourly) simultaneous measurements of GPP (non gap-filled) and photosynthetically active 143 radiation (PAR, total and diffuse) for at least 3 consecutive years. Only two Canadian sites meet the criteria: Groundhog River (CA-Gro, 82.2°W, 48.2°N), a mixed forest (MF), and 144 Quebec Mature Boreal Forest Site (CA-Qfo, 73.4°W, 49.7°N), an evergreen needleleaf forest 145 (ENF). At the two selected sites, we calculate the Pearson's correlation coefficients between 146 half-hourly GPP and different components of PAR. In total, we select 2432 and 3201 pairs of 147 GPP and PAR measurements at CA-Gro and CA-Qfo, respectively. We then apply 148 instantaneous Level 2 Collection 6 of AOD pixels at 3-km resolution retrieved by the 149 Moderate Resolution Imaging Spectroradiometer (MODIS, https://ladsweb.nascom.nasa.gov/) 150

151 onboard the Aqua and Terra satellites (Levy et al., 2013). The MODIS 3-km AOD product has been fully validated against ground-based sun photometers at both global (Remer et al., 152 153 2013) and urban/suburban (Munchak et al., 2013) scales. Strada et al. (2015) used ground-154 based AOD observations from the Aerosol Robotic Network (AERONET) near AMF sites to 155 validate the sampling technique of MODIS 3-km AOD product. They found high correlations 156 of 0.89-0.98 and regression slopes from 0.89 to 1.03 for daily AOD between AERONET and 157 MODIS at four AMF sites. For this study, the validation against ground-based AOD observations was not possible because no AERONET stations exist near to the selected AMF 158 159 sites.

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Every day, MODIS satellite sensors pass a specific region between 10:00 and 14:00 Local 161 162 Time (LT), leaving patchy signals around the AmeriFlux sites. Most of MODIS AOD data at high latitudes are available only in boreal summer; as a result, we narrow our explorations of 163 the GPP-AOD relationships to the noontime (10:00-14:00 LT) from June to August. The 164 chosen noontime window limits the contributions that confounding factors such as low solar 165 angles and high diffuse fraction may have on the amount of diffuse PAR and plant 166 productivity (Niyogi et al., 2004). For each summer day, we select instantaneous MODIS 3-167 168 km AOD pixels that are (a) located within a distance of 0.03° (about 3 km) from the targeted AMF site and (b) "quasi-coincident" with AMF data, which are available each half-hour. 169 170 Because of the unavoidable temporal differences between MODIS overpass and AMF data availability, we name this selection "quasi-coincident". A cloud mask applied to the MODIS 171 172 retrieval procedure conveniently filters out cloudy instants and should reduce the effect of clouds in the scattering process. We calculate both the correlation and regression coefficients 173 174 between "quasi-coincident" GPP and AOD at the selected sites. Negative GPP is considered 175 as a missing value. To further reduce the influence of cloud cover, we discard instants (both 176 AMF and MODIS data) when precipitation is non-zero. In total, we select 65 pairs of GPP 177 and AOD at CA-Gro site and another 59 pairs at CA-Qfo site. The GPP-AOD sampling pairs are much fewer than GPP-PAR, because we select instants when both instantaneous AOD 178 179 and GPP data are available. In addition, AOD is screened for clear instants to exclude the impacts of clouds. 180

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182 **2.2 Wildfire emissions**

Wildfire emissions used in climate modeling are calculated as the product of area burned, 184 fuel consumption, and emission factors. To predict area burned, we build stepwise 185 186 regressions for area burned in 12 boreal ecoregions (Yue et al., 2015). Observed area burned 187 aggregated from inter-agency fire reports is used as the predictand. Predictors are selected 188 from 44 ($5 \times 6 + 7 \times 2$) variables including five meteorological parameters (mean and maximum 189 temperature, relative humidity, precipitation, and geopotential height at 500 hPa) of six 190 different time intervals (winter, spring, summer, autumn, fire season (May-October), and the whole year), as well as the mean and maximum values of 7 fire indexes from the CFWI 191 system during fire season. We consider the impacts of antecedent factors on current fire 192 193 activity by including all above variables at the same year and those in the previous two years, making a total of 132 (44×3) factors. The final formats of regression are different among 194 195 ecoregions, depending on the selection of the factors that contribute the maximum observed 196 variance in predictand but remain the minimum collinearity among predictors. These 197 regression functions are then driven with output from 13 Coupled Model Intercomparison Project Phase 3 (CMIP3) climate models under A1B scenario (Meehl et al., 2007) to predict 198 199 area burned at present day (1981-2000) and midcentury (2046-2065). In the A1B scenario, CO₂ concentration is projected to 532 ppm by the year 2050, similar to the value of 541 ppm 200 201 in IPCC RCP8.5 scenario (van Vuuren et al., 2011) archived for the Coupled Model 202 Intercomparison Project Phase 5 (CMIP5).

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We derive $1^{\circ} \times 1^{\circ}$ gridded area burned based on the prediction for each ecoregion following 204 205 the approach by Yue et al. (2015). Temporally, the annual area burned estimated with regressions is first converted to monthly area burned using the mean seasonality for each 206 207 boreal ecoregion during 1980-2009. Spatially, large fires tend to burn in ecosystems where 208 historical fires are frequent because of favorable conditions (Keane et al., 2008). In each 1°×1° 209 grid square, we calculate the frequency of large fires (>1000 ha) during 1980-2009; these 210 fires account for about 85% of total area burned in boreal North America. We arbitrarily attribute 85% of area burned within each ecoregion to a number of fires with fixed size of 211 1000 ha. We then allocate these large fires among the $1^{\circ} \times 1^{\circ}$ grid cells based on the observed 212 spatial probability of large fires. For example, if one grid box (named grid 'A') bears 1% of 213 large fires (>1000 ha) within an ecoregion at present day, the same grid will bear the same 214 215 possibility for large fires in the future. On the other hand, fuel availability limits reburning and fire spread during the forest return interval, suggesting that current burning will decrease 216

217 the possibility of future fires in the same location. To consider such impact, we scale the observed probabilities by the fraction remaining unburned in each grid box, and then use this 218 219 modified probability distribution to allocate large fires for the remaining months. For 220 example, if present-day fires have consumed 20% of the total area within the grid 'A', then 221 the possibility of large fire will be 0.8% (1%×0.8, instead of 1%) for this grid. Finally, we 222 disaggregate the remaining 15% of area burned into fires 10 ha in size, and randomly 223 distribute these fires across all grid boxes in the ecoregion. With this method, we derive the gridded area burned for boreal North America by eliminating reburning issues. Sensitivity 224 tests show that specifying different area burned to the large fires (100 or 10 000 ha rather 225 than 1000 ha) yields < 1 % changes in predicted biomass burned, suggesting that this 226 227 approach is not sensitive to the presumed fire size in the allocation procedure.

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Fuel consumption, the dry mass burned per fire area, is the product of fuel load and burning 229 severity. For fuel load in Alaska, we use 1-km inventory from the US Forest Service (USFS) 230 Fuel Characteristic Classification System (FCCS, McKenzie et al., 2007). For fuel load in 231 232 Canada, we use a 1-km fuel type map from the Canadian Fire Behavior Prediction (FBP) 233 system (Nadeau et al., 2005), combined with fuel-bed definition from the FCCS. Burning 234 severity, the fraction of fuel load burned by fires, is calculated with the USFS CONSUME 235 model 3.0 following the approach described in Val Martin et al. (2012). With both fuel load 236 and burning severity, we derive fuel consumption and further calculate biomass burned in boreal North America with the predicted area burned. As in Amiro et al. (2009) and Yue et al. 237 238 (2015), we apply constant fuel load for both present day and midcentury because opposite and uncertain factors influence future projections (Kurz et al., 2008; Heyder et al., 2011; 239 240 Friend et al., 2014; Kim et al., 2017). Instead, we consider changes in burning severity due to 241 perturbations in fuel moisture as indicated by CFWI indexes (Yue et al., 2015). On average, 242 we estimate a 9% increase in fuel consumption over boreal North America by the midcentury, because higher temperature and lower precipitation result in a future with drier fuel load 243 (Flannigan et al., 2016). 244

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Fire emissions for a specific species are then estimated as the product between biomass burned and the corresponding emission factor, which is adopted from measurements by Andreae and Merlet (2001) except for NO_x . We use the average value of 1.6 g NO per Kg dry mass burned (DM) from six studies as NOx emission factor, because the number of 3.0 g NO per Kg DM reported in Andreae and Merlet (2001) is much higher than that of 1.1 g NO per
Kg DM from field observations (Alvarado et al., 2010). Based on projected area burned and
observation-based fuel consumption and emission factors, we derive fire emissions of NO_x,
carbon monoxide (CO), non-methane volatile organic compounds (NMVOCs, Alkenes and

Alkanes), NH₃, SO₂, black (BC) and organic carbon (OC) in the present day and midcentury.

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256 2.3 NASA ModelE2-YIBs model

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The NASA ModelE2-YIBs is an interactive climate-carbon-chemistry model, which couples 258 259 the chemistry-climate model NASA ModelE2 (Schmidt et al., 2014) and the YIBs vegetation model (Yue and Unger, 2015). NASA ModelE2 is a general circulation model with 260 horizontal resolution of 2°×2.5° latitude by longitude and 40 vertical layers up to 0.1 hPa. It 261 dynamically simulates both the physical (emissions, transport, and deposition) and chemical 262 263 (production, conversion, and loss) processes of gas-phase chemistry (NO_x, HO_x, O_x, CO, CH₄, and NMVOCs), aerosols (sulfate, nitrate, ammonium, BC, OC, dust, and sea salt), and their 264 265 interactions. In the model, oxidants influence the photochemical formation of secondary aerosol species (e.g., sulfate, nitrate, and biogenic secondary organic aerosol), in turn, 266 267 aerosols alter photolysis rates and influence the online gas-phase chemistry. Size-dependent 268 optical parameters computed from Mie scattering, including extinction coefficient, single 269 scattering albedo, and asymmetry parameters, are applied for each aerosol type (Schmidt et al., 2014). The model also considers interactions between climate and atmospheric 270 271 components. Simulated climate affects formation, transport, and deposition of atmospheric components, in turn, both O₃ and aerosols influence climate by altering radiation, temperature, 272 273 precipitation, and other climatic variables. Both observation-based evaluations and multi-274 model inter-comparisons indicate that ModelE2 demonstrates skill in simulating climatology 275 (Schmidt et al., 2014), soil moisture (Fig. S1), radiation (Wild et al., 2013), atmospheric 276 composition (Shindell et al., 2013b), and radiative effects (Shindell et al., 2013a).

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YIBs is a process-based vegetation model that dynamically simulates changes in leaf area
index (LAI) through carbon assimilation, respiration, and allocation for prescribed PFTs.
Coupled photosynthesis-stomatal conductance is simulated with the Farquhar-Ball-Berry
scheme (Farquhar et al., 1980; Ball et al., 1987). Leaf-level photosynthesis is upscaled to
canopy level by separating diffuse and direct light for sunlit and shaded leaves (Spitters,

1986). Plant respiration considers thermal dependence as well as acclimation to temperature 283 (Atkin and Tjoelker, 2003). Soil respiration is calculated based on the carbon flows among 12 284 285 biogeochemical pools (Schaefer et al., 2008). Net carbon uptake is allocated among leaves, 286 stems, and roots to support leaf development and plant growth (Cox, 2001). The YIBs model 287 has been benchmarked against in situ GPP from 145 eddy covariance flux tower sites and 288 satellite retrievals of LAI and phenology (Yue and Unger, 2015). An interactive flux-based 289 O₃ damage scheme proposed by Sitch et al. (2007) is applied to quantify the photosynthetic responses to ambient O₃ (Yue and Unger, 2014). For this scheme, O₃ damaging level is 290 dependent on excess O_3 stomatal flux within leaves, which is a function of ambient O_3 291 concentration, boundary layer resistance, and stomatal resistance. Reduction of 292 photosynthesis is calculated on the basis of plant functional types (PFTs), each of which 293 294 bears a range of low-to-high sensitivities to O₃ uptake.

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297 2.4 Simulations

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299 Using the NASA ModelE2-YIBs model, we perform 6 time-slice simulations, three for 300 present-day (2010s) and three for midcentury (2050s), with atmosphere-only configuration to explore the impacts of fire emissions on NPP in boreal North America (Table 1). Simulations 301 302 F10CTRL and F50CTRL turn off all fire emissions as well as O₃ vegetation damage for the 2010s and 2050s, respectively. However, climatic feedbacks of aerosols from other sources 303 304 (both natural and anthropogenic) and related photosynthetic responses are included. Simulations F10AERO and F50AERO consider the responses of plant productivity to 305 306 perturbations in radiation and meteorology caused by aerosols, including emissions from 307 wildfires and other sources, but do not include any O₃ vegetation damage. In contrast, 308 simulations F10O3 and F50O3 calculate offline O₃ damage based on the simulated O₃ from 309 all sources including fire emissions. For these simulations, reductions of GPP are calculated twice with either low or high O₃ sensitivity. However, both of these GPP changes are not fed 310 back into the model to influence carbon allocation and tree growth. Plant respiration is 311 changing in response to meteorological perturbations, either due to climate change or aerosol 312 radiative effects. We assume no impact of O₃ damage to plant respiration and examine 313 vegetation NPP, the net carbon uptake by biosphere, for the current study. The difference 314 between AERO and CTRL runs isolates the impacts of fire aerosols on NPP, and the 315

difference between O3 and CTRL runs isolates O₃ vegetation damage caused by fire and nonfire emission sources.

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All simulations are conducted for 20 years and outputs for the last 15 years are used for 319 320 analyses. The simulations apply sea surface temperatures (SSTs) and sea ice distributions 321 from previous NASA GISS experiments under the IPCC RCP8.5 scenario (van Vuuren et al., 322 2011). Decadal average monthly-varying SST and sea ice of 2006-2015 are used as boundary conditions for present-day (2010s) runs while that of 2046-2055 are used for future (2050s) 323 324 runs. In the RCP8.5 scenario, global average SST increases by 0.62 °C while sea ice area decreases by 13.8% at the midcentury compared to the present-day level. Decadal average 325 326 well-mixed greenhouse gas concentrations and anthropogenic emissions of short-lived 327 species, both at present day and midcentury, are adopted from the RCP8.5 scenario (Table 2). 328 The enhancement of CO₂ will affect climate (through longwave absorption) and ecosystem 329 productivity (through CO₂ fertilization), but not the fire activity and related emissions 330 directly. Natural emissions of soil and lightning NO_x, biogenic volatile organic compounds 331 (BVOC), dust, and sea salt are climate-sensitive and simulated interactively. The YIBs vegetation model cannot simulates changes in PFT fractions. The RCP8.5 land cover change 332 333 dataset shows limited changes in land cover fractions between 2010s and 2050s (Oleson et al., 2010). For example, relative to the 2010s, a maximum gain of 5% is predicted for grassland 334 335 in the 2050s, resulting from a 1% loss in deciduous forest and another 1% loss in needleleaf 336 forest over boreal North America. As a result, a land cover dataset derived from satellite 337 retrievals (Hansen et al., 2003) is applied as boundary conditions for both the 2010s and 2050s. 338

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To evaluate the simulated GPP responses to changes in diffuse radiation, we perform site-340 341 level simulations using standalone YIBs model, which is driven with observed hourly 342 meteorology (including temperature, relative humidity, surface pressure, wind speed, and soil moisture) and both diffuse and direct PAR at sites CA-Gro and CA-Qfo. To isolate the 343 344 impact of individual aerosol-induced climatic perturbations on NPP, we perform 10 345 sensitivity experiments using the offline YIBs model driven with offline meteorology 346 simulated by ModelE2-YIBs model (Table 3). For example, the offline run Y10 CTRL is driven with variables from the online simulation of F10CTRL (Table 1). The run Y10 TAS 347 348 adopts the same forcing as Y10 CTRL except for temperature, which is simulated by the climate simulation of F10AERO. In this case, we quantify the NPP responses to individual 349

and/or combined climate feedback (mainly in temperature, radiation, and soil moisture) by
fire aerosols. Each offline run is conducted for 12 years and the last 10 years are used for
analyses.

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354 2.5 Observation datasets

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356 We use observations to evaluated GPP, AOD, and O₃ in boreal North America simulated by ModelE2-YIBs. For GPP, we use a benchmark data product upscaled from FLUXNET eddy 357 covariance data using an ensemble of regression trees (Jung et al., 2009). For AOD 358 observations, we use satellite retrieval at 550 nm from Terra MODIS Level 3 data product. 359 360 For O₃, gridded datasets are not available. We use site-level observations from 81 U.S. sites at the Clean Air Status and Trends Network (CASTNET, https://www.epa.gov/castnet) and 361 362 202 Canadian sites at the National Air Pollution Surveillance (NAPS, http://www.ec.gc.ca/rnspa-naps/) program. All datasets are averaged over the 2008-2012 363 period to represent present-day climatological conditions. Gridded datasets are interpolated to 364 the same $2^{\circ} \times 2.5^{\circ}$ resolution as ModelE2-YIBs model. 365

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368 3 Results

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370 3.1 Observed GPP-AOD relationships

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Positive correlations between GPP and diffuse PAR are found at the two boreal sites (Figs 372 373 2b-2c). The magnitude of diffuse PAR is similar for these sites, possibly because they are 374 located at similar latitudes (Fig. 2a). GPP values at CA-Gro are generally higher than that at 375 CA-Qfo, likely because deciduous broadleaf forest (DBF) has higher photosynthetic rates. Consequently, the slope of regression between GPP and PAR_{dif} is higher at CA-Gro than that 376 at CA-Qfo, suggesting that GPP of DBF (or MF) is more sensitive to changes in diffuse PAR 377 than that of ENF. We find almost zero correlation between GPP and PAR_{dir} at the two sites 378 379 (Table 4), indicating that photosynthesis is in general light-saturated for sunlit leaves at these sites during boreal summer noontime. As a result, modest reductions in direct light by 380 381 aerosols will not decrease GPP of the whole canopy.

With satellite-based AOD, we find positive correlations between GPP and AOD at both sites (Figs 2d-2e). However, the slope of regression between GPP and AOD is lower (and not significant) at CA-Gro compared with that at CA-Qfo, opposite to the GPP-PAR_{dif} regressions. The cause of such discrepancy might be related to the limitation of data availability. For the same reason, the GPP-AOD correlation is insignificant at CA-Gro site. On average, GPP sensitivity (denoted as mean \pm range) is estimated as $3.5 \pm 1.1 \mu mol m^{-2} s^{-1}$ per unit AOD at lower latitudes of boreal regions in the summer.

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391 3.2 Model evaluations

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Simulated summer GPP shows high values in mid-western Canada (Alberta and 393 394 Saskatchewan) and the Southeast (Ontario) (Fig. 3a). Forest GPP at high latitudes is low because of the cool weather and light limitation there. Simulated GPP reasonably captures the 395 spatial distribution with a high correlation coefficient of 0.77 ($p \ll 0.01$) and relatively small 396 biases within 20% of the data product. Simulated AOD reproduces the observed spatial 397 pattern including the high values in boreal forests (Fig. 3b). In contrast to the MODIS 398 observations, predicted AOD is relatively uniform over the West with a background value of 399 400 ~0.1. This discrepancy explains the low correlation coefficient (R = 0.25, p < 0.01) between the model and MODIS data. The simulation fails to capture the high values in the west, 401 402 possibly due to a climate model underestimation of biogenic secondary organic aerosol, which may be an important contribution over the western boreal forest. Simulated maximum 403 404 daily 8-hour average (MDA8) [O₃] shows low values in boreal North America and high values in the western and eastern U.S. (Fig. 4a). This pattern is consistent with surface 405 406 observations (Fig. 4b), but the model overestimates the measured surface O_3 by 22%. The 407 Canadian measurement sites are located near the southern boundary, and as a result do not 408 represent the average state over the vast boreal region at higher latitudes.

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With the Sitch et al. (2007) scheme, the YIBs model simulates reasonable GPP responses to [O₃] in North America (Yue and Unger, 2014; Yue et al., 2016). Generally, damage to GPP increases with the enhancement of ambient [O₃], but with varied sensitivities for different plant species (see Fig. 6 of Yue and Unger (2014)). In responses to the same level of [O₃], predicted O₃ damages are higher for deciduous trees than that for needleleaf trees, consistent with observations from meta-analyses (Wittig et al., 2007). The model also reproduces 416 observed light responses of GPP to diffuse radiation in boreal regions. With the site-level 417 simulations, we evaluate the modeled GPP-PAR_{dif} relationships at the hourly (instead of halfhourly) time step during summer. For 1342 pairs of GPP and PAR_{dif} at the site CA-Gro, the 418 419 observed correlation coefficient is 0.42 and regression slope is 0.011, while the results for the 420 simulation are 0.60 and 0.014, respectively. At the site CA-Qfo, the observations yield a 421 correlation coefficient of 0.46 and regression slope of 0.007 for 1777 pairs of GPP and PAR_{dif}. The simulated correlation is 0.61 and the regression is 0.011 at the same site. The 422 423 GPP sensitivity to PAR_{dif} in the model is slightly higher than that of the available observations, likely because the latter are affected by additional non-meteorological abiotic 424 factors. To remove the influences of compound factors other than radiation, we follow the 425 426 approach of Mercado et al. (2009) to discriminate GPP responses to 'diffuse' and 'direct' 427 components of PAR at the two sites (Fig. 5). The model successfully reproduces the observed GPP-to-PAR sensitivities. Increase in PAR boosts GPP, but the efficiency is much higher for 428 429 diffuse light than that for direct light, suggesting that increase of diffuse radiation is a benefit for plant growth. 430

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- 432 **3.3 Simulation of wildfire O₃ and aerosols**
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During 1980-2009, wildfire is observed to burn 2.76×10^6 ha and 156.3 Tg DM every year 434 over boreal North America. Similarly, the ensemble prediction with fire regression models 435 estimates present-day area burned of 2.88×10^6 ha yr⁻¹ and biomass burned of 160.2 Tg DM 436 yr⁻¹ (Yue et al., 2015). By the midcentury, area burned is projected to increase by 77% (to 437 5.10×10^6 ha yr⁻¹) in boreal North America, mainly because of the higher temperature in 438 future fire seasons. Consequently, biomass burned increases by 93% (to 308.6 Tg DM yr⁻¹) 439 because fuel consumption also increases by 9% on average in a drier climate (Yue et al., 440 441 2015). Enhanced fire emissions increase concentrations of surface O₃ and column AOD, 442 especially over Alaska and central Canada (Fig. 6). The maximum centers of air pollutants are collocated for O₃ and AOD but with unproportional magnitudes, suggesting non-linear 443 444 conversion among fire emission species as well as the interactions with natural emission 445 sources (e.g., lightning/soil NO_x and BVOC). On average, wildfire emissions contribute $7.1 \pm$ 3.1% (2.1 ± 0.9 ppbv) to surface O₃ and $25.7 \pm 2.4\%$ (0.03 ± 0.003) to AOD in the summer 446 over boreal North America in the present day. By midcentury, these ratios increase 447 significantly to $12.8 \pm 2.8\%$ (4.2 ± 0.9 ppbv) for O₃ and $36.7 \pm 2.0\%$ (0.05 ± 0.003) for AOD. 448

450 **3.4 Simulation of fire pollution impacts on NPP**

451

Surface O₃, including both fire and non-fire emissions (Table 2), causes limited (1-2%) 452 453 damages to summer GPP in boreal North America (Fig. 7). The most significant damage is 454 predicted over eastern U.S., where observed $[O_3]$ is high over vast forest ecosystems (Fig. 4). 455 In the western U.S., [O₃] is also high but the O₃-induced GPP reduction is trivial because low stomatal conductance in the semi-arid ecosystems limits O3 uptake there (Yue and Unger, 456 457 2014). Over boreal North America, dominant PFTs are ENF (accounting for 44% of total vegetation cover) and tundra (treated as shrubland, accounting for 41% of total vegetation 458 459 cover). Both species have shown relatively high O₃ tolerance with a damaging threshold of 40 ppbv as calculated with Sitch's scheme (Yue and Unger, 2014). For boreal regions, the 460 461 mean $[O_3]$ of 28 ppbv (Fig. 4a) is much lower than this damaging threshold, explaining why the excess O₃ stomatal flux (the flux causing damages) is low there (Fig. 8). Statistics in Yue 462 et al. (2015) show that maximum daily 8-hour average (MDA8) [O₃] with fire contributions 463 can be higher than 40 ppbv in Alaska and Canada. However, such episodes appear at 95 464 percentile for present day and 90 percentile for midcentury, suggesting that O₃ vegetation 465 damage is rare in boreal North America and fire-induced O3 enhancement does not 466 467 exacerbate such damages. Therefore, we do not consider O₃ damage effects further.

468

Fire aerosols cause significant perturbations in shortwave radiation at surface (Fig. 9). The 469 470 direct light is largely attenuated especially over Alaska and central Canada, where fire aerosols are most abundant (Fig. 6). In contrast, diffuse light widely increases due to particle 471 scattering. In the present day, the average reduction of 5.6 W m⁻² in the direct light 472 component is in part offset by the enhancement of 2.6 W m^{-2} in the diffuse light component. 473 leading to a net reduction of 3.0 W m^{-2} in solar radiation over boreal North America. By the 474 midcentury, a stronger reduction of 9.5 W m^{-2} in direct light is accompanied by an increase of 475 4.0 W m⁻² in diffuse light, resulting in a net reduction of 5.5 W m⁻² in solar radiation. Fire-476 induced BC aerosols strongly absorb solar radiation in the atmospheric column (Figs 10a-477 10b). On average, fire aerosols absorb 1.5 W m⁻² in the present day and 2.6 W m⁻² by the 478 479 midcentury.

481 Atmospheric circulation patterns respond to the aerosol-induced radiative perturbations (Figs 10c-10d). Surface radiative cooling and atmospheric heating together increase air stability 482 483 and induce anomalous subsidence. In the present day, such descending motion is confined to 484 55-68°N, accompanied by a rising motion at 52-55°N (Fig. 10c). As a result, fire aerosols 485 induce surface warming at higher latitudes but cooling at lower latitudes in boreal regions 486 (Fig. 11a). Meanwhile, precipitation is inhibited by the subsidence in northwestern Canada 487 but is promoted by the rising motion in the Southwest (Fig. 11c). By the midcentury, the range of subsidence expands southward to 42°N (Fig. 10d) due to strengthened atmospheric 488 heating (Fig. 10b). The downward convection of warm air offsets surface radiative cooling 489 (Fig. 9b), leading to a significant warming in the Southwest (Fig. 11b). The expanded 490 491 subsidence further inhibits precipitation in vast domain of Canada (Fig. 11d). Soil moisture is 492 closely related to rainfall and as a result exhibits dipole changes (drier north and wetter south) 493 in the present day (Fig. 11e) but widespread reductions (Fig. 11f) by the midcentury.

494

495 In response to the climatic effects of fire aerosols, boreal NPP shows distinct changes 496 between the present day and midcentury (Fig. 12). Such changes in NPP are a consequence of 497 changes in GPP and autotrophic respiration (Fig. S2). Variations in plant respiration resemble 498 those of GPP, because higher photosynthesis leads to faster leaf/tissue development, resulting larger maintenance and growth respiration. In the 2010s, forest NPP increases by 5-15% in 499 500 Alaska and southern Canada, but decreases by 5-10% in northern and eastern Canada. This pattern of NPP changes (Δ NPP) is connected to the climatic effects of aerosols, especially 501 502 changes in soil moisture (Fig. 11). The correlation between ΔNPP (Fig. 12a) and changes in 503 soil moisture (Fig. 11e) reaches R = 0.56 (n = 356), much higher than the values of R = -0.11504 for temperature change (Fig. 11a) and R = 0.22 for precipitation change (Fig. 11c). At the continental scale, the patchy responses of NPP offset each other. Since the dominant fraction 505 of carbon uptake occurs in southern Canada (Fig. 3a), where positive NPP change is 506 predicted (Fig. 12a), wildfire aerosols enhance the total NPP by 72 Tg C yr⁻¹ in the present 507 day (Table 5). In contrast, increased wildfire emissions in the 2050s inhibit precipitation (Fig. 508 11d) and decrease soil moisture in boreal North America (Fig. 11f), leading to widespread 509 NPP reductions and a total NPP loss of 118 Tg C yr⁻¹ (Fig. 12b, Table 5). 510

- 511
- 512

513 4 Discussion

515 4.1 Roles of aerosol climatic feedback

516

The contrasting sign of NPP responses in the present day and midcentury are closely related 517 to the aerosol-induced surface climatic feedback. Sensitivity experiments using offline YIBs 518 519 model (Table 3) allowed assessment of the impacts of individual changes in the major meteorological drivers, including temperature, radiation (diffuse and direct), and soil 520 521 moisture (Table 5). The offline simulations driven with changes in all three variables yield Δ NPP of 126 Tg C yr⁻¹ for the 2010s and -97 Tg C yr⁻¹ for the 2050s. These values are 522 different from the online simulations, which predict ΔNPP of 72 Tg C yr⁻¹ for the 2010s and -523 118 Tg C yr⁻¹ for the 2050s. Missing of other aerosol climatic feedbacks in the offline model, 524 for example, changes in relative humidity, surface pressure, soil temperature, and turbulence 525 526 momentum, may cause such discrepancy between the online and offline simulations. Seasonal analyses show that summertime $\triangle NPP$ is 99 Tg C at present day and -95 Tg C at 527 528 midcentury, dominating the NPP changes all through the year, because both wildfire 529 emissions and ecosystem photosynthesis maximize in boreal summer.

530

531 Observations show that aerosols can promote plant photosynthesis through increasing diffuse 532 radiation (Niyogi et al., 2004; Cirino et al., 2014; Strada et al., 2015). Our analyses with ground data also show positive correlations between GPP and PAR_{dif} (Fig. 2 and Table 4), 533 and the model reproduces observed GPP responses to perturbations in direct and diffuse PAR 534 (Fig. 5). Wildfire aerosols enhance diffuse radiation by 2.6 W m⁻² (1.7%) at present day and 535 4.0 W m⁻² (2.3%) at midcentury in boreal North America (Fig. 9). With these changes, 536 simulated NPP increases by 8 Tg C yr⁻¹ at the 2010s and 14 Tg C yr⁻¹ at the 2050s (Table 5). 537 Near the two AmeriFlux sites (Fig. 2a), wildfires increase local AOD by 0.03 (Fig. 6c). 538 Meanwhile, we estimate that summer average (00:00-24:00) GPP increases by 0.04 μ mol m⁻² 539 s⁻¹ in the same region due to aerosol diffuse fertilization effects (DFE) based on the results of 540 (Y10 PAR – Y10 CTRL). This change suggests a simulated GPP sensitivity of 1.2 μ mol m⁻² 541 s^{-1} (22%) per unit AOD. Observed GPP sensitivity to AOD at the two sites are 2.3 (19%) and 542 4.5 µmol m⁻² s⁻¹ (58%) per unit AOD, respectively (Figs 2d-2e). The absolute value of GPP 543 sensitivity from simulations is much smaller than that of observations, because the former is 544 545 for 24-h average while the latter is only for noontime (10:00-14:00). The relative change of 22% in YIBs model falls within the observed range of 19-58%. 546

The estimated NPP changes of 8 Tg C yr⁻¹ by the radiative effects of boreal fire aerosols are 548 much weaker than the enhancement of 78-156 Tg C yr⁻¹ by fires in Amazon basin (Rap et al., 549 2015). There are at least two reasons for such a difference in the DFE between boreal and 550 551 Amazon fire aerosols. First, wildfire emissions and associated impacts on radiation are much smaller in boreal regions. Wildfires in Alaska and Canada directly emit 68 Tg C yr⁻¹ at the 552 2010s, resulting in enhancement of summer AOD by 35% and diffuse radiation by 1.7%. 553 These boreal emissions are much smaller than the \sim 240 Tg C yr⁻¹ in Amazon basin (van der 554 Werf et al., 2010), where fires enhances regional PM2.5 concentrations by 85% and diffuse 555 556 radiation by 6.2% in dry seasons (Rap et al., 2015). Second, larger solar insolation in lower 557 latitudes allows stronger DFE for the same unit change of diffuse radiation. In our prediction, most of NPP changes occur at high latitudes of boreal regions (Fig. 12), where total 558 559 insolation is not so abundant as that at the tropical areas. Consequently, decline of direct 560 radiation in boreal regions more likely converts the light availability of sunlit leaves from light-saturation to light-limitation, offsetting the benefit from enhanced diffuse radiation for 561 shaded leaves. For this study, we do not find GPP reduction by the decline of direct light at 562 the two Ameriflux sites (Table 4), possibly because these sites are located at middle latitudes 563 (<50°N). In the future, more observations at higher latitudes (> 55°N) are required to explore 564 the sensitivity of GPP to AOD at the light-limited conditions. 565

566

Simulations have shown that absorbing aerosols can cause regional drought by increasing air 567 568 stability (Liu, 2005; Cook et al., 2009; Tosca et al., 2010). Our results confirm such tendency but with varied range of hydrological responses depending on the magnitude of wildfire 569 570 emissions (Figs 11c-11f). Observations suggest that precipitation (and the associated soil 571 moisture) is the dominant driver of the changes in GPP over North America, especially for 572 the domain of cropland (Beer et al., 2010). Sensitivity experiments with offline YIBs model 573 show that changes in soil moisture account for 82.5% of Δ NPP at present day and 70.5% of Δ NPP at midcentury (Table 5). These results suggest that aerosol-induced changes in soil 574 575 water availability, instead of temperature and radiation, dominantly contribute to the changes 576 of boreal NPP, consistent with observational and experimental results (Ma et al., 2012; Girardin et al., 2016; Chen et al., 2017). 577

578

579 4.2 Limitations and uncertainties

In this study, we examine the interactions among climate change, fire activity, air pollution, 581 582 and ecosystem productivity. To reduce the complexity of the interactions, we focus on the most likely dominant feedback and thus main chain of events: "climate \rightarrow fire \rightarrow pollution \rightarrow 583 584 biosphere' (Fig. 1). However, our choice of feedback analysis does not mean that the 585 interplay of other processes is unimportant. For example, climate-induced changes in vegetation cover/types can influence fire activity by alteration of fuel load, and air pollution 586 587 by BVOC emissions (climate \rightarrow biosphere \rightarrow fire/pollution). In addition, other feedbacks may 588 amplify ecosystem responses but are not considered. For example, the drought caused by fire 589 aerosols in the midcentury (Fig. 11) may help increase fire activity (fire \rightarrow pollution \rightarrow 590 climate \rightarrow fire). Furthermore, we apply fixed SSTs in the climate simulations because reliable 591 ocean heat fluxes for the future world were not available. Many previous studies have 592 investigated regional aerosol-climate feedbacks without ocean responses. For example, Cook 593 et al. (2009) found that dust-climate-vegetation feedback promotes drought in U.S., with a 594 climate model driven by prescribed SSTs. Similarly, Liu (2005) found fire aerosols enhance 595 regional drought using a regional climate model, which even ignores the feedback between 596 local climate and large-scale circulation. While we do concede that our experimental design is not a complete assessment of all known processes and feedbacks, within these limitations, 597 598 this study for the first time quantifies the indirect impacts of wildfire on long-range 599 ecosystem productivity under climate change.

600

We use the ensemble projected fire emissions from Yue et al. (2015). Area burned is 601 602 predicted based on the simulated meteorology from multiple climate models. Such an 603 approach may help reduce model uncertainties in climatic responses to CO₂ changes (Collins et al., 2013; Kirtman et al., 2014), but cannot remove the possible biases in the selection of 604 605 climate scenarios and fire models. All predictions in Yue et al. (2015) are performed under 606 the IPCC A1B scenario. With two different scenarios, A2 of high emissions and B2 of low 607 emissions, Balshi et al. (2009) showed that future area burned in boreal North America 608 increases at a similar rate until the 2050s, after which area burned in A2 scenario increases 609 much faster than that in B2 scenario. On average, boreal area burned in Balshi et al. (2009) 610 increases by ~160% at 2051-2060 compared with 2001-2010, much higher than the change of 611 66% in Yue et al. (2015). In contrast, Amiro et al. (2009) predicted that boreal area burned at the $2 \times CO_2$ scenario increases only by 34% relative to the $1 \times CO_2$ scenario. This ratio is only 612

613 half of the estimate in Yue et al. (2015), which compared results between periods with $1.44 \times CO_2$ and $1 \times CO_2$. The discrepancies among these studies are more likely attributed to 614 615 the differences in fire models. Although both Amiro et al. (2009) and Yue et al. (2015) 616 developed fire-weather regressions in boreal ecoregions, the former study did not include 617 geopotential height at 500 hPa and surface relative humidity as predictors, which make 618 dominant contributions to area burned changes in the latter study. On the other hand, Balshi 619 et al. (2009) developed nonlinear regressions between area burned and climate at grid scale, which helps retain extreme values at both the temporal and spatial domain. Compared to 620 previous estimates, Yue et al. (2015) predicted median increases in future fire emissions over 621 622 boreal North America.

623

624 We apply constant land cover and fuel load for both present day and midcentury, but we 625 estimate an increase in fuel consumption due to changes in fuel moisture. Future projection of 626 boreal fuel load is highly uncertain because of multiple contrasting influences. For example, using a dynamic global vegetation model (DGVM) and an ensemble of climate change 627 projections, Heyder et al. (2011) predicted a large-scale dieback in boreal-temperate forests 628 629 due to increased heat and drought stress in the coming decades. On the contrary, projections 630 using multiple DGVMs show a widespread increase in boreal vegetation carbon under the global warming scenario with CO₂ fertilization of photosynthesis (Friend et al., 2014). In 631 632 addition, compound factors such as greenhouse gas mitigation (Kim et al., 2017), pine beetle 633 outbreak (Kurz et al., 2008), and fire management (Doerr and Santin, 2016) may exert varied 634 impacts on future vegetation and fuel load. Although we apply constant fuel load, we consider changes of fuel moisture because warmer climate states tend to dry fuel and increase 635 636 fuel consumption (Flannigan et al., 2016). With constant fuel load but climate-driven fuel 637 moisture, we calculate a 9% increase in boreal fuel consumption by the midcentury (Yue et 638 al., 2015). Although such increment is higher than the prediction of 2-5% by Amiro et al. 639 (2009) for a doubled- CO_2 climate, the consumption-induced uncertainty for fire emission is likely limited because changes in area burned are much more profound. 640

641

Predicted surface $[O_3]$ is much higher than observations over boreal North America (Fig. 4). This bias does not affect main conclusions of this study, because predicted O_3 causes limited damages to boreal GPP even with the overestimated $[O_3]$ (Fig. 7). The result confirms that fire-induced O_3 vegetation damage is negligible in boreal North America. For aerosols, the 646 model captures reasonable spatial pattern of AOD but with a background value of ~ 0.1 647 outside fire-prone regions, where the observed AOD is usually 0.1-0.2 (Fig. 3). This 648 discrepancy may be related to the insufficient representations of physical and chemical 649 processes in the model, but may also result from the retrieval biases in MODIS data due to 650 the poor surface conditions (Liu et al., 2005) and small AOD variations (Vachon et al., 2004) 651 at high latitudes.

652

Simulated aerosol climatic effects depend on radiative and physical processes implemented in 653 the climate model. We find that present-day boreal fire aerosols on average absorb 1.5 W m^{-2} 654 in the atmosphere (Fig. 10), which is much smaller than the value of 20.5 ± 9.3 W m⁻² for 655 fires in equatorial Asia (Tosca et al., 2010). This is because boreal fires enhance AOD only 656 657 by 0.03 while tropical fires increase AOD by ~0.4. Previous modeling studies showed that fire plumes induce regional and downwind drought through enhanced atmospheric stability 658 (Feingold et al., 2005; Tosca et al., 2010; Liu et al., 2014). Most of these results were based 659 on the direct and/or semi-direct radiative effects of fire aerosols. Inclusion of the indirect 660 aerosol effect may further inhibit precipitation and amplify drought, but may also introduce 661 additional uncertainties for the simulations. The fire-drought interaction may promote fire 662 activity, especially in a warmer climate. Ignoring this interaction may underestimate future 663 area burned and the consequent emissions. 664

665

666 4.3 Implications

667

Inverse modeling studies have shown that the land ecosystems of boreal North America are 668 carbon neutral in the present day, with the estimated land-to-air carbon flux from -270 ± 130 669 Tg C yr⁻¹ to 300 ± 500 Tg C yr⁻¹ (Gurney et al., 2002; Rodenbeck et al., 2003; Baker et al., 670 2006; Jacobson et al., 2007; Deng et al., 2014). Here, we reveal a missing land carbon source 671 672 due to future wildfire pollution, taking into account full coupling among fire activity, climate change, air pollution, and the carbon cycle. Fire pollution aerosol increases boreal NPP by 72 673 Tg C yr⁻¹ in the present day, comparable to the direct carbon loss of 68 Tg C yr⁻¹ from 674 wildfire CO₂ emissions (product of biomass burned and CO₂ emission factors). By 675 midcentury, increasing fire emissions instead cause a NPP reduction of 118 Tg C yr⁻¹ due to 676 677 the amplified drought. Although NPP is not a direct indicator of the land carbon sink, reduction of NPP is always accompanied with the decline of net ecosystem exchange (NEE) 678

and the enhanced carbon loss. In combination with the enhanced carbon emission of 130 Tg C yr⁻¹, future boreal wildfire presents an increasing threat to the regional carbon balance and global warming mitigation. Furthermore, the NPP reductions are mostly located in southern Canada, where cropland is the dominant ecosystem, newly exposing the future wildfirerelated air pollution risk to food production.

684

685 Our analyses of fire pollution effects on boreal North American productivity may not be representative for other boreal ecosystems and/or on the global scale. There is substantial 686 variability in plant species, topography, and climatology across different boreal regions. Such 687 differences indicate distinct GPP sensitivities as well as fire characteristics. At lower latitudes, 688 where anthropogenic pollution emissions are more abundant, ambient ozone concentrations 689 may have exceeded damaging thresholds for most plant species. In those regions, additional 690 691 ozone from a fire plume may cause more profound impacts on photosynthesis than our estimate for boreal North America. For example, Amazonian fire is predicted to reduce forest 692 NPP by 230 Tg C yr⁻¹ through the generation of surface ozone (Pacifico et al., 2015). 693 694 Meanwhile, solar radiation is more abundant at lower latitudes, indicating more efficient increases in photosynthesis through aerosol DFE because the sunlit leaves receive saturated 695 696 direct light in those regions. As shown in Beer et al. (2010), partial correlations between GPP 697 and solar radiation are positive in boreal regions but negative over the subtropics/tropics, 698 suggesting that light extinction by fire aerosols has contrasting impacts on plant photosynthesis in the high versus low latitudes. Further simulations and analyses are required 699 700 to understand the net impacts of ozone and aerosols from biomass burning on the global 701 carbon cycle.

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Youth Talents Plan". Nadine Unger acknowledges funding support from The University of
Exeter.

710 711 712 Table 1. Online simulations with ModelE2-YIBs climate model ^a 713

Simulations	SST	$[CO_2]$	Emissions	Fires	O ₃ effect	Aerosol effect
F10O3	2010s	2010s	2010s	2010s	Yes	No
F10AERO	2010s	2010s	2010s	2010s	No	Yes
F10CTRL	2010s	2010s	2010s	No	No	Yes
F50O3	2050s	2050s	2050s	2050s	Yes	No
F50AERO	2050s	2050s	2050s	2050s	No	Yes
F50CTRL	2050s	2050s	2050s	No	No	Yes

714

 a Values of SST, [CO₂], and emissions are adopted from RCP8.5 scenario, with the average

of 2006-2015 for the 2010s and that of 2046-2055 for the 2050s. For fire emissions, values at
the 2010s are predicted based on meteorology for 1981-2000 and those at the 2050s are for
2046-2065.

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Table 2. Emissions from wildfires and non-fire sources over boreal North America

Graning	Fire emission	ons (Tg yr ⁻¹)	Non-fire emiss	sions (Tg yr ⁻¹)
species	2010s	2050s	2010s	2050s
NO _x ^a	0.39	0.74	2.43	2.08
СО	15.7	28.8	5.9	4.0
SO ₂ ^a	0.12	0.22	1.95	1.28
NH ₃	0.22	0.40	0.80	1.15
BC	0.08	0.16	0.03	0.01
OC	1.10	2.04	0.04	0.02
NMVOC	0.39	1.34	0.49	0.30
BVOC ^b	N/A	N/A	15.3	15.1

^a Natural emissions are included for NO_x (lightning and soil) and SO₂ (volcano). ^b ModelE2-YIBs calculates BVOC emissions using photosynthesis-dependent scheme implemented by Unger et al. (2013).

Table 3. Simulations with YIBs vegetation model driven by offline meteorology from
ModelE2-YIBs climate model

Simulations	Base forcing	Temperature	PAR	Soil moisture
Y10_CTRL	F10CTRL			
Y10_ALL	F10CTRL	F10AERO	F10AERO	F10AERO
Y10_TAS	F10CTRL	F10AERO		
Y10_PAR	F10CTRL		F10AERO	
Y10_SLM	F10CTRL			F10AERO
Y50_CTRL	F50CTRL			
Y50_ALL	F50CTRL	F50AERO	F50AERO	F50AERO
Y50_TAS	F50CTRL	F50AERO		
Y50_PAR	F50CTRL		F50AERO	
Y50_SLM	F50CTRL			F50AERO

743
744 Table 4. Pearson's correlation coefficients for GPP-PAR and GPP-AOD relationships at
745 Ameriflux (AMF) sites ^a

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Site	Period ^b	Pearson's R					
Site	I chida	GPP-PAR	GPP-PAR _{dir}	GPP-PAR _{dif}	GPP-AOD	AOD-PAR _{dif}	AOD-PAR _{dir}
CA-Gro	2004-2013	0.19 (2432)	-0.01 (2432)	0.42 (2432)	0.15 (65)	0.60 (65)	-0.52 (65)
CA-Qfo	2003-2014	0.16 (3201)	-0.04 (3201)	0.45 (3201)	0.36 (59)	0.91 (34)	-0.80 (34)

747

^a Both GPP and PAR (direct PAR_{dir} and diffuse PAR_{dif}) data are adopted from site-level AMF measurements. AOD data are adopted from instantaneous MODIS Aqua and Terra 3-km retrievals. Correlations are calculated for quasi-coincident AMF and MODIS data over summer noontime (June-August, 10:00-14:00 Local Time). The sampling number for each correlation is denoted in brackets. Significant (p<0.05) correlation coefficients are bolded. ^b For CA-Gro site, diffuse PAR observations of 2005-2009 have been discarded because of poor calibration, as documented on the AMF website.

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Table 5. Changes in NPP (Tg C yr⁻¹) caused by composite and individual climatic effects of
fire aerosols

	2010s	2050s
Online ^a	72	-118
Offline total ^b	126	-97
Temperature	11	-22
Radiation	8	14
Soil moisture	104	-86

^a Online results are calculated using the ModelE2-YIBs model with (F10AERO – F10CTRL)
 for the 2010s and (F50AERO – F50CTRL) for the 2050s.

^b Offline results are calculated with the YIBs model driven with individual or combined
 changes in temperature, radiation, and soil moisture.

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- organic compound (NMVOC)) and associated BVOC changes causes direct damage to plantphotosynthesis.



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Figure 2. Relationships between (b, c) GPP and diffuse PAR and (d, e) GPP and MODIS 1170 AOD at (a) two boreal sites: Groundhog River (Gro) and Quebec Mature Boreal Forest Site 1171 1172 (Qfo). The two sites are from the AmeriFlux network in Canada and are dominated by mixed forest (MF at Gro) and evergreen needleleaf forest (ENF at Qfo) (Table 1). Data cover 1173 summer days (June-August). AmeriFlux diffuse PAR and GPP (in μ mol m⁻² s⁻¹) are half-1174 hourly observations (10:00-14:00 LT). Instantaneous MODIS Aqua and Terra 3-km AOD are 1175 selected in a time span centered on AmeriFlux record time. For each plot: the red line 1176 indicates the regression line, black lines depict the 1- σ interval; the regression slope and 1177 correlation coefficient are both included for each site (in bold if statistically significant at 95% 1178 1179 confidence level). Blue dots in (b, c) show instants when MODIS Aqua and Terra 3-km 1180 AODs overlap AmeriFlux data.

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Figure 3. Evaluation of simulated summer (a) GPP and (b) AOD at 550 nm with (c, d)
observations. Simulation results are from F10AERO (Table 1). Each point on the (e, f) scatter
plot represents one grid square in boreal North America. The number of points (n),
correlation coefficient (r), and relative bias (b) for the evaluation are presented on the plot.



Figure 4. Evaluation of simulated summer surface maximum daily 8-hour average [O₃] with observations for 2008-2012. Observations are collected from 81 U.S. sites at the Clean Air Status and Trends Network (CASTNET) and 202 Canadian sites at the National Air Pollution Surveillance (NAPS) program. The number of points (n), correlation coefficient (r), and mean bias (b) for the evaluation are presented on the plot. Values over Canada and Alaska are denoted with blue points.

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1242Figure 5. Observed (blue) and simulated (red) response of GPP to diffuse (square) and direct1243(triangle) PAR at boreal sites (a) CA-Gro (2004-2013) and (b) CA-Qfo (2004-2010).1244Observations and simulations are split into 'diffuse' and 'direct' conditions if the diffuse1245fraction is >0.8 and < 0.2, respectively. Data points are then averaged over PAR bins of 30 W</td>1246m⁻² with error bars indicating one standard deviation of GPP for each bin.



Figure 6. Changes in summer (a, b) $[O_3]$ and (c, d) AOD at 550 nm induced by wildfire emissions in (a, c) the 2010s and (b, d) the 2050s over boreal North America. Only significant changes (p < 0.05) are shown.



Figure 7. Simulated O₃ damages to summer GPP in North America. Results shown are from simulations with (a, b) low and (c, d) high O₃ sensitivities for (a, c) 2010 and (b, d) 2050.
Simulated [O₃] includes contributions from both wildfire and non-fire emissions. Results for 2010 are derived as (F10O3/F10CTRL-1)×100%. Results for 2050 are derived as (F50O3/F50CTRL-1)×100%.



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Figure 8. Simulated summertime O_3 stomatal fluxes in boreal North America. Results shown are the (a, b) mean and (c, d) excess flux at (a, c) 2010 and (b, d) 2050. Simulated $[O_3]$ includes contributions from both wildfire and non-fire emissions. Excess O_3 stomatal flux is calculated as the difference between the stomatal flux and a PFT-specific threshold as defined in Sitch et al. (2007).

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(a) Δ SW_tot by fire aerosol in 2010



Figure 9. Changes in surface radiative fluxes induced by wildfire aerosols in boreal North America. Results shown are for the changes in summertime (June-August) (a, b) total, (c, d) direct, and (e, f) diffuse solar radiation at surface caused by aerosols from wildfire emissions at (a, c, e) present day and (b, d, f) midcentury. Significant changes (p < 0.05) are marked with black dots. Results for 2010 are calculated as (F10AERO - F10CTRL). Results for 2050 are calculated as (F50AERO - F50CTRL).



Figure 10. Predicted (a, b) absorption of shortwave radiation and (c, d) perturbations in vertical velocity by wildfire aerosols at (a, c) present day and (b, d) midcentury. The absorption of shortwave radiation is calculated as the differences of radiative perturbations between top of atmosphere and surface. Vertical velocity is calculated as the longitudinal average between 105°W and 112.5°W (two blue lines in a). Positive (negative) values indicate descending (rising) motion. Results for the 2010s are calculated as (F10AERO -F10CTRL). Results for the 2050s are calculated as (F50AERO - F50CTRL). Significant changes (p < 0.05) in (a, b) are indicated as black points.



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Figure 11. Predicted changes in summertime (a, b) surface air temperature, (c, d) precipitation, and (e, f) soil water content at surface caused by aerosols from wildfire emissions at (a, c, e) present day and (b, d, f) midcentury. Results for temperature and precipitation are shown as absolute changes. Results for soil water are shown as relative changes. Results for the 2010s are calculated as (F10AERO - F10CTRL). Results for the 2050s are calculated as (F50AERO - F50CTRL). Significant changes (p<0.05) are marked with black dots.





Figure 12. Predicted percentage changes in summer NPP caused by wildfire aerosols at (a) present day and (b) midcentury. Results for the 2010s are calculated as (F10AERO/F10CTRL -1) × 100%. Results for the 2050s are calculated as (F50AERO/F50CTRL - 1)×100%. Significant changes (p<0.05) are marked with black dots.